



2023 Annual Report



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Printed on March 31st, 2024



I. Names, Titles, Contact telephone numbers, and E-mail addresses of the Spokesperson and Deputy Spokesperson:

Spokeperson name: Deputy Spokeperson name:

Lee-Cheng Liu Chih-Jung Chang

Title: Title:

Chairman and President Senior Vice-President

Tel: +886-2-7708-0123 Tel: +886-2-7708-0123

E-mail: IR@eirgenix.com E-mail: IR@eirgenix.com

II. Address and Telephone of Headquarters, Branches and Plant:

<u>Address</u> <u>Telephone</u>

Headquarters Xizhi No. 101, Lane 169, Kangning St., Xizhi Dist, +886-2-7708-0123

New Taipei City 22180

Branches No.168, Sec. 1, Shengyi Rd., Zhubei +886-3-620-5088

Zhubei City, Hsinchu County 302

III. Firm name, Address, Website URL and Telephone number of Stock Transfer Agent:

KGI Securities, Department of Stock Agency

Address: 5F, No. 2, Sec. 1, Chongqing S. Rd., Zhongzheng Dist., Taipei City 100

Tel: +886-2-2389-2999

Website URL: https://www.kgi.com.tw

IV. Firm name, Address, Website URL, Telephone number, and the name of the CPA who attested the most recent year's financial report

Name of the CPA: Sheng-Wei Deng, Yu-Fang Yen\

Firm name: PricewaterhouseCoopers Taiwan

Address: 27F, No. 333, Sec. 1, Keelung Rd., Xinyi Dist., Taipei City 110

Website URL: http://www.pwc.tw Tel: +886-2-2729-6666

V. The name of any exchanges where the company's securities are traded offshore, and the method by which to access information on said offshore securities: None.

VI. Company Website URL: http://www.eirgenix.com



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Auditors' Report

2023 Independent Financial Statements and Independent

Auditors' Report



I. Letter to Shareholders

Dear Shareholders,

1. 2023 Business Result

(1) Business plan implementing results

EirGenix was established on December 21, 2012 and listed in the market on June 28, 2019. It is a biotechnology and medical company focusing on biosimilars, drug discovery, and biopharmaceutical Contract Development and Manufacturing Organization (CDMO). The revenue was NT\$1,022,653 thousand in 2023 and NT\$1,481,017 thousand in 2022. The difference was mainly due to (1) deferred recognition of the milestone payments as a result of a delay in overseas medicine certificate, (2) facility expansion and equipment re-validation in Zhubei, and (3) the decrease in the market demand in CDMO. EirGenix holds the critical technology of biotechnological drug development and manufacture and is able to provide differentiated services with high value-added. Once production line expansion and upgrade have been completed, the growth momentum of revenues will resume. The consistent and stable operating income can cover part of the development expense for biosimilars. Various drug development projects are being implemented successively as planned. EirGenix's financial and business condition will rise substantially after obtaining the medicine certificate for mass production.

(2) Research and development status

A. Establish competitive and complete production line development strategies:

- (A) EirGenix is currently developing the product for the treatment of HER2+ breast cancer. The dual-target treatment with Pertuzumab in combination with Trastuzumab for late-stage HER2+ breast cancer is gradually being used for early-stage breast cancer. EG1206A is one of the biosimilar leaders in the Pertuzumab market. This will also boost the market share of EG12014.
- (B) The exclusive licensee of EG12014 (Trastuzumab Biosimilar), Sandoz, maintains close communication with the FDA (Food and Drug Administration). Sandoz also works with EirGenix and suppliers for improvements based on the FDA's opinions in the shortest timeframe possible and files a resubmission after the response. Within six months after the resubmission, the FDA will provide the review results in response to the application for approval. Meanwhile, the license of the



- EC (European Commission) and TFDA (Taiwan Food and Drug Administration) has been approved.
- (C) Phase III clinical trial of EG1206A (Pertuzumab Biosimilar) is expected to apply in 2024.
- (D) The antibody-drug conjugate (ADC) EG12043 (TSY0110), jointly developed by the Company and Formosa Pharmaceuticals, expected to apply for Phase I clinical trial in 2024.
- B. Outstanding development and manufacture technology of biotechnological drugs:
 - (A) EirGenix's Zhubei plant has passed the review by the FDA and obtained an EIR (Establishment Inspection Report) before the drug launch.
 - (B) The CDMO contracts signed in 2023 reached a total value of NT\$1.1 billion (US\$36 million). The CAGR (compound annual growth rate) was 27.3% in 2017-2023.
 - (C) In 2023, the mammalian capacity reached 25,500L and the microbial capacity reached 150 L. Building B at the Zhubei plant is expected to be completed in 2026, to increase the microbial capacity to 1,500 L. Meanwhile, a three-stage expansion of the mammalian plant which has 150,000 L capacity, is under planning at Ciaotou Science Park, Kaohsiung.
 - (D) Granted Accreditation Certificate of Foreign Drug Manufacturer by Japan MHLW, with the accreditation category of "biological products" and effective date from October 24, 2022 to October 30, 2027.

C. Affirmation on business performance:

- (A) Received the approval letter from TFDA that the API Trastuzumab has obtained the license and the DMF number.
- (B) The Phase I clinical trial of the biosimilar EG1206A has met its primary endpoint and showed Pharmacokinetic biosimilarity.
- (C) TD.
- (D) Won the highest honor of "New Taipei City Family Friendly Work Equality Measures" in 2023.
- (E) The biosimilar drug, "EIRGASUN vial 150 mg", has been approved by National Health Insurance Administration to be enrolled in the reimbursement system.
- (F) Received a positive CHMP opinion for the biosimilar drug EG12014 licensed.



- (G) Top 5% among TPEx-listed companies in the 9th Corporate Governance Evaluation.
- (H) Received the approval letter from EC for EG12014 licensed.
- (I) EIRGASUN vial 150 mg won the Golden award of National Pharmaceutical Technology Research Development Award in 2023.

(3) Financial revenue and expenditure and profitability analysis

The annual operating incomes are NT\$ 1,022,653 thousand dollars, which are mainly contributed by CDMO business and cooperative development revenue. The gross profit is NT\$ 236,741 thousand dollars with a 23% gross margin rate. The major expenditures in 2023 were biosimilars development and research expenses. The reason for that is because the products are still in the development stage and require more investments for research and development funds, such as clinical study expenses, research and development material expenses, and research and development staff salaries. CDMO sales and other revenues are still unable to fully cover the research and development expenditures mentioned previously at this point, which is the main reason that caused EirGenix's loss. The investment of research and development expenditures now is to accumulate the energy for future profit growth after the product launches.

Unit: %

Item	Year	2023	2022
Financial	Debt Ratio	10.26	9.83
Structure	Long Term Funds to property, plant, and equipment	313.25	426.10
Calvamay	Current Ratio	977.92	1,133.94
Solvency	Quick Ratio	868.41	1,015.88
	Rate of return on assets	(7.88)	(0.93)
Duafitability	Rate of return on equity	(8.84)	(1.09)
Profitability	Net Profit Margin	(89.49)	(7.80)
	Earnings per share (NTD)	(\$3.00)	(\$0.38)

(4) Budget implementation status

EirGenix had only set up an internal budget goal for 2023 and did not disclose the financial forecast to the public. The overall budget implementation has met the goal.

2. 2024 Business Plan Summary



(1) Business policy

EirGenix's business policy is to maintain the sustainable growth since its establishment. It came up with three major service items after considering the three factors of the sales and developing time of drugs, risk value, and potential returns, three stages of the business focus have been set: 1. Contract Development and Manufacturing Organization (CDMO); 2. Biosimilar Development, and 3. Me too and Novel biologics development to make the best of EirGenix's cGMP production factory, equipment, and high-end technology human resources.

(2) Estimated sales, and its basis, and important production and sales policy.

EirGenix's biosimilars in development are still in the developing stage. The main revenue resource comes from Contract Development and Manufacturing Organization and authorized product collaborations. The senior management team proposes the overall goal and strategy, and the research and development team make various development project plans. The project schedule for plan implementation and sales projection is made by feasibility analysis, market potential and financial evaluation.

3. EirGenix's future development strategy

- (1) The short-term development strategy is "Build up the foundation and move forward step by step." The strategy plans for products in development and CDMO sales & marketing development are as follows:
 - A. EG12014 approved by the FDA and other countries in Aisa.
 - B. EG12014 (HERWENDA® Sandoz | EIRGASUN® EirGenix) market launch
 - C. EG1206A submit the application for Phase III trials.
 - D. Application for EG12043 (TSY0110) clinical trials (IND).
 - E. EG1211X pre-clinical preparation completed.
 - F. Expansion of Building B at Zhubei plant to increase the microbial capacity to 1,500 L in 2026.
- (2) The medium- and long-term development strategy is "Products are developing and launching one after another to promote stable growth in revenue. The strategy plans for products in development and CDMO sales development are as follows:
 - A. New dosage forms or new drug delivery systems of biosimilars: development of Trastuzumab high-concentration subcutaneous doses; planning for the development of EG12014+EG1206A dual-targeting high-concentration subcutaneous doses. The successful development of high-concentration



subcutaneous doses will strengthen the product market share of these products and enable EirGenix as the primary supplier of biosimilar drugs for the treatment of HER2+ breast cancer.

B. Developing the biosimilar for the treatment of blood cancer are currently ongoing. According to the development schedule, one new product will be introduced to the market each one to two years starting in 2027. Hence, a three-stage expansion of the mammalian capacity by 150,000L is under planning at Ciaotou Science Park, Kaohsiung. The new capacity can be used to manufacture in-house developed drugs and accept customers' orders for commercial and scale production.

4. Effects by the external competitive environment, legal environment, and overall business environment

The mission of EirGenix at the beginning is to provide high-quality and cost-effective Contract Development and Manufacturing Organization and develop biosimilars with commercial values. The medium to long-term goal is focusing on Niche Biologics development to benefit the human and the society and improve the life quality. EirGenix insists on making the technology first with excellent quality as the foundation and be responsible for customer's success. The goal is to become an international biotechnology and medicine company that begins in Taiwan and focuses on the global market.

We would like to thank all of the shareholders, customers, and collaborating business partners for encouraging and supporting us, as well as the contribution and hard work from our employees. Together it brings prosperity and constant growth for EirGenix.

EirGenix, Inc.

Chairman & President: Lee-Cheng Liu

Head of Accounting Department: Hsiu-Chuan Yang



II. Company Profile

1. Date of Incorporation

December 21st, 2012.

2. Company History

- 2012 EirGenix Inc. was incorporated as a company limited by shares and registered under the provisions of the Company Act of the Republic of China.
- On March 15th, 2013, EirGenix, Inc., Formosa Laboratories, Inc., and Development Center for Biotechnology (DCB) signed a joint venture agreement. EirGenix Inc. (EirGenix) obtained the management rights and completed the transfer of all technologies, R&D, and production personnel in April 2013. Meanwhile EirGenix inherited the existing pilot plant and the R&D core, competencies including cell line development, production process development, protein characterization, quality control, and two Taiwan FDA certified cGMP facilities one for mammalian cells and one for microbial.
 - Completed capital injection in November, with the capital reaching NT\$ 540 million.
- 2014 Granted PIC/S GMP certificate by Taiwan FDA.
- 2015 Completed capital injection with the capital reaching NT\$ 790 million.
 - Received the Gold Prize for "Biomedical and New Agricultural Industry Award" in 2015.
- 2016 Initiated EG12014 Phase I clinical trial in Europe.
 - Completed capital injection, with the capital reaching NT\$ 1.0097 billion.
 - Completed IPO and publicly listed in TPEx Emerging Stock Board.
 - Initiation construction of the new PIC/S GMP bio-pharmaceutical facility with commercial mass-production scale situated in the Zhubei Biomedical Park at the end of 2016.
- Nominated for the Best Process Technology and received Grand Winner of
 Best Bioprocess Excellence in Taiwan by Biologics Manufacturing Asia
 (BMA). Received the excellence award for Antibody Drug Conjugate
 platform. Earned international recognition in bioprocess technology.
 - EG12014 met primary endpoint, bioequivalence, after the completion of Phase I clinical trial in Europe.
 - Granted Accreditation Certificate of Foreign Drug Manufacturer by Japan MHLW, with the accreditation category of "biological products" and effective date from October 31st, 2017 to October 30th, 2022. During the effective period the biological products manufactured by EirGenix's



- designated facility is allowed can be launched in Japan.
- Received "2017 Biomarker Industry Potential Benchmark Award" by Taiwan Bio Industry Organization.
- Received the "Asia's Best CMO (Contract Manufacturing Organization)
 Award" in Asia-Pacific Bioprocessing Excellence Awards 2018.
 - Ranked 145th in Deloitte Technology Fast 500 Asia Pacific.
 - "Trastuzumab biosimilar EG12014" won the 17th Taiwan FDA "Pharmaceutical Technology & Research Development Bronze Award."
 - Completed twice capital injection, with the capital reaching NT\$ 1.490229 billion.
 - First patient enrolled in Phase III clinical trial of the proprietary EG12014.
 - Received the Opinion on Successful and Marketable Development of Product or Technology in Scientific and Technological Industry issued by the Industrial Development Bureau (IDB), Ministry of Economic Affairs.
 - EG12014 won the 15th National Innovation Award-Enterprise Innovation Award.
- EirGenix, Inc. held the opening ceremony to commemorate the launch of the new "Protein Drug Commercial Production Plant" in Hsinchu (Zhubei)
 Biomedical Park.
 - Won the Grand Winner of Best Bioprocess Excellence in Taiwan Award in Singapore for the 3rd consecutive year.
 - Granted approval by 11 regulatory agencies including the United States, Taiwan, Georgia, Russia, Belarus, South Korea, India, Ukraine, Chile, South Africa, and Colombia to initiate EG12014 Phase 3 clinical trial since 2018.
 - Won the 6th National Industrial Innovation Award-Excellent Innovation Enterprise of the Ministry of Economic Affairs.
 - In April 2019, EirGenix Inc. signed a global licensing agreement with Sandoz AG, a global leader in generics and biosimilars. The licensing agreement authorized Sandoz AG to the exclusive commercial rights of EirGenix's EG12014 (Trastuzumab biosimilar) in all global markets except mainland China and Taiwan. The licensing agreement includes a signing fee and milestone payments, and additional royalty payment in the authorized markets after product launch.
 - Completed the Initial Public Offering listing.
 - Established EirGenix Europe GmbH subsidiary in Germany.
 - Won the New Technology Award of "2019 Taipei Biotech Awards".



- Won the subsequent award of National Innovation Award, Enterprise Innovation Award Continuation Award-Innovation Excellence Award.
- Completed capital injection, with the capital reaching NT\$ 1.691204 billion
- The independently administered Pharmaceuticals and Medical Devices Agency (PMDA), under Japan's Ministry of Health, Labour, and Welfare, carried out an on-site inspection of EirGenix's biopharmaceutical manufacturing facility from September 9th to September 12th, 2019. On February 3rd, 2020, EirGenix received PMDA's official approval in its issued GMP Compliance Inspection Result Notification, proclaiming EirGenix's compliance with relevant Japanese regulations regarding the quality, effectiveness, and safety of pharmaceutical manufacturing, which represented a remarkable milestone for EirGenix as the first GMP biopharmaceutical manufacturing facility in Taiwan to receive the authority's approval; not only the only one in both sides of the Taiwan Straits but also one of the few biopharmaceutical manufacturers in Asia receiving Japan's PMDA approval.
 - 807 patients enrolled in EirGenix's Phase III clinical trial of EG12014.
 - Received 2020 Bioprocessing Excellence Award in Greater China Region.
 - Completed capital injection, with the capital reaching NT\$ 2.048565 billion.
 - Completed neoadjuvant treatment and surgery of the last patient for the Phase III clinical trial of the breast cancer biosimilar EG12014.
 - Won the 17th National Innovation Award.
- The Phase III clinical trial of EG12014 showed equivalent efficacy in regards to its clinical response.
 - Completed capital injection, with the capital reaching NT\$ 2.430389 billion.
 - EirGenix's Xizhi site has been certified by Taiwan FDA as the GMP production facility for commercial biopharmaceutical drug substances. Zhubei site has been inspected and approved by Taiwan FDA as the GMP pilot production facility for biopharmaceutical drug substances.
 - Won the Globalizing Award of "2021 Taipei Biotech Awards".
 - Completed capital injection (Private placement), with the capital reaching NT\$ 3.002317 billion.
- EMA has officially accepted the review of the MAA submitted by Sandoz AG (exclusive partner of EirGenix) for trastuzumab biosimilar EG12014.
 - EirGenix, Inc., has officially submitted the biosimilar drug EIRGASUN of 150 mg powder for concentrate for solution for infusion presentation NDA



to the TFDA.

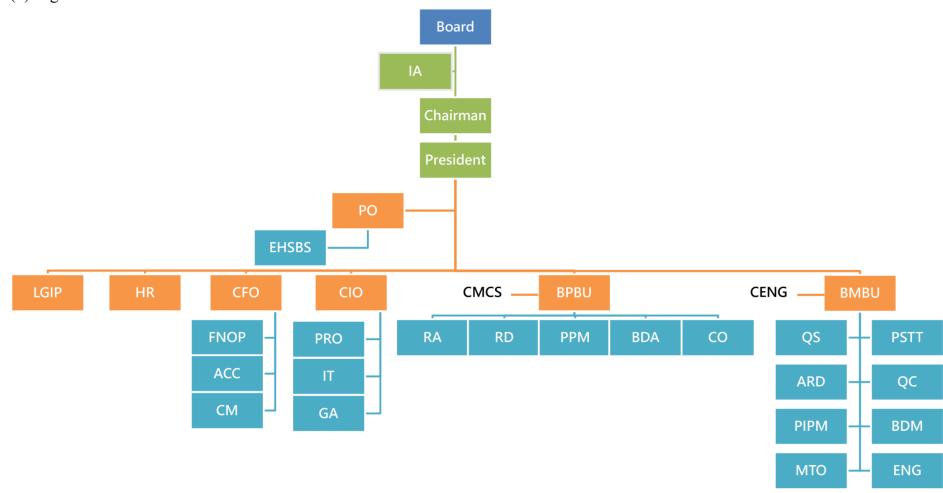
- EirGenix has officially submitted for the Phase I PK biosimilarity clinical study of developmental product EG1206A (proposed Pertuzumab biosimilar) in Europe.
- Granted Accreditation Certificate of Foreign Drug Manufacturer by Japan MHLW, with the accreditation category of "biological products" and effective date from October 24, 2022, to October 30, 2027.
- Best Bioprocessing Awards in Taiwan and the Greater China issued by Biologics Manufacturing Asia (BMA).
- National Innovation and Advancement Award (EG12014 and CRM197).
- Top 5% among TPEx-listed companies in the 8th Corporate Governance Evaluation.
- EirGenix's Zhubei plant has passed the review by the FDA and obtained an EIR before the drug launch.
 - The DMF registration for the API of EG74032 carrier protein was obtained from the FDA.
 - Top 5% among TPEx-listed companies in the 9th Corporate Governance Evaluation.
 - Received the approval letter from TFDA that the API Trastuzumab has obtained the license and the DMF number.
 - The Phase I clinical trial of the biosimilar EG1206A has met its primary endpoint and showed Pharmacokinetic biosimilarity.
 - Received the approval letter from Ministry of Health and Welfare for the biosimilar drug EIRGASUN 150 mg powder for concentrate for infusion.
 - Won the highest honor of "New Taipei City Family Friendly Work Equality Measures" in 2023.
 - The biosimilar drug, "EIRGASUN vial 150 mg", has been approved by National Health Insurance Administration to be enrolled in the reimbursement system.
 - Received a positive CHMP opinion for the biosimilar drug EG12014 licensed.
 - Received the approval letter from EC for EG12014 licensed.
 - EIRGASUN vial 150 mg won the Golden award of National Pharmaceutical Technology Research Development Award in 2023.



III. Corporate Governance Report

1. Organization

(1) Organization Chart





(2) Major Corporate Functions

Department	Functions
President	Formulate the corporate business philosophy, policies, strategies, and major investment plans.
President Office	Internal and external administrative communication and necessary contact.
Internal Auditing	Inspection and review of the Company's internal control system with adequacy in design and effectiveness in operation. Audit the integrity of financial information and establish internal risk assessment and management mechanisms.
Environmental safety and Health & Biosafety	Responsible for labor safety and health, industrial waste disposal, fire precautions, and factory area management.
Legal & Intellectual Property	Responsible for managing intellectual property rights, treatment of legal affairs, and compliance with domestic and foreign laws and regulations.
Human Resources	Responsible for providing a suitable working environment for colleagues through various activities of recruitment, hiring, training, and retaining via HR.
Finance Operations	Responsible for annual budget preparation, long-term and short-term financial forecast, financial analysis, fundraising, investing, M&A, stock affairs, public announcement, investor relations, public relations, and convening the Board of Directors/Shareholders' Meeting.
Cash Management	Responsible for banking matters, cash operation, working capital management, capital dispatch, and bank financing.
Accounting	Responsible for accounting affairs related businesses, financial statement preparation, tax planning, supplementary explanation for competent authorities, the administrative remedy of tax, and other related businesses.
Procurement	Outsource raw materials, equipment, and project and purchase the general materials/packing materials. Develop suppliers and collect goods data. Draw up, coordinate, formulate and manage the domestic and foreign sales contracts. Process the import and export operations. Analyze and plan the strategic purchasing.
Information Technology	Establish and maintain the office information infrastructure. Plan and maintain the information hardware and troubleshooting. Plan and manage the application software and troubleshooting.
General Affairs	Be responsible for internal and external administrative communication, necessary contact, and treatment of general affairs in the office.



Department	Functions									
CMC Strategy	Provide regulatory CMC strategy, planning and lead the cross-functional sub-teams for the assigned projects/products. Ensures effective communication of CMC regulatory strategy, risks, and overall plans to leadership and teams. Leads cross-functional teams responsible for the preparation of CMC regulatory submissions and responses for assigned projects/products. Coach and review persuasive briefing documents, submission sections, and responses.									
Regulatory Affairs	Provide product lifecycle strategy and perform activities related to regulatory maintenance. Ensure alignment and compliance with local and regional registration requirements. Ensure the timely submissions including CONS, IND, BLA, MAA, etc. Provide regular updates regarding the FDA/EMA/TFDA guidance and regulation. Provide the regulatory filing strategy for post-approval CMC changes. Provide regulatory support for other departments.									
Product Portfolio Management	Discuss and formulate a self-owned product development strategy and plan a product development schedule and budget with relevant departments. Supervise the overall progress of project development and coordinate cross-departmental technical discussion and work communication. Manage and control project risk and coordinate various departments to prepare relevant contingency measures. Manage the stakeholders of product projects and ensure good communication with internal teams, strategic partners, external consultants, and outsourcing manufacturers. Assist in business development related to self-owned products. Assist in administrative affairs related to self-owned products.									
Research & Development	According to the evaluation of the key quality attributes for internal products, establish the optimal cell line and develop the upstream cell culture and downstream purification process. Compile the upstream and downstream process parameters as well as quality analysis results of internal products and transfer the technology to the corresponding unit for GMP production. Write and integrate research reports and documents generated from internal product development as the basis for applying the drug certificate.									
Business Development & Alliance	Responsible for ex-Taiwan out-licensing of proprietary biosimilars and biologics Responsible for the business discussions with potential co-development and alliance partners									



Department	Functions
	Responsible for the coordination and evaluation of potential merger & acquisition and corporate investment projects Responsible for collecting market intelligence and market evaluation. To attend related conferences/events and have meeting with potential partners
Commercial Operation	Plan and implement drug marketing in Taiwan. Be responsible for the selection and follow-up implementation of new products in Taiwan and assistance in global connection and coordination.
CENG	Strategy planning for production capacity Guiding and directing the construction project
Quality System	Review and verify the effective plan, product, process, equipment changes, or other changes to determine whether effective changes need to be re-implemented. Establish an appropriate quality management system and internal GMP audit and training plan. Labor safety and health, industrial waste disposal, fire control measures, and factory management.
	Upstream and downstream process development, optimization and scale-up production testing of monoclonal antibodies and recombinant protein drugs
Process Sciences	Development of final protein drug fill finish and drug product (DP)
& Technology Transfer	technology Development and establishment of microbial expression platform and development of production process Transfer of upstream and downstream process technology of protein drugs to assist the smooth progress of cGMP manufacturing
Analytical	Analytical support in process development, stability study, protein
Research and	structure characterization, method development of physicochemical
Development	analysis, biochemical and cell-based assays during product lifecycle
Quality Control	Conduct method validation, perform GMP sample testing, including raw materials, in-process samples, DS, and DP. Conduct equipment qualification, and environmental monitoring.
Principal Investigator & Project Management	Be responsible for internal and external coordination, communication, and management of the implementation contents and administrative affairs of each stage of the project. Establish a project management process and supervision and management mechanism. Be responsible for contract fulfillment and assist the finance department in confirming the revenue based on the percentage of completion method.



Department	Functions
Business Development & Marketing	Be responsible for the expansion of the corporate business and the establishment and development of relationships with new and existing clients. Be responsible for writing the quotation. Be responsible for external and internal technical discussions and client demand confirmation before signing the contract. Plan and implement domestic and foreign publicity and exhibition work, and regularly update the Company's website. Be responsible for receiving and visiting domestic and foreign clients and related businesses. Be responsible for regular discussions and business support with overseas business colleagues.
Manufacturing & Technical Operation	GMP production. Manage the in-and-out storage and ship-out of raw materials, cell bank, and products. Scale up the process and transfer the technology.
Process& Facility Engineering	Be responsible for GMP plant system monitoring, quality maintenance, cleaning, and equipment maintenance. Plant construction project and equipment planning of production line.

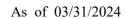


- 2. Information on the company's directors, supervisors, general manager, assistant general managers, deputy assistant general managers, and the supervisors of all the company's divisions and branch units
 - (1) Directors

A. Information of Directors

As of 03/31/2024

																A	s or u	3/31/	202 4
Title	Name	Age/Gender	Nationality/ Place of Incorporation	Date First Elected	Date Elected	Term (Year)	Shareholdir when Electe	ed	Curren	ling	Shareho	olding	Sharehold Nomin		Experience (Education)	Other Position	or Sup Spous Deg	Executives, D or Supervisors Spouses or wit Degrees of K Title Name	
Chairman	Lee-Cheng Liu	M 71~75	R.O.C	2012/12/20	2022/6/10	3	Shares 2,286,884	0.75	Shares 2,440,984	0.80	Shares 203,608	0.07	581,600 (Note)		 Columbia University Ph D, Chemical Engineering & Applied Chemistry President and COO of AnGes Inc. Process Development Department Manager, Novartis Inc. 	 President & CEO, EirGenix, Inc. Executive Director, Taiwan Bio Industry Organization Industry Consultant, Forward BioT Venture Capital. 	None		e None
Director	National Development Fund, Executive Yuan	-	R.O.C	2013/6/14	2022/6/10	3	15,288,860	5.03	15,288,860	4.99	0	0	0	0		 Director, Genovate Biotechnology Co., Ltd. Director, Taiwan Biotech Co., Ltd. Director, ScinoPharm Taiwan., Ltd. Director, Taiwan Flower Biotechnology Co., Ltd. Director, United Biomedical, Inc., Asia. Director, Adimmune Corporation. Director, TaiGen Biotechnology Holdings, Ltd. Director, PharmaEssentia Corporation. Director, PharmaEngine, Inc. Director, TaiAn Technologies Corp. Director, Intech Biopharm Corporation. Director, Point Robotics Holding Limited 	None	Non	e None





																A	s of 03		
Title	Name	Age/Gender	Nationality/ Place of	Date First Elected	Date Elected	Term (Year)	Shareholdin when Electe		Current Sharehold		Spouse & Shareho		Shareholdi Nomin		Experience (Education)	Other Position	or Supe Spouse	ervisor es or w	Directors s Who are ithin Two
			Incorporation				Shares	%	Shares	%	Shares	%	Shares	%	_				Kinship Relation
							Sildies	70	Shares	70	Shares	70	Shares	70	-	 Director, Locus Cell Corporation. Director, MetaTech (AP) Inc. Director, Wellell Inc. Director, TaiMed Biologics, Inc. Director, Taiwan Bio- Manufacturing Corporation 	Title	Name	Relation
	Representative: Hsiu-Hui Chen	F 51~60	R.O.C	2016/9/13	2022/6/10	3	0	0	0	0	0	0	0	0	 Ph.D., National Taiwan University Department of Agricultural Chemistry Researcher, Yi-cheng Biotech Inc. Postdoctoral Fellow, Institute of Plant and Microbial Biology, Academia Sinica 	 Vice President, Development Center for Biotechnology Director, Genovate Biotechnology Co., 	None	None	None
	Formosa Laboratories, Inc.	-	R.O.C	2013/6/14	2022/6/10	3	18,845,818	6.21	18,552,818	6.06	0	0	0	0	-	 Director, Formosa Pharmaceuticals, Inc. Director, A.R.Z Taiwan Ltd. Director, Epione Investment Cayman Ltd. Director, Epione Pharmaceuticals, Inc. 	None	None	None
Director	Representative: Cheng-Yu Cheng	M 71~75	R.O.C	2013/6/14	2022/6/10	3	0	0	0	0	0	0	0	0	 Ph.D., University of California, San Francisco Postdoctoral Fellow, Massachusetts Institute of Technology Research, DuPont de Nemours, Inc. Professor, National Taiwan University Department of Pharmacy Chairman, L. C. United Chemical Co., Ltd. 			None	None

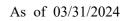


As of 03/31/2024

Title	Name	Age/Gender	Nationality/ Place of Incorporation	Date First Elected	Date Elected	Term (Year)	Shareholdir when Electe		Current Sharehold Shares		Spouse & Sharehol		Shareholdii Nomine Shares		Experience (Education)	Other Position	or Sup Spous Deg	Executives, Director Supervisors What Spouses or within Degrees of Kins Title Name Rel	
	Yao-Hwa Glass Co., Ltd, Management Commission	-	R.O.C	2019/6/12	2022/6/10	3			13,078,082		0	0	0	0	- Director, TaiGen Biotechnology Holdings, Ltd.	 Director, Adimmune Corporation Director, Locus Cell Corporation Director, Taiwan Bio- Manufacturing Corporation Director, ZHI KANG Venture Capital Investment Company, Ltd. 			None
Director	Representative: Ku-Sung Weng	M 51~60	R.O.C	2022/6/10	2022/6/10	3	0	0	0	0	0	0	0	0	 M.S., Chemical Engineering, National Tsing Hua University Director, Printing Technology Research Institute Director, SAR Technology Inc. 	 Deputy Director, Consumer Goods and Chemical Industries Division, Industrial Development Administration, Ministry of Economic Affairs - Director, Stone & Resource Industry R&D Center 	None	None	None
Director	Foxconn Technology Co., Ltd.	-	R.O.C	2022/6/10	2022/6/10	3	27,500,000	9.05	27,500,000	8.98	0	0	0	0	-	- Director and Supervisor of Hua- Zhun Investment Co., Ltd.	None	None	None

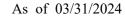


	Representative: Chun- Fu Lu	M 51~60	R.O.C	2023/1/10 2023/1/10	(Note)		0	0	0	0	0	0	 Master of EMBA program, Chinese University of Hong Kong Master of EMBA program, National Sun Yat-sen University CFO and spokesperson of Foxsemicon Integrated Technology Inc. 	- Chairman, Foxconn Technology Co. Ltd Director, Hon Fujin Precision Industry (Taiyuan) Co., Ltd Director, Hon Fujin Precision Industry (Jincheng) Co., Ltd Director, Q-Run Holdings Limited - Director, Sotera Wireless, Inc Director, FTC Japan Co., Ltd Director, Zap Medical System Ltd Chairman, Precision Healthcare Co., Ltd Chairman, Ultimate Aluminum Magnesium Technology Co., Ltd Legal representative and President, FTC Technology Vietnam Company Limited - Director, Foxconn Technology Pte. Ltd Director, Atkinson Holdings Limited - Director, Eastern Star Limited - Director, Foxconn Precision Components Holding Company Limited - Director, Gold Glory International Limited - Director, High Tempo International Limited - Director, Precious Star International Limited - Director, Precious Star International Limited - Director, Precious Star International Limited - Director, Q-Run Far East Corporation - Director, Topfry Industrial Limited - Director, Topfry Industrial Limited - Director, World Trade Trading Limited - Director, World Trade Trading Limited - Director, World Trade Trading Limited - Director, Refront IoMT Corp.	ne None N	Ione
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Title	Name	Age/Gender	Nationality/ Place of Incorporation	Date First Elected	Date Elected	Term (Year)	Shareholdin when Elect	ed	Current Sharehold	ing	Shareho	lding	Shareholdir Nomine	ee	Experience (Education)	Other Position	or Sup Spous Deg	ervisors es or wi rees of	Directors Who are thin Two Kinship
	Representative: Yu-Ting Chen	F 31~40	R.O.C	2022/9/7	2022/9/7	(Note)	Shares 0	0	Shares 0	0	Shares 0	0	Shares 0	0	 MBA in Finance, National Taiwan University Special Assistant to CIO, Hon Hai Precision Industry Co., Ltd. 	 Senior Investment Manager, GTM Management Co., Ltd. Director, Retain Biotech Corp. Director, YongLin Healthcare Foundation Director, YL Capital Ltd. 	None		None
Independent Director	Ming-Thaur Chang	M 71~75	R.O.C	2016/9/13	2022/6/10	3	0	0	0	0	0	0	0	0	 Rutgers University, NJ, USA MBA GM, CTBC Bank Tokyo Branch Chief Rep. (Taipei) United Commercial Bank EVP, Cosmos/KGI Commercial Bank Independent Director, Kaison Green Energy Technology Co., Ltd. 	- Independent Director, DBS Bank (Taiwan) Ltd.	None	None	None
Independent Director	Po-Chih Chen	M 71~75	R.O.C	2022/6/10	2022/6/10	3	0	0	0	0	0	0	0	0	 Ph.D. in Economics, National Taiwan University Chairman, Taiwan Thinktank National Policy Advisor to the President Economic Advisor to the President Director, Central Bank of the Republic of China 	 Senior Advisors to the President Honorary Chairman, Taiwan Thinktank Emeritus Professor, National Taiwan University 		None	None





Title	Name	Age/Gender	Nationality/ Place of Incorporation	Date First Elected	Date Elected	Term (Year)	Shareholdir when Electe	ed	Current Sharehold	ing	Sharehol	lding	Shareholdi Nomin	ee	Experience (Education)		or Supe Spouses Degre	rvisors s or with ees of K	
Independent Director	Fu-Shiow Yin	F 71~75	R.O.C	2016/9/13	2022/6/10	3	Shares 0	0	Shares 0	0	Shares 0	0	Shares 0	0	 Ph.D., Rutgers, the State University of New Jersey, USA Independent Director, PharmaEngine, Inc. Director, TaiGen Biotechnology Holdings, Ltd. Director, Reber Genetics Co., Ltd. Independent Director, Pac-Link Bio Ventures Member of Independent Investment Committee, Boston Life Science Venture Co., IBT Management Corp. Consultant, Department of Economic Development, Taipei City Government Science Advisor, Department of Industrial Technology, Ministry of Economic Affairs 	Co., Ltd.			None
Independent Director	Ming-Shen Chen	M 61~70	R.O.C	2016/9/13	2022/6/10	3	0	0	0	0	0	0	0	0	- Ph.D., Michigan State University, Finance.	 Professor of Finance at National Taiwan University. Director, ROC Foundation for Autistic Children And Adults in Taiwan 		None	None

Note:

- a. The stock shares delivered to the trust account, and the vested conditions in the restricted stock issuance method will be reversed according to the vested ratio.
- b. Director Yu-Ting Chen and Director Chun- Fu Lu, the representatives of Foxconn Technology Co., Ltd., was on board on 2022/09/07 and 2023/01/10.
- c. Where the chairman of the Board of Directors and the general manager or person of an equivalent position (the highest level manager) of the company are the same person, spouses, or relatives within the first degree of kinship, an explanation, reasonableness, necessity thereof, and the measures adopted in response thereto (for example, increase the number of independent directors, and there should be more than half of the directors not serving as employees or managers, etc.) shall be given.
 - The Shareholders Meeting elected the 5th term of the Board on 2022/6/10. The Board nominated Director Lee-Cheng Liu as the Chairman of the Board at the unanimous consent of the Directors on the same day.
 - It is necessary to establish four seats for Independent Directors for the Chairman acting also in the capacity as the President. The Company has complied with applicable law in this regard. The number of Independent Directors and representatives of shareholders from the public sector occupied more than half of the seats of the 5th Board that the monitoring capacity is sound.
 - On the basis of the current stage of the operation and management of the Company, and in consideration of the development in the future, the 5th Board nominated Director and President Lee-Cheng Liu as the Chairman of the Company for the full-range operation of the Company.
 - For the proper pursuit of corporate governance and strengthening the independence of the Board, the 5th Board also requested the Company to select the right candidate to act as the President of the Company at the right time.
- d. If a director's experience is related to their current position, such as having worked at the accounting firm or its affiliate during the disclosure period, the title and position shall be specified: None.



B. Major shareholders of the institutional shareholder

Name of Institutional Shareholders	Major Shareholders	%						
National Development Fund, Executive Yuan	Government Agencies							
	Cheng-Yu Cheng	6.44						
	Japan Securities Finance Co., Ltd.	3.02						
	Li Hsiu-Hui	2.55						
	Moraga Inc.	2.22						
	J.P. MORGAN SECURITIES PLC	2.10						
	Cathay Life Insurance Company, Ltd.	2.07						
Formosa Laboratories, Inc.	Ding Li Development Limited	2.02						
	MITSUBISHI UFJ MORGAN							
	STANLEY SECURITIES CO.,	2.00						
	LTD EQUITY TRADING							
	Standard Chartered Bank -	1.05						
	administrator JPMorgan	1.95						
	UBS Europe SE	1.93						
V II C1 C I.1	The Yao-Hwa Co., Ltd. Management							
Yao-Hwa Glass Co., Ltd,	Commission is a management commis	ssion						
Management Commission	managed by the Ministry of Economic Affairs.							
	Hon Hai Precision Industry Co.,	9.88						
	Ltd.	9.00						
	BaoXin International Investment	8.92						
	Co., Ltd.	0.92						
	Hyield Venture Capital Co., Ltd.	6.01						
	XinSheng Investment Co., Ltd.	4.70						
	HongYuan International Investment	2.41						
Foxconn Technology Co., Ltd.	Co. Ltd.	2.41						
	HongQi International Investment	2.25						
	Co., Ltd.	4.43						
	Standard Chartered Bank as	2.11						
	custodian of LGT	2.11						
	Citibank Custody Investment							
	Account of the Norwegian Central	1.11						
	Bank							



Name of Institutional Shareholders	Major Shareholders	%
	Vanguard Emerging Markets Stock	
	Index Fund, A Series of Vanguard	1.09
	International Equity Index Funds	
	JPMorgan in custody for Vanguard	
	Total International Stock Index	1.04
	Fund, a series of Vanguard Star	1.04
	Funds	

C. Major shareholders of the Company's major institutional shareholders

March 31, 2024

Name of Institutional	Major Shareholders	%
Mayaaa Iyaa	Xiu-Hui Li	64.28
Moraga Inc.	Wen-Jing Lin	7.14
Cathay Life Insurance	Cathay Financial Holding Co., Ltd.	100
Ding Li Development Ltd	Ding-Wu Hu	100
	Gou, Tai-Ming	12.56
	Citibank Hosting Government of Singapore Investment Account	2.78
	New Labor Pension Fund	1.64
	LGT Bank AG	1.20
	JPMorgan Chase Hosting Vanguard Developing Markets Index Fund	1.19
Hon Hai Precision Industry	JPMorgan Chase Bank Hosting Vanguard STAR Developed Markets Index Fund	1.14
Co., Ltd.	Citibank Hosting Norges Bank Investment Account	1.06
	Citibank Hosting Hon Hai Precision Industry Co., Ltd. Depositary Receipts Account	1.06
	Yuanta Taiwan Excellence 50 in custody with CTBC Bank	0.89
	Standard Chartered Bank's I Shares Emerging Markets ETF Investment Account	0.82
BaoXin International Investment Co., Ltd.	Hon Hai Precision Industry Co., Ltd.	100



Name of Institutional	Major Shareholders	%
Hariald Wanter Carried Ca	Hon Hai Precision Industry Co., Ltd.	97.95
Hyield Venture Capital Co.,	BaoXin International Investment Co.,	2.05
Ltd.	Ltd.	2.05
XinSheng Investment Co., Ltd.	Hopetown Properties Ltd.	100
HongYuan International	Han Hai Duaniaian Induator Ca I td	100
Investment Co. Ltd.	Hon Hai Precision Industry Co., Ltd.	100
HongQi International	Henry Hei Done is in the desertion Co. Head	100
Investment Co., Ltd.	Hon Hai Precision Industry Co., Ltd.	100



D.Disclosure of information as professional qualifications and independent status of directors and independent directors

	Qualification	Professional qualifications and experience	Independent status	Number of Other Public Companies in Whic
Name		•	1	Concurrently Servin as an Independent Director
Chairman	Lee-Cheng Liu	 Columbia University Ph D, Chemical Engineering & Applied Chemistry President and COO of AnGes Inc. Process Development Department Manager, Novartis Inc. President and CEO of EirGenix, Inc. Executive Director, Taiwan Bio Industry Organization None of the circumstances in the subparagraphs of Article 30 of the Company Act. 	N/A	0
Director	National Development Fund, Executive Yuan Representative: Hsiu-Hui Chen	 Ph.D., National Taiwan University Department of Agricultural Chemistry Postdoctoral Fellow, Institute of Plant and Microbial 	N/A	0
Director	Formosa Laboratories, Inc.	 Ph.D., University of California, San Francisco. Research, DuPont de Nemours, Inc. Professor, National Taiwan University Department of Pharmacy. Chairman, L. C. United Chemical Corporation. Chairman & President, Formosa Laboratories, Inc. None of the circumstances in the subparagraphs of Article 30 of the Company Act. 	N/A	0
Director	Yao-Hwa Glass Co., Ltd, Management Commission Representative: Ku-Sung Weng	 M.S., Chemical Engineering, National Tsing Hua University Director, Printing Technology Research Institute Director, SAR Technology Inc. Deputy Director, Consumer Goods and Chemical Industries Division, Industrial Development Administration, Ministry of Economic Affairs. Director, Stone & Resource Industry R&D Center None of the circumstances in the subparagraphs of Article 30 of the Company Act. 	N/A	0
Director		 Master of EMBA program, Chinese University of Hong Kong CFO and spokesperson, Foxsemicon Integrated Technology Inc. Chairman, Foxconn Technology Co. Ltd. None of the circumstances in the subparagraphs of Article 30 of the Company Act. 	N/A	0
Director	Foxconn Technology Co., Ltd. Representative: Yu-Ting Chen	 MBA in Finance, National Taiwan University Special Assistant to CIO, Hon Hai Precision Industry Co., Ltd. Senior Investment Manager, GTM Management Co., Ltd. None of the circumstances in the subparagraphs of Article 30 of the Company Act. 	N/A	0
Independent Director	Ming-Thaur Chang	 Rutgers University, NJ, USA MBA GM, CTBC Bank Tokyo Branch (2001-2003) Chief Rep. (Taipei) United Commercial Bank (2004-2010) EVP, Cosmos/KGI Commercial Bank (2010-2014) Independent Director, DBS Bank (Taiwan) Ltd. (2020-Now) 	Independent Directors are qualified for independence and competency.	1



Name	Qualification	Professional qualifications and experience	Independent status	Number of Other Public Companies in Which the Individual is Concurrently Serving as an Independent Director
		Member of Audit Committee. Have work experience in the area of commerce, law, finance, or accounting, or otherwise necessary for the business of the company. None of the circumstances in the subparagraphs of Article 30 of the Company Act.		
Independent Director	Po-Chih Chen	 Ph.D. in Economics, National Taiwan University National Policy Advisor to the President (2003-2006) Economic Advisors to the President (2003-2006) Senior Advisors to the President (2016-Now) Honorary Chairman, Taiwan Thinktank (2012-Now) Emeritus Professor, National Taiwan University Member of Audit Committee. Have work experience in the area of commerce, law, finance, or accounting, or otherwise necessary for the business of the company. None of the circumstances in the subparagraphs of Article 30 of the Company Act. 		0
Independent Director	Fu-Shiow Yin	 Ph.D., Rutgers, the State University of New Jersey, USA Master of Agriculture Chemistry, National Taiwan University Independent Director, PharmaEngine, Inc. (2011-2019) Independent Director, Pac-Link BioVentures Member of Independent Investment Committee, Boston Life Science Venture Co., IBT Management Corp. (2009-2014) Consultant, Department of Economic Development, Taipei City Government. (2011-2014) Independent Director, Foresee Pharmaceuticals Co., Ltd. (2016-Now) Member of Audit Committee. Have work experience in the area of commerce, law, finance, or accounting, or otherwise necessary for the business of the company. None of the circumstances in the subparagraphs of Article 30 of 		1
Independent Director	Ming-Shen Chen	the Company Act. - Ph.D., Michigan State University, Finance. - Professor of Finance at National Taiwan University. (2005-Now) Member of Audit Committee. Have work experience in the area of commerce, law, finance, or accounting, or otherwise necessary for the business of the company. None of the circumstances in the subparagraphs of Article 30 of the Company Act.		0

During the two years before being elected or during the term of office, an independent director of a public company may not have been or be any of the following:

- An employee of the company or any of its affiliates.
- A director or supervisor of the company or any of its affiliates.
- A natural-person shareholder who holds shares, together with those held by the person's spouse, minor children, or held by the person under others' names, in an aggregate of one percent or more of the total number of issued shares of the company or ranking in the top 10 in holdings.



- A director, supervisor, or employee of a corporate shareholder that directly holds five percent or more of the total number of issued shares of the company, or that ranks among the top five in shareholdings, or that designates its representative to serve as a director or supervisor of the company under Article 27, paragraph 1 or 2 of the Company Act.
- A director, supervisor, officer, or shareholder holding five percent or more of the shares, of a specified company or institution that has a financial or business relationship with the company.
- The amounts of the pay received by the independent director for any services such as business, legal, financial, or accounting services provided to the Company or any affiliate thereof within the past 2 years.

Unit: NT\$ thousands

				O III τ τ τ ψ tillo tip till tip
Independent Director	Remuneration	2022	2023	01/01/2024~03/31/2024
Mina Thoma Chana	Base Compensation	854	960	240
Ming-Thaur Chang	Allowances	50	30	10
D. Chil. Ch.	Base Compensation	536	960	240
Po-Chih Chen	Allowances	30	25	10
En Chiam Via	Base Compensation	854	960	240
Fu-Shiow Yin	Allowances	50	30	10
Ming-Shen Chen	Base Compensation	854	960	240
	Allowances	50	30	10

E. Diversity and independence of the Board of Directors

(1) Diversity of the Board of Directors:

Based on the policy of diversification and strengthening of corporate governance in order to promote the sound development of the Company's board composition and structure, the nomination of candidates for directors of the Company shall be adopted the candidate nomination system in accordance with the provisions of the Company's Articles of Incorporation. Each candidate's academic qualifications, work experience, professional background, integrity or relevant professional qualifications, and others are evaluated and considered. After the Board of Directors passed the resolution, the proposed nominees will be submitted to the Shareholders Meeting for election. With regard to the board composition, it is advisable that the number of the directors who concurrently serve as the managers of the Company should not exceed one-third of the board seats. In addition, the Company has, based on its own operations, operational patterns and developmental needs, formulated appropriate diversification policies including but not limited to the following:

The Board of Directors guides the operations strategy, supervises management, and examines the operations of corporate governance, while exercising duties and powers in accordance with laws and regulations and resolutions adopted by the shareholders' meeting and being responsible to the Company, shareholders, and employees, to further improve the Company's operating performance.

The Company has adopted a candidate nomination system for all directors as per the Articles of Incorporation and the Rules of Election of Directors to evaluate candidates based on their education and experience, and the shareholders' meeting elects and appoints candidates from the list of candidates. Relevant regulations are disclosed on the Company's website and Market Observation Post System (MOPS). The Company has formulated a board diversity policy according to the operating model and development needs, including basic qualifications, professional backgrounds, and industry experience, to ensure directors' suitability, independence, and professionalism. There should not be over one-third of directors who serve as managers concurrently on the Board, and directors should be equipped with the qualities in two aspects below:

- (A) Basic criteria and values: Gender, age, nationality, and culture.
- (B) Professional knowledge and skills: Professional backgrounds (such as law, accounting, industry, finances, marketing, or technology) as well as professional skills and industry experience needed for performing duties.

To achieve the corporate governance goals, the Board as a whole should be equipped with the capabilities below:

(A) Operational judgment.



- (B) Accounting and financial analysis skills.
- (C) Business management capability.
- (D) Risk control and crisis management capabilities.

More than half of the directors on the Company's Board should have the industry experience and overall planning, leadership, and management capabilities. There are currently 10 directors on the 5th Board of Directors. Among them, four are from the professional biotechnology background. All directors have experience in business, finances, and accounting as well as overall planning, leadership, and management capabilities, while possessing the professional knowledge and skills needed to perform their duties and actively participating in Board meetings and exchanging opinions with management to make business decisions. The following details our overall quantitative data and the composition of the Board based on the board diversity policy:

Title	Name	Gender	Age	Nationality/ Place of Incorporation	Professional biotechnology background	Experience in business, finances, and accounting	Overall planning, leadership, and management capabilities	Possession of college lecturer qualifications or professional and national technical certification
Chairman	Lee-Cheng Liu	M	> 60	R.O.C	✓	✓	✓	
Director	Hsiu-Hui Chen	F	< 60	R.O.C	✓	✓	✓	
Director	Cheng-Yu Cheng	M	> 60	R.O.C	√	✓	✓	✓
Director	Ku-Sung Weng	M	< 60	R.O.C		✓	✓	
Director	Chun-Fu Lu	M	< 60	R.O.C		✓	✓	
Director	Yu-Ting Chen	F	< 40	R.O.C		✓		
Independent Director	Ming-Thaur Chang	M	> 60	R.O.C		✓	✓	
Independent Director	Po-Chih Chen	M	> 60	R.O.C		✓	✓	✓
Independent Director	Fu-Shiow Yin	F	> 60	R.O.C	✓	✓	✓	
Independent Director	Ming-Shen Chen	M	> 60	R.O.C		✓	✓	✓

- There is only one director who also serves as an employee at the Company, accounting for 10%.
- There are four independent directors, accounting for 40% of the total, and the term of office of independent directors should not exceed nine years.
- There are three female directors, accounting for 30%, and seven male ones, accounting for 70%. To implement the gender equality, the goals is the proportion of female director over 30% in the future.
- There are six directors who are over 60 years old, accounting for 60%, three who are 51–60 years old, accounting for 30%, and one who are 31–40 years old, accounting for 10%.
- There are four directors from the professional biotechnology background, accounting for 40%.
- There are three with professional teaching qualifications and professional certifications, accounting for 30%.
- There are nine with overall planning, management, and leadership capabilities, accounting for 90%.
- All directors are Taiwanese citizens, and many directors possess knowledge of and experience in international business.

(2) Independence of the Board of Directors:

The Board of Directors of the Company consists of ten directors, of which four are independent directors, accounting for 40%. None of the circumstances prescribed in paragraph 3 and paragraph 4, Article 26-3 of the Securities Exchange Act exist among the directors and independent directors. No spousal relationship or familial relationship within the second degree of kinship exist between any directors. The Board of Directors of the Company is independent (Please refer to this Annual Report - Disclosure of information on professional qualifications of directors and independence of independent directors).



(2) Information on the company's directors, supervisors, general manager, assistant general managers, deputy assistant general managers, and the supervisors of all the company's divisions and branch units

March 31, 2023; Unit: Shares; %

Title	Name	Gender	Nationality	Date Effective (Note)	Sharehol	ding	Spouse & Sharehol		Sharehol by Nom Arranger	inee	Experience (Education)	Other Position	Spou Two	Managers who a Spouses or Within Two Degrees of Kinship	
				(11010)	Shares	%	Shares	%	Shares	%			Title	Name	Relation
- President -CEO of BioPharma Business Unit	Lee-Cheng Liu	M	R.O.C	2013/04/01	2,440,984	0.80	203,608	0.07	581,600	0.19	 Ph.D., Chemical Engineering & Applied Chemistry, Columbia University President and COO of AnGes Inc. Process Development Department Manager, Novartis Inc. 	 Executive Director, Taiwan Bio Industry Organization Industry Consultant, Forward BioT Venture Capital. 	None	None	None
-CEO of BioManufacturing Business Unit -Senior Vice President	Chih-Jung Chang	M	R.O.C	2013/04/01	1,029,384	0.34	100,000	0.03	66,564	0.02	 Ph.D., Chemistry, National Taiwan University. Ex-Director of PM for Oncology, TTY Biopharm 	- Director, TFBS Bioscience Inc.	None	None	None
-CFO -Manager of Corporate Governance -Vice President	Hsiu-Chuan Yang	F	R.O.C	2016/05/03	456,294	0.15	0	0	64,703	0.02	 Master of Accounting, University of New Haven General Manager, JIATE Excelsior Co., Ltd. V.P., Arich Enterprise Co., Ltd. 	None	None	None	None
-CENG -Executive Director	Shang-Chung Ju	M	R.O.C	2013/04/01	459,338	0.15	0	0	32,171	0.01	 Ph.D., Chemical Engineering, National Taiwan University Ex-Head of Production at DCB BPPF 	None	None	None	None
-CMC Strategy Lead -Executive Director	Ae-Ning Lin	F	R.O.C	2013/04/01	496,707	0.16	0	0	29,771	0.01	 Ph.D., Chemistry and Biochemistry, University of Maryland College Park. Ex-head of Purification and Protein Characterizations at DCB BPPF 	None	None	None	None
-Executive Director	Ching-Ying Chen	F	R.O.C	2021/06/07	49,275	0.02	0	0	42,371	0.01	 Master of Chemical Engineering, National Taiwan University of Science and Technology V.P., MYCENAX Biotech Inc. Manager, R&D Department, Taiwan Advance Bio-Pharmaceutical Inc. Manager, DCB 	None	None	None	None



Title	Name	Gender	Nationality	Date Effective (Note)	Shareholding		Spouse & Minor Shareholding		Shareholding by Nominee Arrangement		Experience (Education)	Other Position	Managers who are Spouses or Within Two Degrees of Kinship		
					Shares	%	Shares	%	Shares	%			Title	Name	Relation
-Executive Director	Ren-Yo Forng	М	R.O.C U.S.A	2021/07/05	28,124	0.01	0	0	39,371	0.01	 Ph.D., Laboratory of Microbial and Biochemical Sciences, Georgia State University Scientific Director, Amgen Inc. Head of QC Micro, Site Microbiologist, AstraZeneca biologics 	None	None	None	None
-Executive Director	Ywan-Feng Li	F	R.O.C	2022/04/18	13,036	0	0	0	36,248	0.01	 Ph.D., Biology, University of North Carolina at Chapel Hill, USA Vice President, Medical, Clinical & Regulatory Center, United Biopharma Division of pharmaceutical science, Center for drug evaluation-Taiwan 	None	None	None	None
-Senior Director	Tsan-Hui Wu	M	R.O.C	2017/05/01	226,346	0.07	0	0	16,249	0.01	- Ph.D., Biochemistry, National Taiwan University	None	None	None	None
-Senior Director	Hwei-Rung Wang	F	R.O.C U.S.A	2022/09/26	10,313	0.00	0	0	845	0.00	 Ph.D., Material Science and Engineering, Univ. of Michigan Director, Drug Delivery and Device Development, Alexion Principal Engineer and Senior Engineer, Biogen Idec Principal Engineer, Amgen 	None	None	None	None
-Director	Chung-Huan Lin	M	R.O.C	2019/01/02	80,198	0.03	0	0	23,348	0.01	- MBA, Case Western Reserve University - Sr. BD Manager, ScinoPharm Taiwan - BD Manager, TWi Pharma	None	None	None	None
-Director	Yu-Wen Liu	F	R.O.C	2019/05/20	110,103	0.04	0	0	14,598	0.00	- MBA, Business, St. U. of New York, New Paltz - Manager, China Productivity Center	None	None	None	None



Title	Name	Gender	Nationality	Date Effective (Note)	Shareholding		Spouse & Minor Shareholding		Shareholding by Nominee Arrangement		Experience (Education)	Other Position	Managers who are Spouses or Within Two Degrees of Kinship		
					Shares	%	Shares	%	Shares	%			Title	Name	Relation
-Director	Tsung-Chih Wang	M	R.O.C	2020/08/03	30,448	0.01	578	0.00	14,598	0.00	 MS, Taipei Medical University Marketing Manager, Marketing, Novartis Sandoz Pricing Strategy Manager, Pfizer 	None	None	None	None
-Director	Ming-Tao Pai	M	R.O.C	2020/12/01	53,198	0.02	0	0	20,848	0.01	- Ph.D., National Tsing Hua University	None	None	None	None
-Director	Chih-Yuan Ma	М	R.O.C	2022/12/16	24	0.00	0	0	302	0.00	 Ph.D., Institute of Basic Medical Sciences, National Cheng Kung University Associate Director, PMO Assistant Director, PM Department, WuXi Biologics Assistant Supervisor, Pipeline Management, MYCENAX Biotech Inc. 	None	None	None	None
-Director	Sz-Wei Wu	М	R.O.C	2023/07/03	4,078	0.00	0	0	408	0.00	 Ph.D., National Taiwan University Sr. Director, Adagene Inc. Director, WuXi Biologics Manager, CHO Pharma Sr. Application specialist, ThermoFisher Scientific Postdoctoral Fellow, Academia Sinica 	None	None	None	None

Note:

- a. Date effective is the date which be appointed as the position, not the actual date of on duty.
- b. The stock shares delivered to the trust account, and the vested conditions in the restricted stock issuance method will be reversed according to the vested ratio.
- c. If a manager's experience is related to their current position, such as having worked at the accounting firm or its affiliate during the disclosure period, the title and position shall be specified: None.
- (3) Where the chairperson of the board of directors and the general manager or person of an equivalent post (the highest-level manager) of a company are the same person, spouses, or relatives within the first degree of kinship, an explanation shall be given of the reason for, reasonableness, necessity thereof, and the measures adopted in response thereto:
 - The Shareholders Meeting elected the 5th term of the Board on 2022/6/10. The Board nominated Director Lee-Cheng Liu as the Chairman of the Board at the unanimous consent of the Directors on the same day.
 - It is necessary to establish four seats for Independent Directors for the Chairman acting also in the capacity as the President. The Company has complied with applicable law in this regard. The number



of Independent Directors and representatives of shareholders from the public sector occupied more than half of the seats of the 5th Board that the monitoring capacity is sound.

On the basis of the current stage of the operation and management of the Company, and in consideration of the development in the future, the 5th Board nominated Director and President Lee-Cheng Liu as the Chairman of the Company for the full-range operation of the Company.

For the proper pursuit of corporate governance and strengthening the independence of the Board, the 5th Board also requested the Company to select the right candidate to act as the President of the Company at the right time.

3. Remuneration of Directors, Supervisors, President, and Vice Presidents

(1) Remuneration of Directors

A. Remuneration of Directors (including Independent Directors) in 2023

Unit: NT\$ thousands; %

					Remune					Remu	of Total neration		Di		t Remuneration Received by ors Who are Also Employees					Con	io of Total appensation	D amount in n
		,	mpensation (A)			Compe	ectors nsation(C)		rances (D)	Inco	+D) to Net me (%)	Allowa	onuses, and inces (E)		• ` ′	Emplo	yee Con	npensati	on (G)		C+D+E+F+G) Income (%)	Remuneration from ventures other than
Title	Name	The company	All companies in the consolidate d financial statements	1 ,	All companies in the consolidate d financial statements		All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements		Stock	consol finar staten	the idated neial nents	The	Companies in the consolidated financial statements	
Chairman Lee-Cheng l	Liu	0	0	0	0	0	0	30	30	30 (0.003)	30 (0.003)	37,408 (Note)	37,408 (Note)	0	0	0	0	0	0	37,438 (4.09)	37,438 (4.09)	0
Director National Dev Fund, Execu		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Director Rep Hsiu-Hui Ch		0	0	0	0	0	0	30	30	30 (0.003)	30 (0.003)	0	0	0	0	0	0	0	0	30 (0.003)	30 (0.003)	0
Director Formosa Lal	boratories, Inc.	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Director Rep Cheng-Yu C		0	0	0	0	0	0	25	25	25 (0.003)	25 (0.003)	0	0	0	0	0	0	0	0	25 (0.003)	25 (0.003)	0
	lass Co., Ltd, t Commission	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Director Rep Ku-Sung We		0	0	0	0	0	0	25	25	25 (0.003)	25 (0.003)	0	0	0	0	0	0	0	0	25 (0.003)	25 (0.003)	0
Director Foxconn Tec Ltd.	chnology Co.,	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0



Unit: NT\$ thousands: %

																					Omi. IV	15 thousands; %		
					Remun					Remui	of Total neration		Rele Di	vant Remu rectors Wh	neration Re o are Also	ceived b Employe	ees			Ratio of Total Compensation		Daman anatian		
		Base Con	mpensation (A)	Severan	ce Pay (B)	Dir Compe	rectors nsation(C)	Allow	ances (D)		+D) to Net ne (%)		onuses, and ances (E)	Severano	ce Pay (F)	Employee Con		mpensation (G)			C+D+E+F+G) Income (%)	Remuneration from ventures other than		
Title	Name	Name		The company	All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements		All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements	The company	All companies in the consolidate d financial statements	The co	mpany	consol finar stater	the idated icial	The company	Companies in the consolidated financial statements	
Director Rep Yu-Ting Che		0	0	0	0	0	0	25	25	25 (0.003)	25 (0.003)	0	0	0	0	0	0	0	0	25 (0.003)	25 (0.003)	0		
Director Rep Chun-Fu Lu	resentative	0	0	0	0	0	0	25	25	25 (0.003)	25 (0.003)	0	0	0	0	0	0	0	0	25 (0.003)	25 (0.003)	0		
Former Direct Representative Jih-Luh Tang	ve	0	0	0	0	0	0	5	5	5 (0.0005)	5 (0.0005)	0	0	0	0	0	0	0	0	5 (0.0005)	5 (0.0005)	0		
Independent Ming-Thaur		960	960	0	0	0	0	30	30	990 (0.11)	990 (0.11)	0	0	0	0	0	0	0	0	990 (0.11)	990 (0.11)	0		
Independent Po-Chih Che	Director n	960	960	0	0	0	0	25	25	985 (0.11)	985 (0.11)	0	0	0	0	0	0	0	0	985 (0.11)	985 (0.11)	0		
Independent Fu-Shiow Yi		960	960	0	0	0	0	30	30	990 (0.11)	990 (0.11)	0	0	0	0	0	0	0	0	990 (0.11)	990 (0.11)	0		
Independent Ming-Shen C		960	960	0	0	0	0	30	30	990 (0.11)	990 (0.11)	0	0	0	0	0	0	0	0	990 (0.11)	990 (0.11)	0		

a. Salary, Bonuses, and Allowances (E) including the share-based payment.

If the Company has net profit in this fiscal year, the Company shall set aside 3% (inclusive) or less of its profits as bonus to Directors. The distribution of director remuneration shall be heard by over two-thirds of the Board of Directors, be voted in favor for implementation by over one-half of the directors present and represented, and also be reported at the shareholders' meeting.

The Company did not pay any director remuneration during the previous two years. Directors only received traffic allowances for conducting businesses. Independent directors receive fixed emoluments for performing businesses. The aforesaid traffic allowances and emoluments for conducting businesses have been reviewed by Remuneration Committee and approved by the Board of Directors.

c. In addition to the above remuneration, director remuneration shall be disclosed as follows when received from companies included in the consolidated financial statements in the most recent year to compensate directors for their services, such as being independent contractors: None.

b. Please describe the policy, system, standard, and structure of remuneration to independent directors, and the correlation between duties, risk, and time input with the amount of remuneration:



B. Range of Remuneration for Directors (including Independent Directors)

B. Range of Remuneration for Directors (including In	dependent Directors)	Name of	Directors			
	Total of (A+B+C+D)	Total of (A+B-	+C+D+E+F+G)		
Range of Remuneration	The company	Companies in the consolidated financial statements (H)	The company	Companies in the consolidated financial statements (I)		
	Chairman - Lee-Cheng Liu					
	Directors		Directors			
	Representative: Hsiu		Representative: Hsiu-			
	- Formosa Laboratorie	s, Inc.	- Formosa Laboratories			
	Representative: Cher	g-Yu Cheng	Representative: Chen	g-Yu Cheng		
	- Yao-Hwa Glass Co.,	Ltd, Management	- Yao-Hwa Glass Co.,	Ltd, Management		
	Commission		Commission			
Less than NT\$ 1,000,000	Representative: Ku-S	lung Weng	Representative: Ku-Sung Weng			
	- Foxconn Technology	Co., Ltd.	- Foxconn Technology Co., Ltd.			
	Representative: Yu-7	ing Chen	Representative: Yu-T	ing Chen		
	Representative: Chur	ı-Fu Lu	Representative: Chun	-Fu Lu		
	Former Director Rep	resentative: Jih-Luh Tang	Former Director Repo	esentative: Jih-Luh Tang		
	Independent Directors		Independent Directors			
	- Ming-Thaur Chang		- Ming-Thaur Chang			
	- Po-Chih Chen		- Po-Chih Chen			
	- Fu-Shiow Yin		- Fu-Shiow Yin			
	- Ming-Shen Chen		- Ming-Shen Chen			
NT\$1,000,000 (Included) ~ NT\$2,000,000 (Not included)	-	-	-	-		
NT\$2,000,000 (Included) ~ NT\$3,500,000 (Not included)	-	-	-	-		
NT\$3,500,000 (Included) ~ NT\$5,000,000 (Not included)	-	-	-	-		
NT\$5,000,000 (Included) ~ NT\$10,000,000 (Not included)	-	-	-	-		
NT\$10,000,000 (Included) ~ NT\$15,000,000 (Not included)	-	-	-	-		
NT\$15,000,000 (Included)~ NT\$30,000,000 (Not included)	-	-	-	-		
NT\$30,000,000 (Included) ~ NT\$50,000,000 (Not included)			Chairman			
1\(\partial \pi 50,000,000\) (included) \(\sim 1\(\pi 50,000,000\) (included)	-	-	- Lee-Cheng Liu			
NT\$50,000,000 (Included) ~ NT\$100,000,000 (Not included)	-	-	-	-		
Greater than or equal to NT\$100,000,000	-	-	-	-		
Total			1			

(2) Remuneration of Supervisors

EirGenix, Inc. has set up the Audit Committee on June 12th, 2018.

(3) Remuneration of the President and Vice President

A. Remuneration of the President and Vice Presidents in 2023

Unit: NT\$ thousands

														T \$ tilousalius
Title		Salary(A)		Severance Pay (B)		Bonuses and Allowances (C)		Employee Compensation (D)			sation	compo (A+B+C	±100 net	Remuneration from ventures
	Name	The company	All companies in the consolidated financial statements	The	All companies in the consolidated financial statements	The company	All companies in the consolidated financial statements	com	he pany Stock	consol final states	the lidated ncial ments	The	All companies in the consolidated financial statements	other than subsidiaries or from the parent company
President	Lee-Cheng Liu													
Senior Vice President	Chih-Jung Chang	17,058	17,058	216	216	32,986	32,986	0	0	0	0	50,260 (5.49)	50,260 (5.49)	0
Vice President	Hsiu-Chuan Yang													

Note:

a. Remuneration of the President and Vice Presidents in 2023 includes the expenses of share-based payment.



B. Range of Remuneration for President and Vice President

Domas of Domas mortion	Name of President	and Vice Presidents		
Range of Remuneration	The company	Companies in the consolidated		
Less than NT\$ 1,000,000	-	-		
NT\$1,000,000 (Included)~ NT\$2,000,000(Not included)				
NT\$2,000,000 (Included)~ NT\$3,500,000(Not included)	-	-		
NT\$3,500,000 (Included) ~ NT\$5,000,000(Not included)	-	-		
NT\$5,000,000 (Included) ~ NT\$10,000,000(Not included)	Chih-Jung Chang,	Hsiu-Chuan Yang		
NT\$10,000,000 (Included) ~ NT\$15,000,000(Not included)	-	-		
NT\$15,000,000 (Included) ~ NT\$30,000,000(Not included)	-	-		
NT\$30,000,000 (Included) ~ NT\$50,000,000(Not included)	Lee-Ch	eng Liu		
NT\$50,000,000 (Included) ~ NT\$100,000,000(Not included)	-	-		
Greater than or equal to NT\$100,000,000	-	-		
Total	3	3		

C. Managerial officers with the top five highest remuneration amounts

Unit: NT\$ thousands

		Sal	ary(A)	Severano	ce Pay (B)		ses and ances (C)	Emplo	yee Con	npensati	on (D)	comp (A+B+0	of total pensation C+D) to net ome (%)	on from
Title	Name	The company	All companies in the consolidate d financial		All companies in the consolidate d financial	The	ea	The co	ompany	in conso fina	npanies the lidated ncial ments	The	ea	ventures other than subsidiaries or from the parent company
			statements		statements		financial statements	Cash	Stock	Cash	Stock		financial statements	
President	Lee-Cheng Liu	9,235	9,235	-	-	28,173	28,173	0	0	0	0	37,408 (4.09)	37,408 (4.09)	0
Senior Vice President	Chih-Jung Chang	4,240	4,240	108	108	2,384	2,384	0	0	0	0	6,732 (0.74)	6,732 (0.74)	0
Vice President	Hsiu-Chuan Yang	3,583	3,583	108	108	2,429	2,429	0	0	0	0	6,120 (0.67)	6,120 (0.67)	0
Executive Director	Ren-Yo Forng	3,576	3,576	108	108	4,990	4,990	0	0	0	0	8,674 (0.95)	8,674 (0.95)	0
Executive Director	Shang-Chung Ju	3,179	3,179	108	108	1,690	1,690	0	0	0	0	4,977 (0.54)	4,977 (0.54)	0

Note: Bonuses and Allowances include the expenses of share-based payment.



- (4) Employee Profit Sharing Granted to Management Team: None.
- (5) Comparison of Remuneration for Directors, Supervisors, President and Vice Presidents in the Most Recent Two Fiscal Years and Remuneration Policy for Directors, Supervisors, President and Vice President
 - A. The ratio of total remuneration paid by the Company and by all companies included in the consolidated financial statements for the two most recent fiscal years to directors, supervisors, president, and vice presidents of the Company, to the net income.

Item	Ratio of total remuneration paid to directors, supervisors, president, and vice presidents to net income (%)									
		2022	2023							
Title	Total remuneration	Companies in the consolidated financial statements	Total remuneration	Companies in the consolidated financial statements						
Directors	3,948	(3.42)	4,120	(0.45)						
President and Vice President	51,395	(44.48)	50,260	(5.49)						

- B. The policies, standards, and portfolios for the payment of remuneration, the procedures for determining remuneration, and the correlation with risks and business performance.
 - (A) Remuneration for Directors, President and Vice Presidents

Suppose the Company has the net profit in this fiscal year. In that case, the Company shall set aside between 1% to 5% of its profits as a bonus to the Company's employees and set aside 3% (inclusive) or less of its profits as a bonus to Directors. The distribution of bonus to employees may be made by way of cash or shares by the resolution of the Board of Directors. The employees may include certain employees of the subsidiaries who meet the conditions prescribed by the Company. Over two-thirds shall hear the distribution of employee remuneration and director remuneration of the Board of Directors, be voted in favor of implementation by over one-half of the directors present and represented and be reported at the shareholders' meeting.

The Company shall first offset its losses in previous years that have not been previously offset, and then set aside annual profits as a bonus to the Company's employees and set aside annual profits as a bonus to Directors.

EirGenix did not pay any director remuneration during the previous two years. Directors only received traffic allowances for conducting business. Former Chairman and independent directors receive fixed emoluments for performing businesses. The remaining remuneration to directors is the salary of the current Chairman as an employee. The aforesaid traffic allowances and emoluments for conducting businesses have been reviewed by Remuneration Committee and approved by the Board of Directors.

The President and Vice Presidents of the Company are remunerated in commensurate with their position, contribution to the Company and with reference to industry standards subject to the review of the Remuneration Committee and reporting to the Board for final approval. No remuneration has been appropriated by EirGenix as remuneration to employees in the last 2 years.

(B) Association with operation performance and risks in the future

The remunerations to the Directors and managers will be determined on the basis of the operation of the Company, the operation risk and development in the future with reference to industry standards and the assigned duties and contribution, and the association with the operating performance at a significant level. The management and the Remuneration Committee of the Company will review the remuneration level at regular intervals for appropriate adjustment for a proper balance between risk control and sustainable development of the Company.



4. Implementation of Corporate Governance

(1) Operations of the Board of Directors:

A total of <u>6</u> (A) meetings of the Board of Directors was held in 2023. The attendance of directors was as follows:

Title	Name	Actual Attendance (B)	By Proxy	Attendance Rate (%) (B/A)	Remarks
Chairman	Lee-Cheng Liu	6	0	100	
Director	National Development Fund, Executive Yuan Representative: Hsiu-Hui Chen	6	0	100	
Director	Formosa Laboratories, Inc. Representative: Cheng-Yu Cheng	6	0	100	
Director	Yao-Hwa Glass Co., Ltd, Management Commission Representative: Ku-Sung Weng	5	1	83.33	
Director	Foxconn Technology Co., Ltd. Representative: Chun-Fu Lu	6	0	100	
Director	Foxconn Technology Co., Ltd. Representative: Yu-Ting Chen	5	1	83.33	
Independent Director	Ming-Thaur Chang	6	0	100	
Independent Director	Po-Chih Chen	5	1	83.33	
Independent Director	Fu-Shiow Yin	6	0	100	
Independent Director	Ming-Shen Chen	6	0	100	

A total of $\underline{2}$ (A) meetings of the Board of Directors was held by the end of March 2024. The attendance of directors was as follows:

Title	Name	Actual Attendance (B)	By Proxy	Attendance Rate (%) (B/A)	Remarks
Chairman	Lee-Cheng Liu	2	0	100	-
Director	National Development Fund, Executive Yuan Representative: Hsiu-Hui Chen	2	0	100	-
Director	Formosa Laboratories, Inc. Representative: Cheng-Yu Cheng	2	0	100	-
Director	Yao-Hwa Glass Co., Ltd, Management Commission Representative: Ku-Sung Weng	2	0	100	-
Director	Foxconn Technology Co., Ltd. Representative: Chun-Fu Lu	1	1	50	
Director	Foxconn Technology Co., Ltd. Representative: Yu-Ting Chen	2	0	100	-
Independent Director	Ming-Thaur Chang	2	0	100	-
Independent Director	Po-Chih Chen	2	0	100	-
Independent Director	Fu-Shiow Yin	2	0	100	-
Independent Director	Ming-Shen Chen	2	0	100	-



Other mentionable items:

A. In any of the following circumstances, the dates of the meetings, sessions, contents of motion, all independent directors' opinions and the company's response to independent directors' opinion should be specified:

(A). The circumstances referred to in Article 14-3 of the Securities and Exchange Act:

Date of Meeting/ Term of Board of Directors	Contents of Motion	Independent Director's Opinion	The Company's Response to Independent Director's Opinion
2023/03/10 The 8th meeting of the 5th board	 a. Approved the motion of the ratification of the assessment of the independence and competence of the CPAs retained as external Auditors. b. Approved the motion of the ratification of the appointment of CPAs as external auditors and the remuneration to the CPAs. c. Approved the certified public accounts previously approved as the Independent Auditors, the CPA office and affiliates to render non-assurance services to the Company and the subsidiaries. d. Adoption of the 2023 Employee Restricted Stock Awards. e. Approved the Company will raise capital through private placements of common shares. f. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of directors and their representatives. g. Adoption of the Issuance of 2022 Employee Stock Options. 	None	Not applicable
2023/05/10 The 9th meeting of the 5th board	a. Adoption of the Issuance of 2022 Employee Stock Options.		
2023/08/08 The 10th meeting of the 5th board	a. Adoption of the Issuance of 2022 Employee Stock Options.b. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of directors and their representatives.		
2023/11/09 The 11th meeting of the 5th board	 a. Approved the establishment of a subsidiary in the United States based on the operational development needs. b. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". c. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) d. Approved to update the 2023 1st and 2nd Employee Restricted Stock Awards to employees. e. Approved to grant 2022 1st Employee Restricted Stock Awards to employees. f. Approved to grant 2023 2nd Employee Restricted Stock Awards to employees. g. Approved to sign the Amended and Restated License Agreement for the biosimilar EG12014 with Sandoz AG. h. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of directors and their representatives. 		
2023/12/22	a. Approved to formulate "Rules Governing Financial and Business Matters Between this Corporation and its Affiliated Enterprises" and amend		



Date of Meeting/ Term of Board of Directors	Contents of Motion	Independent Director's Opinion	The Company's Response to Independent Director's Opinion
The 12th	"Management of Affiliated Enterprises Transaction".		
meeting of the	b. Approved the amendment of "written accounting systems".		
5th board	c. Approved the amendment of Implementation Report for the Sound		
	Business Plan and estimation of Income Statement.		
	d. Approved the subscription of the AmMax Bio, Inc. cash capital increase.		
	based on potential business collaboration opportunities and financial		
	investment considerations.		
	e. Approved to grant 2022 Employee Stock Options to employees.		
	f. Approved to grant 2023 1st Employee Restricted Stock Awards to		
	employees		
	a. Approved the CPAs replacement due to PricewaterhouseCoopers Taiwan		
	internal organization adjustment.		
	b. Approved the motion of the ratification of the assessment of the		
2024/03/08	independence and competence of the CPAs retained as external Auditors.		
The 13th	c. Approved the motion of the ratification of the appointment of CPAs as		
meeting of the	external auditors and the remuneration to the CPAs.		
5th board	d. Adoption of the 2024 1st Employee Restricted Stock Awards.		
	e. Approved the Company will raise capital through private placements of common shares.		
	f. Approved to sign the contract with clinical CRO and the relevant		
	companies for Phase III clinical trial of the EG1206A.		

(B). In addition to said circumstances, any other matter about which an independent director expresses an objection or reservation that has been included in records or stated in writing: None.

B. If there is Directors' avoidance of motions in conflict of interest, the Directors' names, contents of motions, causes for avoidance, and voting should be specified:

Date of Meeting	Name	Meeting Agenda	Causes for avoidance	Result of Voting	
2023/01/16	Lee-Cheng Liu	a. Approved the motion of distribution of year-end bonuses for the managers.		<u> </u>	
2022/02/10	Lee-Cheng Liu	 a. Adoption of the 2023 1st and 2nd Employee Restricted Stock Awards. 	Excused from		
2023/03/10	Chun Fu Lu Yu-Ting Chen	a. Release the prohibition on directors or representatives of directors from participation in competitive business.	discussion and resolution of this		
2023/08/08	Hsiu-Hui Chen	a. Release the prohibition on directors or representatives of directors from participation in competitive business.	agenda item puto paragraph 3	of	
2022/11/00	Lee-Cheng Liu	 a. Adoption of the 2023 2nd Employee Restricted Stock Awards. 	Article 206 of Company Act.		
2023/11/09	Chun Fu Lu	a. Release the prohibition on directors or representatives of directors from participation in competitive business.			



C. Conducting Evaluations of Board Performance

		Evaluation		Evaluation content
period		method		Evaluation content
January 1st— December 31st, 2023	Board of Directors, individual board member, and functional committees (including Remuneration Committee and Audit Committee)	method Internal self- evaluation of the board of directors and self- evaluation of directors	a. b. c.	Performance evaluation of the board of directors: Include the degree of participation in the company's operations, the decision-making quality of the board of directors, the composition and structure of the board of directors, the selection and continuous education of directors, internal control, etc. Performance evaluation of individual directors: Include grasping the company's objectives and tasks, recognition of directors' responsibilities, level of participation in the company's operations, internal relationship management and communication, expertise and continuous education of directors, internal control, etc. Performance evaluation of functional committees: the degree of participation in the company's operations, the recognition of functional committee's responsibilities, the decision-making quality of functional committees, the composition of functional committees and the selection of members, internal control, etc. The Board, Boardmembers, and functional
October 1st,2021— September 30th, 2022	Conducting Evaluations of Board Performance	Appointment of the external professional institutions: Taiwan Investor	a. b. c. d. e.	committees (including Remuneration Committee and Audit Committee) received a self-assessment score over 90. The organization and professional development of the Board. Quality of decision-making of the Board. The performance result of the Board. Internal control and risk management. Level of participation in corporate social responsibility by the Board.
	1st— December 31st, 2023 October 1st,2021— September	Directors, individual board member, and functional committees (including Remuneration Committee and Audit Committee) October Conducting Evaluations of September Board	Directors, individual board member, and functional committees (including Remuneration Committee and Audit Committee) October 1st,2021— Evaluations of September 30th, 2022 Performance individual board of directors and self-evaluation of directors	Directors, individual board board of directors and self-functional committees (including Remuneration Committee and Audit Committee) October September Board Board Professional Board Sult, 2022 Performance Professional Board Investor Relations Directors, evaluation of the board of directors and self-evaluation of directors and self-evaluation of directors directors and self-evaluation of directors directors c. Taiwan d. Investor e. Relations

- D. An evaluation of targets for strengthening of the functions of the board during the current and immediately preceding fiscal years (e.g., the establishment of the Audit Committee, improvement of information disclosure transparency) and measures taken toward achievement:
 - (A) The board of directors is composed of ten directors (including four independent directors) with rich academic and industry experience. The board of directors follows the "Standards of the Board of Directors' Procedures" and the "Management Measures for the Operation of the Board of Directors' Procedures", and regularly reviews and discusses various business development plans.
 - (B) The PwC accountants attended the board meeting, provided taxation and legal information and suggestions, and reported the quarterly financial report inspection results to the directors.
 - (C) EirGenix establishes Audit Committee and to strengthen the effectiveness of internal control. In addition to helping independent directors fully understand the financial report review, they can also conduct substantive supervision through the



- disclosure and exchange of financial information and build an appropriate and comprehensive risk management supervision mechanism.
- (D) EirGenix has elected 4 Independent Director and set up a Remuneration Committee, and an Audit Committee. In 2022, set up the Corporate Governance Committee.
- (E) EirGenix has set up the spokesman and vice spokesman system and disclosed the financial and business information on the Market Observation Post System and the corporate website according to laws and regulations.

(2) Operation of Audit Committee

A. Information of Audit Committee Operation:

A total of $\underline{5}$ (A) meetings of the Audit Committee were held in 2023. The attendance of independent directors was as follows:

Title	Name	Attendance in Person(B)	By Proxy	Attendance rate (%)
Independent Director	Ming-Thaur Chang	5	0	100
Independent Director	Po-Chih Chen	5	0	100
Independent Director	Fu-Shiow Yin	5	0	100
Independent Director	Ming-Shen Chen	5	0	100

A total of $\underline{1}$ (A) meetings of the Audit Committee was held by the end of March 2024. The attendance of directors was as follows:

Title	Name	Attendance in Person(B)	By Proxy	Attendance rate (%)
Independent Director	Ming-Thaur Chang	1	0	100
Independent Director	Po-Chih Chen	1	0	100
Independent Director	Fu-Shiow Yin	1	0	100
Independent Director	Ming-Shen Chen	1	0	100

Other mentionable items:

- (A) In any of the following circumstances, the dates of the meetings, sessions, contents of motion, audit committee's resolutions, and the company's response to the audit committee's opinion should be specified:
 - a. The circumstances referred to in Article 14-5 of the Securities and Exchange Act:

Date and Term of Meeting	Contents of Motion	Audit Committee's Resolutions	The Company's Response to Audit Committee's Opinion
	a. Accept 2022 Financial Statements and Business Report.	Consent	Approved as
	b. Ratification of the 2022 Deficit Offset Proposal.		proposed
	c. Approved the motion of issuance of the 2022 Declaration of Internal		
	Control System of the Company.		
	d. Approved the motion of the ratification of the assessment of the		
	independence and competence of the CPAs retained as external Auditors.		
2023/03/10	e. Approved the motion of the ratification of the appointment of CPAs as		
The 7th meeting	external auditors and the remuneration to the CPAs.		
of the 3rd term	f. Approved the CPAs previously approved as the Independent Auditors, the		
	CPA firms, and affiliates to render non-assurance services to the		
	Company and the subsidiaries.		
	g. Approved the amendment to the Article of the Company.		
	h. Approved the amendment to the Regulations Governing Procedure for		
	Board of Directors Meetings.		
	i. Adoption of the 2023 1 st and 2 nd Employee Restricted Stock Awards.		



Date and Term of Meeting	Contents of Motion	Audit Committee's Resolutions	The Company's Response to Audit Committee's Opinion
2023/05/10 The 8th meeting	 j. Approved the Company will raise capital through private placements of common shares. k. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of directors and their representatives. l. Approved to grant 2022 Employee Stock Options to employees. a. Accept 2023 Q1 Financial Statements. b. Approved to grant 2022 Employee Stock Options to employees. 		
The 8th meeting of the 3rd term			
2023/08/08 The 9th meeting of the 3rd term	 a. Accept 2023 Q2 Financial Statements. b. Approved to formulate "Operational Procedure for Preparation and Validation of the Sustainability Report". c. Approved to grant 2022 Employee Stock Options to employees. d. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of directors and their representatives. 		
2023/11/09 The 10th meeting of the 3rd term	 a. Accept 2023 Q3 Financial Statements. b. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". c. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) d. Approved to grant 2022 1st Employee Restricted Stock Awards to employees. e. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of directors and their representatives. 		
2023/12/22 The 11th meeting of the 3rd term	 a. Approved to formulate "Rules Governing Financial and Business Matters Between this Corporation and its Affiliated Enterprises" and amend "Management of Affiliated Enterprises Transaction". b. Approved the amendment of "Written Accounting Systems". c. Approved the Internal Audit Plan for the fiscal year 2024. d. Approved the operation plan for the fiscal year 2024. e. Approved the amendment of Implementation Report for the Sound Business Plan and estimation of Income Statement. f. Approved to grant 2022 Employee Stock Options to employees. g. Approved to grant 2023 1st Employee Restricted Stock Awards to employees 		
2024/03/08 The 12th meeting of the 3rd term	 a. Accept 2023 Financial Statements and Business Report. b. Ratification of the 2023 Deficit Offset Proposal. c. Approved the CPAs replacement due to PricewaterhouseCoopers Taiwan internal organization adjustment. d. Approved the motion of the ratification of the assessment of the independence and competence of the CPAs retained as external Auditors. e. Approved the motion of the ratification of the appointment of CPAs as external auditors and the remuneration to the CPAs. f. Approved the motion of issuance of the 2023 Declaration of Internal 		



			The
Date and Term of Meeting	Contents of Motion	Audit Committee's Resolutions	Company's Response to Audit Committee's Opinion
	Control System of the Company. g. Adoption of the 2024 1 st Employee Restricted Stock Awards.		
	h. Approved the Company will raise capital through private placements of common shares.		

b.Other matters which were not approved by the Audit Committee but were approved by two-thirds or more of all directors: None.

- (B) If there are independent directors' avoidance of motions in conflict of interest, the directors' names, contents of motion, causes for avoidance, and voting should be specified: None.
- (C) Communications between the independent directors, the Company's chief internal auditor, and CPAs (e.g., the material items, methods and results of audits of corporate finance or operations, etc.):

For the implementation of the supervision mechanism, the internal audit reports are submitted to the board of directors and management for review on a regular basis by the audit unit and also delivered and notified to independent directors according to regulations. The Company's execution of the internal control system is compliant with regulations and will be continually followed up. When a board of directors meeting is held, attending directors with opinions may have discussion and communication. If there's any material violation or any likelihood of material damage to the company, related personnel shall promptly prepare and present a report and notify the independent directors.

The Company invites CPA, independent directors, and internal auditors to the communication meeting at least once a year. CPA will report the Company's financial position and auditing process to independent directors and then listen to the voice of independent directors and the chief internal auditor.

The Company has invited the Independent Auditors to sit in the meetings of the Audit Committee and the Board as observers at least 4 times as year (2023/3/10, 2023/5/10, 2023/8/8,2023/11/19) for engagement in discussion and review the audit or review result of the quarterly reports and annual financial statements, significant accounting standards and interpretation, applicable legal rules governing securities, and tax law. The internal auditor communicated in the meeting with the independent directors and the CPAs on 2023/11/9, and the topic is the explanation of the risk assessment and key audit issues of the Company in terms of the AQI (Audit Quality Indicators). The conclusion suggested closer attention to key audit issues. In addition to the key point of AQI, independent director suggested the sharing the information through the CPAs to guide the company to attach importance to shareholder's right and ESG.

- (D) The powers of the Committee are as follows:
 - a. The adoption of or amendments to the internal control system pursuant to Article 14-1 of the Securities and Exchange Act.
 - b. Assessment of the effectiveness of the internal control system.
 - c. The adoption or amendment, pursuant to Article 36-1 of the Securities and Exchange Act of the procedures for handling financial or business activities of a material nature, such as acquisition or disposal of assets, derivatives trading, loaning of funds to others, and endorsements or guarantees for others.
 - d. Matters in which a director is an interested party.
 - e. Asset transactions or derivatives trading of a material nature.
 - f. Loans of funds, endorsements, or provision of guarantees of a material nature.
 - g. The offering, issuance, or private placement of equity-type securities.
 - h. The hiring or dismissal of a certified public accountant, or their compensation.
 - i. Annual and semi-annual financial reports.
 - j. Other material matters as may be required by this Corporation or by the competent authority.



B. EirGenix established the Audit Committee to replace supervisors on June 12, 2018

(3) Corporate Governance Implementation Status and Deviations from "the Corporate Governance Best-Practice Principles

for TWSE/TPEx Listed Companies".

for TW	SE/TPEx Listed Companies"	•			
				Implementation Status	Deviations from
	Evaluation Item	Y	N	Abstract Illustration	"Corporate Governance Best-Practice Principles for TWSE/TPEx Listed Companies" and Reasons
the Corpo Principles Governan	company establish and disclose brate Governance Best-Practice is based on "Corporate ace Best-Practice Principles for PEx Listed Companies"?	√		EirGenix has formulated the Corporate Governance Best Practice Principles and Corporate Social Responsibility Best Practice Principles; and EirGenix has operated in accordance with the institutional regulations of internal control and internal audit, and also formulated institutional measures such as Regulations Governing the Acquisition and Disposal of Assets, Procedures for Endorsements and Guarantees, Procedures for Loaning Funds to Others, Rules of Procedure for the Board of Directors Meetings, Management of Procedure for the Board of Directors, Procedure for Election of Directors, Rules of Procedure for Shareholders Meetings and Codes of Ethical Conduct, with the goal of implementing Corporate Governance.	Corporate Governance Best-Practice Principles for TWSE/TPEx Listed Companies".
rights (1) Does the coperating poperating poperating disputes, and are considered to the constant of the const	ompany establish an internal procedure to deal with ers' suggestions, doubts, and litigations, and implement the procedure?	√		EirGenix has established a spokesman and vice spokesman system and has dedicated personnel responsible for disclosing corporate information and handling shareholders' suggestions and doubts to ensure shareholders' rights.	Corporate Governance Best-Practice Principles
(2) Does the c	ompany possess the list of its reholders as well as the ultimate those shares?	√		EirGenix has regularly collected the list of shareholders according to the list of shareholders obtained by the stock affair agency on the book closure date of EirGenix and maintains good interaction with major shareholders to further collect the list of ultimate controllers.	Corporate Governance Best-Practice Principles
the risk ma	ompany establish and execute anagement and firewall system conglomerate structure?	√		EirGenix has established a German subsidiary and has also established risk control mechanisms such as Management of Related Party Transactions, Measures for Management of Transactions with Related Party, Specific Companies and Group Enterprises, internal control, and internal audit system, which are regularly reviewed and handled in accordance with regulations	Corporate Governance Best-Practice Principles for TWSE/TPEx Listed Companies"
	ompany establish internal rules iders trading with undisclosed n?	✓		EirGenix has established the administrative measures for preventing insider trading and Codes of Ethical Conduct from forbidding insiders from acquiring private interests or competing with EirGenix with undisclosed information.	Corporate Governance



			Implementation Status	Deviations from
Evaluation Item	Y	N	Abstract Illustration	"Corporate Governance Best-Practice Principles for TWSE/TPEx Listed Companies" and Reasons
3. Composition and Responsibilities of the Board of Director(1) Does the Board develop and implement a diversified policy for the composition of its members?	√		Please refer to this Annual Report – E. Diversity and independence of the Board of Directors.	Compliant with "the Corporate Governance Best-Practice Principles for TWSE/TPEx Listed Companies"
(2) Does the company voluntarily establish other functional committees in addition to the Remuneration Committee and the Audit Committee?	√		EirGenix has set up a Remuneration Committee in 2016, an Audit Committee in 2018, and Corporate Governance Committee in 2022. EirGenix will set up other types of functional committees as required by operational development.	Corporate Governance
(3) Does the company establish a standard to measure the performance of the Board and implement it annually, and are performance evaluation results submitted to the Board of Directors and referenced when determining the remuneration of individual directors and nominations for reelection?			On November 11, 2020, the Board of Directors formulated the performance evaluation method for the Board of Directors, specifying that external evaluation shall be carried out at least once every three years. EirGenix conducts performance evaluations regularly every year. As recently as March 8, 2024, the Board of Directors submitted a 2023 internal self-assessment of the Board of Directors, assessing 45 items around the degree of participation in the corporate operation, improvement in the decision-making quality of the Board of Directors, the composition, and structure of the Board of Directors, the election of directors and their continuing education, and internal controls with an average score of more than 90 points, good performance and no major matters to be improved. The Company has retained Taiwan Investor Relations Institute to evaluate the Board in performing its function. The evaluation period was 2021/10/1 to 2022/9/30 and the reporting day was 2022/11/11. The summary of the report has been presented to the Boards on 2023/3/10. The overall condition and recommendation are specified as follows: A. Establishment of the functional committees for the "Sustainable Development Committee": Assist the Board in the ongoing advocacy and intensification of sustainable development and corporate governance pertinent to corporate social responsibility and review the allocation of corporate sources and performance from a higher altitude, and how to systematically connect with and present the ESG Sustainability Report. These will be essential for vitalizing the function of the Board in monitoring and bolstering management mechanisms. B. Early scheduling of the Board meetings and the key issues of the agenda of the year:	Corporate Governance Best-Practice Principles for TWSE/TPEx Listed Companies"



			Implementation Status	Deviations from
			-	"Corporate Governance
Evaluation Item		. .		Best-Practice Principles for TWSE/TPEx Listed
	Y	N	Abstract Illustration	Companies" and
				Reasons
			There are far too many motions proposed in the sessions	
			of the Board and these motions always entail professional	
			content. The early planning of the schedule allows the	
			Directors who do not participate in the routine operation of	
			the Company understand the operation strategy, policy and	
			progress of the Company systematically in full-range and	
			helps to enhance the functional performance of the Board.	
			C. In other words, it should be high time to prepare the 2022	
			consolidated financial statements and the schedule in	
			compliance with applicable laws (presentation of	
			unaudited financial statements to the Board for resolution	
			within 75 days after the conclusion of the fiscal year).	G 11
(4) Does the company regularly evaluate the	√		The Audit Committee of the Company consult the AQIs	-
independence of CPAs?			annually for the assessment of the independence and	-
			competence of the certified public accountants (CPAs) retained as Independent Auditors and present the assessment	_
			report to the Board. There are 17 AQIs including the	
			following dimensions:	Companies
			A. The CPAs do not have any direct or indirect financial	
			interest with the Company, and do not share any benefit.	
			B. There is no undue related between the CPAs and the	
			Group in terms of conflict of interest, financing, or	
			acceptance of kickback.	
			C. The CPAs do not hold any stock or other securities	
			issued by the Group.	
			D. The CPAs do not hold any concurrent positions of the	
			Group.	
			E. No CPA has ever been retained by the Company for	
			more than 7 years and the audit fee is justifiable.	
			F. Annual declaration of the independence of CPAs.	
			G. The CPAs have not been subject to disciplinary action	
			or administrative penalty.	
			H. The stability of audit and taxation staff, provide quality	
			service, conforming to time limit requirement and	
			update information on applicable laws. I. The communication between CPAs and the Directors	
			and Management is good.	
			J. The CPAs have presented recommendation to the	
			system and internal control of the Group and assess and	
			monitor the inherent and potential risks.	
			K. The Board and the Audit Committee assessed the	
			independence of Sheng-Wei Teng and Yu-Fang Yan,	



						Implementation Status	Deviation	ns from
							"Corpora	te Governance
	.						Best-Prac	ctice Principles
	Eva	luation Item	Y	N		Abstract Illustration	for TWS	E/TPEx Listed
			1	1			Compani	es" and
							Reasons	
					CPAs	of PwC Taiwan on 2024/03/08, and confirmed that		
					they a	re conforming to the aforementioned requirements		
					in ind	ependence and competence and are retained as our		
					Indep	endent Auditors for certification.		
4.	Does the com	npany appoint a suitable	✓		On Ma	y 12, 2021, the Board of Directors approved the	Complia	nt with "the
	number of co	mpetent personnel and a			appointme	ent of Chief Financial Officer Hsiu-Chuan Yang,	Corporat	e Governance
	supervisor res	sponsible for corporate			who has n	nore than three years of experience in the position	Best-Prac	ctice Principles
	governance m	natters (including but not			of head	of finance and stock affairs in public issuing	for TWS	SE/TPEx Listed
	limited to pro	viding information for			companie	s, as the head of corporate governance to protect	Compani	es"
	directors and	supervisors to perform their			the equition	es of shareholders, strengthen the functions of the		
	functions, ass	sisting directors and			Board of	Directors, and be responsible for affairs related to		
	supervisors w	vith compliance, handling			corporate	governance jointly with the Finance Department.		
	work related	to meetings of the board of			The he	ad of corporate governance main duties includes		
	directors and	the shareholders' meetings,			the follow	ing items:		
	and producing	g minutes of board meetings			A. Handli	ng matters relating to board meetings and		
	and sharehold	ders' meetings)?			shareho	olders meetings according to laws.		
					B. Assisti	ng in onboarding and continuous development of		
				directors. C. Furnishing information required for business execution by directors.				
					D. Assisti	ng directors with legal compliance.		
						to the Board on review result of the Independent		
					•	ors at the time of their nomination, election, and		
						ity within the term of office.		
						sing the change in Directors.		
						matters set out in the articles or corporation or		
					contrac	•		
Man	ager of Corpo	rate Governance Directors' t	traini	ing r				
Date	<u> </u>	Learning institutions		01	-	Course Title		Hours
2023	3/04/13	Taiwan Academy of Bankir	ng an	d Fi	nance	Corporate Governance Lectures	,	3
2023	3/08/02	Chinese Association of Bus	iness	s and	l Intangible	Measuring Sustainable Value, the Key to ESG: C	orporate	3
2022	,, oo, o <u>u</u>	Assets Valuation				Sustainability Rating Analysis		
2023	3/08/09	Taipei Exchange				Orientation of the Insider Equity for TPEx and E	merging	3
						Stock Market Listed Company The Practice of Corporate Mergers and Acquis	itions at	
2023	3/08/16	Taiwan Investor Relations I	nstit	ute		Home and Abroad	inons at	3
5.	Does the com	npany establish a	√		EirGen	ix has set up the spokesman and vice spokesman	Complia	nt with "the
communication channel and build a				system an	d disclosed the financial and business information	Corporat	e Governance	
	designated section on its website for				on the Ma	arket Observation Post System and the corporate	Best-Prac	ctice Principles
	stakeholders (including but not limited to				website a	according to laws and regulations, designated	for TWS	SE/TPEx Listed
	shareholders,	employees, customers, and			dedicated	personnel responsible for properly responding to	Compani	es"
	suppliers), as	well as handle all the issues			important	issues regarding corporate social responsibility		
	,				-	by stakeholders, and set up a stakeholder's area on		
						^		



		Implementation Status			Deviations from
				1	"Corporate Governance
					Best-Practice Principles
	Evaluation Item	Y	N	Abstract Illustration	for TWSE/TPEx Listed
					Companies" and
					Reasons
	they care for in terms of corporate social			the corporate website to maintain a good and smooth	
	responsibilities			communication channel.	
6.	Does the company appoint a professional	✓		EirGenix has appointed a professional stock affair agency	Compliant with "the
	shareholder service agency to deal with			to handle the shareholders' meeting and stock affairs as the	Corporate Governance
	shareholder affairs			Agency Department of KGI Securities (Stock) Company	Best-Practice Principles
				(Address: 5th Floor, No.2, Section 1, Chongqing South	for TWSE/TPEx Listed
				Road, Taipei City, 100, Tel: (02)2389-2999).	Companies"
7.	Information Disclosure			The website of EirGenix is www.EirGenix.com, on which	Compliant with "the
(1)	Does the company have a corporate	✓		the corporate governance and financial business information	Corporate Governance
	website to disclose both financial			is disclosed in Chinese and English versions.	Best-Practice Principles
	standings and the status of corporate				for TWSE/TPEx Listed
	governance?				Companies"
(2)	Does the company have other information	✓		The website of EirGenix is equipped with a language	Compliant with "the
	disclosure channels (e.g., building an			switching interface, including Chinese and English versions;	Corporate Governance
	English website, appointing designated			there is also the spokesman and acting spokesman system	Best-Practice Principles
	people to handle information collection			and special personnel responsible for collecting and	for TWSE/TPEx Listed
	and disclosure, creating a spokesman			disclosing the corporate information. In addition, relevant	Companies"
	system, webcasting investor conferences)?			information about EirGenix's participation in the Investor	
				Conference has been published on the Market Observation	
				Post System and the corporate website in accordance with	
				regulations.	
(3)	Does the company announce and report		√	EirGenix has announced and reported the financial	Plan to announce and
	annual financial statements within two			reports for the first, second and third quarters and the	report annual financial
	months after the end of each fiscal year			operating conditions for each month in advance before the	
	and announce and report Q1, Q2, and Q3			prescribed time limit; and has not announced and reported	
	financial statements, as well as monthly			the annual financial report within two months after the end	each fiscal year.
	operation results, before the prescribed			of the accounting year.	
	time limit?				
8.	Is there any other important information	√		(1) Employee rights and employee care:	Compliant with "the
	to facilitate a better understanding of the			EirGenix has regularly held all-staff communication	
	company's corporate governance			meetings and Management and Labor Council to exchange	
	practices (e.g., including but not limited			opinions with employees, and also learned about the needs	
	to employee rights, employee wellness,			of employees in a timely manner through multiple	•
	investor relations, supplier relations,			mechanisms such as communication, educational training,	
	rights of stakeholders, directors' and			and incentive.	
	supervisors' training records, the	√		(2) Investor relations and stakeholder rights:	Compliant with "the
	implementation of risk management			In addition to disclosing the financial and business	
	policies and risk evaluation measures,			information in accordance with laws and regulations,	
	the implementation of customer			EirGenix has also established the spokesman and vice	
	relations policies, and purchasing			spokesman system and special personnel responsible for	Companies"
	insurance for directors and supervisors)?			maintaining good investor relations and stakeholder rights.	



			Implementation Status	Deviations from
Evaluation Item	Y	N	Abstract Illustration	"Corporate Governance Best-Practice Principles for TWSE/TPEx Listed Companies" and Reasons
(4) Directors' training records in 2023:	√		(3) Supplier relationship: EirGenix has set up a supplier management policy whose cooperation with suppliers complies with laws and regulations, and contracts to safeguard the rights of both parties.	Best-Practice Principles
Analysis, The Practice of Corporate Merg B. Director-Hsiu-Hui Chen: Corporate Gove Compliance with Laws in Corporate Gove C. Director-Cheng-Yu Cheng: The Duties of Corporate Financial Information and Corp D. Director- Ku-Sung Weng: The Duties of I to ESG: Corporate Sustainability Rating A E. Director- Chun Fu Lu: Measuring Sustain Analysis, The Practice of Corporate Merg F. Director-Yu-Ting Chen: The Practice of C Foreign Corporation (CFC) and Global Ar G. Independent Director-Ming-Thaur Chang: Corporate Sustainability Rating Analysis. H. Independent Director- Po-Chih Chen: Mer Rating Analysis, Orientation of the Inside I. Independent Director-Fu-Shiow Yin: Mea Rating Analysis, The Practice of Corporate	ers arman rman Erna Loya Anal able ers a Corp nti- Grea asur r Equasuri te M	and Ance Lance. yalty and Ance Sand Ance Cax From the Constant of the Constant of the Cax From Sand Sand Sand Sand Sand Sand Sand Sand	and Care of Directors, Analysis and Decision-making of overnance with Securities Regulations. Ind Care of Directors, Measuring Sustainable Value, the Key use, the Key to ESG: Corporate Sustainability Rating Acquisitions at Home and Abroad. Mergers and Acquisitions at Home and Abroad, Controlled Evasion. Douse Gas Inventory and Application, the Key to ESG: Sustainable Value, the Key to ESG: Corporate Sustainability for TPEx and Emerging Stock Market Listed Company. Justainable Value, the Key to ESG: Corporate Sustainability are and Acquisitions at Home and Abroad.	Corporate Governance Best-Practice Principles for TWSE/TPEx Listed Companies"
J. Independent Director-Ming-Shen Chen: C Development Action Plans for TWSE and			(5) Status of implementation of risk management policies and risk measurement standards: EirGenix has set up the risk management policies and procedures and regularly submitted them to the Board of Directors; EirGenix has operated in accordance with laws and regulations, corporate management measures, and various internal control systems, and carried out various risk assessments and controls. Report to the status of risk management at regular intervals of the year: the last report to the Board was presented on 2023/12/22.	for TWSE/TPEx Listed Companies" Compliant with "the Corporate Governance Best-Practice Principles for TWSE/TPEx Listed



			Implementation Status	Deviations from
				"Corporate Governance
Evaluation Item				Best-Practice Principles
Evaluation Item	Y	N	Abstract Illustration	for TWSE/TPEx Listed
				Companies" and
				Reasons
	✓		(7) Status of EirGenix purchasing liability insurance	Compliant with "the
			for Directors:	Corporate Governance
			In the Articles of Incorporation, it has been stated clearly	Best-Practice Principles
			that, within the term of the Directors, EirGenix shall	for TWSE/TPEx Listed
			purchase liability insurance for the compensation liabilities	Companies"
			of the Directors in accordance with the law in the scope of	
			their business. EirGenix has underwritten the Directors'	
			Liability Insurance of US\$ 5 million to Fubon Insurance Co.,	
			Ltd In the future, in addition to continuing underwriting the	
			insurance according to regulations, EirGenix will adjust the	
			insured amount in due course according to operation needs	
			to provide appropriate coverage.	

^{9.} Please explain the improvements which have been made in accordance with the results of the Corporate Governance Evaluation System released by the Corporate Governance Center, Taiwan Stock Exchange, and provide the priority enhancement measures.

EirGenix has been ranked among the top 5% of publicly listed companies by the Taipei Exchange Corporate Governance Evaluations. EirGenix will evaluate the feasibility of the strategies in the current year and future through the items that have not passed the evaluation every year in the future, obtain a balance between the policy development of the competent authority and the development of the company, and immediately promote the implementation plan for the items that can be improved at this stage.



(4) Composition, Responsibilities and Operations of the Remuneration Committee

A. Members of Remuneration Committee

Title	Criteria Name	Professional Qualification / Work Experience	Independence Criteria	Number of Other Public Companies in Which the Individual is Concurrently Serving as a Remuneration Committee Member
Independent Director	Ming-Thaur Chang			0
Independent Director	Po-Chih Chen	Please refer to In	nformation of	0
Independent Director	Fu-Shiow Yin	Directors.		1
Independent Director	Ming-Shen Chen			0

B. Information of Remuneration Committee Operation

- (A) Total members of EirGenix's Remuneration Committee are four people.
- (B) The remuneration committee shall exercise the care of a good administrator in faithfully performing the official powers listed below and shall submit its recommendations for deliberation by the board of directors.
 - i. Prescribe and periodically review the Remuneration Committee Charter.
 - ii. Prescribe and periodically review the performance review and remuneration policy, system, standards, and structure for directors and managerial officers.
 - iii. Periodically evaluate and prescribe the remuneration of directors and managerial officers.
- (C) The current term of the Remuneration Committee is from August 11, 2022, until June 9, 2025.

A total of <u>4</u> (A) Remuneration Committee meetings were held in 2023. The attendance record of the Remuneration Committee members was as follows:

Title	Name	Attendance in Person(B)	By proxy	Attendance Rate (%) 【B/A】
Convener	Ming-Thaur Chang	4	0	100
Committee Member	Po-Chih Chen	3	1	75
Committee Member	Fu-Shiow Yin	4	0	100
Committee Member	Ming-Shen Chen	4	0	100

Note: Independent director Po-Chih Chen was newly appointed on 2022/6/10 and should attend 6 meetings.

A total of $\underline{1}$ (A) meetings of the Board of Directors was held by the end of March 2024. The attendance of directors was as follows:

Title	Name	Attendance in Person(B)	By Proxy	Attendance rate (%)
Independent Director	Ming-Thaur Chang	1	0	100
Independent Director	Po-Chih Chen	1	0	100
Independent Director	Fu-Shiow Yin	1	0	100
Independent Director	Ming-Shen Chen	1	0	100

Other mentionable items:

- a. If the board of directors declines to adopt or modifies a recommendation of the remuneration committee, it should specify the date of the meeting, session, the content of the motion, resolution by the board of directors, and the Company's response to the remuneration committee's opinion (e.g., the remuneration passed by the Board of Directors exceeds the recommendation of the remuneration committee, the circumstances and cause for the difference shall be specified): None.
- b. Resolutions of the remuneration committee objected to by members or expressed reservations and recorded or declared in writing, the date of the meeting, session, the content of the motion, all members' opinions, and the response to members' opinion should be specified: None.
- c. All members of the Company's Remuneration Committee the attended the committee meetings at least twice a year, with a total attendance of 100% and regularly review the policies, systems, standards, and structures for performance evaluation and remuneration to directors and managers as at the 14th meeting convened by the 2nd the Remuneration Committee, to comply with the existing system. Its regular review is based on three major aspects: 1. to ensure external competitiveness, it formulates the salary structure for the senior management with reference to the salary levels in the same industry to



- enhance the Company's competitive advantage; 2. it evaluates the values of their work according to their contribution and abilities based on their responsibilities and positions to ensure fairness in the organization; 3. it rewards them for their special performance and links senior managers' remuneration with the Company's business performance to ensure individual fairness and the organization's competitiveness. The objectives of this salary policy are reviewed based on fairness, reasonableness, motivation, finance, and market competitiveness.
- d. If the Company has net profit in this fiscal year, the Company shall set aside between 1% to 5% of its profits as bonus to employees of the Company and set aside 3% (inclusive) or less of its profits as bonus to Directors. The distribution of bonus to employees may be made by way of cash or shares by the resolution of the Board of Directors. The employees may include certain employees of the subsidiaries who meet the conditions prescribed by the Company. The distribution of employee remuneration and director remuneration shall be heard by over two-thirds of the Board of Directors, be voted in favor for implementation by over one-half of the directors present and represented, and also be reported at the shareholders' meeting. The Company shall first offset its losses in previous years that have not been previously offset, and then set aside annual profits as a bonus to employees of the Company and set aside annual profits as a bonus to Directors.
- e. The performance evaluation of directors and senior managers is linked to their remuneration and their remuneration is determined with reference to the payment standard in the industry and the salary at each job level, while based on their performance and contribution, responsibilities, continuous learning, realization of the Company's core value, leadership and management abilities, training ability, and business goal achievement rate, financial position (such as revenue or achieving status of after-tax net income target), and the progress of self-developed products (such as launch and sales of EG12014 or international factory inspection and certification). It regularly evaluates the performance achievement and reviews the remuneration policy in a timely manner.

C. Nominating Committee: None.

- (5) Fulfillment of ESG and Deviations from the "Corporate Social Responsibility Best Practice Principles for TWSE/TPEx Listed Companies" and disclosure of Climate-Related Information of the company
 - A. Fulfillment of ESG and Deviations from the "Corporate Social Responsibility Best Practice Principles for TWSE/TPEx Listed Companies"

			Implementation Status	Deviations from "the Corporate Social
Evaluation Item	Y	N	Abstract Explanation	Responsibility Best- Practice Principles for TWSE/TPEx Listed Companies" and Reasons
1. Does the company assess ESG risks associated with its operations based on the principle of materiality and establish related risk management policies or strategies?	✓		The Board of Directors delegates the President to integrate the sustainable development concept into the Company's business strategy and lead the finance, human resources, R&D, production, and other departments to promote the Company's core spirit, namely empathy, integrity, responsibility, and global vision, while implementing corporate governance, employee care, environmental sustainability, and social charity projects, on a long-term and systematic basis. The Department of Finance has been responsible for the integration of relevant sustainable development mechanism since the Company was established and recently reported on the implementation status to the Board of Directors on March 8, 2024. The management team reports on the progress of the financial business and devises and regularly reviews business strategy at each Board meeting.	None.



			Implementation Status Deviation Corpora				
Evaluation Item			N	Abstract Explanation	Responsibility Best- Practice Principles for TWSE/TPEx Listed Companies" and Reasons		
				Former Chairman Chung-Hur Lee was appointed the Chief Corporate Sustainable Development Officer and established the Greenhouse Gas Inspection Committee in 2022; planned to complete greenhouse gas inspection and confirmation in 2023, which is ahead of schedule.			
2. Does the company assess ESG risks associated with its operations based on the principle of materiality, and establish related risk management policies or strategies?			The Company's and our subsidiaries' main operational sites and sustainable development performance are within the boundaries of risk assessment. Our risk management organization evaluates the concern about and impact of corporate sustainability and risks in the aspects of environment, society, and corporate governance as per the GRI Standards, the Company's business characteristics, and factors of internal and external environments and stakeholders. The management team formulates management policies after discussions to reinforce our business advantages and risk control. Each operating unit completed the planning of implementation of risk countermeasures and reported them to the Board of Directors.		None.		
Environment	Environment and Management	(2)	laur Gra acci Obt 202 Eir dev	reditation category of "biological products" and effective date from 2022/1 ains ISO14001 (Certificate No. ARES/TW/I2211076E), the effective date 5/11/23. Genix has made great efforts in energy conservation and sustainal elopment and has incorporated the concept of green building into the plant Genix obtained the Green Building Certificate (Green Building Certificate).	MHLW, with the 0/24 to 2027/10/30. from 2022/11/24 to the environmental in Zhubei. In 2020,		
Social	Safe Working Environment Products and services comply with relevant laws and international standards	(2)	Obt date Arra drill Eir	ains ISO45001 Occupational Health and Safety (Certificate No. OHS75 of from 2021/11/09 to 2024/11/08. It and the employee health examination and holds public health and safety to and the education training relevant GMP regularly. Genix has designated exclusive personnel responsible for client contacts, and the processing standards to regularly supervise the implementation restrement and strengthen service processes.	raining, firefighting		
Corporate Governance	Legal Strengthen the functions of directors	We ensure that our personnel duly comply with laws and regulations by establishing a governant organization and implementing the internal control system. (1) Keep updated on the revision of laws and regulations, review various internal norms, and follow laws and regulations to reduce the risk of violations. (2) EirGenix has underwritten the Directors' Liability Insurance of US\$ 5 million to Fubon Insurance Co., Ltd. In the future, in addition to continuing underwriting the insurance according to regulations, EirGenix will adjust the insured amount in due course according to operation needs to provide appropriate coverage.					



						Implementation	Status		Deviations from "the Corporate Social		
	Evaluation Item			N		Abstract Ex	xplanation		Responsibility Best- Practice Principles for TWSE/TPEx Listed Companies" and Reasons		
		Stakeholder rights		in a resp	a timely manner to conses. Genix has establis	identify the issues various stakeholders are concerned about and include the timely manner through the positive interaction with them, while proonses. Senix has established the stakeholders' communication mailbox IR@I gned the spokesperson and vice spokesperson to be the external communication.					
3. (1)	Does the environm	mental issues company establish proper nental management based on the ristics of their industries?	✓		As a profess has established implemented ther official approval Notification, procregulations regal pharmaceutical milestone for Eir facility to receive passed the review	sional drug R&D are perfect environments. On February 3, 2 in its issued GM claiming EirGenix's dearding the quality manufacturing, where the authority's approximately by the FDA and passed the review by	ental management 2020, EirGenix recompliance Instrumental compliance with recompliance and the coval. EirGenix's Zobtained an EIR	apany, EirGenix t systems and ceived PMDA's spection Result elevant Japanese and safety of a remarkable manufacturing Chubei plant has before the drug	None.		
(2)	utilize al efficientl materials	company endeavor to l resources more y and use renewable s which have a low impact avironment?	✓		EirGenix bei industry, which be environmental lo adhered to relev policies and been utilization.	None.					
(3)	potential climate c present a and take	company evaluate the risks and opportunities in change with regard to the nd future of its business appropriate action to climate change issues?	✓		business operation raw material costs to devise measinformation discluthe aim of minir	ont leads all employ ons, with a focus on s, and increased gree sures for develop osure, energy mana mizing the impact of I reports regularly to	environmental regenhouse gas (GHG) ment green build gement, and resource of our operating a	gulations, rising a) emissions and ldings, carbon arce reuse, with activities on the	None.		
(4)	of its grewater conweight or years and energy endioxide reduction	enhouse gas emissions, insumption, and the total of waste in the last two dimplement policies on efficiency and carbon reduction, greenhouse gas in, water reduction, or inagement?	✓		Year 2023 2022 Year 2023 2022 Year	Weight Hazardous waste 7.3638 7.9286	Vater Consumption 95,893.97 73,953.05 of Waste Non-hazardous waste 49.0784 75.4426 nhouse Gas Emiss	Unit: tons Unit: mt Total 56.4422 83.3712 Unit: mt	None.		
					2023		13,837.61 11,594.94				



			Implementation Status	Deviations from "the Corporate Social
				Responsibility Best-
Evaluation Item	Y N		Abstract Explanation	Practice Principles for
		N		TWSE/TPEx Listed
			Companies" and	
				Reasons

Policy on energy conservation and carbon reduction, GHG reduction, water consumption reduction, or other waste management:

- (1) The primary source of power consumption for the Company is electricity. The Company will save energy through (1) continued power monitoring and control system, (2) voluntary management of power consumption units, (3) introduction of energy efficient and green equipment, and (4) effective adjustment of production scheduling to enhance the efficient use of power and avoid unnecessary waste.
- (2) In responding to global climate change, stabilization of water supply emerged as an issue confronting all countries. The Company seeks to pursue its corporate social responsibility and respond to the issue of global water shortage through (1) the continued water consumption monitoring and control system, (2) recycling and reuse of water emitted from pure water system, (3) effective adjustment of production scheduling to reduce the water productivity intensity (total water consumption volume/production value at US million) with the expectation of tackling the challenge from climate change in joint action with enterprises of the world.
- (3) Regularly review climate change-related regulations, establish an internal mechanism for greenhouse gas inventory, and integrate carbon emission information.
- (4) EirGenix belongs to the pharmaceutical research and development industry, which basically does not use materials with high impact on environmental load; Moreover, since its establishment, EirGenix has adhered to relevant government environmental protection laws and policies and been committed to improving the efficiency of resource utilization. Mitigate the impacts of products on the environment and engage in joint venture with academic units in the development of material for disposable items that could possibly be recycled and reused.
- (5) Management of toxic substances: duly observe the rules and regulations governing toxic and chemical items of concern and make all functional units staffed with toxic substances management personnel. In addition, the Company keeps track on the volume of toxic substances in the operation with proper marking of the storage zones and operation areas. All these facilities and areas will be controlled by locking.
- 4. Social issues
- (1) Does the company formulate appropriate management policies and procedures according to relevant regulations and the International Bill of Human Rights?

In order to fulfill the corporate social responsibility and implement the protection of human rights, with reference to the principles enshrined in international human rights conventions such as the Universal Declaration of Human Rights and the United Nations Guiding Principles on Business and Human Rights, EirGenix has respected the basic internationally-recognized human rights and formulated human rights policies applicable to EirGenix, to prevent violations of human rights, provide reasonable and safe workplaces and enable the current colleagues to obtain reasonable and dignified treatment.

None.

The Company's human rights policy and specific management program are as follows:

(1) Diversity, inclusion, and equal opportunity:

In terms of recruitment, remuneration and benefits, training, performance evaluation, promotion, resignation, or retirement, the Company treats all employees and job applicants equally regardless of their socioeconomic status, age, gender, sexual orientation, marriage, family status, disabilities, race, religion, appearance, nationality, language, political affiliation, or pregnancy. We also provide effective and appropriate grievance mechanisms and diverse communication channels to avoid situations that endanger employees' rights and interests, thereby achieving equal employment.

(2) Against forced labor and child labor:

To ensure compliance with corporate social responsibility and ethical standards, the Company's regulations on normal working hours and extended working hours, leave, paid leave, and other types of leave are in compliance with labor laws. We do not force employees to perform labor services. The Company complies with the local regulations on the minimum working age and does not employ child workers.

(3) Physical and psychological health, work balance, and a safe work environment:

The Company attaches great importance to safety and health in the workplace for employees to work in a healthy, safe, and humane environment with a healthy body and mind. The Company encourages employees to participate in health promotion activities and set up their own clubs to bond



Evaluation Item			Implementation Status	Deviations from "the Corporate Social
	Y N			Responsibility Best-
		Abstract Explanation	Practice Principles for TWSE/TPEx Listed	
		Abstract Explanation	Companies" and	
				Reasons

through club activities. In addition to holding the year-end party, cycling, and basketball games to balance their life and help them bond, the Company has installed fitness equipment for them to use after work.

- (4) The Company has established the policies for the prevention of sexual harassment at workplace and the regulations governing reporting on complaints to protect the employees, dispatched personnel and applicants for jobs from sexual harassment and provide them a workplace free of such harassment. The Company also adopts proper measures to prevent, correct, punish and respond any misconduct of this kind and protect the right and privacy of the complainants.
- (5) The company values the opinions and thoughts of all circles and devotes itself to providing open and transparent channels. The company has complaint telephone, mailbox, quarterly labor management meetings. Employees may reflect various problems regarding organization, system and working environment through different channels to carry out our diversified voice response and valuation.
- (2) Does the company have reasonable \checkmark EirGenix has formulated and implemented reasonable employee None. employee benefit measures welfare measures, which can be detailed in the explanation of V. Labor (including salaries, leave, and other Relations of this annual report. benefits), and do business EirGenix has also appropriately reflected the operating performance or results reflect on performance and results in the salaries of employees, has set up employee salaries? bonuses associated with the performance target achievement of employees, departments, and company, and has also issued employee stock options associated with in-service seniority, restricted stock awards associated to the corporate objectives at various stages, and cash capital increase to retain employee stock options, so as to share the corporate operation performance with employees. (3) Does the company provide a EirGenix has attached great importance to providing a healthy and None. healthy and safe working safe working environment, regularly organized public safety and health, environment and organize training GMP-related educational training, arranged physical examination and on health and safety for its group insurance for employees to ensure the safety and health of employees on a regular basis? employees, and also provides the COVID-19 rapid test kit for employee to reduce the infection risk. EirGenix obtains ISO45001 Occupational Health and Safety (Certificate No. OHS751791) in 2021.

No occupational accident took place in 2023 and 2022. Occupational security education and training over the past two years:

	I						
Site	Year	Number of training sessions	Number of attendees for the training				
771 1 '	2023	982	1,905				
Zhubei	2022	777	1,313				
W. 1.	2023	515	1,066				
Xizhi	2022	383	629				

(1)Employee safety and health:

- A. On the first day when employees enter the Company, they will receive the first-day training; the Company will hold safety education and training for at least 3 hours each time at least twice a year. The training mainly covers fire escape drills, emergency drills for poisoning disasters, basic knowledge of occupational safety, and chemical classification management.
- B. We provide adequate personal protective equipment according to the needs in the work environment.
- C. Arrange monthly health check-ins by nurses for colleagues, and also quarterly visits by occupational physicians to the workplace for health consultations. These initiatives provide colleagues with advice on health matters and care for their physical and mental well-being. Additionally, regular health seminars are scheduled, along with weekly distribution of health newsletters via email, fostering a healthy and comfortable work environment for colleagues.



			Implementation Status	Deviations from "the
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				Companies" and
				Reasons

- D. Each employee undergoes a comprehensive health examination every two years, while employees engaged in special operations undergo a special health examination annually in accordance with the Occupational Safety and Health Act.
- E. Regular occupational safety and GMP (Good Manufacturing Practices) related educational training sessions are held to enhance the safety and health awareness and capabilities of all staff in the factory.

(2) Work environment:

- A. Every six months, regular workplace environment monitoring is conducted for employees and the work environment. Testing items include illumination, carbon dioxide levels, noise, high temperatures, chemical operations, etc., ensuring that colleagues operate in a safe and harmless environment.
- B. Daily inspections of the entire factory area are conducted, along with scheduled audits and periodic inquiries to staff about areas needing improvement, eliminating hazards and uncertainties, and providing colleagues with a safe environment.
- C. Reserved parking spaces for female colleagues who are pregnant or within one year postpartum are provided to create a friendly environment. Additionally, lactation rooms are set up within the factory, with allotted time for breastfeeding, ensuring peace of mind and dedicated space for breastfeeding mothers during work hours.

To provide an account of the number of fire incidents, casualties, and the ratio of casualties to the total number of employees for the current year, as well as the corresponding improvement measures in response to fires: No related incidents occurred

WCI	as the corresponding improvement mea	sures in response to fires. No related incidents occurred	
(4)	Does the company provide its	The employees have been performing to achieve their annual	None.
	employees with career	targets based on their personal strength. The supervisors will also	
	development and training sessions?	provide advice and guidance; EirGenix organized internal education and	
		training regularly and encouraged employees to participate in external	
		educational training or continue their studies to enhance their self-	
		ability. In September 2017, EirGenix established the "EIRGer's Learning	
		Center," planning diversified internal training courses every year. The	
		key learning focus is on professional, technical orientation,	
		supplemented by management and core functions.	
(5)	Do the company's products and	EirGenix will follow the relevant regulations and international	None.
	services comply with relevant laws	standards on the advertisement, labeling of products and services,	
	and international standards in	customer health and safety, and client privacy When the self-owned	
	relation to customer health and	products come into the market, EirGenix will formulate the customer	
	safety, customer privacy, and	protection policies and appealing procedures; In addition, for CDMO of	
	marketing and labeling of products	bio-pharmaceuticals, EirGenix has designated exclusive personnel	
	and services, and are relevant	responsible for client contacts, and the responsible unit has set up the	
	consumer protection and grievance	processing standards to regularly supervise the implementation results,	
	procedure policies implemented?	implement product improvement and strengthen service processes.	
(6)	Does the company implement	EirGenix has established the procedures for the assessment and	None.
	supplier management policies,	management of suppliers in 2020 and specify in the contracts with the	
	requiring suppliers to observe	suppliers that all shall duly observe applicable laws governing labor,	
	relevant regulations on	human right, environmental protection, safety and health, and	
	environmental protection,	environment and the society, or provide related declarations for the	
	occupational health, and safety, or	upgrade of sustainable development of the enterprises as the goal. In	
	labor and human rights?	addition, the Company also includes environmental, occupational safety	
		and health management, technical skills and supply capacity evaluation	
		as management measures for the screening of contractors.	



				Implementation Status	Deviations from "the Corporate Social
					Responsibility Best-
	Evaluation Item				Practice Principles for
		Y	N	Abstract Explanation	TWSE/TPEx Listed
					Companies" and
					Reasons
5.	Does the company reference	√		EirGenix published the Corporate Social Responsibility Report	None.
	internationally accepted			based on the GRI Standard in 2023 and uploaded to MOPS and company	
	reporting standards or			website. EirGenix will continue the compilation of the report in 2024	
	guidelines and prepare			and will appoint a third-party accreditation institution for assurance	
	reports that disclose non-			inspection.	
	financial information of the				
	company, such as corporate				
	social responsibility reports?				
	Do the reports above obtain assurance from a third-party				
	verification unit?				

6. Describe the difference, if any, between actual practice and the corporate social responsibility principles if the company has implemented such principles based on the Corporate Social Responsibility Best Practice Principles for TWSE/TPEx Listed Companies:

EirGenix has formulated the corporate social responsibility principles in accordance with the Corporate Social Responsibility Best Practice Principles for TWSE/TPEx Listed Companies, and EirGenix has operated in accordance with relevant laws and regulations without significant difference

7. Other useful information for explaining the status of corporate social responsibility practices:

With the corporate spirit indicators of Empathy, Integrity, Responsibility, and Global Vision, all employees of EirGenix shall follow this indicator spirit to practice corporate social responsibility.

• The specific benefits of investing in energy-efficient or green energy-related environmentally sustainable machinery and equipment, as well as investing in country's green energy peripheral industries, are as follows

Investment equipment	Investment amount	Investment benefits
Chiller Unit	NT\$ 4,220 thousands	 In 2023, the Company invested in the construction of a new chiller unit, opting for equipment rated at performance level 1, which compared to previous installations using performance level 2 equipment: Energy Savings: 175,173 kWh per year Reducing energy consumption by 630,622.8 million joules Reduction of 86.71 metric tons CO2e

B. Climate-Related Information of TWSE/TPEx Listed Company

(A) Implementation of Climate-Related Information

Item	Implementation status
Describe the board of directors' and management's oversight and governance of climate-related risks and opportunities.	The board of directors has authorized the general manager to integrate sustainable development into the business strategy, guiding each department with corporate core values such as Empathy, Integrity, Responsibility, and Global Vision. This initiative aims to promote corporate governance, employee wellbeing, environmental sustainability, and social welfare. Company employees are expected to adhere to these values in order to fulfill their corporate social responsibility. In active cooperation with Taiwan government commitment to achieving net-zero carbon emissions by 2050, EirGenix officially adopted the Task Force on Climate-related Financial Disclosures (TCFD) and established the TCFD Risk Management Task Force starting in 2023. The company follows the four frameworks of TCFD, conducting discussions on climate governance, strategy, risk management, and goal setting. Additionally, climate-related issues are incorporated into the risk management process. The TCFD Risk Management Task Force will hold regular meetings to monitor, assess, and discuss climate risks. It



Item	Implementation status						
	will also provide an annual report to the Board of Directors on the regulation, assessment, and						
	implementation outcomes related to climate risks.						
2. Describe how the identified climate risks and opportunities affect the business, strategy, and finances of the business (short, medium, and long term).	EirGenix adheres to the TCFD framework in order to identify risks and opportunities that affect its business, strategy, and financial planning. The relevant departments define and list these risks and opportunities. EirGenix considers the climate scenarios RCP 2.6, RCP 4.5, and RCP 8.5 as defined by the Intergovernmental Panel on Climate Change (IPCC). The Company conducts risk assessments to evaluate transition risks, acute physical risks, and chronic physical risks. These assessments identify and analyze climate risks and opportunities within the company's operational scope, taking into account short, medium, and long-term perspectives.						
3. Describe the financial impact of extreme weather events and transformative actions.	Based on observation data from the weather stations of the Central Weather Administration, the average annual temperature in Taiwan has risen by approximately 1.6°C over the past 110 years (1911-2020). Furthermore, there has been an accelerating trend of warming in the past 50 and 30 years. Based on the scenario which are the temperatures in different regions of Taiwan are expected to continue increasing in the future. Under the worst-case scenario of global warming (SSP5-8.5), the average temperature in the middle and end of the 21st century may rise by more than 1.8°C and 3.4°C, respectively. Under the ideal mitigation scenario (SSP1-2.6), the temperature may increase by 1.3°C and 1.4°C to simulate the potential negative impacts on our company due to extreme weather events in the mid-21st century (2050), and the results are as follows:						
	Scenario Potential Climate Impacts						
	 RCP 2.6 The average annual temperature is projected to increase by over 2.2°C, potentially impacting the temperature within the factory and its surrounding environment, thereby affecting production efficiency. Consequently, it is imperative to invest in improvement equipment. A rise in rainfall can potentially cause flooding, particularly with a nearly 10% increase in maximum rainfall. Insufficient drainage facilities near the factory area may result in the flooding of factory buildings or damage to raw materials, finished products, and equipment. 						
	 RCP 4.5 The average annual temperature is projected to increase by 2.5°C and 2.8°C, potentially impacting the temperature within the factory and its surrounding environment, thereby affecting production efficiency. Consequently, it is imperative to invest in improvement equipment. Moreover, with the longer duration of high temperatures in recent summers, it may be necessary to enhance ventilation and air conditioning systems to safeguard employees from heatstroke. However, this will lead to increased electricity expenses and equipment maintenance costs. The increase in average rainfall may lead to an increase in flooding. Currently, the annual average rainfall in the EirGenix factories' areas has increased by approximately 9 to 11.1%. Poor drainage facilities near the factories may result in flooding of the premises or damage to raw materials, finished products, and machinery. 						
	 RCP 8.5 There is a possibility that the average annual temperature increase may exceed 3.2°C, which could result in a continuous temperature rise. It is essential to consistently enhance the air conditioning in the factory buildings. Rising annual average temperatures could potentially reduce the frequency of typhoons and increase the likelihood of droughts. In the event of extreme weather conditions, the country where EirGenix is situated may be prone to flooding. This could result in transportation disruptions, impacting the commute of personnel and potentially causing injuries. 						
4. Describe how climate risk identification, assessment, and management processes are integrated into the overall risk management system.	The Board of Directors at EirGenix formulated the "Risk Management Policy and Procedures" in 2020. This policy serves as the highest guiding principle for our risk management and aims to identify potential risks (including market, liquidity, operational, hazards, and legal risks) that may impact our operations and profitability. It provides a reference for formulating operational strategies. Additionally, we take appropriate preventive measures for risk warnings to enhance our ability to respond to risk events and minimize their impact on our business operations. Every year, the responsible units identify the relevant risk factors and analyze various risks to assess their potential impact on operations. Risk control measures						



	Item	Implementation status
		are subsequently developed to ensure that the risks remain within manageable and acceptable limits. A report on the implementation of risk management is then submitted to the board of directors. EirGenix oversees climate-related action plans through the TCFD Risk Management Task Force, which comprises a chairman, convener, multiple departments, and external professional advisory consultants. Under the guidance of the TCFD Risk Management Task Force, department managers and colleagues are evaluating industry characteristics and operational conditions to assess the potential impact of different risks and opportunities on our operations. The board of directors should receive an annual report on the status of risk management operations and execution, which should also include discussions on climate change issues.
5.	If scenario analysis is used to assess resilience to climate change risks, the scenarios, parameters, assumptions, analysis factors and major financial impacts used should be described.	According to Sustainable Development Roadmap for TWSE/TPEx Listed Companies, the assessment will be done on 2029 expectedly due to the company's capital is lower than NT\$5 billion.
6.	If there is a transition plan for managing climate-related risks, describe the content of the plan, and the indicators and targets used to identify and manage physical risks and transition risks.	
7.	If internal carbon pricing is used as a planning tool, the basis for setting the price should be stated.	
8.	If climate-related targets have been set, the activities covered, the scope of greenhouse gas emissions, the planning horizon, and the progress achieved each year should be specified. If carbon credits or renewable energy certificates (RECs) are used to achieve relevant targets, the source and quantity of carbon credits or RECs to be offset should be specified.	
9.	Greenhouse gas inventory and assurance status and reduction targets, strategy, and concrete action plan	Please refer to the following explanations.

(B) Greenhouse Gas Inventory and Assurance Status for the Most Recent 2 Fiscal Years

i. Greenhouse Gas Inventory Information

Greenhouse Gas Inventory Information for the most recent 2 fiscal years as the following table, and the Scope 1, Scope 2

are the information of EirGenix Inc. and EirGenix Europe GmbH.

N/	Sec	ope 1	Sec	ope 2	Assurance	Assurance Opinion
Year	Emission Volume (mt CO2e)	Intensity (mt CO2e/NT\$ million)	Emission Volume (mt CO2e)	Intensity (mt CO2e/NT\$ million)	Institutions C BSI Rea As	Opinion
2022	933.418	0.00063	8631.333	0.00582	BSI	Reasonable Assurance
2023	1090.1268	0.00106	10348.4132	0.01011	NA	NA



In the disclosure of total greenhouse gas emissions by the company, 9564.751 metric tons of CO2e (accounting for 82.5% of total emissions) have been assured by the assurance institutions in accordance with ISO 14064-3:2019 assurance standards. The assurance opinion is reasonable assurance.

ii. Greenhouse Gas Assurance Information

As of 2023, the greenhouse gas emissions inventory for both 2022 and 2023 has been completed. The inventory for the year 2022 was externally assured by BSI in December 2023, in accordance with the ISO 14064-3:2019 assurance standard. The assurance results provide reasonable assurance. Due to Financial Supervisory Commission requirements, the assurance is scheduled for completion in 2026 based on the company's capital.

(C) Greenhouse Gas Reduction Targets, Strategy, and Concrete Action Plan

EirGenix is a professional pharmaceutical research and production company that has implemented a comprehensive environmental management system. In order to fulfill its corporate social responsibility and strive for environmental sustainability, EirGenix prioritizes "energy conservation and carbon reduction." The company is currently in an expansion phase, using 2022 as the base year. Once the expansion is completed, EirGenix will gradually reduce energy intensity and minimize resource and energy waste. To achieve this goal, the company has established three key performance indicators for "Electricity sage," "Water resources," and "Waste," and is actively promoting environmental sustainability initiatives. The company introduced greenhouse gas inventory in 2023 to monitor the Company's greenhouse gas emissions. The company's carbon neutrality roadmap as follow.

	Carbo	n Neutrality Roadmap	
Time	Short-term (~2025)	Mid-term (2025~2030)	Long-term (2030~2050)
Reduction Targets	Gradually reduce after the expansion is completed.	Reduce 10% compare with 2022	To achieve carbon neutrality
Strategy, and Concrete Action Plan	 Obtain ISO 14001:2015 Environmental Management System Certification for Biotechnology Testing and Analysis Implementation of ISO 14064-1 greenhouse gas inventory counseling and planning and verification By 2025, the proportion of renewable energy will reach 1%. Promote low-carbon manufacturing and consistently review the reduction of carbon emission intensity. EirGenix's Zhubei A plant obtained the Green Building Certificate. Improve energy efficiency to attain an annual energy- saving performance of 1%. Actively engaged in a net- zero green lifestyle. 	 Obtain Certification for ISO 50001 Energy Management System. Gradually increase the utilization of renewable energy to reach 6% by the year 2030. EirGenix's Zhubei new B plant obtained the Green Building Certificate. Continuously enhance energy management to attain an annual energy-saving efficiency of 1%. Implement low carbon supplier management. Evaluate waste management policies and eco-friendly packaging materials. 	 Gradually increase the utilization of renewable energy to reach 10% by the year 2050. Continued focus on carbon rights, carbon sink, and renewable energy. Implement a green supplier management system and measure sustainability indicators. Participate in climate advocacy organizations or alliances to collectively promote environmental sustainability.



(6) Fulfillment of Ethical Corporate Management and Deviations from the "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies"

				Implementation Status	Deviations from the "Ethical
Evaluation Item		Y	N	Abstract Illustration	Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies" and Reasons
1. (1)	Establishment of ethical corporate management policies and programs Does the company have a Board-approved ethical corporate management policy and stated in its regulations and external correspondence the ethical corporate management policy and practices, as well as the active commitment of the Board of Directors and management towards enforcement of such policy?	✓		The Board of Directors of EirGenix has passed the establishment of the Ethical Corporate Management Best Practice Principles as well as Procedures and Guidelines of Conduct for Integrity Management to express the policies of integrity operation. The Board of Directors and the management team have also actively implemented integrity management and clearly expressed the policies and practices of integrity management in the corporate regulations and external business contracts.	Compliant with "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies."
(2)	Does the company have mechanisms in place to assess the risk of unethical conduct and perform regular analysis and assessment of business activities with a higher risk of unethical conduct within the scope of business? Does the company implement programs to prevent unethical conduct based on the above and ensure the programs cover at least the matters described in Paragraph 2, Article 7 of the Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies?			EirGenix has formulated the Ethical Corporate Management Best Practice Principles, Procedures, and Guidelines of Conduct for Integrity Management, Codes of Ethical Conduct, and Procedures of Administrative Measures for Preventing Insider Trading. The Legal Department and Audit Department have also regularly reported to the Board of Directors on the status of implementation and irregularly checked, analyzed, and evaluated the operating activities within the business scope that have a high risk of dishonest behavior.	Compliant with "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies."

- A. When engaging in commercial activities, directors, managers, employees, and mandataries of the company or persons having substantial control over such companies shall not directly or indirectly offer, promise to offer, request or accept any improper benefits, nor commit unethical acts including breach of ethics, illegal acts, or breach of fiduciary duty for purposes of acquiring or maintaining benefits.
- B. The company shall establish a risk assessment mechanism against unethical conduct, analyze and assess on a regular basis business activity within their business scope which are at a higher risk of being involved in unethical conduct, and establish prevention programs accordingly and review their adequacy and effectiveness on a regular basis.
 - The company to refer to prevailing domestic and foreign standards or guidelines in establishing the prevention programs, which shall at least include preventive measures against the following:
 - (A) Offering and acceptance of bribes.



				Implementation Status	Deviations from the "Ethical						
	Evaluation Item		N	Abstract Illustration	Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies" and Reasons						
	(B) Illegal political donations.										
	(C) Improper charitable donations or sponsorship.										
	(D) Offering or acceptance of unreasonable presents or hospitality, or other improper benefits.										
	(E) Misappropriation of trade secrets and infringement of trademark rights, patent rights, copyrights, and other intellectual property rights.										
	(F) Engaging in unfair compe	titiv	e pra	ctices.							
	(G) Damage directly or indirect	ctly	caus	ed to the rights or interests, health, or safety of consumers or other stal	keholders in the course of						
	research and development	, pro	cure	ment, manufacture, provision, or sale of products and services.							
(3)	Does the company provide	✓		EirGenix has formulated the Ethical Corporate Management	Compliant with "Ethical						
	clearly the operating			Best Practice Principles, Procedures and Guidelines of Conduct for	Corporate Management Best						
	procedures, code of conduct,			Integrity Management, Employee Working Principles, Codes of	Practice Principles for						
	disciplinary actions, and			Ethical Conduct, and Administrative Measures for Preventing	TWSE/TPEx Listed						
	appeal procedures in the			Insider Trading, set up a disciplinary and appealing system for	Companies".						
	programs against unethical			violations, regularly conducted review and correction, and							
	conduct? Does the company			implemented and advocated operating activities to prevent risks of							
	enforce the programs above			dishonest behavior.							
	effectively and perform										
	regular reviews and										
	amendments?										
2.	Fulfill operations integrity										
	policy.	√									
(1)	Does the company evaluate			EirGenix has conducted its business activities in a fair and	Compliant with "Ethical						
	business partners' ethical			transparent manner. Before business activities, EirGenix has	Corporate Management Best						
	records and include ethics-			avoided dealings with trading partners who have dishonest	Practice Principles for						
	related clauses in business			behaviors, with the terms of cooperation stated in the contract.	TWSE/TPEx Listed						
(2)	contracts?				Companies."						
(2)	Does the company have a unit	V		EirGenix has set up a dedicated unit under the Board of	Compliant with "Ethical						
	responsible for ethical			Directors to promote corporate integrity management as the Legal	Corporate Management Best						
	corporate management on a			Department, which is responsible for formulating and supervising	Practice Principles for						
	full-time basis under the Board			the implementation of integrity management policies and	TWSE/TPEx Listed Companies."						
	of Directors, which reports the ethical corporate management			prevention plans, handling and reporting the breach of integrity that	Companies.						
	policy and programs against			may be found in the internal control audit in accordance with							
	unethical conduct regularly (at			relevant laws and regulations, and ensuring that the corporate integrity management policies can be implemented and reported to							
	least once a year) to the Board			the Board of Directors regularly every year, with the latest reporting							
	of Directors while overseeing			date of March 8, 2024.							
	such operations?			date of March 6, 2024.							
(3)	Does the company establish	√		EirGenix has formulated the Ethical Corporate Management	Compliant with "Ethical						
	policies to prevent conflicts of			Best Practice Principles, Procedures and Guidelines of Conduct for	Corporate Management Best						
	interest and provide			Integrity Management, Employee Working Principles, Codes of	Practice Principles for						
	appropriate communication			Ethical Conduct, and Administrative Measures for Preventing	TWSE/TPEx Listed						
	channels, and implement it?			Insider Trading, and set up whistle blower policy with a designated	Companies."						
	•			email for employees putting a stop on all unethical immoral or	-						
				illegal work.							



				Implementation Status	Deviations from the "Ethical Corporate Management Best
	Evaluation Item		N	Abstract Illustration	Practice Principles for TWSE/TPEx Listed Companies" and Reasons
(4)	Does the company have effective accounting and internal control systems in place to implement ethical corporate management? Does the internal audit unit follow the results of unethical conduct risk assessments and devise audit plans to audit the systems accordingly to prevent unethical conduct, or hire outside accountants to perform the audits?			EirGenix has established effective systems for both accounting and internal control, and the internal audit unit has also conducted audits on a regular basis and reported to the Board of Directors and the audit committee every time; it has also appointed CPAs to carry out the audit.	Compliant with "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies."
(5)	Does the company regularly hold internal and external educational trainings on operational integrity?	✓		EirGenix has regularly held all-staff communication meetings and internal educational training to make employees understand the corporate spirit indicators and the corporate culture of integrity management and encouraged employees to participate in external educational training.	Compliant with "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies."
3.	Operation of the integrity channel				
(1)	Does the company establish both a reward/punishment system and an integrity hotline? Can the accused be reached by an appropriate person for follow-up	✓		EirGenix has formulated the Ethical Corporate Management Best Practice Principles and Guidelines of Conduct for Integrity Management. In case of any breach of integrity, employees can report it to the heads of department, Legal Department or Audit Department at any time through the reporting email address or in any form.	Compliant with "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies."
(2)	Does the company have in place standard operating procedures for investigating accusation cases, as well as follow-up actions and relevant post-investigation confidentiality measures?	\		EirGenix has formulated the Ethical Corporate Management Best Practice Principles and Guidelines of Conduct for Integrity Management, provided smooth reporting channels, and implemented the principle of confidentiality.	Compliant with "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies."
(3)	Does the company provide proper whistleblower protection?	✓		EirGenix keeps the contents of reporting on breach of integrity management confidential and protects the whistleblower from improper disposal due to reporting.	Compliant with "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies."
4.	Strengthening information disclosure Does the company disclose its ethical corporate management policies and the results of its implementation on the	✓		EirGenix has disclosed the Ethical Corporate Management Best Practice Principles and Guidelines of Conduct for Integrity Management and information related to integrity management on	Compliant with "Ethical Corporate Management Best Practice Principles for TWSE/TPEx Listed Companies."



	Implementation Status			Deviations from the "Ethical
				Corporate Management Best
Evaluation Item	Y N Abstrac			Practice Principles for
		Abstract Illustration	TWSE/TPEx Listed Companies"	
				and Reasons
company's website and MOPS?			the Market Observation Post System, annual report, and corporate	
			website.	

- 5. If the company has established the ethical corporate management policies based on the Ethical Corporate Management Best-Practice Principles for TWSE/TPEx Listed Companies, please describe any discrepancy between the policies and their implementation:
 - EirGenix has formulated the Ethical Corporate Management Best Practice Principles and Guidelines of Conduct for Integrity Management in accordance with the Integrity Management Principles for TWSE/TPEx Listed Companies, with no difference between the actual operation and the principles.
- 6. Other important information to facilitate a better understanding of the company's ethical corporate management policies (e.g., review and amend its policies):
 - (1) EirGenix has formulated the Ethical Corporate Management Best Practice Principles and Guidelines of Conduct for Integrity Management, which will be amended as appropriate according to the operational development.
 - (2) All employees have signed the Declaration of Ethic Code of Conduct and Business Integrity.
 - (3) Education on applicable laws and important notice of insider trade and equity holding of insider for the Directors and the management will be provided every month. The Directors have completed their training of the company operation or related training for 6 hours in 2023.
 - (4) Required training for all employees on laws and prevention of insider trade for one hour on 2023/08/15 and 2023/08/22, respectively.
 - (5) Legal Affairs, Audit Office and Finance Department provide information on applicable laws and case studies to the Directors, managers, and employees from time to time to realize ethical corporate management and the prevention of insider trade. Related rules and regulations have been disclosed at the intranet and external official website of the Company.
 - (6) Legal Affairs and Audit Office conduct audit on respective functional departments at random, and report to the Board of the status, conduct analysis and control of business activities at high risk of unethical practices within the scope of operation of the Company.



(7) If the company has adopted corporate governance best-practice principles or related bylaws, disclose how these are to be searched:

EirGenix has instituted related rules and regulations in accordance with the "Corporate Governance Best Practice Principles for TWSE/TPEx Listed Companies" and the requirements of the competent authority of securities and exchange. For further information, visit the official website of the Company for inquiry of the content of related rules and regulations.

(8) Other Important Information Regarding Corporate Governance:

All directors, managers, and insider newly assumed office would be released the updated version of the Regulations Governing the Equity Ownership of Insiders of Companies listed at TPEx and Emerging Stock Market" compiled by Taipei Exchange. Information on the amendment to the regulations will be announced and promoted in the Board meeting and the executive meeting every year for the insiders to abide by.

The Company has established the Corporate Governance Committee organized by the Chairman and four Independent Directors. The purpose is to strengthen corporate governance and upgrade the performance of the Board. This committee shall perform the following functions:

- A. Review the institution and amendment to corporate governance systems such as the Corporate Governance Best Practice Principles of the Company.
- B. Monitor and supervise the practice of corporate governance of the Company.
- C. Monitor and supervise the Company in the participation of corporate governance evaluation.
- D. Evaluate the performance of the Board, the committees and the Directors, the independence of the Independent Directors, and present the evaluation result to the Board.
- E. Assess the channels for the gathering of information for the Board, and the quality and timing of the information received by the Directors.
- F. Monitor the governance relations between the Company and its subsidiaries and other affiliates.
- G. Other materiality as required by the Company or the competent authority.

The tenure for the members of the 1st term of the committee started on 2022/12/28 and ends on 2025/06/09. The Committee convened once at the end of March 2024 for discussion on the amendment to the Corporate Governance Best Practice Principles.

Title	Name	Main expertise	Attendance in Person	Attendance Rate (%)
Convener	Ming-Shen Chen	Finance · Corporate Governance	1	100
Committee Member	Lee-Cheng Liu	Operation management · Biotechnology · Corporate Governance	1	100
Committee Member	Ming-Thaur Chang	Finance	1	100
Committee Member	Po-Chih Chen	Economist	1	100
Committee Member	Fu-Shiow Yin	Biotechnology	1	100

- (9) Disclosures Required for the Implementation of the Internal Control System:
 - A. Statement of Internal Control System: Please refer to appendix 1.
 - B. If CPA has been hired to carry out a special audit of the Internal Control System, the company shall furnish the CPA audit report: None.
- (10) If there has been any legal penalty against the company or its internal personnel, or any disciplinary penalty by the company against its internal personnel for violation of the internal control system, during the most recent fiscal year or during the current fiscal year up to the publication date of the annual report, where the result of such penalty could have a material effect on shareholder equity or securities prices, the annual report shall disclose the penalty, the main shortcomings, and condition of improvement: None.



(11) Material resolutions of a shareholders meeting or a board of directors meeting during the most recent fiscal year (2023) or during the current fiscal year up to the date of publication of the annual report:

Item & Date	Major Resolutions of Shareholders' Meeting/ Board of directors
Board	
Meeting 2023/01/16	A. Approved the motion of distribution of year-end bonuses for the managers.
	A. Accept 2022 Financial Statements and Business Report.
	B. Ratification of the 2022 Deficit Offset Proposal.
	C. Approved the motion of issuance of the Declaration of Internal Control System of the Company.
	D. Approved the motion of the ratification of the assessment of the independence and competence of the CPAs retained as external Auditors.
	E. Approved the motion of the ratification of the appointment of CPAs as external auditors and the remuneration to the CPAs.
	F. Approved the certified public accounts previously approved as the Independent Auditors, the CPA office
	and affiliates to render non-assurance services to the Company and the subsidiaries.
	G. Approved the application to Far Eastern International Bank for the loan.
	H. Approved the extension to Shanghai Commercial & Savings Bank for the loan.
	I. Approved the extension to Chang Hwa commercial bank for the loan.
	J. Approved the extension to Cathay United bank for the loan.
	K. Approved the amendment to the "Article of the Company".
Board	L. Approved the amendment to the "Regulations Governing Procedure for Board of Directors Meetings".
Meeting	M. Approved the amendment to the "Sustainable Development Best Practice Principles".
2023/03/10	N. Approved the amendment to the "Corporate Governance Best Practice Principles".
	O. Approved the base date of employee stock option into common stocks capital increase.
	P. Approved the base date of cancellation of the restricted stock award.
	Q. Adoption of the 2023 1 st and 2 nd Employee Restricted Stock Awards.
	R. Approved the discontinue the Private Security Offering Approved by the 2022 Shareholders' Meeting.
	S. Approved the Company will raise capital through private placements of common shares.
	T. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of directors and their representatives.
	U. Approved the motion of the agenda and related matters of the Shareholders' Meeting of 2023.
	V. Approved salary policies, regulations, standards, and structure.
	W. Approved to grant 2022 Employee Stock Options to employees.
	X. Approved to grant 2022 1st Employee Restricted Stock Awards to employees.
	Y. Approve the annual adjustment to the salary of the executive.
	Z. Approved the annual adjustment to the salary and the motion for distribution of year-end bonuses for the
	managers at a European subsidiary.
Board Meeting 2023/05/10	A. Accept 2023 Q1 Financial Statements.
	B. Approved the application to Yuanta Commercial Bank for the loan.
	C. Approved the base date of employee stock option into common stocks capital increase.
	D. Approved to grant 2022 Employee Stock Options to employees.
	Proposed Resolutions:
	A. Accept 2022 Financial Statements and Business Report.
Shareholders	Implementation review: Financial Statements have been announced on 2023/03/16 and Business Report
Meeting	has been announced on 2023/04/28.
2023/05/31	B. Ratification of the 2022 Deficit Offset Proposal.
	Implementation review: It has been announced on 2023/05/31 after the shareholders' meeting.
	C. Adjustment of the Utilization Plan for Capital Injection by Private Placement.



Item & Date	Major Resolutions of Shareholders' Meeting/ Board of directors
	Implementation review: It has been announced by every quarter.
	Matters for discussion:
	A. Approved the amendment to the Article of the Company.
	Implementation review: Approval No. 11230107610 dated July 16, 2023.
	B. Approved the Issuance of Employee Restricted Stock Awards.
	Implementation review: Implementation completed in accordance with the resolution of the
	Shareholders Meeting. Approval No. 1123589721 dated Oct. 27, 2023.
	C. Approved the Company will raise capital through private placements of common shares.
	Implementation review: As the termination date is near, in consideration of working capital and market
	status, the said private placement shall not be renewed and continued.
	D. Release the Prohibition on Directors or Representatives of Directors from Participation in Competitive
	Business.
	Implementation review: Implementation completed in accordance with the resolution of the
	Shareholders' Meeting.
	A. Accept 2023 Q2 Financial Statements.
	B. Approved the extension to The Shanghai Commercial & Savings Bank for the loan.
	C. Approved the extension to CTBC Bank for the loan.
	D. Approved the extension to Mega Bank for the loan.
Board	E. Approved the extension to Bank of Panshin for the loan.
Meeting	F. Approved to formulate "Operational Procedure for Preparation and Validation of the Sustainability
2023/08/08	Report".
	G. Approved the base date of employee stock option into common stocks capital increase.
	H. Approved to grant 2022 Employee Stock Options to employees.
	I. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of
	directors and their representatives.
	A. Accept 2023 Q3 Financial Statements.
	B. Approved the extension to Taichung Commercial Bank for the loan.
	C A managed the establishment of a subsidiance in the Highest Cotates bear down the enough and development
	C. Approved the establishment of a subsidiary in the United States based on the operational development
	needs.
	needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and
	needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form".
	needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget
Board	needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase)
Board Meeting	needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) F. Approved the base date of employee stock option into common stocks capital increase.
Meeting	needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) F. Approved the base date of employee stock option into common stocks capital increase. G. Approved the base date of cancellation of the restricted stock award.
Meeting	needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) F. Approved the base date of employee stock option into common stocks capital increase. G. Approved the base date of cancellation of the restricted stock award. H. Approved the donation of EIRGASUN to NTUH for academic research.
Meeting	 needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) F. Approved the base date of employee stock option into common stocks capital increase. G. Approved the base date of cancellation of the restricted stock award. H. Approved the donation of EIRGASUN to NTUH for academic research. I. Approved to update the 2023 1st and 2nd Employee Restricted Stock Awards to employees.
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Meeting	 needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) F. Approved the base date of employee stock option into common stocks capital increase. G. Approved the base date of cancellation of the restricted stock award. H. Approved the donation of EIRGASUN to NTUH for academic research. I. Approved to update the 2023 1st and 2nd Employee Restricted Stock Awards to employees. J. Approved to grant 2022 1st Employee Restricted Stock Awards to employees. K. Approved to grant 2023 2nd Employee Restricted Stock Awards to employees. L. Approved to sign the Amended and Restated License Agreement for the biosimilar EG12014 with Sandoz AG. M. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of
Meeting 2023/11/09	needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) F. Approved the base date of employee stock option into common stocks capital increase. G. Approved the base date of cancellation of the restricted stock award. H. Approved the donation of EIRGASUN to NTUH for academic research. I. Approved to update the 2023 1st and 2nd Employee Restricted Stock Awards to employees. J. Approved to grant 2022 1st Employee Restricted Stock Awards to employees. K. Approved to grant 2023 2nd Employee Restricted Stock Awards to employees. L. Approved to sign the Amended and Restated License Agreement for the biosimilar EG12014 with Sandoz AG. M. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of directors and their representatives.
Meeting 2023/11/09 Board	 needs. D. Approved the amendment of "IA-18 Implementation of authorization" and "deputy systems and Approval Authority Form". E. Approved the Production Line Expansion and Production Equipment at Zhubei Facility. (Budget Increase) F. Approved the base date of employee stock option into common stocks capital increase. G. Approved the base date of cancellation of the restricted stock award. H. Approved the donation of EIRGASUN to NTUH for academic research. I. Approved to update the 2023 1st and 2nd Employee Restricted Stock Awards to employees. J. Approved to grant 2022 1st Employee Restricted Stock Awards to employees. K. Approved to grant 2023 2nd Employee Restricted Stock Awards to employees. L. Approved to sign the Amended and Restated License Agreement for the biosimilar EG12014 with Sandoz AG. M. Approved the proposal to the shareholders meeting to lift the restrictions on the non-competition of
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tem & Date	Major Resolutions of Shareholders' Meeting/ Board of directors
	D. Approved the Internal Audit Plan for the fiscal year 2024.
	E. Approved the operation plan for fiscal year 2024.
	F. Approved the amendment of Implementation Report for the Sound Business Plan and estimation of
	Income Statement.
	G. Approved the subscription of the AmMax Bio, Inc. cash capital increase. based on potential
	business collaboration opportunities and financial investment considerations.
	H. Approved the base date of employee stock option into common stocks capital increase.
	I. Approved to grant 2022 Employee Stock Options to employees.
	J. Approved to grant 2023 1st Employee Restricted Stock Awards to employees.
Board	
Meeting	A. Approved the motion of distribution of year-end bonuses for the managers.
2024/01/25	
	A. Accept 2023 Financial Statements and Business Report.
	B. Ratification of the 2023 Deficit Offset Proposal.
	C. Approved the CPAs replacement due to PricewaterhouseCoopers Taiwan internal organization
	adjustment.
	D. Approved the motion of the ratification of the assessment of the independence and competence of
	the CPAs retained as external Auditors.
	E. Approved the motion of the ratification of the appointment of CPAs as external auditors and the
	remuneration to the CPAs.
	F. Approved the motion of issuance of the 2023 Declaration of Internal Control System of the
	Company.
	G. Approved the amendment to the "Rules of Procedure for Shareholders Meetings".
Board	H. Approved the amendment to the "Regulations Governing Procedure for Board of Directors
Meeting	Meetings".
2024/03/08	I. Approved the amendment to the "Audit Committee Charter".
	J. Approved the base date of employee stock option into common stocks capital increase.
	K. Adoption of the 2024 1st Employee Restricted Stock Awards.
	L. Approved the discontinue the Private Security Offering Approved by the 2023 Shareholders'
	Meeting.
	M. Approved the Company will raise capital through private placements of common shares.
	N. Approved the extension to The Shanghai Commercial & Savings Bank for the loan.
	O. Approved the application to Taiwan Business Bank for the 10-year capital expenditure loan and
	total amount within NT\$1.974 billion due to the financial needs of the expansion of Zhubei facility.
	P. Approved to sign the contract with clinical CRO and the relevant companies for Phase III clinical
	trial of the EG1206A.
	Q. Approved the motion of the agenda and related matters of the Shareholders' Meeting of 2024.

- Where, during the most recent fiscal year or during the current fiscal year up to the date of publication of the annual report, a director or supervisor has expressed a dissenting opinion with respect to a material resolution passed by the board of directors and said dissenting opinion had been recorded or prepared as a written declaration, disclose the principal content thereof: None.
- (13) A summary of resignations and dismissals, during the most recent fiscal year or during the current fiscal year up to the date of publication of the annual report, of the company's chairperson, general manager, chief accounting officer, chief financial officer, chief internal auditor, chief corporate governance officer, and chief research and development officer: None.



5.Information Regarding the Company's Audit Fee and Independence

(1) The non-audit fees paid to the certified public accountant, to the accounting firm of the certified public accountant, and/or to any affiliated enterprise of such accounting firm, the amounts of both audit and non-audit fees as well as details of non-audit services:

Unit: NT\$ thousands

Accounting Firm	Name of CPAs	Period Covered by CPA's Audit	Audit Fee	Non-audit Fee	Total
PricewaterhouseCoopers	Sheng-Wei Teng	January 1st,2023 to	3,050	927	3,977
Taiwan.	Yu-Fang Yen	December 31st,2023	3,030	921	3,977

Details of non-audit services:

- Business Income Tax Audit, NT\$350,000.
- ESG Report Assurance, NT\$340,000
- Others, NT\$237,000.
- (2) When the company changes its accounting firm and the audit fees paid for the fiscal year in which such change took place are lower than those for the previous fiscal year, the amounts of the audit fees before and after the change and the reasons: None.
- (3) When the audit fees paid for the current fiscal year are lower than those for the previous fiscal year by 10 percent or more, the reduction in the amount of audit fees, reduction percentage, and reason(s) therefor: None.

6.Replacement of CPA

(1) Regarding the former CPA:

rung the former CIA.	Doto	of received the notification for ren	lagaman	t. Folymory 20th 200	24				
Replacement Date		Date of received the notification for replacement: February 20 th , 2024 Date of approval by Board of Directors: March 8 th , 2024							
				,	irron intomol				
Replacement reasons and explanations		CPAs replacement due to Prization adjustment.	ncewate	mouseCoopers 1a	iwan internal				
Describe whether the	Stati	IS	Parties	CPA	The Company				
Company terminated, or the	Tern	ination of appointment		-	-				
CPA did not accept the	No le	onger accepted (continued)		-	-				
appointment		intment							
Other issues (except for unqua years	lified i	ssues) in the audit reports within th	e last two	None.					
		Accounting principles or practices							
		Disclosure of Financial Statements							
Differences with the	Y	Audit scope or steps							
company		Others							
	N	N 🗸							
	Rem	Remarks/specify details: None							
Other Revealed Matters				None.					

(2) Regarding the successor CPA:

0		
	Accounting Firm	PricewaterhouseCoopers Taiwan
	Name of CPA	Shu-Fen Yu and Yu-Fang Yen
	Date of appointment	Date of received the notification for replacement: February 20 th , 2024 Date of approval by Board of Directors: March 8 th , 2024



Consultation results and	
opinions on accounting	
treatments or principles with	
respect to specified	Nama
transactions and the	None.
company's financial reports	
that the CPA might issue	
prior to the engagement.	
Succeeding CPA's written	
opinion of disagreement	None.
toward the former CPA	

- (3) The company shall mail to the former certified public accountant a copy of the disclosures it is making pursuant to item A and to (c) of the here preceding item and advise the accountant of the need to respond by mail within 10 days should the accountant disagree. The company shall disclose the content of the reply letter from the former certified public accountant: None.
- 7. The company's chairperson, general manager, or any managerial officer in charge of finance or accounting matters has in the most recent year held a position at the accounting firm of its certified public accountant or at an affiliated enterprise of such accounting firm: None.
- 8. Any transfer of equity interests and/or pledge of or change in equity interests (during the most recent fiscal year or during the current fiscal year up to the date of publication of the annual report) by a director, supervisor, managerial officer, or shareholder with a stake of more than 10 percent during the most recent fiscal year or during the current fiscal year up to the date of publication of the annual report:
 - (1) Net Change in Shareholding by Directors, Management and Shareholders with 10% Shareholdings or More

Unit: Shares

		20	023	2024/01/01~2024/03/31		
Title	Name	Net Change in Shareholding	Net Change in Shares Pledged	Net Change in Shareholding	Net Change in Shares Pledged	
Chairman/ President	Lee-Cheng Liu	181,600	0	(27,500)	0	
D:	National Development Fund, Executive Yuan	0	0	0	0	
Director	Representative: Hsiu-Hui Chen	0	0	0	0	
	Formosa Laboratories, Inc.	(30,000)	(500,000)	0	0	
Director	Representative: Cheng-Yu Cheng	0	0	0	0	
D:	Yao-Hwa Glass Co., Ltd, Management Commission	0	0	0	0	
Director	Representative Ku-Sung Weng	0	0	0	0	
	Foxconn Technology Co., Ltd.	0	0	0	0	
Director	Representative: Chun- Fu Lu	0	0	0	0	
	Representative Yu-Ting Chen	0	0	0	0	
Independent Director	Ming-Thaur Chang	0	0	0	0	
Independent Director	Po-Chih Chen	0	0	0	0	



Unit: Shares

		20	023	2024/01/01	~2024/03/31
Title	Name	Net Change in Shareholding	Net Change in Shares Pledged	Net Change in Shareholding	Net Change in Shares Pledged
Independent Director	Fu-Shiow Yin	0	0	0	0
Independent Director	Ming-Shen Chen	0	0	0	0
Manager	Chih-Jung Chang	(103,354)	0	0	0
Manager	Hsiu-Chuan Yang	37,035	0	0	0
Manager	Shang-Chung Ju	28,511	0	0	0
Manager	Ae-Ning Lin	33,111	0	0	0
Manager	Ching-Ying Chen	18,711	0	0	0
Manager	Ren-Yo Forng	18,711	0	0	0
Manager	Ywan-Feng Li	34,036	0	(21,000)	0
Manager	Tsan-Hui Wu	14,741	14,741 0		0
Manager	Hwei-Rung Wang	10,313	0	0	0
Manager	Chung-Huan Lin	(2,509)	0	0	0
Manager	Yu-Wen Liu	15,991	0	0	0
Manager	Tsung-Chih Wang	15,741	0	0	0
Manager	Ming-Tao Pai	8,491	0	0	0
Manager	Chih-Yuan Ma	24	0	0	0
Manager	Sz-Wei Wu (Note 1)	4,078	0	0	0
Manager	Yi-Yun Ciou (Note 2)	0	0	-	-

Note 1: Sz-Wei Wu was on board on July 10, 2023. Note 2: Yi-Yun Ciou resigned on October 4, 2023.

- (2) Information of Stock Trade: The counterparties of equity transfer are not related parties.
- (3) Information of Stock Pledge: The counterparties of share pledges are not related parties.

9.Relationship information, if among the company's 10 largest shareholders any one is a related party or a relative within the second degree of kinship of another.

April 1, 2024; Unit: Shares; %

Name	Current Shareholding		Spouse's/ minor's Shareholding		Shareholding by Nominee Arrangement		Name and Relationship Between the Company's Top Ten Shareholders, or Spouses or Relatives Within Two Degrees	
	Shares	%	Shares	%	Shares	%	Name	Relationship
Foxconn Technology Co., Ltd.	27,500,000	8.98	0	0	0	0	Yonglin Capital Holding Co., Ltd.	Chairman
Representative: Jun-Fu Lu	0	0	0	0	0	0	N/A	N/A
Yonglin Capital Holding Co., Ltd.	26,500,000	8.65	0	0	0	0	Foxconn Technology Co., Ltd.	Chairman
Representative: Kai-Lin Huang	0	0	0	0	0	0	N/A	N/A



Name	Current Shareholding		Spouse's/ minor's Shareholding		Shareholding by Nominee Arrangement		Name and Relationship Between the Company's Top Ten Shareholders, or Spouses or Relatives Within Two Degrees	
	Shares	%	Shares	%	Shares	%	Name	Relationship
Formosa Laboratories, Inc.	18,552,818	6.06	0	0	0	0	N/A	N/A
Representative: Cheng-Yu Cheng	0	0	0	0	0	0	N/A	N/A
National Development Fund, Executive Yuan	15,288,860	4.99	0	0	0	0	N/A	N/A
Convener: Ming-Hsin Kung, Minister, National Development Council	0	0	0	0	0	0	N/A	N/A
Yao-Hwa Glass Co., Ltd, Management Commission	13,078,082	4.27	0	0	0	0	N/A	N/A
Representative: Chuan-Neng Lin	0	0	0	0	0	0	N/A	N/A
Wen-Ming Pan	11,001,123	3.59	0	0	0	0	N/A	N/A
Taiwania Capital Buffalo II Bioventures, LP	6,970,286	2.28	0	0	0	0	N/A	N/A
Representative: Taiwania Capital Biotechnology Corporation	0	0	0	0	0	0	N/A	N/A
Development Center for Biotechnology	4,506,484	1.47	0	0	0	0	N/A	N/A
Representative: Shiing-Jer Twu	0	0	0	0	0	0	N/A	N/A
CTBC Venture Capital Co., Ltd.	3,781,414	1.23	0	0	0	0	N/A	N/A
Representative: Zhi-Gang Wang	0	0	0	0	0	0	N/A	N/A
Lee-Cheng Liu	2,440,984	0.80	0	0	0	0	N/A	N/A

10. The total number of shares and total equity stake held in any single enterprise by the company, its directors and supervisors, managerial officers, and any companies controlled either directly or indirectly by the company:

Unit: Shares; %

Affiliated Enterprises	Ownership b	y the Company	Direct or Indirect Owr	Total Ownership		
1	Shares	%	Shares	%	Shares	%
EirGenix Europe GmbH	(Note)	100%	0	0	(Note)	100%

Note: As a limited liability company, there are no shares.



IV. Capital Overview

1. Source of Capital

(1) Source of Capital

		Authorized C		Paid-in	Capital	Remark		
Month/ Year	Par Value(NT\$)	Shares	Amount (Unit: NT\$)	Shares	Amount (Unit: NT\$)	Sources of Capital	Capital Increased by Assets Other than Cash	Other
Apr. 2023	10	400,000,000	4,000,000	304,491,454	3,044,915	Exercising employee stock option 274,250 shares Issuing Restricted Stock Awards 59,290 shares Deregistering Restricted Stock Awards 79,004 shares	None	Approval No. 11230051460 dated Apr. 10, 2023
May 2023	10	400,000,000	4,000,000	304,667,204	3,046,672	Exercising employee stock option 175,750 shares	None	Approval No. 11230091640 dated May 31, 2023
Sept. 2023	10	400,000,000	4,000,000	304,838,454	3,048,385	Exercising employee stock option 171,250 shares	None	Approval No. 11230166720 dated Sept. 4, 2023
Dec. 2023	10	400,000,000	4,000,000	305,946,124	3,059,461	Exercising employee stock option 71,250 shares Issuing Restricted Stock Awards 324,820 shares Issuing Restricted Stock Awards 825,562 shares Deregistering Restricted Stock Awards 113,962 shares	None	Approval No. 11230226640 dated Dec. 6, 2023
Jan. 2024	10	400,000,000	4,000,000	306,032,124	3,606,321	Exercising employee stock option 60,500 shares Issuing Restricted Stock Awards 25,500 shares	None	Approval No. 11330003230 dated Jan. 16, 2024

(2) Type of Stock

April 1, 2024; Unit: Shares

CI T		Authorized Capital		Remarks		
Share Type Issued Sha	Issued Shares	Un-issued Shares	Un-issued Shares	Remarks		
Common Share	306,231,649	93,768,351		TPEx Listed Stock Private Placement 55,000,000 shares		

(3) Information for Shelf Registration: None.

2. Structure of Shareholders

As of April 1, 2023; Unit: Person; Shares; %

Shareholders' Structure Numbers	Government Agencies	Financial Institutions	Other Juridical Persons	Domestic Natural Persons	Foreign Institutions & Natural Persons	Total
Number of Shareholders	2	1	107	30,802	130	31,042
Shareholding (shares)	16,939,860	1,160,000	118,433,583	155,409,802	14,288,404	306,231,649
Percentage	5.53	0.38	38.67	50.75	4.67	100



3. Shareholding Distribution Status

(1) Shareholding Distribution Status

As of April 1, 2024; Unit: Person; Shares; %

Class of Shareholding	Number of Shareholders	Shareholding	Percentage
1~999	6,399	1,280,067	0.42
1,000~5,000	20,037	37,913,637	12.38
5,001~10,000	2,269	17,408,241	5.68
10,001~15,000	737	9,353,666	3.05
15,001~20,000	430	7,832,441	2.56
20,001~30,000	430	10,794,845	3.53
30,001~40,000	214	7,549,628	2.47
40,001~50,000	119	5,442,974	1.78
50,001~100,000	216	15,350,738	5.01
100,001~200,000	90	12,305,533	4.02
200,001~400,000	46	12,265,755	4.01
400,001~600,000	19	9,296,271	3.04
600,001~800,000	6	4,097,581	1.34
800,001~1,000,000	8	7,085,639	2.31
1,000,001 or over	22	148,254,633	48.40
Total	31,042	306,231,649	100

(2) Preferred Shares: None.

4.List of Major Shareholders

As of April 1, 2024; Unit: Shares; %

115 011 15111 1, 202 1, 011111 01111100, 70			
Shares	Percentage		
27,500,000	8.98		
26,500,000	8.65		
18,552,818	6.06		
15,288,860	4.99		
13,078,082	4.27		
11,001,123	3.59		
6,970,286	2.28		
4,506,484	1.47		
3,781,414	1.23		
2,440,984	0.80		
	Shares 27,500,000 26,500,000 18,552,818 15,288,860 13,078,082 11,001,123 6,970,286 4,506,484 3,781,414		



5. Share prices for the Past 2 Fiscal Years, together with the Company's Net Worth Per Share, Earnings Per Share, Dividends Per Share, and Related Information

Unit: NT\$; Thousands of Shares

Item		Year	2022	2023	2024 Until Mar. 31 st
Market	Н	ighest Market Price	158.5	134.5	103
Price per	L	owest Market Price	71.6	90.5	85.2
Share	A	verage Market Price	120.54	114.43	92.78
Net Worth	I	Before Distribution	35.10	32.73	-
per Share			35.10	32.73	-
Earnings	Weighted Average Shares		303,258	304,888	-
per Share	Dilu	Diluted Earnings Per Share		(3.00)	-
		Cash Dividends	0	0	-
Dividends	Stock	Dividends from Retained Earnings	0	0	-
per Share	Dividend Distribution	Dividends from Capital Surplus	0	0	-
	Accumulated Undistributed Dividends		-	-	-
	Price / Earnings Ratio (Note)		N/A	N/A	-
Return on Investment	Pr	ice / Dividend Ratio	N/A	N/A	-
III v estillellt	Casl	h Dividend Yield Rate	N/A	N/A	-

6. Dividend Policy and Implementation Status

(1) The Dividend Policy Adopted in the Company's Articles of Incorporation

In accordance with Article 25 and Article 25-1 of EirGenix's Articles of Incorporation.

"Article 25: If the Company has net profit in this fiscal year, the Company shall set aside between 1% to 5% of its profits as a bonus to employees of the Company and set aside 3% (inclusive) or less of its profits as a bonus to Directors. The distribution of bonuses to employees may be made by way of cash or shares by the resolution of the Board of Directors. The employees may include certain employees of the subsidiaries who meet the conditions prescribed by the Company. The distribution of employee remuneration and director remuneration shall be heard by over two-thirds of the Board of Directors, be voted in favor for implementation by over one-half of the directors present and represented, and also be reported at the shareholders' meeting. The Company shall first offset its losses in previous years that have not been previously offset, and then set aside annual profits as a bonus to employees of the Company and set aside annual profits as a bonus to Directors. "

"Article 25-1: If the Company has earnings at the end of the fiscal year, the Company shall first pay all relevant taxes, offset its losses in previous years, and set aside a legal capital reserve at 10% of the net profit, until the accumulated paidin capital reserve has equaled the total capital of the Company; then set aside a special capital reserve in accordance with relevant laws or regulations or as requested by the authorities in charge. The board of directors may propose the distribution for approval in the shareholders' meeting. The company has the surplus profit distributable as dividends and bonuses to shareholders of no less than 50% of its net income and shall be a resolution adopted by a majority of the shareholders present who represent two-thirds or more of the total number of its outstanding shares of the company. As the Company is in the growing stage, the dividend distribution may take the form of a cash dividend and/or stock dividends and shall take into consideration the Company's capital expenditures, R&D plan, future expansion plans, and financial structure and funds requirement for sustainable development needs, etc. The cash dividends may not be less than 10% of the total dividend amount. However, the actual distribution ratio is still subject to the resolution of the shareholders meeting."

(2) The Dividend Distributions Proposed at the Shareholders' Meeting

The proposal for distribution was passed at the Meeting of the Board of Directors. In this proposal, due to net losses of 2023, none of the cash dividend and none of the stock dividend will be discussed at the annual shareholders' meeting.



(3) If a material change in dividend policy is expected, provide an explanation: None.

7. Effect upon business performance and earnings per share of any stock dividend distribution proposed or adopted at the most recent shareholders' meeting: None.

8. Compensation of employees, directors, and supervisors

(1) The percentages or ranges with respect to employee, director, and supervisor compensation, as set forth in the company's articles of incorporation.

Please refer to 6. Dividend Policy and Implementation Status.

(2) The basis for estimating the amount of employee, director, and supervisor compensation, for calculating the number of shares to be distributed as employee compensation, and the accounting treatment of the discrepancy, if any, between the actual distributed amount and the estimated figure, for the current period:

The amount of payment in the past will be taken as the foundation for the estimation of the amount.

If there is a significant change in the amount resolved by the Board to pay after the end of the fiscal year, the amount of change will be recognized as the expense of the year.

If there are still further changes at the time of the decision of the Shareholders Meeting, proceed to the accounting principle of change and enter as adjustment of the year under the resolution of the Shareholders Meeting.

(3) Information on any approval by the board of directors of the distribution of compensation:

None.

- A. The amount of any employee compensation distributed in cash or stocks and compensation for directors and supervisors. If there is any discrepancy between that amount and the estimated figure for the fiscal year these expenses are recognized, the discrepancy, its cause, and the status of treatment shall be disclosed: None.
- B. The amount of any employee compensation distributed in stocks, and the size of that amount as a percentage of the sum of the after-tax net income stated in the parent company only financial reports or individual financial reports for the current period and total employee compensation: None.
- (4) The actual distribution of employee, director, and supervisor compensation for the previous fiscal year (with an indication of the number of shares, monetary amount, and stock price, of the shares distributed), and, if there is any discrepancy between the actual distribution and the recognized employee, director, or supervisor compensation, additionally the discrepancy, cause, and how it is treated: None.

9. Status of a Company Repurchasing its own Shares: None.

10. Corporate bond

- (1) Status of Corporate bond: None.
- (2) Information of Convertible Bond: None.
- (3) Exchangeable Bond: None.
- (4) Shelf Registration: None.
- (5) Bond with Warrants: None.

11. Preferred Shares: None.

12. Global Depository Receipts: None.



13. Employee Share Subscription Warrants

(1)Status of Employee Share Subscription Warrants (as of March 31, 2024)

Type of Stock Option	2014 2nd Employee Share Subscription Warrants					
Regulatory approval date and units issued	2016/07/19 2,100					
Issue date	2015/07/01 2015/07/01 2015/07/06					
Units issued	1,270	130	80			
Number of shares still available for issuance	Each unit	can subscribe 1,000 comm	on shares.			
Option shares to be issued as a percentage of outstanding shares	0.42 %	0.04 %	0.03 %			
Exercising Period	2016/07/01 ~ 2025/06/30	2016/07/06 ~ 2025/07/05				
Conversion measures		Issue new common shares				
Conditional conversion periods and percentages	For every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.					
Converted shares	982,250 shares	92,500 shares	80,000 shares			
Exercised amount	NT\$ 14,733,750	NT\$ 1,850,000	NT\$ 1,600,000			
Number of shares yet to be converted	48,250 shares	5,000 shares	0 share			
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 15 NT\$ 20 -					
Unexercised shares as a percentage of total issued shares	0.02% -					
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.					

Type of Stock Option	2014 2nd Employee Share Subscription Warrants				
Regulatory approval date and units issued	2016/07/19 2,100				
Issue date	2015/07/15 2015/07/19 2015/07/26 2015/08/17				
TT '	10	30	20	10	
Units issued	Eac	h unit can subscribe	1,000 common share	es.	
Number of shares still available for issuance	0				
Option shares to be issued as a percentage of outstanding shares	0.003%	0.01%	0.01%	0.003%	
Exercising Period	2016/07/15 ~2025/07/14	2016/07/19~ 2025/07/18	2016/07/26~ 2025/07/25	2016/08/17~ 2025/08/16	
Conversion measures		Issue new con	nmon shares.		
Conditional conversion periods and percentages	For every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.				
Converted shares	10,000 shares	15,000 shares	10,000 shares	1,875 shares	
Exercised amount	NT\$ 200,000	NT\$ 300,000	NT\$ 200,000	NT\$ 37,500	
Number of shares yet to be converted	0 share	0 share	10,000 shares	0 share	



Adjusted exercise price for those					
who have yet to exercise their	-	-	NT\$ 20	-	
rights					
Unexercised shares as a			0.003%		
percentage of total issued shares	-	-	0.00370	-	
Impact on possible dilution of	The issuance of employee stock option certificate may create the common				
Impact on possible dilution of shareholdings	interests of the Co	ders, which shall be p	ositive effects to		
snareholdings	the rights of shareholders.				

Type of Stock Option	2014 2nd Employee Share Subscription Warrants					
Regulatory approval date and		2016/				
units issued		2,1				
Issue date	2015/08/20	2015/08/31	2015/09/29	2015/11/10		
Units issued	20	60	20	30		
Cliffs Issued	Eacl	n unit can subscribe	1,000 common sha	ires.		
Number of shares still available for issuance		()			
Option shares to be issued as a percentage of outstanding shares	0.01%	0.02%	0.01 %	0.01%		
Exercising Period	2016/08/20 ~2025/08/19	2016/08/31~ 2025/08/30	2016/09/29~ 2025/09/28	2016/11/10~ 2025/11/09		
Conversion measures		Issue new con	nmon shares.			
Conditional conversion periods and percentages	su	bscription proportion	ne accumulated max on will increase 25% n be exercised after	6.		
Converted shares	20,000 shares	40,000 shares	10,000 shares	22,500 shares		
Exercised amount	NT\$ 400,000	NT\$ 800,000	NT\$ 200,000	NT\$ 450,000		
Number of shares yet to be converted	0 share	0 shares	0 share	0 share		
Adjusted exercise price for those who have yet to exercise their rights			-	-		
Unexercised shares as a percentage of total issued shares	-	-	-	-		
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.					

Type of Stock Option	2014 2nd Employee Share Subscription Warrants					
Regulatory approval date and		2016/7/19				
units issued	2,100					
Issue date	2015/12/01 2015/12/14 2015/12/21					
Their issued	5	20	25			
Units issued	Each unit c	an subscribe 1,000 comm	on shares.			
Number of shares still available for issuance	0					
Option shares to be issued as a percentage of outstanding shares	0.002% 0.01% 0.01%					
Exercising Period	2016/12/01~ 2025/11/30	2016/12/14~ 2025/12/13	2016/12/21~ 2025/12/20			
Conversion measures	I	ssue new common shares.				
Conditional conversion periods and percentages	For every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.					
Converted shares	5,000 shares 0 share 25,000 shares					
Exercised amount	NT\$ 100,000	NT\$ 0	NT\$ 500,000			



Number of shares yet to be converted	0 share	0 share	0 share		
Adjusted exercise price for those who have yet to exercise their rights	-	-	-		
Unexercised shares as a percentage of total issued shares	-	-	-		
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.				

Type of Stock Option	2014 2nd Employee Share Subscription Warrants						
Regulatory approval date and units issued	2016/07/19 2,100						
Issue date	2016/01/01						
	30	10	15	25			
Units issued	Eac	ch unit can subscribe	1,000 common share	es.			
Number of shares still available for issuance		0	1				
Option shares to be issued as a percentage of outstanding shares	0.01 %	0.003 %	0.005 %	0.01 %			
Exercising Period	2017/01/01~ 2025/12/31	2017/01/12~ 2026/01/11	2017/01/13~ 2026/01/12	2017/02/14~ 2026/02/13			
Conversion measures		Issue new con	nmon shares.				
Conditional conversion periods and percentages	S	ubscription proportion	ne accumulated maxing on will increase 25%. In the exercised after 4				
Converted shares	3,750 shares	5,000 shares	15,000 shares	18,750 shares			
Exercised amount	NT\$ 75,000	NT\$ 100,000	NT\$ 300,000	NT\$ 375,000			
Number of shares yet to be converted	0 share	0 share	0 share	0 share			
Adjusted exercise price for those who have yet to exercise their rights	-	-	-	-			
Unexercised shares as a percentage of total issued shares							
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.						

Type of Stock Option	2014 2nd Employee Share Subscription Warrants			
Regulatory approval date and	2016/07/19			
units issued	2,100			
Issue date	2016/03/1	2016/03/14		
TT: '4: ' 1	150	15		
Units issued	Each unit can subscribe 1,000 common shares.			
Number of shares still		0		
available for issuance		U		
Option shares to be issued as a				
percentage of	0.05%	0.005 %		
outstanding shares				
Exercising Period	2017/03/01~2026/02/28	2017/03/09~2026/03/08	2017/03/14~2026/03/13	
Conversion measures		Issue new common shares.		
Conditional conversion periods	For every expiration	For every expiration of one year, the accumulated maximum exercise		
and	subscrip	subscription proportion will increase 25%.		
percentages	100% subscription right can be exercised after 4 years.			
Converted shares	37,500 shares	0 share	15,000 shares	



Exercised amount	NT\$ 750,000	-	NT\$ 300,000	
Number of shares yet to be converted	0 share	25,000 shares	0 share	
Adjusted exercise price for				
those who have yet to exercise	-	NT\$ 20	-	
their rights				
Unexercised shares as a		0.008%		
percentage of total issued shares	-	0.00870	-	
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.			

Type of Stock Option	2016 1st Employee Share Subscription Warrants		
Regulatory approval date and units issued	2016/07/19 2,100		
Issue date	2016/05/05	2016/06/01	
TT '. ' 1	45	55	
Units issued	Each unit can subscribe	1,000 common shares.	
Number of shares still available for issuance	()	
Option shares to be issued as a percentage of outstanding shares	0.01%	0.02%	
Exercising Period	2018/05/05~2026/05/04	2018/06/01~2026/05/31	
Conversion measures	Issue new common shares.		
Conditional conversion periods and percentages	50% subscription right can be exercised after 2 years. After 2 years, for every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.		
Converted shares	35,000 shares	40,000 shares	
Exercised amount	NT\$ 1,099,000	NT\$ 1,264,000	
Number of shares yet to be converted	10,000 shares	0 share	
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 29.2	-	
Unexercised shares as a percentage of total issued shares	0.003%		
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.		

Type of Stock Option	2016 2nd Employee Share Subscription Warrants		
Regulatory approval date and	2016/08/30		
units issued	600		
Issue date	2016/10/12	2016/12/29	
Units issued	515	85	
Units issued	Each unit can subscribe 1,000 common shares.		
Number of shares still			
available for issuance	0		
Option shares to be issued as a			
percentage of	0.17 %	0.03 %	
outstanding shares			
Exercising Period	2018/10/12~2026/10/11 2018/12/29~2026/12/28		
Conversion measures	Issue new common shares.		
Conditional conversion periods	50% subscription right can be exercised after 2 years.		
and			



percentages	After 2 years, for every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.			
Converted shares	241,250 shares 40,000 shares			
Exercised amount	NT\$ 7,401,750	NT\$ 1,578,675		
Number of shares yet to be converted	150,000 shares	15,000 shares		
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 29.2	NT\$ 37.5		
Unexercised shares as a percentage of total issued shares	0.05 %	0.005 %		
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.			

Type of Stock Option	2017 1st Employee Share Subscription Warrants			
Regulatory approval date and units issued	2017/05/10 1,700			
Issue date	2017/08/08 2017/12/27 2018/03/2			
** '. ' 1	395	570	175	
Units issued	Each unit	can subscribe 1,000 commo	on shares.	
Number of shares still available for issuance		0		
Option shares to be issued as a percentage of outstanding shares	0.13 %	0.19 %	0.06 %	
Exercising Period	2019/08/08~2027/08/07	2019/12/27~2027/12/26	2020/03/23~2028/03/22	
Conversion measures	Issue new common shares.			
Conditional conversion periods and percentages	50% subscription right can be exercised after 2 years. After 2 years, for every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.			
Converted shares	235,000 shares	330,250 shares	62,000 shares	
Exercised amount	NT\$ 7,161,000	NT\$ 8,620,300	NT\$ 1,516,700	
Number of shares yet to be converted	0 shares	58,500 shares	48,000 shares	
Adjusted exercise price for those who have yet to exercise their rights	-	NT\$ 25	NT\$ 23.5	
Unexercised shares as a percentage of total issued shares	-	0.02 %	0.02 %	
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.			

Type of Stock Option	2017 1st Employee Share Subscription Warrants		
Regulatory approval date and	2018/08/09		
units issued	1,50	00	
Issue date	2019/01/25 2019/05/13		
TT '. ' 1	520	285	
Units issued	Each unit can subscribe 1,000 common shares.		
Number of shares still available for issuance	0		
Option shares to be issued as a percentage of outstanding shares	0.17 %	0.09 %	



Exercising Period	2021/01/25~2029/01/24	2021/05/13~2029/05/12	
Conversion measures	Issue new common shares.		
Conditional conversion periods and percentages	50% subscription right can be exercised after 2 years. After 2 years, for every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.		
Converted shares	275,275 shares 105,750 shares		
Exercised amount	NT\$ 8,120,393	NT\$ 3,641,475	
Number of shares yet to be converted	28,475 shares	93,750 shares	
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 28.7	NT\$ 34.3	
Unexercised shares as a percentage of total issued shares	0.01 %	0.03 %	
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.		

Type of Stock Option	2019 1st Employee Share Subscription Warrants				
Regulatory approval date and units issued	2019/10/29 2,000				
Issue date	2019/11/12 2020/04/15 2020/08/12				
TT '. ' 1	960	775	205		
Units issued	Each unit	can subscribe 1,000 commo	n shares.		
Number of shares still available for issuance		0			
Option shares to be issued as a percentage of outstanding shares	0.32 %	0.26 %	0.07 %		
Exercising Period	2021/11/12~2029/11/11	2022/04/15~2030/04/14	2022/08/12~2030/08/11		
Conversion measures		Issue new common shares.			
Conditional conversion periods and percentages	50% subscription right can be exercised after 2 years. After 2 years, for every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.				
Converted shares	363,750 shares	267,500 shares	96,250 shares		
Exercised amount	NT\$ 9,166,500	NT\$ 7,704,000	NT\$ 4,928,000		
Number of shares yet to be converted	162,500 shares				
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 25.2 NT\$ 28.8 NT\$ 51.2				
Unexercised shares as a percentage of total issued shares	0.05 % 0.03 % 0.02 %				
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.				

Type of Stock Option	2020 1st Employee Share Subscription Warrants			
Regulatory approval date and	2020/11/06			
units issued		3,000		
Issue date	2020/12/23 2021/05/12 2021/08/12 2021/10/01			
TI.:4. : 1	830	315	505	1,185
Units issued	Each unit can subscribe 1,000 common shares.			es.
Number of shares still	0			
available for issuance		U		



Option shares to be issued as a percentage of outstanding shares	0.27 %	0.10 %	0.17 %	0.39 %
Exercising Period	2022/12/23~ 2030/12/22	2023/05/12~ 2031/05/11	2023/08/12~ 2031/08/11	2023/10/01~ 2031/09/30
Conversion measures		Issue new common shares.		
Conditional conversion periods and percentages	50% subscription right can be exercised after 2 years. After 2 years, for every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.			llated maximum 25%.
Converted shares	257,750 shares	0 share	0 share	0 share
Exercised amount	NT\$ 10,851,275	NT\$ 0	NT\$ 0	NT\$ 0
Number of shares yet to be converted	244,750hares	215,000 shares	250,000 shares	700,000 shares
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 42.1	NT\$ 146.4	NT\$ 128.4	NT\$ 117.5
Unexercised shares as a percentage of total issued shares	0.08 %	0.07%	0.08%	0.23%
Impact on possible dilution of shareholdings		mployee stock option mpany and sharehold the rights of s	lers, which shall be p	

Type of Stock Option	2021 1st Employee Share Subscription Warrants			
Regulatory approval date and	2021/10/15			
units issued	3,000			
Issue date	2022/03/22	2022/05/12	2022/08/11	2022/09/08
Units issued	160	225	685	510
Onits issued	Eac	ch unit can subscribe	e 1,000 common shar	es.
Number of shares still available for issuance		()	
Option shares to be issued as a percentage of outstanding shares	0.05 %	0.07 %	0.22 %	0.17 %
Exercising Period	2024/03/22~ 2032/03/21	2024/05/12~ 2032/05/11	2024/08/11~ 2032/08/10	2024/09/08~ 2032/09/07
Conversion measures		Issue new common shares.		
Conditional conversion periods and percentages	50% subscription right can be exercised after 2 years. After 2 years, for every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.			
Converted shares	0 share	0 share	0 share	0 share
Exercised amount	NT\$ 0	NT\$ 0	NT\$ 0	NT\$ 0
Number of shares yet to be converted	80,000 shares	180,000 shares	440,000 shares	315,000 shares
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 93.5	NT\$ 71.6	NT\$ 85.9	NT\$ 118.5
Unexercised shares as a percentage of total issued shares	0.03 %	0.06%	0.14%	0.10%
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.			

Type of Stock Option	2022 1st Employee Share Subscription Warrants			
Regulatory approval date and	2022/09/06			
units issued	4,000			
Issue date	2022/11/08	2023/03/10		



Units issued	615	1,105	
	Each unit can subscribe	1,000 common shares.	
Number of shares still available for issuance Option shares to be issued as a	2,28	80	
percentage of outstanding shares	0.20 %	0.36 %	
Exercising Period	2024/11/08~2032/11/07	2025/03/10~2033/03/09	
Conversion measures	Issue new com	nmon shares.	
Conditional conversion periods and percentages	50% subscription right can After 2 years, for every expiration of o exercise subscription propo 100% subscription right can	one year, the accumulated maximum ortion will increase 25%.	
Converted shares	0 share	0 share	
Exercised amount	NT\$ 0	NT\$ 0	
Number of shares yet to be converted	460,000 shares	935,000 shares	
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 103.5	NT\$ 111.5	
Unexercised shares as a percentage of total issued shares	0.15%	0.31 %	
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to rights of shareholders.		
Type of Stock Option	2022 1st Employee Share	Subscription Warrants	
Regulatory approval date and units issued	2022/0	09/06	
Issue date	2023/05/10	2023/08/08	
Units issued	255	225	
Units issued	Each unit can subscribe	1,000 common shares.	
Number of shares still available for issuance	1,80	00	
Option shares to be issued as a percentage of outstanding shares	0.08%	0.07 %	
Exercising Period	2025/05/10~2033/05/09	2025/08/08~2033/08/07	
Conversion measures	Issue new com	nmon shares.	
Conditional conversion periods and percentages	50% subscription right can After 2 years, for every expiration of a exercise subscription proposition of the can be a subscription o	one year, the accumulated maximum ortion will increase 25%.	
Converted shares	0 share	0 share	
Exercised amount	NT\$ 0	NT\$ 0	
Number of shares yet to be converted	255,000 shares	225,000 shares	
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 120	NT\$ 101.5	
Unexercised shares as a percentage of total issued shares	0.08 %	0.08 %	
Impact on possible dilution of shareholdings	The issuance of employee stock option interests of the Company and shareholder rights of shareholder rights of shareholders.	rs, which shall be positive effects to the	



Type of Stock Option	2022 1st Employee Share Subscription Warrants
Regulatory approval date and	2022/09/06
units issued	4,000
Issue date	2023/12/22
** ** *	270
Units issued	Each unit can subscribe 1,000 common shares.
Number of shares still available for issuance	1,530
Option shares to be issued as a percentage of outstanding shares	0.09 %
Exercising Period	2025/12/22~2033/12/21
Conversion measures	Issue new common shares.
Conditional conversion periods and percentages	50% subscription right can be exercised after 2 years. After 2 years, for every expiration of one year, the accumulated maximum exercise subscription proportion will increase 25%. 100% subscription right can be exercised after 4 years.
Converted shares	0 share
Exercised amount	NT\$ 0
Number of shares yet to be converted	245,000 shares
Adjusted exercise price for those who have yet to exercise their rights	NT\$ 100.5
Unexercised shares as a percentage of total issued shares	0.08 %
Impact on possible dilution of shareholdings	The issuance of employee stock option certificate may create the common interests of the Company and shareholders, which shall be positive effects to the rights of shareholders.



(2) The annual report shall disclose the names of top-level company executives holding employee share subscription warrants and the cumulative number of such warrants exercised by said executives as of the date of publication of the annual report. The annual report shall also disclose the names of the ten employees holding employee subscription warrants authorizing purchase of the most shares, along with the cumulative number of warrants exercised by these ten employees, as of the date of publication of the annual report.

March 31st, 2024

				Option			Exercised				Unexercised	141011 31 , 2021	
	Title	Name	No. of Option Shares	Shares as a Percentage of Shares Issued	No. of Shares Converted	Strike Price (NT\$)	Amount (NT\$)	Converted Shares as a Percentage of Shares issued	No. of Shares Converted	Strike Price (NT\$)	Amount (NT\$)	Converted Shares as a Percentage of Shares issued	
	President	Lee-Cheng Liu											
	Senior Vice President	Chih-Jung Chang									NT\$ 74,343,000		
	Vice President	Hsiu-Chuan Yang	_										
	Executive Director	Ae-Ning Lin											
	Executive Director	Shang-Chung Ju								\$25.2		0.27%	
-	Executive Director	Ching-Ying Chen								\$29.2			
/Iana	Executive Director	Ren-Yo Forng				1,641,000 \$10.2 \$15 \$42.1				\$34.3 \$42.1			
Management	Executive Director	Ywan-Feng Li	2,466,000	0.81%	0.81% 1,641,000 shares \$		NT\$ 17,728,000	0.54%		\$100.5			
ent	Senior Director	Tsan-Hui Wu	shares										
	Senior Director	Hwei-Rung Wang											
	Director	Chung-Huan Lin	_										
	Director	Yu-Wen Liu											
	Director	Tsung-Chih Wang											
	Director	Ming-Tao Pai											
	Director	Chih-Yuan Ma											
	Director	Sz-Wei Wu											
	President of EirGenix Europe GmbH	Thomas Schulze											
	Executive Director of EirGenix Europe GmbH	Barbara Grohmann- Izay									\$15		
	Associate Director	Chien-Hao Chen								\$23.5 \$25.2 \$28.8			
Staff	Associate Director	Wan-Ting Hsieh	590,000		210,000	\$15			272.000				
ff	Associate Director	Chia-Hsin Hsiao	shares	0.19%	218,000 shares	\$25.2 \$28.8	NT\$ 4,794,000	0.07%	373,000 shares	\$42.1 \$111.5	NT\$ 32,353,000	0.12%	
	Associate Director	Ching-Cheng Hsiao				\$42.1				\$117.5			
	Associate Director	Chia-Fang Lin								\$120.0 \$128.4			
	Senior Project Manager	Ryan Lee								V120.1			
	Senior Project Manager	Yen-Ming Peng											
	Senior Manager	An-Chi Fan											



14. Restricted Employee Share

(1) Status of Restricted Employee Share (as of March 31, 2024)

Type of Stock Option	1st Employee Restricted Stock in 2016				
Regulatory approval date					
and shares issued	2,000,000 shares 2016/11/18 2016/08/08				
Issue date					
Units issued Number of shares still	1,659,500 shares	257,500 shares			
available for issuance	0 share				
Strike price	NT\$ 0				
Restricted Stock Awards shares to be issued as a percentage of outstanding shares	0.54 % 0.08 %				
Conditional conversion periods and percentages	Condition A: Company operation performance and employee personal KPI. Achieve a positive income before tax for three quarters, and employee's average KPI shall be over 2.66 for three consecutive years. 33.3% of total shares will be released. Condition B: Employee job tenure and employee personal KPI. Work in EirGenix for ten years, and employee personal average KPI shall be over 2.66 for three consecutive years. 16.7% of total shares will be released. Condition C: Development of biosimilar EG12014 and employee personal KPI Timing I: Complete EG12014 Phase 3 and the employee's average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Timing II: EG12014 Launched to market and employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Condition D: Development of biosimilar EG1206A and employee personal KPI Timing I: Complete EG1206A Phase 1, and employee personal average KPI shall be over 2.66 for three consecutive years. 5.6% of total shares will be released. Timing II: Complete EG1206A out-licensing or Phase 3 and employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Condition E: New plant in Zhubei start running and completes 1,000L or 2*2,000L scale process validation and employee personal KPI. The new plant in Zhubei starts running and completes 1,000L or 2*2,000L scale process validation, and the employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Condition F: Complete IPO in TPEx and employee personal API Complete IPO in TPEx and employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released.				
Restricted Conditions	Please refer to the following table for details.				
Depository methods of new shares	Please refer to the following table for details.				
Handling of an employee's failure to meet the vesting conditions	Please refer to the following table for details.				
Bought-back or canceled new shares of Restricted Stock Awards	654,750 shares	172,000 shares			
Shares of Unrestricted Stock Awards	551,550 shares	39,975 shares			
New shares of Restricted Stock Awards	453,200 shares	45,525 shares			
Percentage of new shares of Restricted Stock Awards to Total Issued Shares (%)	0.15 %				



Impacts on Shareholders' The ratio accounted for by the new shares with restricted rights as above. There is no major Equity impact to the existing shareholders of the Company.

Type of Stock Option	1 st Employee Restricted	1 Stock in 2019			
Regulatory approval date and	2019/12/30				
shares issued	600,000 shares				
Issue date	2020/05/13	2020/12/10			
Units issued	454,500 shares	144,000 shares			
Number of shares still available for issuance	0 share				
Strike price	NT\$ 0				
Restricted Stock Awards shares to be issued as a percentage of outstanding shares	0.15 % 0.05%				
Conditional conversion periods and percentages	 Condition A: Calculated from the working day when relocated to Zhubei branch, after serving for 0.25 years, 0.5 years, 0.75 years and 1 year; 750 shares will be released on each timing. Condition B: Calculated from the working day when relocated to Zhubei branch, after serving for 1.5 years, and 2 years; 2,000 shares will be released on each timing. Condition C Calculated from the working day when relocated to Zhubei branch, after serving for 3 years; 5,000 shares will be released on each timing. 				
Restricted Conditions	Please refer to the following table for details.				
Depository methods of new shares	Please refer to the following table for details.				
Handling of an employee's failure to meet the vesting conditions	Please refer to the following table for details.				
Bought-back or canceled new shares of Restricted Stock Awards	35,250 shares	10,500 share			
Shares of Unrestricted Stock Awards	395,250 shares	67,750 shares			
New shares of Restricted Stock Awards	24,000 shares 65,750 shares				
Percentage of new shares of Restricted Stock Awards to Total Issued Shares (%)	0.01 %				
Impacts on Shareholders' Equity	The ratio accounted for by the new shares with restricted rights as above. There is no major impact to the existing shareholders of the Company.				

Type of Stock Option	2 nd Employee Restricted Stock in 2019			
Regulatory approval date and shares issued	2019/12/30 1,000,000 shares			
Issue date	2020/08/14	2020/12/10		
Units issued	905,700 shares	94,200 shares		
Number of shares still available for issuance	0 share			
Strike price	NT\$ 0			
Restricted Stock Awards shares to be issued as a percentage of outstanding shares	0.30 %	0.03%		
Conditional conversion periods and percentages	Condition A: Company operation performance and employee personal KPI. Achieve a positive income before tax for three quarters, and employee's average KPI shall be over 2.66 for three consecutive years. 33.3% of total shares will be released. Condition B: Employee job tenure and employee personal KPI. Work in EirGenix for ten years, and employee personal average KPI shall be over 2.66 for three consecutive years. 16.7% of total shares will be released. Condition C: Development of biosimilar EG12014 and employee personal KPI Timing I: Complete EG12014 Phase 3 and the employee's average KPI shall be			



	over 2.66 for three consec	cutive years. 11.1% of total shares will be			
	released.	curive years. 11.170 or total shares will be			
		narket and employee personal average KPI			
		ree consecutive years. 11.1% of total shares			
	will be released.	·			
	Condition D: Development of biosimilar EG1206	5A and employee personal KPI			
	Timing I: Complete EG1206A Phas	se 1, and employee personal average KPI			
	shall be over 2.66 for three	ee consecutive years. 5.6% of total shares			
	will be released.				
	Timing II: Complete EG1206A out	e-licensing or Phase 3 and employee personal			
	average KPI shall be ov	er 2.66 for three consecutive years. 11.1% of			
	total shares will be relea	sed.			
	Condition E: New plant in Zhubei start running a	•			
	process validation and employee pe				
	_	ing and completes 1,000L or 2*2,000L scale			
		ee personal average KPI shall be over 2.66			
	for three consecutive years. 11.1% of				
	Condition F: Complete IPO in TPEx and employe	•			
	Complete IPO in TPEx and employee personal average KPI shall be over 2.66				
	for three consecutive years. 10% of total shares will be released				
Restricted Conditions	Please refer to the following table for details.				
Depository methods of new	Please refer to the following table for details.				
Handling of an employee's					
	Please refer to the following table for details.				
conditions					
Bought-back or canceled	272 200 1	0.1			
new shares of Restricted	272,200 shares	0 share			
Stock Awards					
Shares of Unrestricted	253,750 shares	37,550 shares			
Stock Awards					
New shares of Restricted Stock Awards	379,750 shares	56,650 shares			
Percentage of new shares					
of Restricted Stock Awards	0.12 %	0.02 %			
to Total Issued Shares (%)		0.02 /0			
` /	The ratio accounted for by the new shares with re	estricted rights as above. There is no major			
Equity	impact to the existing shareholders of the Compa	· ·			
24		<i>y</i> -			

Type of Stock Option	1 st Employee Restricted Stock in 2021					
Regulatory approval date	2021/09/10					
and shares issued	1,000,000 shares					
Issue date	2021/10/15 2022/01/15 2022/09/08					
Units issued	612,500 shares	184,000 shares	190,000 shares			
Number of shares still available for issuance		0 share				
Strike price		NT\$ 0				
Restricted Stock Awards shares to be issued as a						
percentage of outstanding	0.20 %	0.06%	0.06%			
shares						
Conditional conversion periods and percentages	Condition A: Company operation performance and employee personal KPI. Achieve a positive income before tax for three quarters, and employee's average KPI shall be over 2.66 for three consecutive years. 33.3% of total shares will be released. Condition B: Employee job tenure and employee personal KPI. Work in EirGenix for ten years, and employee personal average KPI shall be over 2.66 for three consecutive years. 16.7% of total shares will be released. Condition C: Development of biosimilar EG12014 and employee personal KPI Timing I: Complete EG12014 Phase 3 and the employee's average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Timing II: EG12014 Launched to market and employee personal average					



years. 11.1% of total rsonal KPI personal average KPI					
shall be over 2.66 for three consecutive years. 5.6% of total shares will be released.					
three consecutive					
L or 2*2,000L scale					
,000L or 2*2,000L					
erage KPI shall be					
ares will be released.					
52,500 shares					
0 share					
137,500 shares					
319,750 shares 120,000 shares 137,500 shares					
0.04 %					
0.0 1 /0					
ove. There is no major					
The ratio accounted for by the new shares with restricted rights as above. There is no major impact to the existing shareholders of the Company.					

Type of Stock Option	1 st Employee Restricted Stock in 2022						
Regulatory approval date and shares issued	2022/09/06 850,000 shares						
Issue date	2022/09/08 2022/11/08 2023/03/10 2023/11/09						
Units issued	62,657 shares	195,137 shares	5,929 shares	324,820 shares			
Number of shares still available for issuance		261,457 shares					
Strike price		NT	\$ 0				
Restricted Stock Awards shares to be issued as a percentage of outstanding shares	0.02 %	0.06%	0.002%	0.11%			
Conditional conversion periods and percentages	The employee must remain employed by the Company on the last date of each vesting period. During the vesting period, the employee may not breach any agreement with the company or violate the Company's employment agreement, service agreement, trust agreement, company governance best practice principles, ethical corporate management best practice principles, work rules, non-compete and non-disclosure agreement of the Company or any agreement with the Company. Specific employee performance metrics and the Company's business performance metrics are met in the Employee Restricted Stock Awards Rules. Condition A: Employees on board at, or before the third quarter of 2022, 100% of shares will be vested at the end of same year. Condition B: Employees on board in the fourth quarter of 2022, 100% of shares will be vested at the end of following year. Condition C: Employees on board between the first quarter to third quarter of 2023, 100% of shares will be vested at the end of same year. Condition D: Shares granted for employees 2021 personal performance and company performance rating over 2.5, will be vested 100%. Condition E: Shares granted for employees 2022 company performance rating over 2.0, and						



	will be vested 100%.					
Restricted Conditions]	Please refer to the following table for details.				
Depository methods of new shares]	Please refer to the following table for details.				
Handling of an employee's failure to meet the vesting conditions		Please refer to the following table for details.				
Bought-back or canceled new shares of Restricted Stock Awards	0 share	3,370 shares	0 share	1,284 shares		
Shares of Unrestricted Stock Awards	62,657 shares	191,181 shares	5,929 shares	289,797 shares		
New shares of Restricted Stock Awards	0 share	586 shares	0 share	33,739 shares		
Percentage of new shares of Restricted Stock Awards to Total Issued Shares (%)		0.0002 %	-	0.01%		
Impacts on Shareholders' Equity	The ratio accounted for by the new shares with restricted rights that have not yet been lifted is a mere 0.012%. There is no major impact to the existing shareholders of the Company.					

Type of Stock Option	2 nd Employee Restricted Stock in 2023					
Regulatory approval date	2023/10/27					
and shares issued	870,000 shares					
Issue date	2023/11/09					
Units issued	825,562 shares					
Number of shares still	44.429 shows					
available for issuance	44,438 shares					
Strike price	NT\$ 0					
Restricted Stock Awards						
shares to be issued as a	0.27 %					
percentage of outstanding	0.27 /0					
shares						
Conditional conversion periods and percentages	The employee must remain employed by the Company on the last date of each vesting period. During the vesting period, the employee may not breach any agreement with the company or violate the Company's employment agreement, service agreement, trust agreement, company governance best practice principles, ethical corporate management best practice principles, work rules, non-compete and non-disclosure agreement of the Company or any agreement with the Company. Specific employee performance metrics and the Company's business performance metrics are met in the Employee Restricted Stock Awards Rules. Condition A: When the annual Key-Performance-Indicator (KPI) of the Company is at least 2.5, the employee will be eligible to receive 100% shares of the Company's restricted stock. Condition B: When EirGenix reaches the break-even milestone for a fiscal year based on an audited income statement, the employee will be eligible to receive 100% shares of Company's restricted stock. Condition C: When the Company and employee's annual Key-Performance-Indicator (KPI) is at least 2.5, the employee will be eligible to receive 100% shares of the Company's restricted stock.					
Restricted Conditions	Please refer to the following table for details.					
Depository methods of new shares	Please refer to the following table for details.					
Handling of an employee's failure to meet the vesting conditions	Please refer to the following table for details.					
Bought-back or canceled new shares of Restricted Stock Awards	0 shares					
Shares of Unrestricted Stock Awards	210,562 shares					
210 111 111 111 111	615,000 shares					



Stock Awards	
Percentage of new shares	
of Restricted	0.2 %
Stock Awards to Total	0.2 70
Issued Shares (%)	
Impacts on Shareholders'	The ratio accounted for by the new shares with restricted rights as above. There is no major
Equity	impact to the existing shareholders of the Company.

Regulatory approval date and shares issued as society approval process stated to the state of th	Type of Stock Option	1 st Employee Restricted Stock in 2023						
Issue date Issue date Issue date Inits issued Number of shares still available for issuance Strike price Restricted Stock Awards shares to be issued as a percentage of outstanding shares to be issued as a percentage of outstanding shares to be issued as a percentage of outstanding shares Condition A: Company operation performance and employee personal KPI. Achieve a positive income before tax for three quarters, and employee's average KPI shall be over 2.66 for three consecutive years, 3.3% of total shares will be released. Condition B: Employee go be tenure and employee personal KPI. Work in Effective ten years, and employee personal RPI. Work in Efficients for ten years, and employee personal API. Work in Efficients for ten years, and employee personal RPI. Work in Efficients for ten years, and employee personal API. Work in Efficients for ten years, and employee personal API. Work in Efficients for ten years, and employee personal API. Work in Efficients for ten years, and employee personal API. Work in Efficients for ten years, and employee personal API. Work in Efficients for ten years, and employee personal API. Work in Efficients for ten years, and employee personal API. Work in Efficients for ten years, and employee personal API. Work in Efficients will be released. Condition E: Development of biosimilar EGI 2014 and employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Timing I: EGI 2014 Launched to market and employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Condition E: New plant in Zhubei start running and completes 1,000L or 2*2,000L scale process validation, and the employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Restricted Conditions Depository methods of new shares Handling of an employee personal everage KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will b	• • • • • • • • • • • • • • • • • • • •	• •						
Issue date	• • • • • • • • • • • • • • • • • • • •							
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Restricted Conditions Depository methods of new shares Handling of an employee's failure to meet the vesting conditions Bought-back or canceled new shares of Restricted Stock Awards Shares of Unrestricted Stock Awards New shares of Restricted Stock Awards Please refer to the following table for details. Please refer to the following table for details. O shares O shares 25,500 shares Percentage of new shares O 01 %	Conditional conversion periods and percentages	income before tax for three quarters, and employee's average KPI shall be over 2.66 for three consecutive years. 33.3% of total shares will be released. Condition B: Employee job tenure and employee personal KPI. Work in EirGenix for ten years, and employee personal average KPI shall be over 2.66 for three consecutive years. 16.7% of total shares will be released. Condition C: Development of biosimilar EG12014 and employee personal KPI Timing I: Complete EG12014 Phase 3 and the employee's average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Timing II: EG12014 Launched to market and employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Condition D: Development of biosimilar EG1206A and employee personal KPI Timing I: Complete EG1206A Phase 1, and employee personal average KPI shall be over 2.66 for three consecutive years. 5.6% of total shares will be released. Timing II: Complete EG1206A out-licensing or Phase 3 and employee personal average KPI shall be over 2.66 for three consecutive years. 11.1% of total shares will be released. Condition E: New plant in Zhubei start running and completes 1,000L or 2*2,000L scale process validation and employee personal KPI. The new plant in Zhubei starts running and completes 1,000L or 2*2,000L scale process validation, and the employee personal average KPI shall be over 2.66 for						
Depository methods of new shares Handling of an employee's failure to meet the vesting conditions Bought-back or canceled new shares of Restricted Stock Awards Shares of Unrestricted Stock Awards New shares of Restricted Stock Awards New shares of Restricted Stock Awards Please refer to the following table for details. O shares 0 shares 25,500 shares Percentage of new shares	Restricted Conditions	-						
Handling of an employee's failure to meet the vesting conditions Bought-back or canceled new shares of Restricted Stock Awards Shares of Unrestricted Stock Awards New shares of Restricted Stock Awards New shares of Restricted Stock Awards Percentage of new shares O shares 25,500 shares	Depository methods of							
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Stock Awards New shares of Restricted Stock Awards Percentage of new shares 0 shares 25,500 shares 0 01 %	new shares of							
Stock Awards Percentage of new shares 0.01 %	Stock Awards	0 shares						
0.01 %	Stock Awards	25,500 shares						
OI ICOUITORA	Percentage of new shares of Restricted	0.01 %						



Stock Awards to Total	
Issued Shares (%)	
Impacts on Shareholders'	The ratio accounted for by the new shares with restricted rights as above. There is no major
Equity	impact to the existing shareholders of the Company.

	Employee Restricted Stock
Restricted Conditions	 During the vesting period, the employee may not sell, pledge, transfer, give to another person, create any encumbrance on, or otherwise dispose of, restricted stock awards Voting right in Shareholders' Meeting: The same as common stock. Dividend: The same as common stock.
Depository methods of new shares	The Employee Restricted Stock issued may be deposited in a security trust account.
Handling of an employee's failure to meet the vesting conditions	 No matter of the Voluntary departure from employment, Dismissal from employment, Retirement, Death or Job position transfer to an affiliate, EirGenix shall buy back and cancel all Restricted Stock Awards. Sufferers of disability due to an occupational accident: EirGenix shall buy back and cancel Restricted Stock Awards unless the permission by the Board. Employees will not have to return the stock dividend or cash dividend occurred by forfeited restricted stock awards Employees who have not reached the vesting conditions: Employees who has subscribed but fail to meet the grant conditions, EirGenix shall buy back and cancel Restricted Stock Awards in accordance with the laws.



(2) List of Executives and the Top 10 Employees Receiving Restricted Stock Awards

March 31st, 2024

			NT C	Receiving		Un	restricted			Rest	ricted	Waren 31 , 2024
	Title	Name Received Restriction Sto	No. of Receiving Restricted Stock Shares	Restricted Stock Shares as a Percentage of Shares issued	No. of Shares Converted	Strike Price	Amount	Converted Shares as a Percentage of Shares issued	No. of Shares Converted	Strike Price	Amount	Converted Shares as a Percentage of Shares issued
	President	Lee-Cheng Liu										
	Senior Vice President	Chih-Jung Chang										
	Vice President	Hsiu-Chuan Yang										
	Executive Director	Shang-Chung Ju										
	Executive Director	Ae-Ning Lin										
Z	Executive Director	Ching-Ying Chen										0.32%
anag	Executive Director	Ren-Yo Forng							969,000 shares	NT\$0	NT\$0	
Management	Senior Director	Tsan-Hui Wu	2,271,000 shares	271,000 hares 0.74%	0.74% 1,302,000 shares	NT\$0	NT\$0 NT\$0	0.43%				
nt	Senior Director	Hwei-Rung Wang	_ Shares									
	Director	Chung-Huan Lin										
	Director	Yu-Wen Liu										
	Director	Tsung-Chih Wang										
	Director	Ming-Tao Pai										
	Director	Chih-Yuan Ma										
	Director	Sz-Wei Wu										
	President of EirGenix Europe GmbH	Thomas Schulze										
	Associate Director	Chia-Hsin Hsiao								NT\$0		0.05%
	Associate Director	Chien-Hao Chen										
S	Senior Manager	Yi-Hsuan Pan										
Staff	Senior Manager	Ying-Chun Chen	348,000	0.11%	202,000	NT\$0	NT\$0	0.01%	146,000 shares		NT\$0	
	Senior Manager	Jui-Chi Lee	shares	21270	shares	2 4 0		3.027.0		- 1240	2 \$ 0	3.02.0
	Senior Manager	Ya-Fen Yang										
	Manager	Chia-Feng Liao										
	Manager	Wen-Yuan Ting										
	Assistant manager	Ya-Han Liu										



15. Issuance of new Shares in Connection with Mergers or Acquisitions or With Acquisitions of Shares of Other Companies: None.

16. The Status of Implementation of Capital Allocation Plans

- (1)Cash Capital Increase in 2020
 - A. Description of the Plan:
 - (A) Date and document reference number for effective registration: Official letter No.1090379952 on January 28th, 2021 for effective registration. Official letter No. 1100134277appoval on March 31st, 2021, to extend the period of the subscription until July 27th, 2021.
 - (B) Total Amount of the Plan: NT\$ 3,202,500,000.
 - (C) Source of Funds: Issuance 35,000,000 new common shares for capital increase. Par value is NT\$ 10 per shares, issuance price is NT\$ 91.5 per share, and the total amount is NT\$ 3,202,500,000.
 - (D) Plan item and Expect implementation progress of fund:

Unit: NT\$ thousands

Total func		Expect implementation progress of Capital Allocation Plans (Note 1)							
Item	Item	2021			2022				
	needed	Q2	Q3	Q4	Q1	Q2	Q3	Q4	
Replenishment of									
working capital	3,202,500	400,000	520,000	550,000	510,000	410,000	410,000	402,500	
(Note 2)									

Note 1: If the fund-raising funds are not in place as expected, then the relevant planned projects will be funded by bank loans, and the borrowed bank loans will be repaid after the fund-raising funds are in place.

Note 2: Replenishment of working capital is used to cover research and development expenditure and funds required for daily operations other than EG62054 and EG12021 (non-HER2).

- (E) Planned benefit: The total amount of capital increase by EirGenix in this plan is NT\$ 3.2025 billion, which will be fully used for replenishment of working capital and R&D expenses except EG62054 and EG12021. Based on the current bank loan interest rate of 1.7970%, it is estimated that EirGenix will save NT\$ 38.366 million in 2021 and NT\$ 57.549 million in interest expense in the future. After the Capital Allocation Plans completes, the plan not only can cover the funds of product developments and daily operations but also can save the interest expenses effectively, improve the financial structure, decrease the dependence on the bank, enhance the flexibility for future capital allocation, and reduce the operation risk. The plan has useful help on the business and operational development of EirGenix.
- (F) Date of entering to MOPS: January 28th, 2021
- (G) Change the content of the plan, the reason for changing, and the benefit of changing: Not Applicable.

B.Implementation status

Unit: NT\$ thousands

Τ.	Implementation status			Advance or delay of plans and the reasons
Item	202	3	Q1	The Company has previously completed
	TT 1.4	Expect	3,202,500	the capital utilization plan for the 2022 Q4,
	Used Amount	Actual	100%	but was delayed in raising capital, to the
Replenishment of		Expect	3,202,500	extent that the use of fund was delayed. As such, an amount of NT\$362,092 thousand
working capital	Actual			was still unconsumed as of December 31,
	Implementation	Actual	100%	2022. Accordingly, the capital utilization plan
				was deferred to 2023 Q1.



A. The impact on shareholders' rights and interests

The fundraising will be completed in the second quarter of 2021, which has no major difference from the expected schedule. The raised funds are used to cover research and development expenses and daily operation funds other than EG62054 and EG12021 (non-HER2), as well as the research and development expenses for investing its products and completing the research and development schedule of each stage of the products, to achieve a sound financial structure, enhance solvency, maintain the working capital requirements needed for its operation and further enhance the competitiveness. In the long run, there should be no significant adverse impact on shareholders' rights and interests.

B. Expect the Implementation Benefit

(A) Replenishment of working capital

The total amount of capital increase by EirGenix in this plan is NT\$ 3.2025 billion, which will be fully used for replenishment of working capital. Through the long-term and stable capital infusion to ensure the company's daily operation. At the same time, it can implement the drug development smoothly and increase the operation scale and value of the company, also can improve the financial structure, and avoid increasing the financing costs to ensure the company operation normally and decrease the operation risk. Based on the current bank loan interest rate of 1.7970%, it is estimated that EirGenix will save NT\$ 38.366 million in 2021 and NT\$ 57.549 million in interest expense in the future.

(B)Improve financial structure:

Unit: NT\$ thousands; %

			Omi: 111 \$\phi\$ thousands, 70
	Year	2020	2021
Item		(Before Capital increase)	(After Capital Increase)
	Current assets	1,494,307	9,070,266
	Total assets	3,835,215	11,440,873
D :	Current liabilities	642,163	703,216
Basic	Total liabilities	1,929,598	1,012,122
Financial	Shareholders' equity	1,905,617	10,428,751
Information	Operation revenue	1,071,838	1,697,359
	Interest expenses	28,500	21,149
	Earnings per share	(5.41)	(0.18)
	Debt Ratio	50.31%	8.85%
Financial	Ratio of long-term capital		
structure	to property, plant and	172.42%	569.09%
	equipment		
G 1	Current ratio	232.70%	1,289.83%
Solvency	Quick ratio	195.19%	1,215.91%

To meet the capital demand for operation, EirGenix conducted this capital increase, with the raised amount of NT\$ 3.2025 billion, which was fully used for replenishment of working capital, to enhance EirGenix's capital, make the financial structure sounder, further reduce EirGenix's operating risks, increase long-term capital stability and enhance market competitiveness. The estimated time of completion for this capital increase is 202Q2, and this capital increase will replenish the working capital. The Ratio of long-term capital to property, plant and equipment will increase from 172.42% to 569.09%; the current ratio and quick ratio increased from 232.70% and 195.19% to 1,289.83% and 1,215.91%; Its financial structure will improve compared with that before the capital increase; can maintain the solvency and the stability of the financial structure at the same time when expanding the scale of operation and the flexibility for future capital allocation will be maintained. If EirGenix had failed to raise funds this time, it would have increased its financial burden. Therefore, the capital increase this time to replenish working capital will help fulfill the operation funs, and its benefits will be reasonable.



(2)Private Placement in 2021

A. Description of the Plan:

- (A) Date and document reference number for effective registration: Official letter No.11001199560 on November 18th, 2021.
- (B) Total Amount of the Plan: NT\$ 5,032,500,000.
- (C) Source of Funds: Issuance 55,000,000 new common shares for capital increase. Par value is NT\$ 10 per shares, issuance price is NT\$ 91.5 per share, and the total amount is NT\$ 5,032,500,000.
- (D) Plan item and Expect implementation progress of fund:

Unit: NT\$

Item	The Usage of funds	Budget Amount
Replenishment of working capital	R&D expenses	3,000,000,000
Building factory	Expansion and building factory	500,000,000
Others	Repay bank loans and replenish horizontal and vertical integration, and other operational funding needs	1,532,500,000

(E) Planned benefit:

- a. To accelerate the product developing efficiency as well as the process of the same drug series to complete the production line. By the comprehensive effect of expanding the market, it can establish EirGenix's unique status in the international biosimilars and CDMO field.
- b. Expand the facility and equipment and increase the production scale to meet the needs of the CDMO business expansion and self-development product commercial operation.
- c. Repay bank loan NT\$316,322,000 and save annual interest expenses roughly about NT\$5,684,000 which calculating under the current EirGenix loan rates of 1.797%. Other unused funds will follow the plan and demonstrate effects continuously.
- d. Co-developed the biosimilar drug TSY0110 (EG12043) of ADC for the treatment of breast cancer with Formosa Pharmaceuticals.
- (F) Date of entering to MOPS: May 4, 2021; May 12, 2022.
- (G) Change the content of the plan, the reason for changing, and the benefit of changing:
 - a. Approved by the board of directors on May 12, 2022, in response to the company's medium and long-term strategic development plan, the adjustments to the capital utilization plan are as follows:

Unit: NT\$

Item	m The Usage of funds	
Replenishment of working capital	R&D expenses	1,016,178,000
Building factory	Expansion and building factory	1,700,000,000
	Repay bank loans and replenish horizontal and vertical integration, and other operational funding needs.	316,322,000
Others	Acquisition or purchase the intangible assets, operation- related assets, and right-of-use assets.	2,000,000,000

- b. Considering the operation and benefits of the company, and avoiding the impact on the shareholder equity, the capital plan and project expenditure situation will be adjusted and changed appropriately according to the needs. It will benefit the company's business and fund while shareholder equity should not be materially affected.
- c. Date of submitting to the shareholders meeting: May 31, 2023.



B.Implementation status

Unit: NT\$

The Usage of funds	Budget Amount	Implementation as of 2024 First Quoter
R&D expenses	1,016,178,000	R&D expenses 397,191,024 and deposit other
•		funds in EirGenix bank accounts.
Expansion and building factory	1,700,000,000	Expansion and building factory 682,214,319 and
Expansion and building factory	1,700,000,000	deposit other funds in EirGenix bank accounts.
Repay bank loans and replenish horizontal and		
vertical integration, and other operational funding	316,322,000	Repay bank loan 316,322,000
needs		
Acquisition or purchase the intangible assets,	2 000 000 000	Acquisition important assets 60,112,501 and
operation- related assets, and right-of-use assets.	2,000,000,000	deposit other funds in EirGenix bank accounts.

C. The impact on shareholders' rights and interests

Boost EirGenix's operating scale, horizontal and vertical integration, and product or market development collaboration, assist EirGenix to improve technology, efficiency, expand the operational scale, and elevate the market status. It has positive benefits in creating EirGenix and shareholder value.

D. Expect the Implementation Benefit

- (A) EirGenix is currently developing the product for the treatment of HER2+ breast cancer. Received the approval letter from Ministry of Health and Welfare for the biosimilar drug EIRGASUN 150 mg powder for concentrate for infusion. Received the approval letter from EC for EG12014 licensed. Preparing the US BLA resubmission for EG12014. The Phase I clinical trial for EG1206A (Pertuzumab Biosimilar) has been completed.
- (B) The second mammalian cell production line for the Zhubei plant phase I facility has been completed. Build microbial cell production line factory for the Zhubei plant phase II facility. The three-stage expansion of the mammalian plant which has 150,000L capacity, is under planning at Ciaotou Science Park, Kaohsiung.
- (C) Repay bank loan NT\$316,322,000 and save annual interest expenses roughly about NT\$5,684,000 which calculating under the current EirGenix loan rates of 1.797%. Other unused funds will follow the plan and demonstrate effects continuously.
- (D) Co-developed the biosimilar drug TSY0110 (EG12043) of ADC for the treatment of breast cancer with Formosa Pharmaceuticals.

(E) Improve financial structure:

Unit: NT\$ thousands; %

Year		2021 Q3	2021 Q4		
Item		(Before Capital increase)	(After Capital Increase)		
Basic Financial Information	Current assets	4,451,420	9,070,266		
	Total assets	6,804,041	11,440,873		
	Current liabilities	862,482	703,216		
	Total liabilities	1,463,501	1,012,122		
	Shareholders' equity	5,340,540	10,428,751		
	Operation revenue	1,273,814	1,697,359		
	Operating cost	401,661	604,305		
	Operating profit	872,153	1,093,054		
	Interest expenses	17,161	21,149		
	Earnings per share	(0.20)	(0.18)		
Financial structure	Debt Ratio	21.51%	8.85%		
	Ratio of long-term capital to property, plant and equipment	318.70%	569.09%		



	Year	2021 Q3	2021 Q4
Item		(Before Capital increase)	(After Capital Increase)
Solvency	Current ratio	516.12%	1,289.83%
	Quick ratio	461.55%	1,215.91%

To meet the capital demand for operation, EirGenix conducted this capital increase, with the raised amount of NT\$ 5.0325 billion, which was fully used for the replenishment of working capital. The private placement can enhance equity capital, make the financial structure sounder, further reduce EirGenix's operating risks, increase long-term capital stability and enhance market competitiveness. The Ratio of long-term capital to property, plant and equipment in 2021 increased from 318.70% to 569.09%; the current ratio and quick ratio increased from 516.12% and 461.55% to 1,289.83% and 1,215.91%; the financial structure improved compared with that before the capital increase; can maintain the solvency and the stability of the financial structure at the same time when expanding the scale of operation and the flexibility for future capital allocation will be maintained. Therefore, the capital increases this time to replenish working capital will help fulfill the operation funds, and its benefits will be reasonable.



V. Operational Highlights

1. Business Activities

(1) Business Scope

- A. Main areas of business operation
 - ①C199990 Other Food Manufacturing Not Elsewhere Classified
 - ②C802041 Drugs and Medicines Manufacturing
 - (3)C802060 Animal Use Medicine Manufacturing
 - **4**C802990 Other Chemical Products Manufacturing
 - (5)F107990 Wholesale of Other Chemical Products
 - ⑥F108021 Wholesale of Drugs and Medicines
 - 7F108031 Wholesale of Drugs, Medical Goods
 - **®F208021** Retail Sale of Drugs and Medicines
 - 9F208031 Retail sale of Medical Equipments
 - (10)F401010 International Trade
 - 11)I199990 Other Consultancy
 - (12)IC01010 Pharmaceuticals Examining Services
 - (13) IG01010 Biotechnology Services
 - (14)IG02010 Research Development Service
 - (15)ZZ99999All business items that are not prohibited or restricted by law, except those that are subject to special approval.

EirGenix is a R&D company for biosimilars and new drugs, provides the biopharmaceutical CDMO (Contract Development & Manufacturing Organization) services, including cell line building platform, process development platform, analytical science, protein identification in PIC/S GMP manufacturing plants, and provides production of drugs for clinical trials, etc.

EirGenix employs a dual-track approach encompassing bio-pharmaceutical CDMO and Product Development, effectively leveraging the company's cGMP production equipment and high-caliber technical expertise. Our company's core competitive advantage lies in simultaneously possessing expertise in two major expression systems: Mammalian cell development and Microbial strain fermentation development. Additionally, we have specialized capabilities in research, manufacturing, and analysis. Through a vertically integrated operating model, we achieve quality control and cost management. In light of the exorbitant prices of biopharmaceuticals from original development companies, which many patients cannot afford, coupled with the escalating burden of healthcare costs on governments worldwide, the mission of our company is: in the short term, to provide customers with high-quality and cost-effective services, while also developing biosimilar products with commercial viability; in the middle to long term, to focus on developing niche biologics that enhance human and societal well-being and improve quality of life. Our ultimate goal is to become an international biopharmaceutical company rooted in Taiwan with a global outlook, dedicated to advancing human health and welfare.

B. Revenue distribution

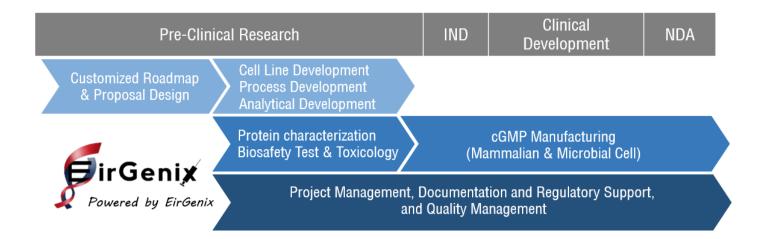
Unit: NT\$ thousands

Year	2022		2023	
Item	Revenue	%	Revenue	%
Service Revenue	757,680	51.16	605,990	59.26
Sales Revenue	461,461	31.16	275,191	26.91
Licensing Revenue	261,876	17.68	141,472	13.83
Total	1,481,017	100	1,022,653	100

The sales are primarily focused on the Asian region, accounting for approximately 65% of revenue, followed by Europe and the Americas, which make up about 35% of revenue.



C. Main products (Service)



Our company has independently developed the following core technologies and six major platforms related to CDMO and the capacity and scale of the CDMO platform:

- (A) Cell line development
- (B) Process development and Scale-up
- (C) Analytical Method development and validation
- (D) Product identification
- (E) cGMP production for clinical trials +stability testing
- (F) CMC (Chemical, Manufacturing, and Control) documents
- (G) The capacity and scale of the CDMO platform

These are described separately as follows:

(A) Cell line development

To accelerate the development, clinical trials, and market entry of next-generation biopharmaceutical products, our company focuses particularly on the first critical technology in the biopharmaceutical development stage- the development of cell lines and strains. This development includes the establishment and optimization of high-yield cell lines and strains, as well as the optimization of culture media and cultivation methods. It also involves the establishment of MCB (Master Cell Bank) and WCB (Working Cell Bank). The key focus at this stage is on selecting the optimal host cells (such as CHO, Sp2/0, NS0, Hybridoma, HEK 293, PER.C6 cells, etc., for animal cells; and E. coli, S. cerevisiae, Pichia, etc., for microbial cells) and optimizing culture media, process development to achieve the highest production (for recombinant protein or monoclonal antibody drugs) and quality of cell lines/strains. The implementation involves inserting the gene into a vector to express the protein, constructing an expression system, and then transfecting it into host cells (adapted to serum-free and suspension culture). Stable and high-yield cell lines are then selected, comparing the stability and quality of production, and establishing MCB and WCB. Simultaneously, suitable or developed cell culture media for production are selected or developed.

(B) Process development and Scale-up

The upstream process development and scale-up primarily focus on the process development and optimization of fed-batch culture, especially on scalability, and manufacturing suitability leading to high density of cell lines/strains during production.

On the other hand, downstream process development emphasizes the recovery and purification processes, viral clearance assay studies, scale-up (currently mainly at 100 liters), formulation development, and providing products/materials to support animal testing studies, reference standards, and quality control (QC) requirements.

(C) Analytical Method development and validation

The development and validation of analytical methods are crucial for product quality confirmation, including:

- (i) Identification: SDS-PAGE, Western blot, IEF, peptide mapping, IEC-HPLC
- (ii) Quantitative determination: BCA/Bradford, A280
- (iii) Purity: SEC-HPLC, RP-HPLC, SDS-PAGE
- (iv) Activity: ELISA, cell-based assay
- (v)Impurity: Host cell DNA, host cell protein, ProA residue, endotoxin, bioburden

These analysis methods will be validated through linearity & range, accuracy, and precision.



(D) Product identification

Protein identification has been increasingly emphasized by regulatory authorities year by year. Our company has established a set of HPLC and LC/MS/MS systems capable of conducting peptide mapping, complete sequence analysis, N-/O-linked carbohydrates analysis, disulfide linkages analysis, oxidation analysis, deamidation analysis, and other post-modifications analysis, as well as analysis of N-/C-terminal variants, secondary and higher-order structures, and other related analytical tasks.

(E)cGMP production for clinical trials and stability testing

The pilot run will be able to provide drugs needed for animal toxicity tests, preliminary stability test data, and reference standard samples, and provide sufficient operating parameters as the basis for GMP production preparation. The GMP production section includes GMP trial production (Engineering Run), GMP production, End of production cell banking and testing, viral clearance assay studies (limited to mammalian cell culture), stability testing, clean validation, and other tasks.

(F) CMC documents

Our company provides comprehensive CMC (Chemistry, Manufacturing, and Control) document services to customers for subsequent product application in clinical trials. At each stage of new drug research, sufficient CMC data should be submitted to ensure appropriate assurance of the new drug's identification, quality, purity, potency, and stability. The required CMC data vary depending on different stages of research, planned testing periods, dosage forms, and other available data. For instance, stability data are necessary throughout all stages of new drug testing to demonstrate that the active pharmaceutical ingredient and drug product remain within acceptable limits during the designated study period. CMC documents can be likened to the manufacturing process's record, serving as a crucial basis for regulatory safety assessments.

(G) The capacity and scale of the CDMO platform

Our company is one of the few in Asia equipped with both upstream mammalian cell and microbial strain fermentation production facilities, along with a comprehensive downstream protein purification system, enabling the production of investigational medicinal products for clinical trials. Within our biopharmaceutical cGMP facilities, we exclusively utilize single-use bioreactors (SUBs) ranging from 50 to 2000 liters. The total production capacity for mammalian cell has reached 25,500 liters and microbial strain 150 liters (will reach 1,500 liters in 2026).

The facilities that have been invested in are outlined below:

Facility	Usage	Highlights
Xizhi Factory	For mammalian cell culture	The production facilities are located on the first floor of the Taiwan Xizhi Plant, including 200L and 1,000L single-use bioreactors (SUBs), purification rooms, final purification rooms, media preparation rooms, buffer preparation rooms, washing rooms, and high-pressure sterilization rooms.
	For microbial cell culture	There is one 30L and one 150L stainless steel bioreactor (SSB) for fermentation.
Zhubei A Factory (in	For mammalian cell culture	• Officially operational since 2019, including 2 units of 1,000L and 4 units of 2,000L single-use bioreactors (SUB).
Hsinchu Biomedical Science Park)		• Subsequently, there will be an expansion with an additional 2 units of 2,000L and another production line consisting of 6 units of 2,000L SUBs, totaling 12 units of 2,000L SUBs. It is estimated that the annual production capacity for monoclonal antibodies could reach 1,000 kilograms.
		 The services provided are applicable from late-stage clinical trials to commercial-scale production after market approval.
Zhubei B Factory (next to A)	Under construction	 Primarily focused on microbial cell production lines, with the establishment of 1500L, 500L, and 75L fermentation tanks, along with 2-3 purification lines. Scheduled for completion and commissioning by 2026.
Ciaotou	Planning	· For mammalian cell culture
Factory		• There are plans for 10 units of 15,000L stainless steel bioreactors (SSB).

EirGenix's cGMP (Current Good Manufacturing Practice) plant (Xizhi plant) has received a PIC/S GMP certificate from TFDA. EirGenix received PMDA's official approval in its issued GMP Compliance Inspection Result Notification, proclaiming EirGenix's compliance with relevant Japanese regulations regarding the quality, effectiveness, and safety of pharmaceutical manufacturing, which represented a remarkable milestone for EirGenix as the GMP biopharmaceutical manufacturing facility to receive the authority's approval in 2020 and 2023. EirGenix's Zhubei plant has passed the review by the FDA and obtained



an EIR before the drug launch. EirGenix passed the review by the TGA in 2023.

D. The new products (services) are planning to development:

(A) Biosimilar drugs

(i) EG12014

EG12014 is EirGenix's first self-developed and successfully marketed Trastuzumab biosimilar (EMA approved trade name HERWENDA, TFDA approved trade name EIRGASUN). Its R&D target HERCEPTIN was acquired by Genentech (merged by Roche in March 2009). obtained marketing approval from the US FDA in September 1998. It is a genetically recombinant monoclonal antibody and is a drug against high-phenotype breast cancer caused by the oncogene (HER2/neu). It is mainly used for the treatment of patients with early breast cancer (EBC), metastatic breast cancer (MBC) and metastatic gastric cancer (mGC) with HER2 overexpression or HER2 gene amplification. In terms of early breast cancer (EBC), it includes: (1) adjuvant therapy through surgery and chemotherapy (preoperative or postoperative). (2) Treat with Doxorubicin and Cyclophosphamide, combined with adjuvant therapy of Paclitaxel or Docetaxel. (3) Adjuvant therapy with Docetaxel and Carboplatin. (4) Preoperative combined with chemotherapy and postoperative adjuvant therapy for the treatment of locally advanced (including inflammation) breast cancer or tumors (>2 cm in diameter); in metastatic breast cancer (MBC): (1) Used alone to treat previously treated breast cancer Metastatic breast cancer that has received more than one chemotherapy; unless the patient is not suitable for Anthracycline or Taxane, the previous chemotherapy should at least include Anthracycline or Taxane. For use in hormone receptor-positive patients who have failed hormonal therapy, unless the patient is not suitable for hormonal therapy. (2) Used in combination with Paclitaxel or Docetaxel for metastatic breast cancer that has not received chemotherapy. (3) Combined with aromatic cyclase inhibitors for hormone receptor-positive metastatic breast cancer; in metastatic gastric cancer (mGC): combined with Capecitabine (or 5-fluorouracil) and Cisplatin for HER2 overexpression that has not received chemotherapy Treatment of metastatic gastric adenocarcinoma (or gastroesophageal junction adenocarcinoma).

(ii) EG1206A

EG1206A is a biosimilar of Pertuzumab. Its research and development target PERJETA has obtained indications for early breast cancer (EBC) and metastatic breast cancer (MBC). In the treatment of metastatic breast cancer (MBC): used in combination with trastuzumab and docetaxel to treat patients with HER2-positive metastatic breast cancer who have not been treated with anti-HER2 or chemotherapy after metastasis. In early breast cancer (EBC): When Pertuzumab is used in combination with chemotherapy drugs for preoperative adjuvant therapy, it is suitable for patients with HER2-positive, locally advanced, inflammatory, or early breast cancer (tumor diameter greater than 2 cm or positive lymph nodes) as a part of a complete treatment prescription for early-stage breast cancer. When used as postoperative adjuvant therapy, it is suitable for patients with early-stage breast cancer who are HER2-positive and have a high risk of recurrence. (According to the results of the APHINITY trial, during postoperative adjuvant therapy, HER2-positive early breast cancer patients with a high risk of recurrence are defined as having lymph node-positive breast cancer.) Pertuzumab has different binding mechanisms for HER2 receptors and can produce double blockade (Dual Blockade) effect. EG1206A is a recombinant humanized monoclonal antibody that targets the extracellular dimerization domain (subregion II) of human epidermal growth factor receptor type 2 (HER2). Therefore, it can block the ligand-dependent heterodimerization of HER2 and other human epidermal growth factor receptor (HER) family members (including EGFR, HER3 and HER4). Inhibits ligandinitiated intracellular signaling through two major signaling pathways: mitogen-activated protein (MAP) kinase and phosphoinositide 3-kinase (PI3K). Inhibition of these signaling pathways will lead to cell growth cessation and apoptosis respectively.

(iii) TSY0110 (EG12043)

EG12043 (TSY0110) is a biosimilar of Trastuzumab Emtansine. It is an ADC (Anti-Drug Conjugate). Its mechanism is an antibody-drug complex that targets HER2. The antibody is human anti-HER2 IgG1 (trastuzumab). The small molecule cytotoxin DM1 is a microtubule inhibitor. After binding to the IV domain of the HER2 receptor, Trastuzumab Emtansine begins to be internalized through the receptor, and the subsequent lysosomal degradation process releases cytotoxic metabolites containing DM1 into the cell. The process of DM1 binding to tubulin will destroy the intracellular microtubule network, leading to cell cycle arrest and apoptosis. In addition, in vitro experiments also show that Trastuzumab Emtansine,



similar to Trastuzumab, also inhibits the function of HER2 receptor signaling, causes antibody-dependent cell-mediated cytotoxicity, and inhibits HER2 extracellular domain shedding in HER2-overexpressing human breast cancer cells. Trastuzumab Emtansine can be used alone to treat HER2-positive metastatic breast cancer in patients who have previously received Trastuzumab and a Taxane drug, or their combination, if they meet the following conditions: have previously received treatment for metastatic cancer or are receiving adjuvant therapy Patients whose cancer relapses during or within 6 months after completing treatment. Early Breast Cancer: Used alone, it is suitable for adjuvant therapy for patients with HER2-positive early breast cancer who still have residual disease after receiving Taxane and Trastuzumab-based lead treatment (neoadjuvant therapy).

(iv) EG1211X

EG1211X is a biosimilar of Atezolizumab. Its R&D target Tecentriq is an Fc-engineered, humanized monoclonal antibody that binds to PD-L1 and blocks its interaction with PD-1 and B7.1 receptors. Atezolizumab is an unglycosylated IgG1 kappa immunoglobulin with a calculated molecular weight of 145 kDa. Its pharmacological effect is that PD-L1 can be expressed on tumor cells and/or tumor-infiltrating immune cells and can inhibit the anti-tumor immune response in the tumor microenvironment. PD-L1 binds to PD-1 and B7.1 receptors on T cells and antigen-presenting cells to inhibit the activity of cytotoxic T cells, T cell proliferation and cytokine production. Atezolizumab is a monoclonal antibody that binds to PD-L1 and blocks its interaction with PD-1 and B7.1 receptors. This releases PD-L1/PD-1-mediated suppression of immune responses, including activation of anti-tumor immune responses without inducing antibody-dependent cellular cytotoxicity. In syngeneic mouse tumor models, blocking PD-L1 activity resulted in reduced tumor growth. Indications span a variety of cancer types: 1. Locally advanced or metastatic urothelial cancer is suitable for the treatment of patients with locally advanced or metastatic urothelial cancer whose disease has worsened after receiving platinum-containing chemotherapy or who are not suitable for cisplatin-containing therapy. 2. Used alone for locally advanced or metastatic non-small cell lung cancer. It is suitable for the treatment of patients with locally advanced or metastatic non-small cell lung cancer whose disease has worsened after receiving platinum-containing chemotherapy. If patients have EGFR or ALK tumor gene abnormalities, they must first be treated with EGFR or ALK inhibitors. If the disease worsens after treatment, Tecentriq can be used. Used in combination with Avastin (bevacizumab), paclitaxel and carboplatin, as a first-line treatment for metastatic non-squamous non-small cell lung cancer without EGFR or ALK tumor gene abnormalities. 3. The combination of Tecentriq and nab-paclitaxel for triple-negative breast cancer is suitable for the treatment of unresectable locally advanced or metastatic triple-negative breast cancer, and the tumor has PD-L1 manifestations (tumor-infiltrating immune cells (IC) \geq 1%) and has not received Chemotherapy is used for patients with metastatic breast cancer. 4. Small cell lung cancer, combined with carboplatin and etoposide, is suitable for the first-line treatment of adults with extensive stage small cell lung cancer. 5. The combined use of hepatocellular carcinoma and bevacizumab is suitable for the treatment of patients with hepatocellular carcinoma who have not received systemic therapy and are unresectable or metastatic, and their liver function is Child-Pugh A.

(v) EG1216X

EG1216X is a biosimilar of Daratumumab. The target of its R&D, DARZALEX, is a humanized IgG1κ monoclonal antibody that can bind to the CD38 antigen. This strain is produced in a mammalian cell line (Chinese Hamster Ovary [CHO]) through recombinant DNA technology. Its pharmacology Mechanism is an IgG1κ human monoclonal antibody (mAb) that binds to the CD38 protein that is highly expressed on the surface of multiple myeloma cells and to varying degrees on other cell types and tissues. CD38 protein has multiple functions, such as receptor-mediated adhesion, signaling and enzyme activity. DARZALEX is suitable for: 1. Patients who have received at least three previous therapies (including a protease inhibitor and an immunomodulator) as a single drug, or whose disease has worsened under treatment with both a protease inhibitor and an immunomodulator (double-refractory to a protease inhibitor and an immunomodulatory agent) in adults with multiple myeloma. 2. Use in combination with lenalidomide plus dexamethasone or with bortezomib plus dexamethasone to treat adult patients with multiple myeloma who have previously received at least one therapy.



(B) New dosage forms and new drug delivery systems of biosimilars

(i) EG13084

EG13084 is a new subcutaneous injection drug that combines Trastuzumab and Pertuzumab. Its R&D target is PHESGO-approved indications for early breast cancer (EBC) and metastatic breast cancer (MBC). In terms of early breast cancer (EBC), when EG13084 is used in combination with chemotherapy drugs for preoperative adjuvant therapy, it is suitable for patients with HER2-positive, locally advanced, inflammatory or early breast cancer (tumor diameter greater than 2cm or positive lymph nodes). Part of a complete treatment prescription. When used as adjuvant treatment after surgery, it is suitable for patients with early-stage breast cancer (EBC) who are HER2-positive and have a high risk of recurrence. (According to the results of the APHINITY clinical trial, in postoperative adjuvant therapy, patients with HER2-positive early breast cancer who have a high risk of recurrence are defined as having lymph node-positive breast cancer.) In terms of metastatic breast cancer (MBC), it is used together with docetaxel to treat metastasis patients with HER2-positive metastatic breast cancer who have not been treated with anti-HER2 or chemotherapy. In addition to the significant clinical benefits of this product, once successfully launched on the market, it will also greatly increase the convenience of breast cancer patients receiving treatment.

(ii) EG7412X

The R&D target of EG7412X is "recombinant human hyaluronidase PH20 (rHuPH20)", which is equivalent to the soluble fragment of human HYAL5. It is genetically engineered in hamster culture cells (CHO) culture containing a DNA plasmid encoding the enzyme. Produced as a type of hyaluronidase (hyaluronidase), its main function is to catalyze the degradation of hyaluronic acid (HA). According to the enzyme reaction products, they are divided into three different types: two eukaryotic endoglycosidase hydrolases and one prokaryotic lyase-type glycosidase. Hyaluronidase catalyzes the hydrolysis of HA, causing the viscosity of HA to decrease, thereby increasing tissue permeability. Therefore, it is often used in medicine together with other drugs, which can accelerate the dispersion and delivery of drugs through subcutaneous absorption.

(C) Special biological drugs

(i) EG74032

EG74032 is CRM197 modified from Diphtheria toxin. It is no longer toxic after modification of amino acids. Therefore, it can be used as a carrier to make a mixed vaccine (Conjugate vaccine) to promote immune effect. CRM197 is a carrier protein that assists vaccine immunity without patent protection. This protein has been widely used in commercial products and clinical development products. In addition to traditional infectious disease vaccines, the field of cancer vaccine research and development has also attracted much attention in recent years. Both R&D units and biotech companies at home and abroad are actively investing in the development of cancer vaccines, hoping to bring a glimmer of hope to various types of cancers for which effective treatments currently do not exist. Early production processes used Corynebacterium diphtheria (Corynebacterium diphtheria) for production, and then carried out downstream recovery, purification and other steps. The yield and recovery rate were usually relatively low. In addition, the acquisition of this strain requires signing an authorization agreement with a specific unit, and the production unit must have a biological protection level that meets the specifications before it can produce.

(2) Industry Overview

A. The Current Status and Development of the Industry

The biopharmaceutical industry can be divided into two categories of drugs based on molecular size: large molecules (Macromolecular) and small molecules. Small molecule drugs have a long history of development and are mostly manufactured through chemical synthesis. Common examples include antibiotics, analgesics, and sedatives. Macromolecular drugs, also known as biologics, have a molecular weight much higher than small molecule drugs. They are primarily produced by genetically modified microorganisms, plants, or animal cells for therapeutic purposes. Common examples include insulin, rheumatoid arthritis (RA) monoclonal antibody therapy drugs, and targeted cancer therapies. Biosimilars are also a category of macromolecular drugs. According to the FDA, biosimilars are defined as follows: A biosimilar is a biologic that is highly similar to another biologic that is already FDA-approved (known as the original biologic). It is both normal and expected for both biosimilars and original biologics to have minor differences between batches of the same medication in terms of the safety, purity, and potency.



Unlike traditional small molecule drugs, biologics have stable chemical structures, larger molecular weights, and complex structures. After approval, biologics, due to their specificity in disease treatment, high safety, and significant efficacy, often become blockbuster drugs shortly after market introduction. With the increasing severity of safety and drug resistance issues caused by chemical drugs, biologics can fill the gaps in the treatment field left by chemical drugs. Their growth rate continues to rise, overwhelming the overall pharmaceutical market trend.

Our company focuses on the research, development, and manufacturing of biopharmaceuticals, with CDMO services as the main focus. We accept commissions from biotechnology and pharmaceutical companies, providing services related to the development and manufacturing of biotech products and biopharmaceuticals. These services include product evaluation and design, overall development and market entry processes, cell lines and strains, processes, culture media required for CMC development to production, clinical trial drugs, raw material production, and developing process of scale-up. Compared to small molecule drugs, the development and manufacturing of biopharmaceuticals have relatively higher entry barriers. In addition to requiring significant investment in infrastructure, the production process is also more complex, with greater difficulty in scaling up processes. The capacity utilization rate of small to medium-sized biopharmaceutical CDMO companies is higher than that of large-scale biopharmaceutical CDMO companies. This is mainly due to the higher flexibility of small to medium-sized CDMO companies in adjusting production capacity, allowing them to meet the diverse production needs of clients.

According to 2023 Market Data Forecast, the biologics CDMO market started from US\$ 14.37 billion in 2024 and estimated grow with 11.52% compound annual growth rate (CAGR) to reach US\$ 24.79 billion in 2029. In order to seize the opportunity of this global trend, EirGenix has been actively expanding its plants and preparing for the market demand in the next 10 years. In addition to the sales of self-developed products, the company has also been actively seizing the contract manufacturing market where biologics manufacturing and demand have significant growth, of which biosimilars are the focus of the fastest growth.

The use of cell types in the manufacturing of biopharmaceuticals also indicates a gradual decrease in microbial cell fermentation, while the use of mammalian cells shows a relative growth trend. This phenomenon is due to the fact that biotech products produced by mammalian cells, such as monoclonal antibodies (mAb), require much higher therapeutic doses compared to protein drugs produced by microbial cells, leading to higher production volumes. Currently, our company's development of biosimilar drugs primarily focuses on mAb. Therefore, we have proactively expanded and established mammalian cell facilities while retaining space for microbial production facilities. We continue to monitor market changes and trends, and will evaluate the timing for establishing microbial cell fermentation facilities. According to survey reports, the global production capacity of mammalian cell systems exceeded demand in 2017. However, most CDMOs remain conservative in their capacity expansion strategies. The increased capacity mainly focuses on Multiple-2,000 SUBs, following a scale-out design concept. The aim is to make the newly added facilities and equipment more flexible and effectively reduce development risks.

According to the 2021 market research report by BioPlan Associates, Inc., nearly 70% of global biopharmaceutical manufacturing capacity is concentrated in Europe and the United States. The main reason for this is the expiration of patents for major biopharmaceuticals worldwide. Additionally, healthcare systems in European and American countries, which serve as benchmarks, are actively seeking high-quality and cost-effective biosimilar drugs to alleviate fiscal pressures on healthcare systems. Our company is currently actively developing biosimilar drugs. Upon completion of the expansion, we will not only be able to meet the market demand for our own products but also offer the remaining capacity to domestic and international clients for contract manufacturing. This strategy will enable us to establish a strong existence in Taiwan and offer the service to clients around the world as well as seize the profit base of competition in the global market.

Figure: 2021 Regional distribution of Global Biomanufacturing Capacity

Region	Regional Capacity, L	Global Capacity %, by Region
US/North America	5,496,290 L	31.7%
Europe	5,982,977 L	34.7%
Asia/Rest of World	5,826,070 L	33.6%
Total Worldwide	~17,300,000 L	100%



(A) Biosimilars

Macromolecule drugs, due to their complex structure, cannot be replicated 100% identical from the original developers. Therefore, for macromolecule drugs developed with reference to marketed biopharmaceuticals, they must be highly similar to the reference product in terms of molecular structure, physical, chemical, and biological properties, with no clinically meaningful differences in safety, quality, and efficacy, as verified by regulatory authorities before being marketed, in order to be termed biosimilar drugs. The development costs and timelines for biosimilar drugs are much higher than for small molecule generic drugs. The main difference from developing new drugs lies in the reverse engineering of cell lines and processes in the front end, ensuring that the product achieves a high degree of similarity to the reference product in molecular structure, physical, chemical, and biological properties. The selection of cell lines and reverse engineering techniques are a highly challenging barrier in the development of biosimilar drugs. Even after process development, biosimilar drugs still need to undergo two stages of human clinical trials. The first stage is a Phase I clinical trial comparing the pharmacokinetics of the drug in the body (bioequivalence), and the second stage compares the efficacy of the biosimilar drug with the reference biopharmaceutical (equivalence). If reliable biomarkers are available, they can also be used as primary clinical endpoints. The development of biosimilar drugs differs from innovative drug development; innovative drug development entails considerable time and cost, especially in late-stage clinical trials, where the failure rate is quite high. Conversely, if a biosimilar product achieves a high degree of similarity and demonstrates bioequivalence in clinical pharmacokinetic studies, the failure rate in Phase III clinical trials is almost negligible.

Since the enactment of The Patient Protection and Affordable Care Act in 2010, the United States (US) has established legislative authority for reviewing biosimilar drugs through the Biologics Price Competition and Innovation (BPCI) Act. As of February 2024, the FDA has approved a total of 46 biosimilar product applications covering 14 different active ingredients. Regulatory authorities in US are actively accelerating the review of biosimilar drugs. Additionally, a landmark decision by the US Supreme Court in June 2017 clarified two key aspects of biosimilar regulation. First, it ruled that the BPCI Act provision requiring biosimilar drug manufacturers to submit their CMC applications to the original biologic manufacturer for review of potential process patent infringement is discretionary rather than mandatory. Second, it provided a definitive ruling on the timing of notification to the original biologic manufacturer by biosimilar manufacturers seeking to launch their products within 180 days, allowing them to notify the original manufacturer before FDA approval. This decision is expected to remove barriers to biosimilar market entry and accelerate their approval process in the US.

After biosimilar are launched, their prices do not decrease as significantly as generic drugs. It is estimated that there will only be a reduction of about 20-35% in the early stages of product launch. However, due to the high cost of treatment, even a slight decrease in price contributes to the overall reduction in healthcare expenditure. Countries are increasing the use of biosimilar drugs to lower medical costs, thereby bringing promising business opportunities, and expected profits to biosimilar drug developers. Based on the experience of using biosimilar drugs in Europe, it can be observed that their price reduction rate is much slower than that of generic drugs. Despite the decrease in unit price, total sales revenue increases. This can be attributed to two main factors: (1) doctors start treating patients proactively before their condition worsens, and (2) patients who could not afford the original biologic drug (reference drug) now have the opportunity to use biosimilar drugs with no clinical differences in safety, quality, and efficacy compared to the reference drug.

The regulatory framework for biosimilar drugs in Europe was established in 2001. Since the approval of the world's first biosimilar drug, Omnitrope®, in April 2006, Europe has approved a total of 76 biosimilar drugs as of February 2023. Among them, the most approved biosimilar drugs are granulocyte colony-stimulating factors (G-CSF), followed by biosimilar drugs referencing Roche's MabThera® (Rituxan®). Other biosimilar drugs referencing Humira®, Herceptin®, Avastin®, Forsteo®, Humalog®, and others are also either approved or pending approval. In addition to the rapid growth of biosimilar drugs in the European and American markets, Japan has also seen an increasing number of biosimilar drugs obtaining marketing approval, with the overall market continuing to grow at an extremely fast pace.

According to the 2021 Evaluate Pharma report, the global oncology drug market is expected to continue its robust growth driven by the emergence of new drugs, primarily immunotherapy drugs, from 2020 to 2026. It is projected to reach \$319 billion by 2026, accounting for 22% of total global drug expenditures. In response to the high cost but remarkable efficacy of newly launched drugs, various countries in Europe and America have actively promoted the extensive use of biosimilar drugs to replace off-patent biologics. This initiative not only helps alleviate the increasingly challenging financial pressures on global healthcare systems, but also accelerates the adoption of next-generation therapeutic drugs. Consequently, advanced countries including those in Europe, America, and Japan have achieved a certain proportion of biosimilar drug utilization in a short period,



indirectly bringing certain benefits to the developers of biosimilar drugs.

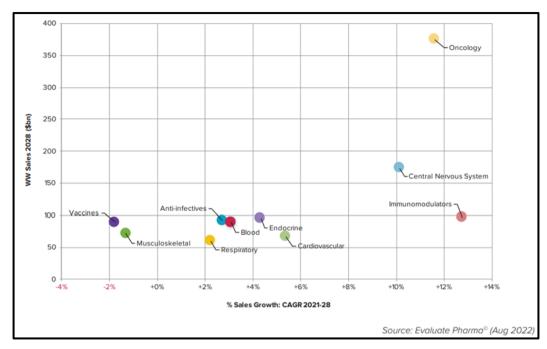


Figure: Top 10 therapy areas in 2028 by market share & sales globally

According to Global Market Insights' 2023 forecast, the global market size of biosimilar drugs reached approximately \$35 billion in 2022 and is expected to continue exponential growth. It is estimated to surpass \$122.9 billion by 2032, with a high CAGR of 13%. This represents a rare surge in the history of pharmaceutical development. The primary market growth opportunities are focused on regions such as the US, which has relatively late-stage biosimilar markets, as well as the Asia-Pacific and Latin America regions, which offer cost-effective growth potential. The increasing prevalence of diabetes worldwide, along with government support, has increased the demand for insulin biosimilars, further driving the adoption of biosimilar drugs.

Affected by the slowing global economic growth and the rapid increase in the elderly population, healthcare systems in various countries are facing increasingly heavy burdens. The prices of biologic drugs, which have significant effects and relatively low side effects but incur high development costs, remain high. Against this backdrop, the adoption of biosimilar drugs, which offer no differences in safety, quality, and efficacy compared to patented biologic drugs but are relatively lower in price, has become one of the effective methods for healthcare systems worldwide to address the current challenges. The momentum generated by the expiration of patents for biologic drugs will also accelerate the replacement of market share held by original biologic drugs with biosimilar drugs.

(B) Antibody drug conjugate (ADC) drugs

ADC, which involve highly cytotoxic small molecules linked to mAb, represent a novel class of drugs with both high specificity and antibody properties. These drugs precisely target malignant tumors with cytotoxic agents in a "targeted" manner, minimizing impact on other normal tissues. The highly cytotoxic small molecules exhibit potent activity in inhibiting cell growth when reaching picomolar concentrations (pM). To align with the trend of developing novel drugs, international CDMO are integrating High Potency Active Pharmaceutical Ingredient (HPAPIs) technology with enhanced capabilities in the production and development of ADCs. Undoubtedly, ADC technology and products have become a trend in the development of novel antibody drugs. Collaborating with CDMOs to become partners for small to medium-sized biotech companies can establish a new market positioning and continue to seize opportunities in the development of biosimilar drugs and antibody-based therapeutics.

The next-generation antibody therapeutics market encompasses ADCs, bispecific antibodies, Fc-fusion antibodies, antibody fragments, and antibody proteins. According to a report by Market Data Forecast in 2023, the ADC market is projected to reach \$18.36 billion in 2023 and is expected to surpass \$55.45 billion by 2028, with a CAGR of 24.74% from 2023 to 2028. As of October 2023, the FDA has approved and successfully launched 15 ADC drugs: Elahere (2022), Tivdak (2021), Zylonta (2021), Aidixi (2021), Blenrep (2020), Trodelvy (2020), Akalux (2020), Enhertu (2019), Padcev (2019), Polivy (2019), Lumoxiti (2018), Besponsa (2017), Mylotarg (2017), Kadcyla (2013), and Adcetris (2011).

Currently, the manufacturing of ADC requires expensive upstream mammalian cell bioreactors and downstream protein purification equipment, along with specialized antibody and chemical drug conjugation technologies, and high-efficiency



cytotoxic small molecule chemical drug manufacturing facilities. These factors have led to most ADC drug manufacturers relying on a few specific CDMOs that can provide various technical services, including monoclonal antibodies, chemical linkers, and cytotoxins. However, only a few companies can offer integrated development services for ADC drugs. It is anticipated that over 10 new commercial ADC products will drive the growth of the entire market in the next decade. Our company is currently collaborating with Formosa Laboratories, Inc., with small molecule technology capabilities to jointly provide this platform service. The collaboration aims to seize the next wave of biotech advancements, potentially setting new milestones for Taiwan in the international market.

B. Upstream and downstream relevance in the industry

The process of developing new drugs generally consists of five stages: drug discovery, preclinical trials, clinical trials, inspection registration, and post-market surveillance. Typically, the entire development process requires decades of effort and significant investment of high-risk funds. Our company and its subsidiaries' CDMO business possess both development and manufacturing capabilities, mastering key technologies in the development and manufacturing of biopharmaceuticals, and possessing international regulatory expertise, enabling us to provide high-value-added differentiated services. The development of biosimilar drugs eliminates the need for drug discovery and preclinical trials but increases the number of product comparative structural analysis testing items, focusing on the CMC part.

Our company's service offerings include cell line development, biopharmaceutical process development and optimization, related analytical method development and validation, compliance with regulatory requirements for quality control, and GMP pilot production. The establishment of expertise in biopharmaceutical CMC and a customer-centric project management mechanism enables us to provide efficient and internationally standardized quality-stable biopharmaceuticals that are safe, effective, and economically viable. Our CDMO business combines development and manufacturing capabilities, mastering key technologies in the development and manufacturing of biopharmaceuticals, and possessing international regulatory expertise, enabling us to provide high-value-added differentiated services.

In 2017, EirGenix was nominated for Best Process Technology by the Biology Manufacturing Asia (BMA), received the Grand Winner of Best Bioprocess Excellence in Taiwan, received the excellence award for Antibody Drug Conjugate platform at the same time, and won international recognition in bioprocess technology. Received the "Asia's Best CMO (Contract Manufacturing Organization) Award" in Asia-Pacific Bioprocessing Excellence Awards 2018. Won the Grand Winner of Best Bioprocess Excellence in Taiwan Award in Singapore again in 2019. Received 2020 Bioprocessing Excellence Award in Greater China Region. Won the Globalizing Award of "2021 Taipei Biotech Awards". Won the Best Bioprocessing Awards in Taiwan and the Greater China issued by Biologics Manufacturing Asia (BMA) and National Innovation and Advancement Award (EG12014 and CRM197) in 2022. EIRGASUN vial 150 mg won the Golden award of National Pharmaceutical Technology Research Development Award in 2023.

In addition, our company has also obtained the Accreditation Certificate of Foreign Drug Manufacturer issued by the Japanese Ministry of Health, Labor and Welfare, with the recognized category being "Biological Products." The certificate is valid from October 24, 2022, to October 30, 2027. During the validity period, biological products manufactured at our Taiwan facility can be sold and marketed in Japan. In 2017, a Japanese client transferred their product, which was already on the market in Japan, from a Japanese CMO to our company for production. After completing technology transfer and process validation, the Pharmaceuticals and Medical Devices Agency (PMDA), an independent administrative agency of the Japanese government, conducted an inspection at our facility in September 2019. The inspection process went smoothly without significant deficiencies, and on February 3, 2020, we received approval from the PMDA. Subsequently, the Japanese client began placing orders for commercial production at our facility and initiated negotiations for long-term supply. Finally, on March 2, 2021, our company signed a long-term supply contract with this Japanese pharmaceutical company, becoming the first Taiwanese manufacturer to supply biopharmaceuticals for a product marketed in Japan. This product is essential for cancer treatment, with a market share in Japan exceeding 30% in its category. Our company is not only the first in the Greater China region but also one of the few in Asia to be audited and approved by the PMDA for biopharmaceutical manufacturing. This certification will enhance the willingness and confidence of Japanese and international biotech companies to entrust us with biopharmaceutical production, aiding in business promotion. In recent years, the demand for biopharmaceutical CDMO services in Japan has been growing. Through our successful track record of selling products in Japan, we are expanding our competitive advantage in the Japanese market, significantly increasing the willingness and confidence of Japanese and international biotech companies to entrust us with biopharmaceutical production. This significant milestone will accelerate the expansion of our CDMO business.



C. Development Trends

(A) Biosimilar drugs

(i) EG12014

EG12014 is a biosimilar of Trastuzumab, and its R&D target HERCEPTIN has global annual sales of 1.79 billion CHF according to Roche's 2023 financial report, of which the European and US markets account for 42%. According to data from the Taiwan Health Insurance Administration, breast cancer ranked second among the top ten cancer health insurance medical expenses in Taiwan in 2022, with drug expenditures of 9.075 billion NTD (the average growth rate from 2018 to 2022 was 7.75%). Among them, the breast cancer target drug HERCEPTIN, Taiwan's health insurance expenditure in 2022 was as high as 1.8 billion NTD. In 2024, the latest health insurance drug price of HERCEPTIN frozen crystal injection form (440 mg) was NT\$43,236 per tube. Currently, the National Health Insurance reimbursed Trastuzumab for patients with early breast cancer (EBC), metastatic breast cancer (MBC) and metastatic gastric cancer (mGC). The number of breast cancer and gastric cancer patients in Taiwan increases every year, and medical expenses also increase accordingly. The latest health insurance price of EirGenix's EIRGASUN 150 mg in 2024 is NT\$11,323/tube, which brings more benefits to more people with a more affordable price and wider treatment benefits. Patients who need to undergo breast cancer treatment can achieve expected clinical efficacy while reducing their medical expenses by using EG12014 (EIRGASUN 150 mg), which has no clinically significant difference in safety, quality and efficacy from HERCEPTIN, to achieve the purpose of truly benefiting the people.

(ii) EG1206A

EG1206A is a biosimilar of Pertuzumab. Since its R&D target PERJETA was launched in the United States in 2013, sales have grown rapidly every year. According to the results of the APHINITY trial of Pertuzumab combined with Trastuzumab and chemotherapy in the treatment of early breast cancer published by Roche in 2022, this combination therapy can reduce the risk of recurrence or death by 23% compared with the control group. From this excellent test results, it is foreseeable that the subsequent product development and therapeutic application of EG1206A will be more extensive. Roche is currently actively promoting the expansion of indications and is expected to continue to expand the treatment scope and market. According to Roche's 2023 annual financial report: the global annual sales of this product still reached 4.06 billion CHF, with an annual growth rate of 9%, and the European and US markets accounted for 54% of revenue contribution.

(iii) TSY0110 (EG12043)

EG12043 (TSY0110) is a biosimilar of Trastuzumab Emtansine. According to Roche's 2023 annual financial report, its R&D target KADCYLA: the global annual sales of this product reached 2.16 billion CHF, with an annual growth rate of 4%, and the European and US markets accounted for 65% of the revenue contribution. According to the 2024 Research and Markets report, the global ADC (Anti-Drug Conjugate) market will be approximately US\$8.8 billion in 2023 and is expected to reach US\$10.7 billion in 2024. It will even reach US\$24.8 billion by 2028 with a compound annual growth rate of 23.4%.

(iv) EG1211X

EG1211X is a biosimilar of Atezolizumab. Its R&D target Tecentriq is an immune checkpoint inhibitor (Immune Checkpoint Inhibitor). According to Roche's 2023 annual financial report, global annual sales reached 3.77 billion CHF, with an annual growth rate of 9%, of which the European and US markets accounted for 73%. As the world's first approved PD-L1 immune checkpoint inhibitor, Roche has invested considerable resources in clinical trials for multiple cancer types, and has successively obtained results for locally advanced or metastatic urothelial cancer, locally advanced or metastatic non-Small cell lung cancer, triple negative breast cancer, small cell lung cancer, hepatocellular carcinoma and other cancer indications. According to a report by IMARC Group, the global immune checkpoint inhibitor market size will reach US\$43.1 billion in 2023, and the market size is expected to reach US\$157.2 billion by 2032. Immune checkpoint inhibitors are drugs that treat cancer by inhibiting specific proteins produced by immune system cells, such as T cells and cancer cells. It can be used alone or in combination with other cancer treatments, including chemotherapy and radiation therapy, to further enhance the effectiveness of the treatment.



(v) EG1216X

Multiple myeloma is a disease caused by the malignant proliferation of plasma cells in the bone marrow. The normal function of plasma cells in the human body is to produce antibodies to resist invasion by foreign germs. Usually the number is not large. When specific germs enter the human body, the production of antibodies will be started. However, the plasma cells of multiple myeloma patients can only produce one type of antibody and cannot resist foreign germs. In addition, excessive production of plasma cells invades the bone marrow cavity, resulting in multiple myeloma patients with low immune function and very vulnerable to infection. Bacterial infection. Diagnosis is often delayed due to atypical symptoms. As the global population continues to age, the incidence rate of myeloma is also increasing year by year. According to 2022 The Lancet Haematology, there are approximately 176,000 newly diagnosed patients worldwide every year, accounting for 14% of blood tumors. Currently, there are more than 700 newly diagnosed cases of myeloma in Taiwan every year. DARZALEX has also driven a substantial growth in its revenue due to the excellent results of its clinical trials. According to J&J's 2023 annual financial report, global sales reached US\$9.7 billion, an increase of 22% from 2022

(B) New dosage forms and new drug delivery systems of biosimilars

(i) EG13084

EG13084 is a new subcutaneous injection drug that combines Trastuzumab and Pertuzumab. Its R&D target is PHESGO-approved indications for early breast cancer (EBC) and metastatic breast cancer (MBC). Since Roche obtained the marketing approval of PHESGO from the EMA and FDA in 2020, it has begun to strategically and actively convert PHESGO in Europe and the US for patients who use Trastuzumab and Pertuzumab in combination. According to Roche's 2023 annual financial report: the global annual sales of this product have to 1.21 billion CHF, with an annual growth rate of 64%. Among them, the European and US markets account for 83%. According to the excellent results of the APHINITY trial (Roche published in 2022, Pertuzumab combined with Trastuzumab and chemotherapy in the treatment of early breast cancer: this combination therapy can reduce the risk of recurrence or death by 23% compared with the control group and), it is foreseeable that EG13084 and EG1206A will jointly expand the treatment scope of patients and continue to expand the market of dual-target therapy in the treatment of HER2 breast cancer

(ii) EG7412X

According to the Hyaluronidase Market Size, Share & Trends Analysis Report 2023-2030 report, the global market value of hyaluronidase (hyaluronidase) in 2023 will be US\$910 million, of which "recombinant human hyaluronidase PH20 (rHuPH20) "Accounting for 23%. Due to its stable purity, the market share is expected to grow more rapidly. It is estimated to grow at a compound growth rate of 9.4% by 2030 and is widely used in new dosage forms of pharmaceuticals.

(C) Special biological drugs

(i) EG74032

EG74032 can be widely used in vaccine products and used as a carrier to make mixed vaccines. A number of vaccines with products such as CRM197 have been launched, and a number of vaccines are also in clinical development. Take Prevnar13 produced by Pfizer as an example. This vaccine combines the capsular antigens of Streptococcus Pneumonia serotypes 1, 3, 4, 5, 6A, 6B, 7F, 9V, 14, 18C, 19A, 19F, and 23F. The sugar suspension is chemically linked to the carrier protein to form a mixed vaccine. It is also used in the clinical development products of many large international pharmaceutical companies (such as Novartis and Mitsubishi Tanabe Pharmaceuticals) to produce multiple types of mixed vaccines, such as Haemophilus type B vaccine, typhoid vaccine or meningitis vaccine, demonstrating its wide application sex.

D. Competition for the Company's Products

Global biotech and pharmaceutical companies are optimistic about the subsequent development potential of the biosimilar drugs and ADC market. International major manufacturers have invested in this market, including biotech giants Amgen and Biogen. Well-known multinational pharmaceutical companies include Sandoz, Pfizer, Merck, and Eli. Lilly et al. In addition to the major international pharmaceutical companies actively exploring the field of biosimilar drugs through joint cooperation or mergers and acquisitions, there are also many small and medium-sized biopharmaceutical companies joining this battle. However, due to the company's size or its own capabilities, they can only strategically Cutting into part of the biosimilar drug value chain, EirGenix is different from other small and medium-sized biopharmaceutical companies. EirGenix's structure



strengthens the company's international competitive advantage on the basis of CDMO. Its niche points are: 1. Through the R&D energy of the Taiwan team, master the cell line development capabilities and keep exclusive technology and manufacturing capabilities in Taiwan; 2. Implement process research and development, as well as product analysis and manufacturing technology through a team with international experience; 3. At the same time Possess Mammalian (mammalian cell line development) and Microbial (microbial cell fermentation) technology platforms; 4. Complete the commercial production base at the Zhubei factory to systematically retain production technology capabilities and provide competitive production costs and profits; 5. Have regulatory experience in international clinical applications and drug approvals. EirGenix is a professional biopharmaceutical company with high concentration of technology and experience. Its business strategy mainly focuses on two major directions: 1. Provide domestic and foreign biopharmaceutical development companies with highquality and market-competitive entrusted process development and production services. EirGenix has cGMP-related facilities that comply with international regulatory standards for clinical/marketed production of biopharmaceuticals. It has two cGMP plants for mammalian cell and microbial fermentation and related technical manpower, which has greatly improved the integrity and quality of the production system for clinical trials and marketed biopharmaceuticals. complementarity. 2. Simultaneously develop high-quality and market-competitive biopharmaceuticals/biosimilar drugs. Through the above two business axes, EirGenix can provide customers with high-quality and cost-effective biotech drug manufacturing services, and jointly develop high-quality and cost-effective biotech drugs with partners to benefit the global medical system and their needs. All patients treated will be the biggest beneficiaries.

(A) Biosimilar drugs

(i) Market Competition Analysis of EG12014

In recent years, although the global sales of Roche, the original developer of HERCEPTIN, have declined year by year due to the entry of biosimilars into the market competition, the global sales of related products developed with its main ingredient Trastuzumab have increased due to the increase in breast cancer patients worldwide and the launch of biosimilars (As of the date of publication, the US FDA has approved 5 items and the EU EMA has approved 7 items), which has improved treatment opportunities for patients and maintained growth. According to the 2024 Research and Markets report, global sales of trastuzumab biosimilars have reached 4.27 billion USD in 2023. With the extensive experience and marketing advantages of Sandoz, a strategic partner of EirGenix, in leading global biosimilars, EG12014 is expected to quickly gain market share after being launched in the European and American markets. In the Taiwan market, where EirGenix is responsible for its own marketing, although 6 Trastuzumab biosimilars have been approved for marketing, only 3 of them (including EIRGASUN 150 mg) have obtained a wider range of National health insurance payment conditions and are substantially competitive. EIRGASUN 150 mg is the only trastuzumab developed and manufactured in Taiwan, and under the premise of the government's policy to promote the stability of the biotechnology industry and supply chain, it is successively entering the procurement list of hospitals at all levels according to the plan, and gradually expanded the revenue.

In addition, in April 2019, EirGenix Inc. signed a licensing and co-development agreement with Sandoz AG, a global leader in generics and biosimilars. The agreement including the up-front payment, milestone by the stage and the royalty payment in the authorized markets after product launch. Sandoz is one of the three major business units of the Novartis Group, one of the top three pharmaceutical companies globally. It holds a leading position in the field of generic drugs and biosimilars. Its products in the biosimilars field accounted for over 22% of the global market, reaching \$1.9 billion in sales by 2020, making it the top player in the industry. With a long history and rich experience in the development and sales of biosimilar drugs and cancer medications, Sandoz aims to achieve \$6 billion in revenue from biosimilar drugs by 2030. This strategic cooperation is expected to improve the global competitiveness of EirGenix's production line, thus expanding EirGenix's overall operating scale and increasing profits, which is of great positive help to financial and business development.

(ii) Market Competition Analysis of EG1206A

Although Roche is currently marketing another subcutaneous injection drug PHESGO that combines Trastuzumab and Pertuzumab globally, which has affected the growth of PERJETA, according to Roche's 2023 annual financial report: the global annual sales of this product still reached 4.06 billion CHF, with an annual growth rate of 9%, and the European and US markets accounted for 54% of revenue contribution. At present, EirGenix's EG1206A R&D progress ranks among the



top three in the global pertuzumab biosimilars, which will be more conducive to seizing the biosimilar market after the expiration of Pertuzumab's patent. The current progress is preparing to enter phase 3 clinical trials, which are expected to be launched in the second half of 2024. Once EG1206A passes Phase III clinical trials and is successfully launched, it will be paired with EG12014 and EG13084 to further strengthen the integrity of the HER2 product portfolio and provide more treatment options for breast cancer patients around the world.

(iii) Market Competition Analysis of TSY0110 (EG12043)

In response to the increasing demand for research and production of anti-drug conjugates and highly cytotoxic/potent substances, many foreign CDMOs and CMOs have begun to expand their service energy, especially for highly cytotoxic/potent active pharmaceutical ingredient (API) and final products. As of March 2024, the U.S. FDA has approved 16 ADCs for marketing, and global pharmaceutical companies are also competing to invest resources. This is also a battleground for the future cancer treatment market. With the cGMP plant, EirGenix has the capability of developing and manufacturing a monoclonal antibody drug process. At the same time, EirGenix forms a strategic alliance with Formosa Laboratories, which has the most experience in developing and manufacturing high-activity raw materials in Taiwan and has experienced antibody drug development and cGMP production talents and international cooperation networks, which is conducive to the development of ADC platform.

technology.

(iv) Market Competition Analysis of EG1211X

Immune checkpoint inhibitors target specific signaling pathways in the immune system to further develop more precise cancer treatments. Compared with traditional chemotherapy, it has the advantages of lower toxicity and fewer side effects, making it better tolerated by patients. The current trend in cancer treatment also focuses on using new sequencing technologies to identify specific genetic mutations in cancer cells, thereby driving the global demand for immune checkpoint inhibitors and becoming the mainstream of current cancer treatment. In the near future, several after the patent of immune checkpoint inhibitors expires, the introduction of biosimilar drugs to provide more affordable treatments with the same efficacy will also be the focus of cancer treatment development.

(v) Market Competition Analysis of EG1216X

J&J currently markets DARZALEX, a breakthrough CD38 monoclonal antibody therapy, in both IV and subcutaneous dosage forms globally, in addition to expanding the market and maintaining its competitive advantage through the patent protection strategy of subcutaneous dosage forms.

(B) New dosage forms and new drug delivery systems of biosimilars

(i) Market Competition Analysis of EG13084

Roche is currently marketing PHESGO replacement Pertuzumab in combination with trastuzumab and chemotherapy globally, emphasizing convenience and maintaining the clinical benefit of this combination therapy in reducing the risk of recurrence or death by 23% compared to the control group. It is foreseeable that the EG13084 that EirGenix focuses on R&D will work with EG1206A to expand the scope of treatment for patients and continue to expand the market of dual-target therapy in HER2 breast cancer treatment, which is also one of the important axes of EirGenix's EG13048 expanding its breast cancer treatment production line.

(ii) Market Competition Analysis of EG7412X

At present, "recombinant human hyaluronidase PH20 (rHuPH20)" is still monopolized by the American company Halozyme with ENHANZE's patented technology and has been licensed to a number of major international pharmaceutical companies for the application of new subcutaneous injection dosage forms of star drugs.

(C) Special biological drugs

(i) Market Competition Analysis of EG74032

At present, there are very few manufacturers with the production capacity of this product in the market. The main manufacturers are as follows:



Company	Location of Manufacturing Plant	Product
SynCon Bio	Netherlands	Production of CRM197 with mutant Diphtheria Bacillus;
Partner	(Amsterdam)	Provision of Prevnar® and Meningites® manufactured by
		Pfizer, and Menveo® manufactured by Novartis
Pfenex,	India	Provision of various specifications of CRM197 raw
Inc.		materials, including raw materials conforming to cGMP
		specifications (CRM197 is expressed with pseudomonas
		fluorescent as host)

EirGenix uses a unique microbial expression system and process to produce high-purity EG74032. Compared with other CRM197 products currently on the market, EirGenix's EG74032 uses a microbial (E. Coli.) expression system and process to produce high-purity EG74032. With a high degree of competitive advantage, EirGenix has completed the development and mass production of the EG74032 process. The current mass production scale can reach 150 liters of fermentation tanks.

(3) An overview of the company's technologies and its research and development work

A. The company's technologies and its research and development work

EirGenix is a professional biopharmaceutical company with a high density of technology and experience, whose business strategy is mainly in two directions:

- (A) Provide domestic and foreign biopharmaceutical development companies with high-quality and market-competitive entrusted process development and production services. EirGenix has cGMP-related facilities that comply with international regulatory standards for clinical/marketed production of biopharmaceuticals. It has two cGMP plants for mammalian cell and microbial fermentation and related technical manpower, which has greatly improved the integrity and quality of the production system for clinical trials and marketed biopharmaceuticals. complementarity.
- (B) Simultaneously develop high-quality and market-competitive biopharmaceuticals/biosimilar drugs.

Through the above two business axes, EirGenix can provide customers with high-quality and cost-effective biotech drug manufacturing services, and jointly develop high-quality and cost-effective biotech drugs with partners to benefit the global medical system and their needs. All patients treated will be the biggest beneficiaries.

B. Successfully Developed Technique or Product

EirGenix and its subsidiary provide contracted process development and production service with high quality and market competitiveness for biopharmaceutical development companies at home and abroad. EirGenix has CGMP-related facilities that conform to international regulations and standards and can be used for biopharmaceuticals' clinical/market production. At the same time, EirGenix has two CGMP plants for mammalian cell and microbial fermentation and related technical manpower, greatly improving the integrity and complementarity of clinical trials and market biotech drug production systems. EirGenix has accumulated technology for many years, continuously introduced international talents and novel equipment to maintain international competitiveness and meet the requirements of international laws and regulations, and continuously carried out technology development.

C. Research and Development Implementation Progress

Item	Indications		Implementation Progress
		•	EirGenix's "Trastuzumab" has obtained API license
			and API master file number from the Food and Drug
			Administration of Taiwan's Ministry of Health and
	Early Breast Cancer (EBC),		Welfare.
EG12014	Metastatic Breast Cancer	•	The biosimilar "EIRGASUN 150 mg" has obtained a
	(MBC) / Metastatic Gastric		marketing approval from Taiwan's Ministry of Health
	Cancer		and Welfare and has been approved by the National
			Health Insurance Agency of the Ministry of Health and
			Welfare to be reimbursed by health insurance benefits.
			The latest health insurance drug price in 2024 was



Item	Indications	Implementation Progress
		NT\$11,323/tube. • HERWENDA-Trastuzumab biosimilar EG12014 (150 mg, for intravenous use) has obtained marketing approval from EC.
EG1206A	Early Breast Cancer (EBC), Metastatic breast Cancer (MBC) /	EirGenix's EG1206A achieved successful phase I clinical results in May 2023 and the current progress is preparing to enter phase III clinical trials, which are expected to be launched in the second half of 2024.
TSY0110 (EG12043)	Breast Cancer /Gastric Cancer	EG12043 (TSY0110) has received positive results from the Scientific Advice Meeting of the European Medicines Agency (EMA); and the Biosimilar Drug Development Type 2 Meeting (BPD Type 2) of the US Food and Drug Administration (FDA) and expect to apply for the phase I clinical trials in second half of 2024.
EG1211X	Locally Advanced or Metastatic Urothelial Cancer, Locally Advanced or Metastatic Non-Small Cell Lung Cancer, Triple-Negative Breast Cancer, Small Cell Lung Cancer, Hepatocellular Carcinoma	Preclinical evaluation trials are currently underway.
EG1216X	Multiple Myeloma	Preclinical evaluation trials are currently underway.
EG13084	Early Breast Cancer (EBC), Metastatic breast Cancer (MBC) /	At present, the development of subcutaneous injection is underway.
EG7412X	Early Breast Cancer (EBC), Metastatic breast Cancer (MBC) /	At present, the development of subcutaneous injection is underway.
EG74032	Conjugate Vaccine	At present, EirGenix has completed the development and pilot run of the EG74032 process, with the current production scale reaching a 150-liter fermentation tank, which has been sold at home and abroad.

D. R&D Expenses As of the date of issuance for the annual report in the latest 5 year

Unit: NT\$ thousands; %

Year Item	2019	2020	2021	2022	2023
R&D Expenses(A)	959,610	1,561,722	893,510	800,144	952,290
Net Operation Revenue(B)	476,085	1,071,838	1,697,359	1,481,017	1,022,653
(A) / (B)	202	146	53	54	93

(4) Long-term and Short-term Development

- A. The short-term development strategy is "Build up the foundation and move forward step by step." The strategy plans for products in development and CDMO sales & marketing development are as follows:
- (A) EG12014 approved by the FDA and other countries in Aisa.
- (B) EG12014 (HERWENDA® Sandoz | EIRGASUN® EirGenix) market launch.
- (C) EG1206A submit the application for Phase III trials.



- (D) Application for EG12043 (TSY0110) clinical trials (IND).
- (E) EG1211X pre-clinical preparation completed.
- (F) Expansion of Building B at Zhubei plant to increase the microbial capacity to 1,500 L in 2026.
- B. The medium and long-term development strategy is "Products are developing and launching one after another to promote stable growth in revenue. The strategy plans for products in development and CDMO sales development are as follows:
- (A) New dosage forms or new drug delivery systems of biosimilars: development of Trastuzumab high-concentration subcutaneous doses; planning for the development of EG12014+EG1206A dual-targeting high-concentration subcutaneous doses. The successful development of high-concentration subcutaneous doses will strengthen the product market share of these products and enable EirGenix as the primary supplier of biosimilar drugs for the treatment of HER2+ breast cancer.
- (B) Developing the biosimilar for the treatment of blood cancer are currently ongoing. According to the development schedule, one new product will be introduced to the market each one to two years starting in 2027. Hence, a three-stage expansion of the mammalian capacity by 150,000L is under planning at Ciaotou Science Park, Kaohsiung. The new capacity can be used to manufacture in-house developed drugs and accept customers' orders for commercial and scale production.

2. Market and Sales Overview

(1) Market Analysis

A. Sales (Service) Region

Unit: NT\$ thousands; %

Year	202	23	2022		
Area	Amount	%	Amount	%	
Taiwan	474,123	46.36	746,845	50.43	
Japan	186,584	18.25	133,023	8.98	
USA and Canada	119,293	11.67	304,969	20.59	
Europe	242,186	23.68	283,328	19.13	
Other Regions	467	0.04	12,853	0.87	
Total	1,022,653	100	1,481,017	100	

B. Market Share

At present, the CDMO business of EirGenix is extremely competitive in the market. In addition to a certain proportion of the market share in Taiwan, EirGenix has actively expanded to Asian markets such as Japan and the mainland; In the future, EirGenix will be committed to increasing the market share of Europe, America, and other regions. The self-developed products of EirGenix have not been sold on the market before, so there is no market share analysis at present.

C. Supply and Demand of the Market and Growth in the Future

(A) Biosimilar drugs

(i) EG12014

EG12014 is a biosimilar of Trastuzumab, and its R&D target HERCEPTIN has global annual sales of 1.79 billion CHF according to Roche's 2023 financial report, of which the European and US markets account for 42%. According to data from the Taiwan Health Insurance Administration, breast cancer ranked second among the top ten cancer health insurance medical expenses in Taiwan in 2022, with drug expenditures of 9.075 billion NTD (the average growth rate from 2018 to 2022 was 7.75%). Among them, the breast cancer target drug HERCEPTIN, Taiwan's health insurance expenditure in 2022 was as high as 1.8 billion NTD. In 2024, the latest health insurance drug price of HERCEPTIN frozen crystal injection form (440 mg) was NT\$43,236 per tube. Currently, the National Health Insurance reimbursed Trastuzumab for patients with early breast cancer (EBC), metastatic breast cancer (MBC) and metastatic gastric cancer (mGC). The number of breast cancer and gastric cancer patients in Taiwan increases every year, and medical expenses also increase accordingly. The latest health insurance price of EirGenix's EIRGASUN 150 mg in 2024 is NT\$11,323/tube, which brings more benefits to more people with a more affordable price and wider treatment benefits. Patients who need to undergo breast cancer



treatment can achieve expected clinical efficacy while reducing their medical expenses by using EG12014 (EIRGASUN 150 mg), which has no clinically significant difference in safety, quality and efficacy from HERCEPTIN, to achieve the purpose of truly benefiting the people.

(ii) EG1206A

EG1206A is a biosimilar of Pertuzumab. Since its R&D target PERJETA was launched in the United States in 2013, sales have grown rapidly every year. According to the results of the APHINITY trial of Pertuzumab combined with Trastuzumab and chemotherapy in the treatment of early breast cancer published by Roche in 2022, this combination therapy can reduce the risk of recurrence or death by 23% compared with the control group. From this excellent test results, it is foreseeable that the subsequent product development and therapeutic application of EG1206A will be more extensive. Roche is currently actively promoting the expansion of indications and is expected to continue to expand the treatment scope and market. Although Roche is currently marketing another subcutaneous injection drug PHESGO that combines Trastuzumab and Pertuzumab globally, which has affected the growth of PERJETA. According to Roche's 2023 annual financial report: the global annual sales of this product still reached 4.06 billion CHF, with an annual growth rate of 9%, and the European and US markets accounted for 54% of revenue contribution.

(iii) TSY0110 (EG12043)

Roche's Trastuzumab emtansine (T-DM1) has been successfully marketed by using Linker to bind Trastuzumab antibody and cytotoxic chemical DM-1. The conjugate (Linker, Connector) of T-DM1 will not affect the antibody-dependent cytotoxic activity and will not interfere with the antibody neutralization activity dominated by HER2. Therefore, T-DM1 not only retains the anticancer effect of Trastuzumab but also enables the powerful cytotoxic drugs attached to it to exert a stronger effect. In Phase III clinical trials (EMILIA trial*) for patients with advanced breast cancer patients of HER2 overexpression and who have used Trastuzumab or Taxane, it was found that patients receiving T-DM1 treatment have a longer and better progression-free survival (T-DM1: 9.6 months; Lapatinib plus capecitabine: 6.4 months)(HR: 0.65; 95%CI: 0.55-0.77; P<0.001) and overall survival (T-DM1: 30.9 months; Lapatinib plus capecitabine: 25.1 months) (HR: 0.68; 95% CI: 0.55-0.85; P<0.001) over patients treated with Lapatinib in combination with Capecitabine. It shows that this kind of treatment is progress that cannot be ignored in both clinical medicine and pharmacology.

As far as the global market is concerned, there are very few companies in the world that can provide ADC development services. EirGenix has formed a strategic alliance with Formosa Laboratories. According to the schedule plan, they will jointly become one of the world's leading manufacturers in the ADC field in the shortest possible time to expand their competitive advantage.

(iv) EG1211X

EG1211X is a biosimilar of Atezolizumab. Its R&D target Tecentriq is an immune checkpoint inhibitor (Immune Checkpoint Inhibitor). According to Roche's 2023 annual financial report, global annual sales reached 3.77 billion CHF, with an annual growth rate of 9%, of which the European and US markets accounted for 73%. As the world's first approved PD-L1 immune checkpoint inhibitor, Roche has invested considerable resources in clinical trials for multiple cancer types, and has successively obtained results for locally advanced or metastatic urothelial cancer, locally advanced or metastatic non- Small cell lung cancer, triple negative breast cancer, small cell lung cancer, hepatocellular carcinoma and other cancer indications. According to a report by IMARC Group, the global immune checkpoint inhibitor market size will reach US\$43.1 billion in 2023, and the market size is expected to reach US\$157.2 billion by 2032. Immune checkpoint inhibitors are drugs that treat cancer by inhibiting specific proteins produced by immune system cells, such as T cells and cancer cells. It can be used alone or in combination with other cancer treatments, including chemotherapy and radiation therapy, to further enhance the effectiveness of the treatment.

(v) EG1216X

Multiple myeloma is a disease caused by the malignant proliferation of plasma cells in the bone marrow. The normal function of plasma cells in the human body is to produce antibodies to resist invasion by foreign germs. Usually the number is not large. When specific germs enter the human body, the production of antibodies will be started. However, the plasma cells of multiple myeloma patients can only produce one type of antibody and cannot resist foreign germs. In addition, excessive production of plasma cells invades the bone marrow cavity, resulting in multiple myeloma patients with low



immune function and very vulnerable to infection. Bacterial infection. Diagnosis is often delayed due to atypical symptoms. As the global population continues to age, the incidence rate of myeloma is also increasing year by year. According to 2022 The Lancet Haematology, there are approximately 176,000 newly diagnosed patients worldwide every year, accounting for 14% of blood tumors. Currently, there are more than 700 newly diagnosed cases of myeloma in Taiwan every year. DARZALEX has also driven a substantial growth in its revenue due to the excellent results of its clinical trials. According to J&J's 2023 annual financial report, global sales reached US\$9.7 billion, an increase of 22% from 2022.

(B) New dosage forms and new drug delivery systems of biosimilars

(i) EG13084

EG13084 is a new subcutaneous injection drug that combines Trastuzumab and Pertuzumab. Its R&D target is PHESGO-approved indications for early breast cancer (EBC) and metastatic breast cancer (MBC). Since Roche obtained the marketing approval of PHESGO from the EMA and FDA in 2020, it has begun to strategically and actively convert PHESGO in Europe and the US for patients who use Trastuzumab and Pertuzumab in combination. According to Roche's 2023 annual financial report: the global annual sales of this product have to 1.21 billion CHF, with an annual growth rate of 64%. Among them, the European and US markets account for 83%. According to the excellent results of the APHINITY trial (Roche published in 2022, Pertuzumab combined with Trastuzumab and chemotherapy in the treatment of early breast cancer: this combination therapy can reduce the risk of recurrence or death by 23% compared with the control group and), it is foreseeable that EG13084 and EG1206A will jointly expand the treatment scope of patients and continue to expand the market of dual-target therapy in the treatment of HER2 breast cancer

(ii) EG7412X

According to the Hyaluronidase Market Size, Share & Trends Analysis Report 2023-2030 report, the global market value of hyaluronidase (hyaluronidase) in 2023 will be US\$910 million, of which "recombinant human hyaluronidase PH20 (rHuPH20) "Accounting for 23%. Due to its stable purity, the market share is expected to grow more rapidly. It is estimated to grow at a compound growth rate of 9.4% by 2030 and is widely used in new dosage forms of pharmaceuticals.

(C) Special biological drugs

(i) EG74032

This product is a carrier protein made of diphtheria toxin through genetic improvement and amino acid replacement.

At present, it has been applied in commercially available vaccines of Pfizer and Novartis, with annual sales of billions of US dollars. In 2015, the global vaccine market value was nearly US\$ 30 billion, while the market value of conjugate vaccines with carrier protein reached up to US\$ 7 billion. These show that the future market for such products is quite amazing.

This protein has been widely used in commercially available products and clinical development products. There are more than 60 completed or ongoing clinical trials with relevant products on ClinicalTrials.gov of the National Institute of Health. Among them, only Novartis has carried out more than 20 clinical trials, which shows that the conjugate vaccine with this carrier protein has a great market.

In addition to the traditional vaccines for infectious disease, the research and development of cancer vaccines have attracted much attention in recent years. Both R&D organizations and biotech companies at home and abroad are actively investing in the development of cancer vaccines, hoping to bring a glimmer of light to all kinds of cancers that have no effective therapy yet.

In the early process, Corynebacterium diphtheria was used for production, followed by downstream recovery, purification, and other steps, and the yield and recovery rate were usually relatively low. In addition, for the acquisition of the strain, a license agreement needs to be signed with a specific organization, and the production organization needs to have a biological protection level that meets the standards before production. Compared with the above processes, the microbial expression platform is used for the EG74032 process developed by EirGenix, which will achieve the effects of a quite high yield and purity as well as immune enhancement. Compared with traditional processes or other products on the market, EirGenix's products will be competitive in quality and price and are expected to be widely used in research and development or marketing products.

EirGenix and its subsidiary CRM197 is expected to occupy a certain market share in the academic and pre-clinical



markets.

D. Competitive niche

(A) The advanced technology platform of EirGenix helps customers shorten the time schedule of biopharmaceutical development.

(i) Cell line development

Customers only need to provide DNA expression sequences or amino acid sequences of proteins, and EirgGenix can complete the establishment of high-expression cell lines. EirgGenix possesses dual-platform cell lines, including the internationally renowned Life Technologies CHO-S cell line and our self-developed EG CHO K1 sv cell line platform. This platform includes cell lines, expression vectors, and culture media. Under normal culture conditions, it can achieve antibody protein yields of 3-10 grams per liter, reducing costs for customers in both drug development and commercial production stages.

(ii) Process development and Scale-up

In terms of cell culture process, with AmbrTM micro bioreactor, EirGenix can simulate the culture conditions of the large-volume bioreactor in a 15 ml test tube and can control and adjust a number of parameters on a small scale to achieve the comprehensive effect of saving time and cost.

In terms of process amplification, at present, many projects, including the customer's products and the EG12014 product within EirGenix, have successfully entered the cGMP plant for the product at 200/4,000 liters or more. EirGenix has mastered the setting of various important parameters in bio-fermentation tanks of various scales in the process amplification of the cGMP plant. Once the culture condition parameters of small-scale fermentation tanks from 2 to 5 liters are available, they can be successfully amplified to a scale of 200, 500, 1,000, to 4,000 liters. This technology platform can save customers the time and various costs required in the process amplification.

(iii) Protein analysis and identification

Due to the characteristics of biopharmaceuticals, in the production process, each batch of products cannot be 100% the same. Various analytical methods are needed to identify the characteristics of the protein drugs produced. In addition, corresponding analytical methods are needed to detect the amino acid sequence of the protein, the purity of the product, impurity produced in the process, the activity of the protein, and the monitoring of microorganisms that may cause biosafety issue. The team of EirGenix has established the full identification, characterization and analytical methods of protein drugs internally to ensure the strictest control of product quality and safety at all stages and reduce the risks caused by unstable product quality in the drug development process.

(iv) Provide a full range of services to meet the needs of customers:

EirGenix also operates mammalian and microbial production facilities that adhere to PIC/S cGMP standards and can offer comprehensive services tailored to clients' requirements. The mammalian cell production line at the Xizhi Facility includes one cell bank production line, two upstream production suites, and one downstream purification production suite. The upstream production suites feature reactors with scales of 50-200-1000 liters and another suite with scales of 50-200 liters, all utilizing single-use bioreactors. In the Xizhi Facility's microbial cell production line, there is one upstream production suite and one downstream purification production suite, equipped with a 20L stainless-steel fermentor and a 100L stainless-steel fermentor in the upstream production line. At the Zhubei Facility, there are two mammalian cell production lines, each equipped with a 200-2000 (2x1000L)-12000 (6x2000L) liters single-use bioreactor upstream production suite and one downstream production suite. The use of single-use bioreactors offers advantages such as reduced cross-contamination and simplified equipment setup. Throughout the drug development phase, our company ensures maximum flexibility and a broad range of options in process development to accommodate the diverse needs of clients' drug development projects.

(B) EirGenix has formed the strategic industrial alliance with Formosa Laboratories in large molecule drugs and small molecular drugs.

At present, Formosa Laboratories, with a number of raw materials and anti-sunburn series active ingredient products marketed all over the world as well as ISO certificate, has successfully passed the GMP plant inspection certification of



Taiwan Department of Health, FDA of the United States, BGV of Germany, EDQM of the European Union and PMDA of Japan, which is a major manufacturer of small molecule raw materials in the world. At present, Formosa Laboratories has built a high-activity raw material production plant for the production of small molecule drugs with high toxicity. In combination with EirGenix's ability to produce large molecule antibody drugs, EirGenix and Formosa Laboratories have jointly established a production platform for ADC, with EirGenix producing antibodies, Formosa Laboratories producing small molecule drugs and carrying out antibody-drug conjugation, and EirGenix carrying out various identification and analysis related to ADC products. The establishment plan of this production platform has been subsidized and supported by the Ministry of Economy's Industry Development Technology Plan. The alliance between EirGenix and Formosa Laboratories has made the establishment of the ADC production platform completed and will make EirGenix one of the few CMO companies in the world that can produce antibody-drug conjugates.

E. Favorable and Unfavorable Factors in the Long Term

(A) Favorable factors

- (i) EirGenix has protein drug development platform technology and a cGMP pilot plant. In coordination with the toxicology laboratory and bio-safety testing laboratory previously established by the Biotechnology Center, EirGenix can integrate the upstream, midstream, and downstream protein drug R&D chains and provide a series of complete technical services.
- (ii) At the same time, EirGenix has rich experience in cell line cloning and microbial process technology development and continues to introduce domestic and foreign experienced and technical talents. Good production and development quality, good manpower quality, low turnover rate, and high work efficiency can shorten the biopharmaceutical development time.
- (iii) The relevant GMP production facilities comply with international regulations (including FDA GMP and PIC/S GMP), which is conducive to obtaining foreign sources of cases. Through business cooperation with strategic alliance partners, CDMO business has expanded rapidly.
- (iv) The protein-drug market continues to grow, and there is still a wide range of therapeutic applications to be developed. Drugs have entered preclinical and Phase I/II clinical trials one after another. There is a high demand for CDMO at this scale at home and abroad. Upstream R&D organizations at home and abroad have invested heavily in the research and development of biopharmaceuticals. The number of pipelines for bio-pharmaceuticals continues to increase. There is an urgent need for mid-stream research and development institutions that can undertake research and development results in order to extend the results to pre-clinical and Phase I/II clinical trials. The demand for microbial fermentation systems is gradually increasing in biopharmaceutical companies at home and abroad. The establishment of a CGMP microbial fermentation system can be applied not only to mature microbial expression systems such as E. coli and Pichia but also to the mass production and development of DNA vaccines.
- (v) The government actively constructs an environment conducive to the development of the bio-pharmaceutical industry, including tax exemption and tax relief, to further enhance the competitiveness of domestic manufacturers.
- (vi) EirGenix's development of biosimilars follows the international guidance, and its quality has the competitive strength of major international factories. With the gradual development of products, active international cooperation will be conducive to the deep roots of the brand in the near future.

(B) Unfavorable factors and countermeasures

(i) There are relatively few CDMO projects in the late development stage and commercial production.

Countermeasures: EirGenix has completed the construction of the first cell line 12,000L (3 sets of 2x2,000 L) on the 3F for commercial scale production in the Hsinchu Biomedical Park. It was put into operation in January 2019 and mainly used for the production of our own products (EG12014 and EG1206A). The second production line 12,000L (3 sets of 2x2,000 L) on the 5F has been launched in November 2023. Currently, we have several projects in hand and are continuously communicating with domestic and foreign biopharmaceutical companies, including biosimilar drug developers, to seek opportunities for late development stage and commercial scale production cooperation. The



commercial microbial production plant (350 L+1,000 L and 2x DSP suites) in Building B of the Hsinchu Park has started construction and is scheduled to be completed and put into operation in 2026. Some production capacity is currently reserved for the future needs of existing customers, and we will continue to communicate with domestic and foreign biopharmaceutical companies to seek projects for late development stage and production of CGT projects as well as DNA plasmids and enzymes related to mRNA.

(ii) Foreign bio-pharmaceutical manufacturers have been actively deployed, and their brand advantages will form pressure.

Countermeasures: EirGenix has established an experienced R&D team to continuously enter the market as early as possible with the development efficiency of new products (Biosimilar, Biobetter, etc.) through the improvement of R&D technology capabilities, passing US FDA, EMA, or Japan PMDA inspections will increase customer confidence, and to reduce the threat of price competition through the expansion of factories under the reduction of production costs. In addition, EirGenix has actively cooperated with local pharmaceutical companies to carry out clinical development, product production, and marketing.

(iii) For biosimilars products, the long R&D time is long and high fund investment make it unfavorable to the P&L.

Countermeasures: EirGenix has evaluated the R&D of a series of new indications related to HER2 with its professional development capability so as to expand the product market effectively and continuously carry out product life cycle management; has also sought strategic alliances and shared with domestic and foreign partners and combined with the capital market to ensure the smooth marketing of products.

- (2) Production Procedures of Main Products
 - A. Major Products and Their Main Uses
 - (A) Biosimilar drugs

(i) EG12014

So far, there are three approved indications of Trastuzumab by the FDA of the United States, including the treatment of patients with early breast cancer (EBC), metastatic breast cancer (MBC), and metastatic gastric cancer (mGC) of HER2 overexpression or HER2 gene amplification. In early breast cancer (EBC), it includes (1) adjuvant therapy after surgery and chemotherapy (preoperative or postoperative). (2) Adjuvant therapy of Doxorubicin and Cyclophosphamide combined with Paclitaxel or Docetaxel. (3) Adjuvant therapy of Docetaxel combined with Carboplatin. (4) Preoperative chemotherapy and postoperative adjuvant therapy are used to treat locally advanced (including inflammatory) breast cancer or tumor (diameter > 2 cm); In metastatic breast cancer (MBC): (1) it is used alone for metastatic breast cancer that has been treated with chemotherapy once or more; Unless the patient is not suitable for using Anthracycline or Taxane, the previous chemotherapy should include at least Anthracycline or Taxane. It is used for patients positive in hormone receptors who have failed in hormone therapy unless the patient is not suitable for hormone therapy. (2) Combined with Paclitaxel or Docetaxel for patients with metastatic breast cancer that has not been treated with chemotherapy. (3) Combined with aromatic cyclase inhibitors for patients with hormone receptor-positive metastatic breast cancer; In metastatic gastric cancer (mGC), combined with capecitabine (or 5-fluorouracil) and cisplatin for the treatment for metastatic gastric adenocarcinoma (or gastroesophageal junction adenocarcinoma) of HER2 over-expression that has not been treated with chemotherapy.

(ii) EG1206A

EG1206A is a biosimilar of Pertuzumab. The reference drug of Pertuzumab, Perjeta, is very promising in clinical efficacy and sales prospect. As the incidence of breast cancer tends to be younger year by year and the 5-year recurrence rate of HER2+ early breast cancer is 17~40%, the utilization rate of EG1206A will continue to grow. It is estimated that the market demand for this drug will gradually increase in the future.



(iii) TSY0110 (EG12043)

Antibody-drug conjugates (ADC) are undoubtedly the most selective anti-cancer therapy for tumors, but their performance is not protruding in drug delivery. Therefore, antibody-drug conjugates (ADC) need to be combined with powerful drugs. At present, ADCs are commonly combined with maytansinoids (T-DM1) and dolastatin analogs (brentuximab vedotin), which act on microtubules and can inhibit microtubule kinetics. Once reaching picomolar concentration (pM), these drugs can show super activity to inhibit cell growth; Therefore, ADC is expected to kill the enemy more effectively by accurately "targeting" these cytotoxic drugs into the anti-cancer battlefield. After T-DM1 passes the examination at an extremely fast speed, the research on ADC becomes hot.

(iv) EG1211X

The R&D target Tecentriq is an Fc-engineered, humanized monoclonal antibody that binds to PD-L1 and blocks its interaction with PD-1 and B7.1 receptors. Atezolizumab is an unglycosylated IgG1 kappa immunoglobulin with a calculated molecular weight of 145 kDa. Its pharmacological effect is that PD-L1 can be expressed on tumor cells and/or tumor-infiltrating immune cells, and can inhibit the anti-tumor immune response in the tumor microenvironment. PD-L1 binds to PD-1 and B7.1 receptors on T cells and antigen-presenting cells to inhibit the activity of cytotoxic T cells, T cell proliferation and cytokine production. Atezolizumab is a monoclonal antibody that binds to PD-L1 and blocks its interaction with PD-1 and B7.1 receptors. This releases PD-L1/PD-1-mediated suppression of immune responses, including activation of anti-tumor immune responses without inducing antibody-dependent cellular cytotoxicity. In syngeneic mouse tumor models, blocking PD-L1 activity resulted in reduced tumor growth. Indications span a variety of cancer types: 1. Locally advanced or metastatic urothelial cancer is suitable for the treatment of patients with locally advanced or metastatic urothelial cancer whose disease has worsened after receiving platinum-containing chemotherapy or who are not suitable for cisplatin-containing therapy. 2. Used alone for locally advanced or metastatic non-small cell lung cancer. It is suitable for the treatment of patients with locally advanced or metastatic non-small cell lung cancer whose disease has worsened after receiving platinum-containing chemotherapy. If patients have EGFR or ALK tumor gene abnormalities, they must first be treated with EGFR or ALK inhibitors. If the disease worsens after treatment, Tecentriq can be used. Used in combination with Avastin (bevacizumab), paclitaxel and carboplatin, as a first-line treatment for metastatic non-squamous non-small cell lung cancer without EGFR or ALK tumor gene abnormalities. 3. The combination of Tecentriq and nabpaclitaxel for triple-negative breast cancer is suitable for the treatment of unresectable locally advanced or metastatic triplenegative breast cancer, and the tumor has PD-L1 manifestations (tumor-infiltrating immune cells (IC) \geq 1%) and has not received Chemotherapy is used for patients with metastatic breast cancer. 4. Small cell lung cancer, combined with carboplatin and etoposide, is suitable for the first-line treatment of adults with extensive stage small cell lung cancer. 5. The combined use of hepatocellular carcinoma and bevacizumab is suitable for the treatment of patients with hepatocellular carcinoma who have not received systemic therapy and are unresectable or metastatic, and their liver function is Child-Pugh A.

(v) EG1216X

The target of its R&D, DARZALEX, is a humanized IgG1 κ monoclonal antibody that can bind to the CD38 antigen. This strain is produced in a mammalian cell line (Chinese Hamster Ovary [CHO]) through recombinant DNA technology. Its pharmacology Mechanism is an IgG1 κ human monoclonal antibody (mAb) that binds to the CD38 protein that is highly expressed on the surface of multiple myeloma cells and to varying degrees on other cell types and tissues. CD38 protein has multiple functions, such as receptor-mediated adhesion, signaling and enzyme activity. DARZALEX is suitable for: 1. Patients who have received at least three previous therapies (including a protease inhibitor and an immunomodulator) as a single drug, or whose disease has worsened under treatment with both a protease inhibitor and an immunomodulator (double-refractory to a protease inhibitor and an immunomodulatory agent) in adults with multiple myeloma. 2. Use in combination with lenalidomide plus dexamethasone or with bortezomib plus dexamethasone to treat adult patients with multiple myeloma who have previously received at least one therapy.



(B) New dosage forms and new drug delivery systems of biosimilars

(i) EG13084

EG13084 is a new subcutaneous injection drug that combines Trastuzumab and Pertuzumab. Its R&D target is PHESGO-approved indications for early breast cancer (EBC) and metastatic breast cancer (MBC). In terms of early breast cancer (EBC), when EG13084 is used in combination with chemotherapy drugs for preoperative adjuvant therapy, it is suitable for patients with HER2-positive, locally advanced, inflammatory or early breast cancer (tumor diameter greater than 2cm or positive lymph nodes). Part of a complete treatment prescription. When used as adjuvant treatment after surgery, it is suitable for patients with early-stage breast cancer (EBC) who are HER2-positive and have a high risk of recurrence. (According to the results of the APHINITY clinical trial, in postoperative adjuvant therapy, patients with HER2-positive early breast cancer who have a high risk of recurrence are defined as having lymph node-positive breast cancer.) In terms of metastatic breast cancer (MBC), it is used together with docetaxel to treat metastasis patients with HER2-positive metastatic breast cancer who have not been treated with anti-HER2 or chemotherapy.

(ii) EG7412X

The R&D target of EG7412X is "recombinant human hyaluronidase PH20 (rHuPH20)", which is equivalent to the soluble fragment of human HYAL5. It is genetically engineered in hamster culture cells (CHO) culture containing a DNA plasmid encoding the enzyme. Produced as a type of hyaluronidase (hyaluronidase), its main function is to catalyze the degradation of hyaluronic acid (HA). According to the enzyme reaction products, they are divided into three different types: two eukaryotic endoglycosidase hydrolases and one prokaryotic lyase-type glycosidase. Hyaluronidase catalyzes the hydrolysis of HA, causing the viscosity of HA to decrease, thereby increasing tissue permeability. Therefore, it is often used in medicine together with other drugs, which can accelerate the dispersion and delivery of drugs through subcutaneous absorption.

(C) Special biological drugs

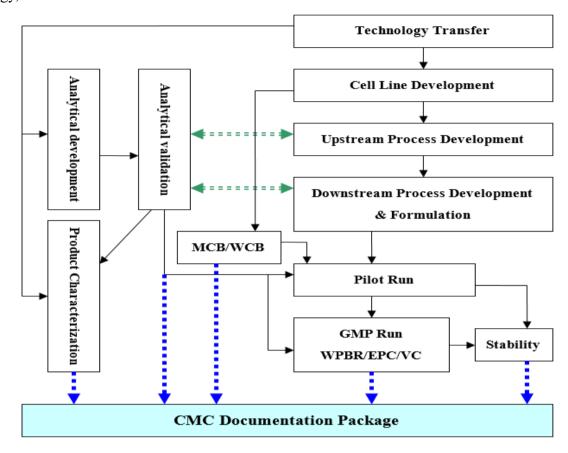
(i) EG74032

It is mainly used in vaccine products and used as a carrier to make conjugate vaccines. There are many commercially available conjugate vaccines, including Pfizer's Prevnar® and Prevnar® 13, Wyeth/Pfizer's Meningitec® and Novartis's Menveo®. At present, as a variety of products have been approved as infectious disease vaccines in Europe and the United States, representing that regulatory organizations have no doubts about the safety and effectiveness of CRM197 in improving immunity, and the subsequent technical and regulatory risks of acting as the raw material of conjugate vaccines are small, much academic research and clinical development projects are in progress.



B. Major Products and Their Production Processes

EirGenix's main core value in the product production process lies in the vertical integration of the upper, middle, and lower reaches of the industrial value chain. From cell line development and cell culture to process development and amplification to downstream product purification and drug stability analysis, EirGenix can master the industrial chain by itself, accurately master technology, and control costs.



Note: MCB/WCB (mother cell bank/working cell bank); WPBR (work production lot report); EPC (end production cell); VC (viral effectiveness)

C. The supply situation for the company's major raw materials

EirGenix's main service items are bio-pharmaceutical contract development & manufacturing organization (CDMO) and self-developed monoclonal antibody bio-similar drugs. The main raw materials are culture medium, buffer solution, chromatographic resin, single-use filter element, packaging materials, etc. The supply sources shall conform to international medical standards. In order to ensure the stable supply of raw materials, an inventory of qualified manufacturers is established to ensure that other supply sources meet the quality requirements.

D. A list of any suppliers and clients accounting for 10 percent or more of the company's total procurement (sales) amount in either of the two most recent fiscal years, the amounts bought from (sold to) each, the percentage of total procurement (sales) accounted for by each, and an explanation of the reason for increases or decreases in the above figures. Where the company is prohibited by contract from revealing the name of a client, or where a trading counterpart is an individual person who is not a related party, it may use a code in place of the actual name.

(A) Major Suppliers Information for the Last Two Calendar Years

As of the date of issuance for the annual report, the business of EirGenix is development of Biosimilars and new drugs, and the major revenue comes from CDMO.

Unit: NT\$ thousands

		2022		2023				
Item	Company	A	Danasant	Relation	Company	A	0/	Relation
	Name	Amount	Percent	With Issuer	Name	Amount	%	With Issuer
1	Merck	135,142	25.48	No	Cytiva	116,665	27.56	No
2	Pall Singapore	115,931	21.86	No	Merck	116,351	27.49	No
3	Life Tech	60,895	11.48	No	Sartorius	49,139	11.61	No
4	Sartorius	58,942	11.11	No	Others	141,153	33.34	No
5	Others	159,523	30.07	No	-	-	-	-



		2023						
Item	Company Name	Amount	Percent	Relation With Issuer	Company Name	Amount	%	Relation With Issuer
6	-	-	-	-	-	-	-	-
	Net Purchases	530,433	100	No	Net Purchases	423,308	100	No

Description of change:

EirGenix and its subsidiary mainly provide biopharmaceutical contract development & manufacturing organization (CDMO) business, and raw materials are mainly purchased according to the progress of each production process. Therefore, the main suppliers set different purchase prices according to the progress of their cases every year.

The industries of EirGenix and its subsidiary are bio-similar drug research and development and CDMO services. The main items purchased are protein ion exchange resin, culture medium, culture bag, filter element, reagent, and general consumables. In order to ensure stable supply quality and consistent comparison basis of experimental data, and some raw materials will be purchased from specific manufacturers according to project requirements. Raw materials cannot be arbitrarily changed in each process stage to avoid affecting test results, so it is a characteristic of the industry that a designated supplier supplies a single source of raw materials. Among the suppliers, Cytiva mainly supplies membrane, liquid handling bag, and mixer bag. Pall Singapore mainly supplies some cell culture medium and purified colloid, and Merck mainly supplies reagents and culture bags. All three companies are internationally renowned biotechnology research and development factories. Apart from good quality and stable supply, they can also provide relevant data and technical support and supporting documents required for drug inspection and registration.

In the development of the Company's products, the applicability of the products is still used to test and screen the required raw materials. Except for some CDMO customers who demand specified materials, the raw materials selected by the Company will be put into production lines only after research and development tests and evaluations. The raw materials of the three major companies are on the candidate list and have no absolute dependence.

Major international biotech factories have a stable supply. If there is any commodity shutdown for the project to be shut down, they will announce in advance and propose alternative commodity schemes and provide sample tests to solve the risk of refueling or material cut-off connection. At present, the newly developed cases of EirGenix and its subsidiary, the medium protein ion exchange resin, colloid, filter, and bag, have been successfully replaced by other brands. The raw materials selected by the Company can be put into the production line only after research and development tests and evaluations. Consider providing customers with better quality and competitive advantages in price, and increasing alternative applicable materials of suppliers, research and development units will also consider using other alternative products or collecting relevant information on products in the market at the initial stage of research and development or testing, so as to cope with and reduce the risk of over-reliance on specific manufacturers.

If there are supply risks in the future, the Company has the ability to select suitable alternative materials from other supply brands through the research and development technology platform to reduce the risks and make the supply risks within the controllable range.

(B) Major Clients Information for the Last Two Calendar Years

Unit: NT\$ thousands

		2022		2023				
Item	Company Name	Amount	Percent	Relation with Issuer	Company Name	Amount	Percent	Relation with Issuer
1	Company MV	514,208	34.72	No	Company G	174,644	17.08	No
2	Company SA	261,876	17.68	No	Company SA	144,479	14.13	No
3	Company BO	168,647	11.39	No	Company MV	125,354	12.26	No
4	Others	536,286	36.21	No	Company HB	113,042	11.05	No
5	-	-	-	_	Company HC	109,281	10.68	No



		2022	2023					
Item	Company Name	Amount	Percent	Relation with Issuer	Company Name	Amount	Percent	Relation with Issuer
6	-	-	-	-	Others	355,853	34.80	No
	Net Sales	1,481,017	100	No	Net Sales	1,022,653	100	No

Description of change:

At present, the main source of income for EirGenix and its subsidiary is the bio-pharmaceutical contract development & manufacturing organization (CDMO). As service income and GMP production are recognized according to various contracts, services provided, or undergoing production progress, the sales number of major sales clients varies according to the progress of their cases each year. In addition, in April 2019, the Company signed a license agreement for the co-development of the breast cancer biosimilar EG12014 (Trastuzumab Biosimilar) with SA. By the end of 2023, the Company had completed the requirements of milestones from Phase I to Phase VI. Therefore, the revenues of contract payment and milestone payment from Phase I to Phase VI were recognized in stages in accordance with standard accounting procedures.

All revenues from the main clients of EirGenix and its subsidiary are presently recognized as the revenue from development and manufacturing services, except for the revenue from the license agreement for the co-development gradually recognized with SA over time. At present, for the development and manufacturing services commissioned by clients, the service contents include the preliminary development work and the back end GMP production, including process verification and verification work. Several clients are close to the listing stage and are also discussing the long-term production of future listing supply. Once the client's products are successfully listed, it is expected to have a long-term stable income from said product supply. As revenue is still growing in the past few years, in the future, with the launch of production plants year by year, the Company will look for potential clients of later products and products to be marketed closely and continue to maintain stable, productive creation and considerable revenue.

At the same time, the continuous expansion of foreign cases is also a medium-term plan to create value. It is obvious that European and American clients are increasing year by year. Due to its characteristics, if a good client relationship is well maintained and quality is ensured, then it is also the keyway to obtain stable considerable revenue. In addition to client maintenance, EirGenix and its subsidiary have also obtained the certification of foreign factories from Japan's Ministry of Health, Laboure, and Welfare. It is expected that in the future, they will gradually obtain internationally important legal certifications from the European Union, the United States, and other countries, which will be more conducive to accelerating the growth of revenue.

E. An indication of the production volume for the two most recent fiscal year

The Company is a CDMO company, and its output value is determined according to the work items of the commissioned case, and there are no products with fixed mass production. The self-developed bio-similar drug products are still in the research and development stage and have not been officially mass-produced and sold, so they are not applicable.

F. An indication of the volume of units sold for the two most recent fiscal year.

Unit: NT\$ thousands

Volume Year		2022				2023			
	Lo	cal Expo		ort	Lo	Local		Export	
Major goods	Quantity	Amount	Quantity	Amount	Quantity	Amount	Quantity	Amount	
Service Revenue	-	330,747	-	426,933	-	350,011	-	255,979	
Sales Revenue	-	416,098	-	45,363	-	124,112	-	151,079	
Licensing Revenue	-	-	-	261,876	-	-	-	141,472	
Total	-	746,845	-	734,172	-	474,123	-	548,530	



Description of change:

The major revenue of EirGenix is a CDMO business, and its output value is determined according to the work items of the commissioned case, and there are no products with fixed mass production. Due to EirGenix increases the contract from current clients and finds new clients to make the performance grow stably. In April 2019, EirGenix Inc. signed a global licensing agreement for EG12014 (Trastuzumab Biosimilar) with Sandoz AG. As of the 2023 Q4, EirGenix has received the signing fee and fulfilled the requirements of its first to sixth milestone; the revenue from the milestone payment will be recognized in stages in accordance with standard accounting procedures.

3. The number of employees employed for the two most recent fiscal years and during the current fiscal year up to the date of publication of the annual report, their average years of service, average age, and education levels.

Unit: Person; age; years; %

Ye	Year		2023	2024 until the end of March
	Management		19	19
F 1	Supervisor	30	31	33
Employees	Staff	358	338	328
	Total	408	388	380
Average Yo	ears of Age	36.38	37.71	37.9
Average Tea	rs of Service	3.84	3.94	3.9
	Ph.D.	7.8	8.8	9.5
F1	Master's	68.6	68	67.1
Education	Bachelor's	23.6	23.2	23.4
	High School	0	0	0



4. Disbursements for Environmental Protection

(1) Any losses suffered by the company in the most recent fiscal year and up to the annual report publication date due to environmental pollution incidents (including any compensation paid and any violations of environmental protection laws or regulations found in environmental inspection, specifying the disposition dates, disposition reference numbers, the articles of law violated, and the content of the dispositions), and disclosing an estimate of possible expenses that could be incurred currently and in the future and measures being or to be taken. If a reasonable estimate cannot be made, an explanation of the facts of why it cannot be made shall be provided:

Since its establishment, EirGenix has been committed to environmental protection, which complies with relevant laws and policies of government on environmental protection. Therefore, the Company has not had any environmental pollution as of the date of issuance for the annual report in the last two years. In the future, the Company will continue to adhere to its consistent philosophy to maintain the best environmental protection results.

5. Labor Relations

(1) List any employee benefit plans, continuing education, training, retirement systems, and the status of their implementation, and the status of labor-management agreements and measures for preserving employees' rights and interests.

A. Employee Reward System

The Company rewards system relates to employee individual performance, his/her contribution to the company, and his/her personal work profession and job levels, as well as the Company's business performance. While the Company is profitable in the current year, 1-5% of the Company annual profit shall be issued to employees as a reward. Employee compensation consists of three parts: salary, bonus, and welfare. Employee salary is related to his/her work profession and position, while the bonus is connected to individual performance, department contribution, and annual company business performance results. As for welfare to employees, it must be not only in compliance with laws and regulations from the government and also designed to meet employee's needs from all aspects. The Company also issues stock-related rewards as welfare, such as Employee Stock Options, Restricted Stock Units, IPO/SPO reserved stock options for employees. All these non-cash rewards to employees are provided to share our accomplishments and also to retain and grow with employees.

B. Workplace diversity and equity

The Company respects the value of diversity and provides all the employees regardless of his/her race, ethnicity, gender, gender identity, sexual orientation, age and socioeconomical background with the equal opportunity on their job rights, compensation, and career development. The indigenous people and with disability are employed by the Company without any discrimination and inequality. In 2023, the new-hired female employee took 43.3%. In the entire company, the female employee is 43.8% and the female managers is 38.6%.

C. Employee Welfare

- (A) In order to promote employee physical and mental health to reach the work-and-life balance, the Company holds an inhouse Employee Welfare Committee according to the Employee Welfare Fund Act. The Company appropriates funds for the Committee to handle welfare issues for employees to promote internal morale and a cozy work environment.
- (B) Employee Leave Policy Superior to the regulated standards of Labor Standards Act and the Regulations of Leave-Taking of Workers
 - (i) Employees are provided with annual personal leave available since his/her first day on board and the paid day leave are higher than the government regulation requirement.
 - (ii) Paid family care leave.
- (C) Other benefits include flexibility of starting and finishing daily working time, wedding leave, funeral leave, hospitalization allowance, maternity allowance, pregnancy leave, employee lunch allowance, department teambuilding feasts, transportation allowance, welfare committee activities, employee outing allowance, and lottery draw-in the annual feast, group insurance, and occupational injury insurance. EirGenix care for the employees and their family members. The health insurance is covered to their family by their options. In the family day event by the Company, employees' family are invited.



D. Training and Development

Starting from the beliefs in lifelong learning, the Company provides learning for positive inter-promotion between work effectiveness, quality, and efficiency. The training program is annually planned to provide employee pre-service and on-the-job training. In addition to the 1st-day training for the newcomer to the Company, it covers three training topics, including professional, leadership and management, and core competency. Through these training courses, the profession of talents, the employee morale to the Company, and the competitiveness of the Company in the global industry are expected to develop in the meantime.

EIRGer's Learning Center is built to shape EirGenix into a learning organization. Also known as ELC, it provides employees with diversified training courses annually. Professional courses take the majority and follow with leadership program and core competency training:

- (A) Experts Program. The training covers professional topics such as cGMP, CMC, biologics, and manufacturing.
- (B) Leadership Program. This program is designed for the current managers and potential supervisors, in which management skills, team building, communication, coaching, strategical thinking, and leadership mindset are provided.
- (C) Common Knowledge Program, as known as core competency training, in which ELC intends to build up morale and teamwork for employees, and also most common knowledge education and training courses are designed to develop employees.

In 2023, ELC offered 15 courses over 70 course hours, with a total of 1364 participants and a total of 4,388 study hours.

E. Retirement Policy

Employees may apply for retirement under any of the following conditions:

- (A) Where the employee attains the age of 55 and has worked for 15 years.
- (B) Where the employee has worked for more than 25 years.
- (C) Where the employee attains the age of 60 and has worked for 10 years.

In compliance with the Labor Pension Act and the "Monthly Contribution Classification of Labor Pension" issued by the government, the Company has the obligations to bear pension contribution amounts for each employee no less than 6% of his/her monthly salary and save in his/her personal pension account. Since the establishment of the Company, two employees have retired, and retirement-related matters have been handled in accordance with the provisions of the Labor Pension Act

The Company has setup an employee benefits trust fund program, which inspires employees through linking long-term benefit plans with the Company's operating performance.

F. Labor-Management Dispute

The Company communicates with employees not only through Town Hall Meeting and Labor-Management Meeting but also through internal emails, office displays, and suggestion boxes for employees to provide their opinions at any time. The Company also meets the needs of employees in a timely manner through communication, education, and incentive mechanisms. The Company has not had any disputes between employers and employees requiring settlements in 2023.

G. Other Employee Rights Mechanism

The Company has a sound system, which sets out various management policies, specifies the employee rights, obligations, and welfare, and regularly reviews and revises the welfare contents to safeguard the rights and interests of all employees.

(2) List any losses suffered by the company in the most recent two fiscal years and up to the annual report publication date due to labor disputes (including any violations of the Labor Standards Act found in labor inspection, specifying the disposition dates, disposition reference numbers, the articles of law violated, the substance of the legal violations, and the content of the dispositions), and disclosing an estimate of possible expenses that could be incurred currently and in the future and measures being or to be taken. If a reasonable estimate cannot be made, an explanation of the facts of why it cannot be made shall be provided:

Since the establishment, EirGenix's labor relations have been harmonious without any loss caused by the labor-management dispute. In the future, both employees and EirGenix should complement each other and grow together to manage the relationships with the heart to avoid the risks of loss caused by the labor-management dispute.



6. Cyber security management

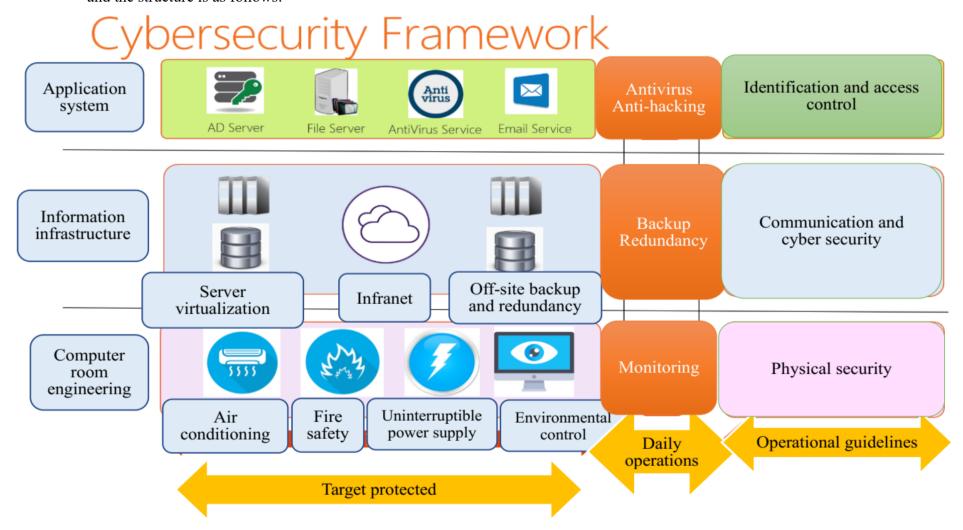
EirGenix has included information security in the annual audit project, regularly reviewed and evaluated security measures, and regularly changed various security settings while updating the system and working with professional vendors to ensure information and network security. Furthermore, to ensure that our information system can continue to provide stable services, we have established various redundancy mechanisms and backup systems and improved relevant processes as appropriate and upgraded computer software and hardware in response. The Information Technology Department often sends information security information to employees via emails and reports information security issues to the Board on March 8, 2024.

EirGenix joined TW-CERT as a member to receive TW-IASC information on information security in real-time. The person in charge of information security and information system updated or adjusted internal information-related equipment, architecture, and operation procedure shortly after considering the risk level, applicability and feasibility to reduce the possibility of severe damage caused by different forms of internal and external information security risks.

(1) Describe the cyber security risk management framework, cyber security policies, concrete management programs, and investments in resources for cyber security management.

A. Cyber Security Risk Management Framework

We have also established an information security risk management framework to reduce the risk of unknown information security threats caused by changes in the internal and external information environment. To reduce the unknown information security risks caused by new information technologies adopted and changes in the external environment, the Information Technology Department is responsible for coordinating information security and relevant matters and formulating internal information security plans. After such plans are approved, the department should conduct information security risk management as per the standard operating procedures, regularly examine internal information security, raise personnel's awareness of information security, and perform information security drills. The Company's information security framework is designed in a layered manner, and the structure is as follows:



B. Cyber Security Policies

It aims to achieve the purpose of sustainable corporate development, ensure the effective operations of the Company's information systems to support the normal operations of various business activities, and ensure continuous operations to minimize operating losses. When all employees of the Company use information-related systems, this information security management policy is used as the basis for management and compliance.

The information system security policy is divided into the aspects below:



- (A) System and regulations: Update relevant information security management regulations, infrastructure, systems, and information security protection technologies in line with relevant laws and regulations and changes in the Company's business and information technologies to maintain the confidentiality, integrity, and availability of our important information systems, and continuously protect information from various threats. The permissions management and changes of the important information systems should be recorded as a basis for auditing.
- (B) Information technology management: Update and evaluate information systems in real-time and execute necessary control measures to ensure the security of data, systems, networks, and information infrastructure.
- (C) Personnel and organization: The Information Technology Department should offer information security education and training to raise internal personnel's awareness of information security and improve their relevant professional skills.
- C. Cyber Security Management Programs and Investments in Resources for Cyber Security Management

The Company actively strengthens the security of the overall information system. Relevant matters, from the information security regulations to the design of information infrastructure, system maintenance and upgrading, professional personnel's training, and raising of employees' awareness of information security, are all included in the scope of information security. We self-examine information security every year to see if relevant systems are aligned with the changes in the environment and make timely adjustments according to needs. We adopted the Taiwan Intellectual Property Management System (TIPS) in 2021 to strengthen the management of the Company's confidential information. Our specific information security management measures implemented are as follows:

Category	Description	Operating method
Permissions management	Personnel and group accounts and verification methods management, permissions management, and system management permissions management	 Personnel accounts management operations should proceed or be changed after an application is filed and approved by responsible managers in accordance with the operating procedures. Each user's use permissions should be immediately revoked after resignation or job change to prevent unauthorized access. Regularly review system-related permissions. Manage system account life cycle and permissions accounts. Adopt multi-factor authentication and designated login to manage important systems.
Access management	Data flow control and auditing, physical equipment access management, audit records, and incident investigation	 Revise data flows into and out of important information systems and keep records of the access for auditing. Conduct physical security protection of the information system console. Analyze audit records and issue automatic warnings of abnormalities. Identify the information security level according to the importance and the degree of risk. Adopt digital rights management technology for important files to control the data flow to avoid unauthorized access.
Threat and risk management	Rate the information risks that may be caused by internal employees, external personnel, and potential vulnerabilities in the systems and take measures to reduce risks	 Standardize the user's computer preset. Launch operating regulations for external vendors to access the Company's information systems. Launch risk assessment procedures for adoption of new technologies. Deploy multiple brands' multi-layer firewalls and cloud email filtering to reduce the chance of external cyber-attacks and intrusion of phishing emails. Strengthen endpoint security, regularly update users' computers, and install antivirus software. Regularly offer information security education and training to improve personnel's awareness of information security.
System integrity and availability management	Maintain the availability and integrity of data and	 The host has been virtualized in a cluster to improve the availability of systems. Adopt large storage devices, regularly automate on-site and off-site backups, and perform recovery tests as planned to ensure the integrity and availability of



Category	Description	Operating method
	systems to resume	systems.
	normal operations in	 Adopt multiple redundancy mechanisms for infrastructure, multiple UPS
	the event of a disaster	systems with automatic generators, N+1 and 1+1 fan coil units, as well as
	or damage	multiple redundancy measures for internal and external network wires and
		equipment to reduce the chance of information service interruption.

Investments in Resources for Cyber Security Management:

- (A) To enhance the confidential and sensitive data access control, our company has implemented Microsoft Azure Information Protection encryption solution. A total of NT\$600,000 was invested from 2022 to 2023, along with engagement of PricewaterhouseCoopers Cybersecurity Consultants to plan the introduction of cybersecurity regulations. It is planned to adopt NIST CSF related specifications over a three-year period.
- (B) In response to the above implementation, over 8 meetings have been held with consultants, attended by external consultants, as well as internal IT and cybersecurity personnel.
- (C) In 2023, there were a total of 8 meetings and discussions on information security-related issues.
- (D) In September 2023, the IT department issued the Information Security Learning Handbook-2023, primarily aimed at establishing concepts of employee information and network security. The target audience includes all employees of the company, who undergo training and assessment after completion.
- (E) The company has also joined TWCERT/CC for cybersecurity information sharing.
- (F) In addition to advocating cybersecurity-related issues in monthly meetings, the IT department promptly informs internal colleagues via communication software and email in case of any special cybersecurity incidents where the severity of the risk may affect the company's information security.
- (2) List any losses suffered by the company in the most recent fiscal year and up to the annual report publication date due to significant cyber security incidents, the possible impacts therefrom, and measures being or to be taken. If a reasonable estimate cannot be made, an explanation of the facts of why it cannot be made shall be provided: None.
- (3) Effects of and Response to Changes in Technology and the Industry Relating to Corporate Finance and Sale
 - A. The government has actively promoted the biotechnology industries in recent years, of which the biopharmaceutical industry has the characteristics of high technical threshold, long R&D cycle, high professional technical demand, and added value. The threshold for the industry is relatively high, so it is not easy to produce drastic changes in a short period of time. Moreover, the Company, with a high degree of professional R&D capability, can closely grasp technological changes and industrial changes and take appropriate countermeasures as needed. As of the date of issuance for the annual report in the latest year, EirGenix and its subsidiary have not had a significant impact on the Company's financial business due to technological changes and industrial changes.
 - B. Considering the rapid changes in information technology and the external environment, to reduce the impact of external changes on our finances, the Company plans to revise relevant information security policies suitable for the operations together with external professional information security consultants aligned with the NIST Cybersecurity Framework (CSF) and relevant standards in the industry. We will implement the revised policies accordingly and review and adjust them regularly as the benchmark for evaluation and judgment when adopting various information systems and services. We've successfully completed the 3rd phase of NIST CSF implementation in the first quarter of 2023.



7. Important Contracts

Type of Agreement		Period	Major Contents	Restrictions
Lease contract	Department Center for Biotechnology	2022/03~2026/12	Lease offices, laboratories, and plants	None.
Lease contract	Hsinchu Science Park Bureau, Ministry of Science and Technology	2016/11~2036/11	Lease the land for plant construction in Biomedical Park.	None.
CDMO	Company HB	2021/02 until project completion	Recombinant protein 1,000L GMP Production.	None.
CDMO	Company G	2021/04 until project completion	Tech transfer, Recombinant protein GMP production	None.
CDMO	Company BO	2021/05 until project completion	Sale of biologic substance for R&D and clinical trials	None.
CDMO	Company AS	2021/09 until IND application	Process development and GMP production	None.
CDMO	Company AP	2022/03 until IND application	Tech transfer, Process development & GMP production	None.
CDMO	Company G	2022/09 until project completion	Perform stability study of recombinant protein	None.
CDMO	Company HB	2023/02 until project completion	Recombinant protein 1,000L GMP Production.	None.
CDMO	Company OM	2023/02 until project completion	Antibody 2,000L GMP Production.	None.
CDMO	Company AD	2023/06 until project completion	Tech transfer, Process development, GMP production and Raw material purchase.	None.
CDMO	Company B	2023/06 until project completion	Recombinant protein GMP production	None.
CDMO	Company HC	2023/06 until project completion	Cell line development, Process development and Antibody GMP production.	None.
CDMO	Company IB	2023/09 until project completion	Cell line development, Process development and Antibody GMP production.	None.
CDMO	Company U	2023/09 until project completion	Antibody 2,000L GMP Production.	None.
CDMO	Company AM	2024/01 until project completion	Cell line development	None.
CDMO	Company BR	2024/01 until project completion	Raw material purchase.	None.
CDMO	Company HB	2024/01 until project completion	Recombinant protein 200L GMP Production.	None.
Co-development	Formosa Pharmaceuticals, Inc.	2022/03~	Co-developed the biosimilar drug TSY0110 (EG12043) of ADC for the treatment of breast cancer.	None.
License agreement	Company SA	2019/04~	Grant the exclusive rights to commercialize the biosimilar EG12014 in license market.	In accordance with that contract
Credit contract	Hua Nan Commercial Bank Ltd	2022/02~2027/03	Establishment of Facility and Production Equipment.	The funds are used to purchase machinery and equipment.



VI. Financial Information

1. Condensed balance sheets and income statements for the past five fiscal years, showing the name of the auditor CPA and the auditor CPA's opinion given

- (1) Condensed Balance Sheet Based on IFRS
 - A. Condensed Balance Sheet- Consolidated

Unit: NT\$ thousands

	Year		Financial	Summary for The	Last Five Years	
Item		2019	2020	2021	2022	2023
Current assets			1,494,307	9,070,266	8,287,878	6,915,506
Property, Plant and Equipment			1,851,850	1,886,824	2,608,848	3,337,685
Intangible assets			33,129	19,553	28,067	28,269
Other assets			455,929	464,230	922,363	881,099
Total assets			3,835,215	11,440,873	11,847,156	11,162,559
Current	Before distribution		642,163	703,216	730,892	707,165
liabilities	After distribution		642,163	703,216	730,892	707,165
Non-current liabil			1,287,435	308,906	433,386	437,931
Total liabilities	Before distribution		1,929,598	1,012,122	1,164,278	1,145,096
	After distribution		1,929,598	1,012,122	1,164,278	1,145,096
Equity attributabl shareholders of the			1,905,617	10,428,751	10,682,878	10,017,463
Capital stock			2,063,751	3,003,845	3,043,358	3,060,516
Capital surplus			2,813,974	10,475,952	7,734,141	7,830,216
Equity attributable to shareholders of the parent	Equity attributable to shareholders of the parent		(2,930,919)	(2,973,500)	(115,540)	(915,208)
Capital stock	Capital stock		(2,930,919)	(2,973,500)	(115,540)	(915,208)
Other equity inter	est	\	(41,189)	(77,546)	20,919	41,939
Treasury stock			-	-	-	-
Non-controlling interest		\	-	-	-	-
Total equity	Before distribution		1,905,617	10,428,751	10,682,878	10,017,463
	After distribution		1,905,617	10,428,751	10,682,878	10,017,463

Note: The financial data for the most recent years has been audited and attested by CPAs.



B. Condensed Balance Sheet- Individual

Unit: NT\$ thousands

	Year		Financial Sun	nmary for The La	ast Five Years	
Item		2019	2020	2021	2022	2023
Current assets		1,048,257	1,491,466	9,064,044	8,269,047	6,909,802
Property, Plant and Eq	uipment	1,878,776	1,851,325	1,885,858	2,607,958	3,337,069
Intangible assets		42,434	32,840 19,553 2		28,067	28,269
Other assets		448,318	456,627	466,522	927,285	888,648
Total assets		3,417,785	3,832,258	11,435,977	11,832,357	11,163,788
Current liabilities	Before distribution	480,325	639,798	698,320	716,093	708,394
Current naomities	After distribution	480,325	639,798	698,320	716,093	708,394
Non-current liabilities		1,082,589	1,286,843	308,906	433,386	437,931
Total liabilities	Before distribution	1,562,914	1,926,641	1,007,226	1,149,479	1,146,325
Total habilities	After distribution	1,562,914	1,926,641	1,007,226	1,149,479	1,146,325
Equity attributable to sthe parent	shareholders of	1,854,871	1,905,617	10,428,751	10,682,878	10,017,463
Capital stock		1,693,041	2,063,751	3,003,845	3,043,358	3,060,516
Capital surplus		2,055,782	2,813,974	10,475,952	7,734,141	7,830,216
Equity attributable to shareholders of the parent Capital stock	Equity attributable to shareholders of the parent	(1,889,249)	(2,930,919)	(2,973,500)	(115,540)	(915,208)
Capital Stock	Capital stock	(1,889,249)	(2,930,919)	(2,973,500)	(115,540)	(915,208)
Other equity interest		(4,703)	(41,189)	(77,546)	20,919	41,939
Treasury stock		-	-	-	-	-
Non-controlling interest		-	-	-	-	-
Total equity	Before distribution	1,854,871	1,905,617	10,428,751	10,682,878	10,017,463
Total equity	After distribution	1,854,871	1,905,617	10,428,751	10,682,878	10,017,463

Note: The financial data for the most recent years has been audited and attested by CPAs.



(2) Condensed Statement of Comprehensive Income – Based on IFRS

A. Condensed Statement of Comprehensive Income- Consolidated

Unit: NT\$ thousands

					111: N I \$ thousands		
Year		Financial Summary for The Last Five Years					
Item	2019	2020	2021	2022	2023		
Operating revenue		1,071,838	1,697,359	1,481,017	1,022,653		
Gross profit		750,667	1,093,054	756,452	236,741		
Income (Loss) from operations		(986,004)	(58,311)	(330,819)	(1,031,977)		
Non-operating income and expenses		(55,319)	17,146	216,504	118,327		
Income (Loss) before tax		(1,041,323)	(41,165)	(114,315)	(913,650)		
Income (Loss) from Continuing Operation		(1,041,670)	(42,581)	(115,540)	(915,208)		
Income (Loss) from Discontinued Operation		-	-	-			
Net income (Loss)		(1,041,670)	(42,581)	(115,540)	(915,208)		
Other comprehensive income (income after tax)		259	5,335	59,311	46,118		
Total comprehensive income (Loss)		(1,041,411)	(37,246)	(56,229)	(869,090)		
Net income attributable to shareholders of the parent		(1,041,670)	(42,581)	(115,540)	(915,208)		
Net income attributable to non- controlling interest		-	-	-	-		
Comprehensive income attributable to Shareholders of the parent		(1,041,411)	(37,246)	(56,229)	(869,090)		
Comprehensive income attributable to non-controlling interest		-	-	-	-		
Earnings per share		(5.41)	(0.18)	(0.38)	(3.00)		

Note: The financial data for the most recent years has been audited and attested by CPAs.



B. Condensed Statement of Comprehensive Income- Individual

Unit: NT\$ thousands

Year	Financial Summary for The Last Five Years					
Item	2019	2020	2021	2022	2023	
Operating revenue	476,085	1,071,838	1,697,359	1,481,017	1,022,653	
Gross profit	254,667	750,667	1,093,054	756,452	236,741	
Income (Loss) from operations	(847,671)	(987,766)	(60,518)	(333,400)	(1,035,394)	
Non-operating income and expenses	(13,254)	(53,557)	18,126	218,198	120,651	
Income (Loss) before tax	(860,925)	(1,041,323)	(42,392)	(115,202)	(914,743)	
Income (Loss) from Continuing Operation	(860,925)	(1,041,670)	(42,581)	(115,540)	(915,208)	
Income (Loss) from Discontinued Operation	-	-	-	-	-	
Net income (Loss)	(860,925)	(1,041,670)	(42,581)	(115,540)	(915,208)	
Other comprehensive income (income after tax)	-	259	5,335	59,311	46,118	
Total comprehensive income (Loss)	(860,925)	(1,041,411)	(37,246)	(56,229)	(869,090)	
Net income attributable to shareholders of the parent	(860,925)	(1,041,670)	(42,581)	(115,540)	(915,208)	
Net income attributable to non- controlling interest	-	-	-	-	-	
Comprehensive income attributable to Shareholders of the parent	(860,925)	(1,041,411)	(37,246)	(56,229)	(869,090)	
Comprehensive income attributable to non-controlling interest	-	-	-	-	-	
Earnings per share	(5.39)	(5.41)	(0.18)	(0.38)	(3.00)	

Note: The financial data for the most recent years has been audited and attested by CPAs.

(3) Auditors' Opinions from 2019 to 2023

Year	CPA	Accounting Firm	Audit Opinion
2019	Shu-Fen Yu, Hui-Chin Tseng	PricewaterhouseCoopers Taiwan	Unmodified Opinion
2020	Sheng-Wei Deng, Shu-Fen Yu	PricewaterhouseCoopers Taiwan	Unmodified Opinion
2021	Sheng-Wei Deng, Yu-Fang Yen	PricewaterhouseCoopers Taiwan	Unmodified Opinion
2022	Sheng-Wei Deng, Yu-Fang Yen	PricewaterhouseCoopers Taiwan	Unmodified Opinion
2023	Sheng-Wei Deng, Yu-Fang Yen	PricewaterhouseCoopers Taiwan	Unmodified Opinion



2. Five-Year Financial Analysis

(1) Consolidated Financial Analysis – Based on IFRS

Year Items for Analysis		Fir	nancial Analysi	s for the Most l	Recent Five Yea	rs
		2019	2020	2021	2022	2023
Financial	Debt Ratio (%)		50.31	8.85	9.83	10.26
structure	Ratio of long-term capital to property, plant and equipment (%)		172.42	569.09	426.10	313.25
	Current ratio (%)		232.70	1,289.83	1,133.94	977.92
Solvency	Quick ratio (%)		195.19	1,215.91	1,015.88	868.41
	Interest coverage ratio		-	-	-	-
	Receivables turnover rate (times)		7.92	19.54	26.16	7.07
	Average collection days for receivables		46.09	18.68	13.95	51.63
	Inventory turnover rate (times)	\	1.98	2.10	1.26	1.05
Operating	Payables turnover rate (times)		11.08	9.47	6.56	7.34
Ability	Average days for sale	\	184.34	173.81	289.68	328.83
	Property, plant and equipment turnover rate (times)		0.57	0.91	0.66	0.34
	Total asset turnover rate (times)		0.30	0.22	0.13	0.09
	Return on assets (%)		(28.10)	(0.34)	(0.93)	(7.88)
	Return on equity (%)	\	(55.40)	(0.69)	(1.09)	(8.84)
Profitability	Ratio of income before tax to paid-in capital (%)		(50.46)	(1.37)	(3.76)	(29.85)
	Profit margin before tax (%)		(97.19)	(2.51)	(7.80)	(89.49)
	Earnings per share (NT\$)		(5.41)	(0.18)	(0.38)	(3.00)
	Cash flow ratio (%)	\	-	-	-	-
	Cash flow adequacy ratio (%)		-	-	-	-
	Cash reinvestment ratio (%)		-	-	-	-
Leverage	Operating leverage		-	-	-	-
Leverage	Financial leverage		-	-	-	-

Analysis of financial ratio differences for the last two years (2022& 2023) (Increase or decrease over 20%):

- 1. Ratio of long-term capital to property, plant and equipment: Mainly because of the capacity expansion of the mammalian cell production line at the 5th floor and the Zhubei plant Phase II facility that there was an increase in the amount of property, plant, and equipment.
- Receivables turnover rate (times) and Average collection days for receivables: Mainly because the collection of the account
 receivables increased at the end of 2023, leading to the decrease in receivables turnover rate and increase in average collection
 days.
- 3. Property, plant and equipment turnover rate (times): Mainly because of the capacity expansion of the mammalian cell production line at the 5th floor and the Zhubei plant Phase II facility that there was an increase in the amount of property, plant, and equipment, leading to the decreased in Property, plant and equipment turnover rate.
- 4. Total asset turnover rate (times): Total assets turnover rate decreased because of the capacity expansion and the increased of R&D expenses.
- 5. Return on assets: Mainly because of the increase of net loss after taxation in 2023 as compared with the same period of 2022.
- 6. Return on equity: Mainly because of the increase of net loss after taxation in 2023 as compared with the same period of 2022.
- 7. Ratio of income before tax to paid-in capital: Mainly because of the increase of net loss after taxation in 2023 as compared with the same period of 2022.
- 8. Profit margin before tax: Mainly because of the decrease of operation revenue and the increase of R&D expenses, leading to the increase in net loss after taxation.

Note: The following calculation formulas shall be listed at the end of this table in the annual report:

A. Financial Structure

- (A) Debt-asset ratio = total liabilities / total assets
- (B) Ratio of long-term capital to property, plant and equipment = (total equity + non-current liabilities) / net worth of property, plant and equipment

B. Solvency

- (A) Current ratio = current assets / current liabilities
- (B) Quick ratio = (current assets inventory prepaid expenses) / current liabilities
- (C) Interest coverage ratio = income before income tax and interest expenses / current interest



C. Operating ability

- (A) Receivables (including accounts receivable and notes receivable arising from business operations) turnover rate = net sales / average receivables (including accounts receivable and notes receivable arising from business operations) for each period
- (B) Average collection days for receivables = 365 / receivables turnover rate
- (C) Inventory turnover rate = cost of sales / average inventory
- (D) Payables (including accounts payable and notes payable arising from business operations) turnover rate = cost of sale / average payables (including accounts payable and notes payable arising from business operations) for each period
- (E) Average days of sale = 365 / inventory turnover rate
- (F) Property, plant and equipment turnover rate = net sales / average net worth of property, plant and equipment
- (G) Total asset turnover rate = net sales / average total assets

D. Profitability

- (A) Return on assets = [net income + interest expenses (1- tax rate)] / average total assets
- (B) Return on equity = net income / average total equity
- (C) Profit margin before tax = net income / net sales
- (D) Earnings per share = (profit and loss attributable to owners of the parent dividends on preferred shares) / weighted average number of issued shares (Note 4)

E. Cash flow

- (A) Cash flow ratio = Net cash flow from operating activities / current liabilities
- (B) Net cash flow adequacy ratio = Net cash flow from operating activities for the most recent five years / (capital expenditures + inventory increase + cash dividend)
- (C) Cash flow reinvestment ratio = (Net cash flow from operating activities cash dividend) / gross property, plant and equipment value + long-term investment + other non-current assets + working capital)

F. Leveraging

- (A) Operating leverage = (net operating revenue variable operating costs and expenses) / operating income
- (B) Financial leverage = operating income / (operating income / interest expenses)



(2) Individual Financial Analysis – Based on IFRS

Year		Financial Analysis for the Most Recent Five Years						
Items for Analysis		2019	2020	2021	2022	2023		
	Debt Ratio (%)	45.73	50.27	8.81	9.71	10.27		
	Ratio of long-term capital to property, plant and equipment (%)	156.35	172.44	569.38	426.24	313.31		
	Current ratio (%)	218.24	233.12	1,297.98	1,154.74	975.42		
Solvency	Quick ratio (%)	112.01	195.54	1,223.59	1,034.37	866.25		
	Interest coverage ratio	-	-	-	-	-		
	Receivables turnover rate (times)	5.20	7.92	19.54	26.16	7.07		
	Average collection days for receivables	70.19	46.09	18.68	13.95	51.63		
	Inventory turnover rate (times)	2.16	1.98	2.10	1.26	1.05		
Operating Ability	Payables turnover rate (times)	17.85	11.08	9.47	6.56	7.34		
1101111	Average days for sale	168.98	184.34	173.81	289.68	328.83		
	Property, plant and equipment turnover rate (times)	0.27	0.57	0.91	0.66	0.34		
	Total asset turnover rate (times)	0.15	0.30	0.22	0.13	0.09		
	Return on assets (%)	(26.47)	(28.11)	(0.34)	(0.93)	(7.89)		
	Return on equity (%)	(43.67)	(55.40)	(0.69)	(1.09)	(8.84)		
	Ratio of income before tax to paid-in capital (%)	(50.85)	(55.46)	(1.41)	(3.79)	(29.89)		
	Profit margin before tax (%)	(180.83)	(97.19)	(2.51)	(7.80)	(89.49)		
	Earnings per share (NT\$)	(5.39)	(5.41)	(0.18)	(0.38)	(3.00)		
	Cash flow ratio (%)	-	-	-	-	-		
	Cash flow adequacy ratio (%)	-	-	-	-	-		
	Cash reinvestment ratio (%)	-	-	-	-	-		
Loverson	Operating leverage	-	-	-	-	-		
Leverage	Financial leverage	-	-	-	-	-		

Analysis of financial ratio differences for the last two years (2022& 2023) (Increase or decrease over 20%):

- 1. Ratio of long-term capital to property, plant and equipment: Mainly because of the capacity expansion of the mammalian cell production line at the 5th floor and the Zhubei plant Phase II facility that there was an increase in the amount of property, plant, and equipment.
- 2. Receivables turnover rate (times) and Average collection days for receivables: Mainly because the collection of the account receivables increased at the end of 2023, leading to the decrease in receivables turnover rate and increase in average collection days.
- 3. Property, plant and equipment turnover rate (times): Mainly because of the capacity expansion of the mammalian cell production line at the 5th floor and the Zhubei plant Phase II facility that there was an increase in the amount of property, plant, and equipment, leading to the decreased in Property, plant and equipment turnover rate.
- 4. Total asset turnover rate (times): Total assets turnover rate decreased because of the capacity expansion and the increased of R&D expenses.
- 5. Return on assets: Mainly because of the increase of net loss after taxation in 2023 as compared with the same period of 2022.
- 6. Return on equity: Mainly because of the increase of net loss after taxation in 2023 as compared with the same period of 2022.
- 7. Ratio of income before tax to paid-in capital: Mainly because of the increase of net loss after taxation in 2023 as compared with the same period of 2022.
- 8. Profit margin before tax: Mainly because of the decrease of operation revenue and the increase of R&D expenses, leading to the increase in net loss after taxation.

Note 1: The following calculation formulas shall be listed at the end of this Table in the annual report:

A. Financial Structure

- (A) Debt-asset ratio = total liabilities / total assets
- (B) Ratio of long-term capital to property, plant and equipment = (total equity + non-current liabilities) / net worth of property, plant and equipment

B. Solvency

- (A) Current ratio = current assets / current liabilities
- $(B)\ Quick\ ratio = (current\ assets-inventory-prepaid\ expenses)\ /\ current\ liabilities$
- (C) Interest coverage ratio = income before income tax and interest expenses / current interest



C. Operating ability

- (A) Receivables (including accounts receivable and notes receivable arising from business operations) turnover rate = net sales / average receivables (including accounts receivable and notes receivable arising from business operations) for each period
- (B) Average collection days for receivables = 365 / receivables turnover rate
- (C) Inventory turnover rate = cost of sales / average inventory
- (D) Payables (including accounts payable and notes payable arising from business operations) turnover rate = cost of sale / average payables (including accounts payable and notes payable arising from business operations) for each period
- (E) Average days of sale = 365 / inventory turnover rate
- (F) Property, plant and equipment turnover rate = net sales / average net worth of property, plant and equipment
- (G) Total asset turnover rate = net sales / average total assets

D. Profitability

- (A) Return on assets = [net income + interest expenses (1- tax rate)] / average total assets
- (B) Return on equity = net income / average total equity
- (C) Profit margin before tax = net income / net sales
- (D) Earnings per share = (profit and loss attributable to owners of the parent dividends on preferred shares) / weighted average number of issued shares (Note 4)

E. Cash flow

- (A) Cash flow ratio = Net cash flow from operating activities / current liabilities
- (B) Net cash flow adequacy ratio = Net cash flow from operating activities for the most recent five years / (capital expenditures + inventory increase + cash dividend)
- (C) Cash flow reinvestment ratio = (Net cash flow from operating activities cash dividend) / gross property, plant and equipment value + long-term investment + other non-current assets + working capital)

F. Leveraging

- (A) Operating leverage = (net operating revenue variable operating costs and expenses) / operating income
- (B) Financial leverage = operating income / (operating income / interest expenses)
- Note 2: When the above formula for calculation of earnings per share is used during measurement, give special attention to the following matters:
 - A. Measurement should be based on the weighted average number of common shares, not the number of issued shares at year end.
 - B. In any case where there is a cash capital increase or treasury stock transaction, the period of time in circulation shall be considered in calculating the weighted average number of shares.
 - C. In the case of capital increase out of earnings or capital surplus, the calculation of earnings per share for the past fiscal year and the fiscal half-year shall be retrospectively adjusted based on the capital increase ratio, without the need to consider the issuance period for the capital increase.
 - D. If the preferred shares are non-convertible cumulative preferred shares, the dividend of the current year (whether issued or not) shall be subtracted from the net profit after tax or added to the net loss after tax. In the case of non-cumulative preferred shares, if there is net profit after tax, dividend on preferred shares shall be subtracted from the net profit after tax; if there is loss, then no adjustment need be made.
- Note 3: Give special attention to the following matters when carrying out cash flow analysis:
 - A. Net cash flow from operating activities means net cash in-flow amounts from operating activities.listed in the statement of cash flows.
 - B. Capital expenditures means the amounts of cash out-flows for annual capital investment.
 - C. Inventory increase will only be entered when the ending balance is larger than the beginning balance. An inventory decrease at year end will be deemed zero for calculation.
 - D. Cash dividend includes cash dividends from both common shares and preferred shares.
 - E. Gross property, plant and equipment value means the total value of property, plant and equipment prior to the subtraction of accumulated depreciation.
- Note 4: Issuers shall separate operating costs and operating expenses by their nature into fixed and variable categories. When estimations or subjective judgments are involved, give special attention to their reasonableness and to maintaining consistency.
- Note 5: In the case of a company whose shares have no par value or have a par value other than NT\$10, for the calculation of the above-mentioned paid-in capital ratio, the ratio of equity attributable to owners of the parent as stated in the balance sheet shall be substituted.



3. Supervisors'/Audit Committee's Report for the Most Recent Year:

Please refer to Appendix 2.

4. Financial statements for the most recent fiscal year, including an auditor's report prepared by a CPA, a two-year comparative balance sheet and income statement, statement of changes in shareholders' equity, cash flow statement, and any attached notes or appendices:

Please refer to Appendix 3.

5. Financial Statements for the Years Ended December 31, 2023 and 2022, and Independent Auditors' Report:

Please refer to Appendix 3.

6. If the company or its affiliates have experienced financial difficulties during the most recent fiscal year or the current fiscal year up to the date of printing of annual report, the annual report shall explain how said difficulties will affect the company's financial situation: None.



VII. Review of Financial Conditions, Financial Performance, and Risk Management

1. Financial Conditions

(1) Consolidated Financial Conditions

Unit: NT\$ thousands; %

Year	2022	2023	Difference		
Item	2022	2025	Amount	%	
Current Assets	8,287,878	6,915,506	(1,372,372)	(17%)	
Property, plant and equipment	2,608,848	3,337,685	728,837	28%	
Right-of-use Assets	325,330	329,236	3,906	1%	
Intangible Assets	28,067	28,269	202	1%	
Other Assets	597,033	551,863	(45,170)	(8%)	
Total Assets	11,847,156	11,162,559	(684,597)	(6%)	
Current Liabilities	730,892	707,165	(23,727)	(3%)	
Non-current Liabilities	433,386	437,931	4,545	1%	
Total Liabilities	1,164,278	1,145,096	(19,182)	(2%)	
Common Stock	3,043,358	3,060,516	17,158	1%	
Capital Surplus	7,734,141	7,830,216	96,075	1%	
Retained Earnings	(115,540)	(915,208)	(799,668)	692%	
Other Adjustments	20,919	41,939	21,020	100%	
Common control equity	-	-	-	-	
Total Shareholders' Equity	10,682,878	10,017,463	(665,415)	(6%)	

The major reason, impact and the response plan of the difference over 20% and the amount over 10 million:

(2) Individual Financial Condition

Unit: NT\$ thousands; %

Year	2022	2023	Difference		
Item	2022	2025	Amount	%	
Current Assets	8,269,047	6,909,802	(1,359,245)	(16%)	
Property, plant and equipment	2,607,958	3,337,069	729,111	28%	
Right-of-use Assets	325,330	329,236	3,906	1%	
Intangible Assets	28,067	28,269	202	1%	
Other Assets	601,955	559,412	(42,543)	(7%)	
Total Assets	11,832,357	11,163,788	(668,569)	(6%)	
Current Liabilities	716,093	708,394	(7,699)	(1%)	
Non-current Liabilities	433,386	437,931	4,545	1%	
Total Liabilities	1,149,479	1,146,325	(3,154)	0%	
Common Stock	3,043,358	3,060,516	17,158	1%	
Capital Surplus	7,734,141	7,830,216	96,075	1%	
Retained Earnings	(115,540)	(915,208)	(799,668)	692%	
Other Adjustments	20,919	41,939	21,020	100%	
Common control equity	-	-	-	-	
Total Shareholders' Equity	10,682,878	10,017,463	(665,415)	(6%)	

^{1.} Property, plant and equipment: Mainly because of the capacity expansion of the mammalian cell production line at the 5th floor and the Zhubei plant Phase II facility that there was an increase in the amount of property, plant, and equipment.

^{2.} Other Adjustments: Mainly because the increase in unrealized gains and losses on financial assets invested in unlisted companies in 2023.



The major reason, impact and the response plan of the difference over 20% and the amount over 10 million:

- 1. Property, plant and equipment: Mainly because of the capacity expansion of the mammalian cell production line at the 5th floor and the Zhubei plant Phase II facility that there was an increase in the amount of property, plant, and equipment.
- 2. Other Adjustments: Mainly because the increase in unrealized gains and losses on financial assets invested in unlisted companies in 2023.
- (3) The main reasons for any material change in the company's financial situation during the past 2 fiscal years, and describe the effect thereof:

The Board of Directors resolution of the establishment of Phase II Facility and production equipment in Hsinchu Biomedical Science Park, the upper limit is NT\$2.468 billion, and the Board of Directors resolution of the establishment R&D laboratory production line and production equipment, the upper limit is NT\$1.1425 billion. The source of funds comes from cash capital increase, equity fund and bank loan. After completion, can accelerate to meet the demand for international CDMO orders, meet the demand for the marketing products of the self-owned product EG12014, and can also be sufficient to provide the drugs for Phase III clinical trial and marketing production demand for subsequent product development. The construction of this new plant will become an efficient engine for the company's rapid growth in the future and promotion of momentum internationally, which will be of positive help to the company's finance and business.

Through the injection capital by private placement, the company can now accelerate the execution of its future strategic planning. For the product development unit, the product pipeline will be expanded to include more biosimilar drug products. For the CDMO unit, the current facility infrastructure will add additional production lines and facilities to handle even more diversified biological products and break into the field of cell and gene therapy, as well as the extension of services to further link upstream, midstream, and downstream development and manufacturing services. Lastly, EirGenix will seek to establish various forms of cooperation with international entities, which include but are not limited to collaborations, strategic alliances, or mergers and acquisitions. Soon EirGenix will become an important hub for biopharmaceutical development and manufacturing on the global stage.



2. Financial Performance

(1) List of Analysis of Financial Performance- Consolidated

Unit: NT\$ thousands; %

Year	2022	2023	Difference	%
Operating revenue	1,481,017	1,022,653	(458,364)	(31%)
Operating costs	724,565	785,912	61,347	8%
Gross profit (loss)	756,452	236,741	(519,711)	(69%)
Operating expenses	1,087,271	1,268,718	181,447	17%
Operating profit (loss)	(330,819)	(1,031,977)	(701,158)	212%
Non-operating income	226,879	130,479	(96,400)	(42%)
Non-operating expenses	(10,375)	(12,152)	(1,777)	17%
Profit (loss) before tax	(114,315)	(913,650)	(799,335)	699%
Income tax expense	(1,225)	(1,558)	(333)	27%
Net Profit (Loss)	(115,540)	(915,208)	(799,668)	692%

The main reason for the major change on Operating revenue, Net operating income (loss), and Income tax expense in currently 2 years:

- 1. Operating revenue and Gross profit: For CDMO business, because of the annual repairment in Xizhi facility and expansion and revalidation of WFI system in Zhubei facility, decreased in market demand and adjusted (postponed) the production schedule based on the client's R&D schedule. In addition, the Company's signing of an authorisation and cooperative development contract of the breast cancer biosimilar EG12014 (Trastuzumab Biosimilar) with Company SA. In 2023, due to the delay of the oversea approval, the recognized revenue decreased.
- 2. Non-operating income: Mainly because the unfavorable movements of the fluctuations in international exchange rate, and lead to the effects on foreign exchange losses.
- 3. Operation loss, loss before tax, and Net loss: Mainly because the decrease in CDMO revenue and licensing revenue.

Note: All the finance data are audited by CPA.

(2) List of Analysis of Financial Performance- Individual

Unit: NT\$ thousands; %

Year	2022	2023	Difference	%
Operating revenue	1,481,017	1,022,653	(458,364)	(31%)
Operating costs	724,565	785,912	61,347	8%
Gross profit (loss)	756,452	236,741	(519,711)	(69%)
Operating expenses	1,089,852	1,272,135	182,283	17%
Operating profit (loss)	(333,400)	(1,035,394)	(701,994)	211%
Non-operating income	228,569	132,821	(95,748)	(42%)
Non-operating expenses	(10,371)	(12,170)	(1,799)	17%
Profit (loss) before tax	(115,202)	(914,743)	(799,541)	694%
Income tax expense	(338)	(465)	(127)	38%
Net Profit (Loss)	(115,540)	(915,208)	(799,668)	692%

The main reason for the major change on Operating revenue, Net operating income (loss), and Income tax expense in currently 2 years:

- 1. Operating revenue and Gross profit: For CDMO business, because of the annual repairment in Xizhi facility and expansion and revalidation of WFI system in Zhubei facility, decreased in market demand and adjusted (postponed) the production schedule based on the client's R&D schedule. In addition, the Company's signing of an authorisation and cooperative development contract of the breast cancer biosimilar EG12014 (Trastuzumab Biosimilar) with Company SA. In 2023, due to the delay of the oversea approval, the recognized revenue decreased.
- 2. Non-operating income: Mainly because the unfavorable movements of the fluctuations in international exchange rate, and lead to the effects on foreign exchange losses.
- 3. Operation loss, loss before tax, and Net loss: Mainly because the decrease in CDMO revenue and licensing revenue.

Note: All the finance data are audited by CPA.



(3) Expected sales volume and its basis:

The self-owned biosimilars and new drugs of EirGenix and its subsidiary are still in the development stage and not commercially available. At present, the main source of revenue is the CDMO business, and the Company will continue to provide customized CDMO services. The management team of EirGenix and its subsidiary puts forward the Company's overall objectives and strategies, and then the research and development team put forward various research and development project plans. After feasibility analysis as well as market sales scale and financial evaluation, the implementation of the research and development plan and the timing of marketing sales are decided.

(4) The possible impact and the response plan for the company's finance and business in the future.

EirGenix and its subsidiary continue to provide bio-drug development technical services and GMP production business and continue to research and develop bio-similar drugs. EirGenix and its subsidiary are financially sound and see no significant adverse impact on the ongoing R&D plan and financial business.

3. Analysis of Cash Flow

(1) Cash Flow Analysis for the Current Year

Unit: NT\$ thousands; %

Year Item	2022	2023	Difference	%
Operating activities	(147,518)	(848,556)	(701,038)	475%
Investing activities	(476,050)	(219,018)	257,032	(54%)
Financing activities	124,788	(6,820)	(131,608)	(105%)

Analysis of change in cash flow:

- 1. Operating activities: Mainly due to the operation revenue lower than 2022. Related to the completion of Phase I clinical trials of the self-developed biosimilar EG1206A, the R&D expense higher than 2022.
- 2. Investing activities: Mainly due to the completion of the expansion of the mammalian cell production line at the 5th floor in Zhubei facility in 2023.
- 3. Financing activities: Mainly due to the bank loan for the expansion of the mammalian cell production line at the 5th floor in Zhubei facility in 2022.

Improve plan for insufficient liquidity: None.

(2) Cash Flow Analysis for the Coming Year (2024)

Unit: NT\$ thousands

Cash and Cash	Estimated Net Cash Flow	Estimated Cash	Cash Surplus	Leverage of C	Cash Deficit
Equivalents, Beginning	from Operating Activities	Outflow (Inflow)	(Deficit)	Lucrostus aut Diana	Eineneine Diene
of Year (1)	(2)	(3)	(1)+(2)-(3)	Investment Plans	Financing Plans
5,053,183	(947,260)	(842,411)	3,263,512	-	-

Analysis of change in cash flow in the next year:

- 1. Operating activities: Mainly due to the continuous investment in R&D costs.
- 2. Investing activities: Mainly due to the payment of the plant expansion and the purchasing of machinery and equipment.
- 3. Financing activities: Mainly due to the bank loan for the expansion of Zhubei plant phase II facility.

Improve plan for insufficient liquidity: None.

4. The effect of major capital expenditures during the most recent fiscal year on company's finance and business operations:

The Board of Directors resolution of the establishment of Phase II Facility and production equipment in Hsinchu Biomedical Science Park, the upper limit is NT\$2.468 billion, and the Board of Directors resolution of the establishment R&D laboratory \ production line and production equipment, the upper limit is NT\$1.1425 billion. After completion, can accelerate to meet the demand for international CDMO orders, meet the demand for the marketing products of the self-owned product, and can also be sufficient to provide Phase III clinical drugs and marketing production demand for subsequent product development. The construction of this new plant will become an efficient engine for the Company's rapid growth in the future and promotion of momentum internationally, which will be of positive help to the company's finance and business.



5. Investment Policy in the Last Year, Main Causes for Profits or Losses, Improvement Plans and Investment Plans for the Coming Year

(1) Investment Policy

The Company's reinvestment in other companies shall be implemented in accordance with the Investment Cycle and Regulations Governing the Acquisition and Disposal of Assets of the internal control system, which shall be discussed and approved by the Board of Directors or Shareholders' Meeting.

(2) Reasons of Investment Gain/Loss and its improving plan:

Year	Item	Recognized Investment Gain/(Loss)	Investment Policy	Reasons of Loss	Improving Plan
2023 Q4	EirGenix Europe GmbH	NTD 2,324 thousand	Development and Research on biotechnology drug and business development.	N/A	N/A

- (3) The investment plans for the coming year: None.
- (4) Investment plan in next year: The Company set up a German subsidiary in the first quarter of 2020, considering future operational needs and implementation of biosimilar clinical trials, of which the benefits will gradually emerge.

6. Analysis of Risk Management

- (1) Effects of Changes in Interest Rates, Foreign Exchange Rates and Inflation on Corporate Finance, and Future Response Measures
 - A. The effect upon the company's profits (losses) of interest rates and response measures to be taken in the future:
 - (A) The effect upon the company's profits (losses)

The Company's interest rate risk mainly comes from long and short-term borrowings from banks; the interest expenses of EirGenix and its subsidiaries amounted to NT\$ 2,300 thousand and NT\$ 1,205 thousand in 2023 and 2022; the increased mainly due to the interest rate increased by Central Bank. However, the company currently has enough operating funds to support. Therefore, the impact of interest rate changes on the Company's profit or loss has gradually decreased.

(B) Response measures to be taken in the future.

EirGenix and its subsidiary will keep abreast of interest rate changes, maintain good interactive communication with banks to obtain preferential interest rates and match up long and short-term capital planning to reduce the overall financing cost of the Company. At present, there is no effect of change in the interest rate on the company's operating results.

- B. The effect upon the company's profits (losses) of exchange rate and response measures to be taken in the future
- (A) The effect upon the company's profits (losses)

EirGenix and its subsidiary mostly denominate receivables and payables in New Taiwan Dollars or important international currencies for current clients and suppliers. The net exchange (losses) gains of the Company for the years 2023 and 2022 amounted to NT\$(9,431) thousand and NT\$ 126,788 thousand respectively, accounting for (0.92%) and 8.56% of the net operating revenues for the respective periods and have not had a significant impact on the Company's operation at present.

(B) Response measures to be taken in the future.

In order to reduce the impact of exchange rate changes on the Company's profit or loss in the future, EirGenix and its subsidiary will collect information on the exchange rate at any time, pay attention to the trends and changes in the exchange rate of major currencies in the international exchange market, grasp the exchange rate trends, and maintain a good interactive relationship with banks so as to obtain more extensive foreign exchange information and more preferential exchange rate quotations.

- C. The effect upon the company's profits (losses) of changes in the inflation rate and response measures to be taken in the future.
- (A) The effect upon the company's profits (losses)

In March 2024, the Chief Accounting Office of the Executive Yuan noted an annual increase rate of 2.14% in the consumer price index. Inflation was slight, and there was no significant impact on the Company's profit or loss.



(B) Response measures to be taken in the future.

In the future, the Company will also continue to track the impact of Inflation on various expenses of the industry and pay attention to market changes at any time as one of the bases for the Company's contingency decisions.

(2) Policies, Main Causes of Gain or Loss and Future Response Measures with Respect to High-risk, High-leveraged Investments, Lending or Endorsement Guarantees, and Derivatives Transactions

EirGenix has formulated the Procedures for Loaning Funds to Others, Procedures for Endorsements and Guarantees, Regulations Governing the Acquisition and Disposal of Assets, and other methods and has followed the specifications. EirGenix and its subsidiary focus on the development of the industry. As of the date of issuance for the annual report in the latest year, EirGenix and its subsidiary have not engaged in high-risk and highly leveraged investment or derivative merchandise transactions and have not lent funds or endorsement guarantees to others.

- (3) Future Research & Development Projects and Corresponding Budget
 - A. Future Research & Development Projects:

 Please refer to this Annual Report V. Operational Highlights-D. The new products (services) are planning to development.
 - B. Expected to Spend on the Research and Development:

 EirGenix and its subsidiary are expected to spend about NT\$ 1,160,000,000 on the research and development of the above products, clinical trials, and the construction of cell line platforms in 2024. The research and development costs of the product development plans will be planned and adjusted according to the actual progress and plan objectives.
- (4) Effects of and Response to Changes in Policies and Regulations Relating to Corporate Finance and Sales

The operation of EirGenix and its subsidiary follows the relevant current laws and regulations at home and abroad, and relevant personnel also pay attention to changes in laws and regulations at any time for the reference of the management echelon. Therefore, the Company can grasp and effectively respond to changes in important policies and laws at home and abroad in real-time. As of the date of issuance for the annual report in the latest year, changes in policies and laws at home and abroad have had no significant adverse impact on the Company's finance and business.

(5) Effects of and Response to Changes in Technology and the Industry Relating to Corporate Finance and Sale

The government has actively promoted the biotechnology industries in recent years, of which the biopharmaceutical industry has the characteristics of high technical threshold, long R&D cycle, high professional technical demand, and added value. The threshold for the industry is relatively high, so it is not easy to produce drastic changes in a short period of time. Moreover, the Company, with a high degree of professional R&D capability, can closely grasp technological changes and industrial changes and take appropriate countermeasures as needed. As of the date of issuance for the annual report in the latest year, EirGenix and its subsidiary have not had a significant impact on the Company's financial business due to technological changes and industrial changes.

Considering the rapid changes in information technology and the external environment, to reduce the impact of external changes on our finances, the Company plans to revise relevant information security policies suitable for the operations together with external professional information security consultants with reference to the NIST Cybersecurity Framework (CSF) and relevant standards in the industry. We will implement the revised policies accordingly and review and adjust them regularly as the benchmark for evaluation and judgment when adopting various information systems and services.

(6) The Impact of Changes in Corporate Image on Corporate Risk Management, and the Company's Response Measures

In recent years, EirGenix has deeply cultivated the Taiwan market, has established good relations with Japanese and Chinese clients, and is actively exploring the European and American markets at present, aiming at the professional technical reputation and good international image. EirGenix has always adhered to the professional and sincere enterprise spirit and implemented it in the daily operation and management of the Company so that the Company's systems and colleagues have sufficient ability to cope with possible enterprise crises and reduce the impact of such risks on the Company's operation. As of the date of issuance for the annual report in the latest year, EirGenix and its subsidiary have not had any negative impact on the Company due to changes in corporate image.

- (7) Expected Benefits from, Risks Relating to and Response to Merger and Acquisition Plans

 As of the date of issuance for the annual report in the latest year, EirGenix didn't have the acquisition plan.
- (8) Expected Benefits from, Risks Relating to and Response to Factory Expansion Plans
 - A. Expected Benefits from Factory Expansion Plans



As the existing Xizhi plant has reached full capacity, at the end of 2016, a new PIC/S GMP biopharmaceutical plant with commercial mass-production scale situated in the Zhubei Biomedical Park broke ground, and a disposable bioreactor (SUB) process was built. It can attract international and domestic clients' demand for large-scale production and contract production of products on the market in the future. Zhubei plant will be responsible for the production of self-owned products and continue to undertake CDMO business.

Currently Zhubei plant phase I facility completed, the production capacity reached 25,500 L. After the completion of Zhubei plant phase II facility, the production capacity of the microbial cell production line will reach 1,500 L. And the third production site, located in the Southern Taiwan Science Park, will expand the mammalian cell production line by 150,000 liters over the next 10 years. After the completion of the construction, it is expected to increase the revenue from the technical service of biopharmaceutical contract development.

B. Risks Relating to and Response to Factory Expansion Plans

The increased production capacity of the new plant will fluctuate with the market of biologics, research and development status, and the receipt of orders for contract development cases. In addition to actively striving for domestic biopharmaceutical contract development orders, the Company will continue to expand and seek overseas orders. It is expected that the expanded commercial plant will be conducive to the development of contract development cases and commission orders for biologics.

- (9) Risks Relating to and Response to Excessive Concentration of Purchasing Sources and Excessive Customer Concentration
 - A. Risks Relating to and Response to Excessive Concentration of Purchasing Sources

Among the top ten suppliers of EirGenix and its subsidiary in the last two years, Merck, Pall Singapore and Cytiva make up over 15%. Because the raw materials for biotechnology research and development products have a high manufacturing technology threshold and strict quality requirements, their suppliers that can be internationally recognized are limited. The Company takes international pharmaceutical companies as its main customers. Therefore, the source of raw materials is internationally renowned international raw material suppliers with stable supply, which is the general trend of the raw material sources for the research and development of most biotechnology companies and pharmaceutical companies in the world. However, the Company keeps an eye on the changing trend of raw material market supply and is committed to actively developing multiple suppliers to reduce the risk of centralized purchase.

B. Risks Relating to and Response to Excessive Customer Concentration

The proportion of the largest trade debtors of EirGenix and its subsidiary in the last two years was 17.08% and 34.72%, respectively. In terms of technical services for biopharmaceuticals, because of its high technical threshold and different characteristics of the developed products, EirGenix, and its subsidiary establish long-term relationships with key clients, with the goal of cooperating in the development of multiple projects or large-scale projects, which is in line with the interests of both parties and the performance of development efficiency. EirGenix has successively developed several stable clients in the past few years and is still continuing to cooperate and establish deep relationships with clients to balance the proportion of sales of individual clients. In the future, the Company will continue to develop clients to reduce the risk of sales concentration.

- (10) Effects of, Risks Relating to and Response to Large Share Transfers or Changes in Shareholdings by Directors, Supervisors, or Shareholders with Shareholdings of over 10%: None.
- (11) Effects of, Risks Relating to and Response to the Changes in Management Rights

As of the date of issuance for the annual report in the latest year, EirGenix and its subsidiary didn't have the situation of Changing management rights.

- (12) Litigation or Non-litigation Matters, List major litigious, non-litigious or administrative disputes that: (1) involve the company and/or any company director, any company supervisor, the general manager, any person with actual responsibility for the firm, any major shareholder holding a stake of greater than 10 percent, and/or any company or companies controlled by the company; and (2) have been concluded by means of a final and unappealable judgment, or are still under litigation. Where such a dispute could materially affect shareholders' equity or the prices of the company's securities, the annual report shall disclose the facts of the dispute, amount of money at stake in the dispute, the date of litigation commencement, the main parties to the dispute, and the status of the dispute as of the date of publication of the annual report: None.
- (13) Other Major Risk and Response



Risks in the development of biosimilars and new drugs and their impact on financial business:

New drug development is a high-risk, time-consuming, and capital-consuming industry. From early research to successful drug marketing, it takes about 10~15 years. If a new drug can successfully enter the drug license examination and be approved for marketing from pre-clinical research through layers of tests, then both the company and investors must realize that the new drug development risk is high, and the investment recovery period is long. The R&D process is long, and huge R&D funds need to be invested from topic selection, process development to clinical trials. If R&D fails, or the net cash flow from operating activities is relatively late, then stable operating revenue will not be smooth, insufficient working capital may occur, and there will be a risk that the new drug R&D plan cannot be completed.

The risk of developing biosimilars is relatively low compared with that of developing new drugs for two reasons. The first concerns whether the reverse engineering technical difficulty of making the product, which is highly similar to the original reference drug in physical, chemical, and biological properties, can be overcome and whether bioequivalence can be achieved in human pharmacokinetic tests (usually Phase I clinical trials). The experience of developing biosimilars in Europe in the past ten years shows that the chances of failure for products to meet the above standards are very small. The second reason is to have sufficient funds for Phase III clinical development and partners for Phase III joint development and sales.

In view of the financial risks in the research and development of biosimilars and new drugs, in addition to generating cash flow by Contract Development & Manufacturing Organization (CDMO) and applying for specialized programs to meet the cash expenditure needs, EirGenix and its subsidiary will also carry out negotiations on the authorization of regional cooperation for products. EirGenix, Inc. has secured a license agreement with Sandoz AG, a global leader in generic and biosimilar drug manufacturing, in April 2019. The signed license agreement grants Sandoz the exclusive rights to globally commercialize EirGenix's proposed trastuzumab biosimilar drug (EG12014). EirGenix will maintain responsibility for the development and manufacturing of the trastuzumab biosimilar, while Sandoz will maintain rights to commercialize the drug upon approval in the authorised regions. Under the terms of the agreement, EirGenix will receive an upfront payment, milestone payments at each stage and profit-sharing royalty on sales of products in the authorized markets. EirGenix, Inc. also undertook the post-marketing production of EG12014. In addition, the Company continues to carry out product life cycle management and evaluate the research and development of new indications related to HER2. By virtue of self-owned specialty and lower development risks, the Company effectively expands the market and life cycle of products so as to continuously increase the market value of products and ensure the Company's ability to continue business development.

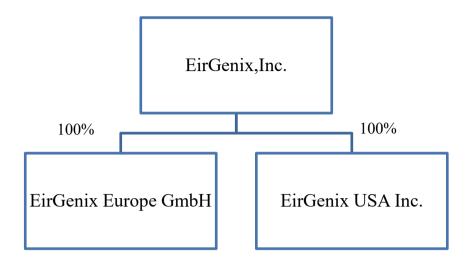
7. Other important matters: None.



VIII. Special Disclosure

1. Information of Affiliated Companies

(1) Investment Structure



(2) Basic information on affiliated enterprises:

Name of Subsidiary	Date of registration	Address	Capital	Main Business Activity
EirGenix Europe GmbH	2020/2/11	Neuhauser Str. 47, 80331 Munchen	EUR25,000	Development and Research on biotechnology drug and business development.
EirGenix USA Inc.	2023/11/28	130 Gates Street, San Francisco, CA 94110	USD100	Consultation of CDMO service

- (3) In Compliance with Article 369-3 of Company Law, it shall be concluded as the existence of the controlling and subordinate relation: Not Applicable.
- (4) The industries covered by the business operated by the affiliates overall. Where connections exist among the businesses operated by individual affiliates, a description of the mutual dealings and division of work among such affiliates should be provided: The major tasks of EirGenix Europe GmbH are managing and executing the clinical trial of the durg development. Its main business activity are development and research on biotechnology drug and business development. The major task of EirGenix USA Inc. is consultation of CDMO service.

(5) Directors, Supervisors and President information on affiliated enterprises:

Name of Subsidiary	Title Name		Shares holding	
E. C E C. III	Director	Lee-Cheng Liu	-	-
EirGenix Europe GmbH	President	Thomas Schulze	-	-
EigConin LICA In a	Chairman and	Las Chana Lin		
EirGenix USA Inc.	President	Lee-Cheng Liu	-	-

(6) Operational information on affiliated enterprises

Dec. 31st, 2023, Unit: NT\$ thousands

Name	Capital	Asset	Liability	Net worth	Revenue	Operating income	Net Gain after tax	EPS
EirGenix Europe GmbH	845	19,273	11,530	7,743	71,764	3,417	2,324	-
EirGenix USA Inc.	-	-	-	-	-	-	-	-

(7) Consolidated Financial Statements of Affiliated Enterprises:

EirGenix's financial information for the 2020Q1 was included in the subsidiary EirGenix Europe GmbH and issued consolidated statements. For the 2023 and 2022, pursuant to "Criteria Governing Preparation of Affiliation Reports, Consolidated Business Reports and Consolidated Financial Statements of Affiliated Enterprises," the entities that are required to be included in the consolidated financial statements of affiliates are the same as the entities required to be included in the consolidated financial statements of parent



and subsidiary companies under International Financial Reporting Standard No. 10. Also, if relevant information that should be disclosed in the consolidated financial statements of affiliates has all been disclosed in the consolidated financial statements of parent and subsidiary companies, it shall not be required to prepare separate consolidated financial statements of affiliates.

- (8) Information of Affiliated Enterprises for Loaning of Funds, Making of Endorsements/Guarantees and Engaging in Derivatives Trading: None.
- (9) Major Trading Matter with Affiliated Enterprises: None.
- (10) Reports on Affiliations: Not Applicable.

2. Private Placement Securities in the Most Recent Years:

Item	2021 First time Private Placement of Securities
	Issue Date (delivery date): 2021/11/30
Securities under	Common Stock
private placement	
Date of resolution and	2021/8/3
approved quantity	55,000,000 shares
Basis and rationale for price setting	The price determination date is based on the board meeting on 2021/10/01. Reference price is the simple average closing price of the common shares of the TWSE listed or TPEx listed company for either the 1, 3, or 5 business days before the price determination date and each of them is 126.5 dollars, 126.67 dollars, and 128.5 dollars. After adjustment for any distribution of stock dividends, cash dividends, or capital reduction, the price is 128.5 dollars; or the simple average closing price of the common shares of the TWSE listed or TPEx listed company for the 30 business days before the price determination date, after adjustment for any distribution of stock dividends, cash dividends, or capital reduction, the price is 128.75 dollars. Select the higher of the above two calculations 128.75 as reference price and actual private placement shall not be lower than 50% of the reference price. The actual private placement price is per share NT\$91.5 which is 71.07% of the reference price: NT\$128.75 and complies with shareholder meetings' decision that no lower than the price based on the pricing principle: at least 50% of the two above-mentioned prices (the higher one). Consult with Hsiu-Luan Lin, Certified Public Accountant from CHAMPiON accounting firm, to issue a submission of the reasonableness for private placement.
Selection method of the placees	The places of the private placement are strategic investors. In accordance with Article 43-6 of the Securities and Exchange Act and Taiwan Finance Certificate (1) No. 0910003455 issued by the Financial Supervisory Commission on June 13, 2002, it states to select those who are beneficial to the long-term development of the Company and improve the operational performance, strengthen competitiveness, and generate benefits for existing shareholders' equity. The purpose of the placees selected this time is to introduce strategic investors. The main targets are strategic investors who have developing experiences in biomedicine and health and can stabilize the Company's equity and capital structure.
The necessary reason for the Private Placement	With the considerations of the timeliness of financing activities and the uncertainty of the capital market, and the benefit for the Company's long term operating development because of the transfer limit of the private placement common share, it plans to conduct the financial activities with the private placement. To accelerate the product development efficiency as well as the process of the same drug series to complete the production line. By the comprehensive effect of expanding the market, it can establish EirGenix's unique status in the international biosimilars and CDMO field. In order to sustain EirGenix's operation and development, it is necessary to conduct private placement to introduce strategic investors by resolution.
Date of payment and completion	2021/10/15
Actual Subscription Price	Per share NT\$91.5



Difference between Actual Subscription price and Reference Price	The actual subs	cribed price	is per share N	Т\$91.5	, 71.07% of th	he reference price: NT\$128.75.		
	Placees	Eligibility (note)	Quantity Subscribed		ionship with Genix, Inc.	Participation in Company Operations		
Information on Placees	Foxconn Technology Co., Ltd. Yonglin Capital Holding Co., Ltd. Hong Wei Investment Co., Ltd.	Note 2.	27,500,000 shares 26,500,000 shares 1,000,000 shares		None	There is no significant change in managerial control within the 1-year period immediately preceding the day on which the board of directors resolves on the private placement and after the introduction of strategic investors through private placement.		
Impact of private	Boost EirGenix's op	erating scale	e, horizontal ar	d verti	cal integration	n, and product or market		
placement on				-		efficiency, expand the operational		
shareholders' equity						ing EirGenix and shareholder value.		
	The Usage of	funds	Budget Am		_	entation as of 2024 First Quarter		
	R&D expenses		NT\$1,016,178,000		R&D expenses NT\$ 397,191,024 and deposit other funds in EirGenix bank accounts.			
	Expansion and building factory NT\$1,700,				Expansion and building factory NT\$			
Use of funds from			1,141,700,00		EirGenix bar			
private placement and progress of proposed plans	Repay bank loans and horizontal and vertice integration, and other operational funding	eal er	NT\$316,322	2,000	Repay bank	loan NT\$316,322,000.		
	Acquisition or purch intangible assets, op related assets, and ri assets.	eration-	NT\$2,000,00	NT\$2,000,000,000		Acquisition important assets NT\$ 60,112,501 and deposit other funds in EirGenix bank accounts.		
Effectiveness of private placement	 EirGenix is currently developing the product for the treatment of HER2+ breast cancer. Received the approval letter from Ministry of Health and Welfare for the biosimilar drug EIRGASUN 150 mg powder for concentrate for infusion. Received the approval letter from EC for EG12014 licensed. Preparing the US BLA resubmission for EG12014. The Phase I clinical trial for EG1206A (Pertuzumab Biosimilar) has been completed. The second mammalian cell production line for the Zhubei plant phase I facility has been completed. Build microbial cell production line factory for the Zhubei plant phase II facility. The three-stage expansion of the mammalian plant which has 150,000L capacity, is under planning at Ciaotou Science Park, Kaohsiung. Repay bank loan NT\$316,322,000 and save annual interest expenses roughly about NT\$5,684,000 which calculating under the current EirGenix loan rates of 1.797%. Other unused funds will follow the plan and demonstrate effects continuously. Co-developed the biosimilar drug TSY0110 (EG12043) of ADC for the treatment of breast cancer with Formosa Pharmaceuticals. Other unused funds will follow the plan and demonstrate effects continuously. 							

- 3. The Shares in the Company Held or Disposed of by Subsidiaries in the Most Recent Years: None.
- 4. Other Matters that Require Additional Description: None.



5. If any of the situations listed in Article 36, paragraph 3, subparagraph 2 of the Securities and Exchange Act, which might materially affect shareholders' equity or the price of the company's securities, has occurred during the most recent fiscal year or during the current fiscal year up to the date of publication of the annual report:

Date	Material Information
2023/01/05	Announcement of EirGenix receives Establishment Inspection Report from US FDA.
2023/04/11	EirGenix, Inc. has received the approval letter from TFDA that the API Trastuzumab has obtained the license and the DMF number.
2023/05/11	The Phase I clinical trial of EirGenix's biosimilar EG1206A has met its primary endpoint and showed Pharmacokinetic biosimilarity.
2023/09/18	Sandoz AG received a positive CHMP opinion for the biosimilar drug EG12014 licensed from EirGenix, Inc.
2023/09/18	EirGenix's biosimilar drug,"Eirgasun vial 150 mg," has been approved by National Health Insurance Administration to be enrolled in the reimbursement system.
2023/11/17	Sandoz AG has received the approval letter from EC for Herwenda-Trastuzumab biosimilar EG12014 (150 mg, for intravenous use) licensed from EirGenix, Inc.
2024/03/08	The Board of Directors resolved to sign the contract with clinical CRO and the relevant companies for Phase III clinical trial of the EG1206A



Appendix 1

EirGenix, Inc.

Statement of Internal Control System

Date: March 8, 2024

Based on the findings of a self-assessment, EirGenix Inc. (hereinafter "the Company") states the following pertaining to its internal control system during year 2023:

- I. The Company is fully aware that establishing, operating, and maintaining an internal control system are the responsibilities of its Board of Directors and managers. The Company has established such a system with an aim to providing reasonable assurance for the achievement of the following objectives: the effectiveness and efficiency of business operation (including profitability, performance, and safeguarding of company assets; the reliability, timeliness, transparency, and regulatory compliance of financial reporting and other related reports; and the compliance with applicable laws, regulations and rulings.
- II. An internal control system has inherent limitations. No matter how perfectly it is designed, an effective internal control system can provide only reasonable assurance of achieving the three above-mentioned objectives. Moreover, the effectiveness of the internal control system may be subjected to changes of environment or circumstances. Nonetheless, the Company's internal control system comprises of self-monitoring mechanisms, and the Company immediately undertakes corrective measures once a deficiency is identified.
- III. The Company assesses the design and operating effectiveness of its internal control system in accordance with the criteria stated in the "Regulations Governing Establishment of Internal Control Systems by Public Companies" (hereinafter referred to as "the Regulations"). The criteria stipulated in the Regulations identify five essential elements of an internal control system based on managerial control process, including (1). Control environment, (2). Risk assessment (3). Control activities, (4). Information and communication, and (5). Monitoring activities. Each essential element further contains several items. Please refer to the Regulations for the aforementioned items.
- IV. The Company has evaluated the design and operating effectiveness of its internal control system according to the aforesaid criteria.
- V. Based on the results of the mentioned assessment above, the Company believes that, as of December 31, 2023, its internal control system, including its supervision and management of subsidiaries, was effective in design and operation and provided reasonable assurance of achievement of operational effectiveness and efficiency, reliability, timeliness, transparency of reporting, and compliance with applicable laws, regulations, and rulings.
- VI. This Statement constitutes an integral part of the Annual Report for the year 2023 and the Prospectus of the Company and will be made public. Any falsehood, concealment, or other



illegality in the content made public will entail legal liability under Articles 20, 32, 171, and 174 in the Securities and Exchange Act.

VII. This Statement has been approved by the Board of Directors in their meeting held on March 8, 2024, with none of the ten attending directors expressing dissenting opinions, and the remainder all affirming the contents of this Statement.

EirGenix Inc.

Chairman and President: Lee-Cheng Liu





EirGenix, Inc.

Audit Committee's Review Report

The Board of Directors has prepared EirGenix's 2023 Business Report, Financial Statement, and Deficit Offset Statement. The CPA Sheng-Wei Deng and Yu-Fang Yen of PricewaterhouseCoopers Taiwan was retained to audit EirGenix's Financial Statement and has issued an audit report relating to the Financial Statement.

The Business Report, Financial Statement, and Deficit Compensation Statement have been reviewed and determined to be correct and accurate by the Audit Committee member of EirGenix. According to relevant requirements of the Securities and Exchange Act and the Company Act, we hereby submit this report.

To

EirGenix, Inc. 2024 Annual Shareholders' Meeting

EirGenix, Inc.

Chairman of Audit Committee: Ming-Thaur Chang

Member of Audit Committee: Po-Chih Chen

Member of Audit Committee: Fu-Shiow Yin

Member of Audit Committee: Ming-Shen Chen

EIRGENIX INC. AND SUBSIDIARIES CONSOLIDATED FINANCIAL STATEMENTS AND INDEPENDENT AUDITORS' REPORT DECEMBER 31, 2023 AND 2022

For the convenience of readers and for information purpose only, the auditors' report and the accompanying financial statements have been translated into English from the original Chinese version prepared and used in the Republic of China. In the event of any discrepancy between the English version and the original Chinese version or any differences in the interpretation of the two versions, the Chinese-language auditors' report and financial statements shall prevail.

EIRGENIX, INC.

<u>Declaration of Consolidated Financial Statements of Affiliated Enterprises</u>

For the year ended December 31, 2023, pursuant to "Criteria Governing Preparation of Affiliation Reports, Consolidated Business Reports and Consolidated Financial Statements of Affiliated Enterprises," the entities that are required to be included in the consolidated financial statements of affiliates are the same as the entities required to be included in the consolidated financial statements of parent and subsidiary companies under International Financial Reporting Standard No. 10. Also, if relevant information that should be disclosed in the consolidated financial statements of affiliates has all been disclosed in the consolidated financial statements of parent and subsidiary companies, it shall not be required to prepare separate consolidated financial statements of affiliates.

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EirGenix Inc.

Representative: Lee-Cheng Liu

March 8, 2024

INDEPENDENT AUDITORS' REPORT TRANSLATED FROM CHINESE

To the Board of Directors and Shareholders of EirGenix Inc.

Opinion

We have audited the accompanying consolidated balance sheets of EirGenix Inc. and subsidiaries (the "Group") as at December 31, 2023 and 2022, and the related consolidated statements of comprehensive income, of changes in equity and of cash flows for the years then ended, and notes to the consolidated financial statements, including a summary of material accounting policies.

In our opinion, the accompanying consolidated financial statements present fairly, in all material respects, the consolidated financial position of the Group as at December 31, 2023 and 2022, and its consolidated financial performance and its consolidated cash flows for the years then ended in accordance with the Regulations Governing the Preparation of Financial Reports by Securities Issuers and the International Financial Reporting Standards, International Accounting Standards, IFRIC Interpretations, and SIC Interpretations that came into effect as endorsed by the Financial Supervisory Commission.

Basis for opinion

We conducted our audits in accordance with the Regulations Governing Financial Statement Audit and Attestation Engagements of Certified Public Accountants and Standards on Auditing of the Republic of China. Our responsibilities under those standards are further described in the Auditors' Responsibilities for the Audit of the Consolidated Financial Statements section of our report. We are independent of the Group in accordance with the Norm of Professional Ethics for Certified Public Accountants of the Republic of China, and we have fulfilled our other ethical responsibilities in accordance with these requirements. We believe that the audit evidence we have obtained is sufficient and appropriate to provide a basis for our opinion.

Key audit matters

Key audit matters are those matters that, in our professional judgment, were of most significance in our audit of the Group's 2023 consolidated financial statements. These matters were addressed in the context of our audit of the consolidated financial statements as a whole and, in forming our opinion thereon, we do not provide a separate opinion on these matters.

The key audit matters for the Group's 2023 consolidated financial statements are stated as follows:

Accuracy of service revenue and authorisation and cooperative development revenue

Description

Refer to Note 4(25) for accounting policy on service revenue and authorisation and cooperative development revenue recognition, Note 5(2) for significant accounting estimates and assumptions, and Note 6(19) for details of operating revenue. The amount of service revenue and authorisation and cooperative development revenue for the year ended December 31, 2023 were NTD 605,990 thousand and NTD 141,472 thousand, respectively.

The Group's service revenue and authorisation and cooperative development revenue primarily arise from offering biopharmaceutical contract development and manufacturing services and authorising intellectual property rights of medicine development to pharmaceutical factory. Revenue is recognised based on the stage of completion at the balance sheet date provided that such transaction amounts can be reliably estimated. Since the information process, recording and maintenance are partially performed manually and the recognition of service revenue and authorisation and cooperative development revenue contains a high degree of uncertainty resulting in a complex calculation process, and revenue recognition is significant to the financial statements, we considered the accuracy of service revenue recognition a key audit matter.

How our audit addressed the matter

We performed the following audit procedures on the above key audit matter:

- 1. Obtained management's accounting policies on the service revenue and authorisation and cooperative development revenue recognition and confirmed that they are reasonable.
- 2. Selected samples and examined the contract in order to confirm whether the judgement made by the management was in line with the contract and generally accepted accounting principles.
- 3. For the performance obligation which was satisfied over time, selected samples and examined each data of contract costs and assessed whether the method and parameters used to measure the completion of performance obligation are reasonable.
- 4. Recalculated the accuracy of amount recognised as revenue and respective timing of recognition.

Impairment assessment of property, plant and equipment

Description

Refer to Note 4(17) for accounting policy on impairment of non-financial assets, Note 5(2) for uncertainty of accounting estimates and assumptions in relation to property, plant and equipment and Note 6(8) for description of property, plant and equipment.

On December 31, 2023, property, plant and equipment amounted to NTD 3,337,685 thousand, which were constructed to extend the production capacity of GMP. The Company assesses at each balance sheet date the fair value or recoverable value of those assets whether there is any indication that they may be impaired based on internal and external information. Since the impairment indication assessment and information and assumptions used to assess recoverable amount of assets have a significant impact to property, plant and equipment, we considered the impairment assessment of property, plant and equipment a key audit matter.

How our audit addressed the matter

We performed the following audit procedures on the above key audit matter:

- 1. Reviewed and assessed the reasonableness of each data in the impairment assessment.
- 2. Assessed the estimation procedure of future cash flows, and checked whether the cash flows listed in the assessment is consistent with operating plans.
- 3. Interviewed management to discuss the Group's operations and reviewed the actual performance of prior years' operating plans in order to understand the Group's intention and ability and ascertained whether there was any significant postponement on research and development.
- 4. Assessed the reasonableness of the significant assumptions adopted in estimating cash flows.

Other matter - Parent company only financial reports

We have audited and expressed an unqualified opinion on the parent company only financial statements of EirGenix Inc. as at and for the years ended December 31, 2023 and 2022.

Responsibilities of management and those charged with governance for the consolidated financial statements

Management is responsible for the preparation and fair presentation of the consolidated financial statements in accordance with the Regulations Governing the Preparation of Financial Reports by Securities Issuers and the International Financial Reporting Standards, International Accounting Standards, IFRIC Interpretations, and SIC Interpretations that came into effect as endorsed by the Financial Supervisory Commission, and for such internal control as management determines is necessary to enable the preparation of consolidated financial statements that are free from material misstatement, whether due to fraud or error.

In preparing the consolidated financial statements, management is responsible for assessing the Group's ability to continue as a going concern, disclosing, as applicable, matters related to going concern and using the going concern basis of accounting unless management either intends to liquidate the Group or to cease operations, or has no realistic alternative but to do so.

Those charged with governance, including the audit committee, are responsible for overseeing the Group's financial reporting process.

Auditors' responsibilities for the audit of the consolidated financial statements

Our objectives are to obtain reasonable assurance about whether the consolidated financial statements as a whole are free from material misstatement, whether due to fraud or error, and to issue an auditors' report that includes our opinion. Reasonable assurance is a high level of assurance, but is not a guarantee that an audit conducted in accordance with the Standards on Auditing of the Republic of China will always detect a material misstatement when it exists. Misstatements can arise from fraud or error and are considered material if, individually or in the aggregate, they could reasonably be expected to influence the economic decisions of users taken on the basis of these consolidated financial statements.

As part of an audit in accordance with the Standards on Auditing of the Republic of China, we exercise professional judgment and professional skepticism throughout the audit. We also:

1. Identify and assess the risks of material misstatement of the consolidated financial statements, whether due to fraud or error, design and perform audit procedures responsive to those risks, and obtain audit evidence that is sufficient and appropriate to provide a basis for our opinion. The risk of not detecting a material misstatement resulting from fraud is higher than for one resulting from error, as fraud may involve collusion, forgery, intentional omissions, misrepresentations, or the override of internal control.

- 2. Obtain an understanding of internal control relevant to the audit in order to design audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the Group's internal control.
- 3. Evaluate the appropriateness of accounting policies used and the reasonableness of accounting estimates and related disclosures made by management.
- 4. Conclude on the appropriateness of management's use of the going concern basis of accounting and, based on the audit evidence obtained, whether a material uncertainty exists related to events or conditions that may cast significant doubt on the Group's ability to continue as a going concern. If we conclude that a material uncertainty exists, we are required to draw attention in our auditors' report to the related disclosures in the consolidated financial statements or, if such disclosures are inadequate, to modify our opinion. Our conclusions are based on the audit evidence obtained up to the date of our auditors' report. However, future events or conditions may cause the Group to cease to continue as a going concern.
- 5. Evaluate the overall presentation, structure and content of the consolidated financial statements, including the disclosures, and whether the consolidated financial statements represent the underlying transactions and events in a manner that achieves fair presentation.
- 6. Obtain sufficient appropriate audit evidence regarding the financial information of the entities or business activities within the Group to express an opinion on the consolidated financial statements. We are responsible for the direction, supervision and performance of the group audit. We remain solely responsible for our audit opinion.

We communicate with those charged with governance regarding, among other matters, the planned scope and timing of the audit and significant audit findings, including any significant deficiencies in internal control that we identify during our audit.

We also provide those charged with governance with a statement that we have complied with relevant ethical requirements regarding independence, and to communicate with them all relationships and other matters that may reasonably be thought to bear on our independence, and where applicable, related safeguards.

From the matters communicated with those charged with governance, we determine those matters that were of most significance in the audit of the consolidated financial statements of the current period and are therefore the key audit matters. We describe these matters in our auditors' report unless law or regulation precludes public disclosure about the matter or when, in extremely rare circumstances, we determine that a matter should not be communicated in our report because the adverse consequences of doing so would reasonably be expected to outweigh the public interest benefits of such communication.

Teng, Sheng-Wei

Yen, Yu-Fang
For and on Behalf of PricewaterhouseCoopers, Taiwan

March 8, 2024

The accompanying consolidated financial statements are not intended to present the financial position and

The accompanying consolidated financial statements are not intended to present the financial position and results of operations and cash flows in accordance with accounting principles generally accepted in countries and jurisdictions other than the Republic of China. The standards, procedures and practices in the Republic of China governing the audit of such financial statements may differ from those generally accepted in countries and jurisdictions other than the Republic of China. Accordingly, the accompanying consolidated financial statements and independent auditors' report are not intended for use by those who are not informed about the accounting principles or auditing standards generally accepted in the Republic of China, and their applications in practice.

As the financial statements are the responsibility of the management, PricewaterhouseCoopers cannot accept any liability for the use of, or reliance on, the English translation or for any errors or misunderstandings that may derive from the translation.

EIRGENIX INC. AND SUBSIDIARIES CONSOLIDATED BALANCE SHEETS DECEMBER 31, 2023 AND 2022

(Expressed in thousands of New Taiwan dollars)

		 December 31, 2023	 December 31, 2022			
	Assets	Notes	 AMOUNT	%	 AMOUNT	%
	Current assets					
1100	Cash and cash equivalents	6(1)	\$ 5,053,183	45	\$ 6,126,885	52
1136	Current financial assets at amortised	6(3)				
	cost		500,000	5	1,000,000	9
1140	Current contract assets	6(19) and 7	293,694	3	234,399	2
1150	Notes receivable, net	6(4)	19	-	-	-
1170	Accounts receivable, net	6(4)	253,390	2	32,782	-
1180	Accounts receivable, net-related	7				
	parties		2,636	-	-	-
1200	Other receivables		20,497	-	24,944	-
1220	Current income tax assets		17,648	-	5,963	-
130X	Inventories	6(5)	680,637	6	739,463	6
1410	Prepayments	6(6)	 93,802	1	 123,442	1
11XX	Total current assets		 6,915,506	62	 8,287,878	70
	Non-current assets					
1510	Non-current financial assets at fair	6(2) and 7				
	value through profit or loss		80,298	1	61,420	1
1517	Non-current financial assets at fair	6(7)				
	value through other comprehensive					
	income		325,887	3	279,325	2
1535	Non-current financial assets at	6(3) and 8				
	amortised cost		40,720	-	41,123	-
1600	Property, plant and equipment, net	6(8), 7 and 8	3,337,685	30	2,608,848	22
1755	Right-of-use assets	6(9)	329,236	3	325,330	3
1780	Intangible assets	6(10)	28,269	-	28,067	-
1990	Other non-current assets	6(8)(11) and 8	 104,958	1	 215,165	2
15XX	Total non-current assets		 4,247,053	38	 3,559,278	30
1XXX	Total assets		\$ 11,162,559	100	\$ 11,847,156	100

(Continued)

EIRGENIX INC. AND SUBSIDIARIES CONSOLIDATED BALANCE SHEETS DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars)

	Tiskillaise and Fanian	Notes		December 31, 2023 AMOUNT	December 31, 2022		
	Liabilities and Equity Current liabilities	Notes	<i></i>	AMOUNI	%	AMOUNT	
2130	Current contract liabilities	6(19) and 7	\$	56,766	- \$	150,475	1
2170	Accounts payable	0(15) and 7	Ψ	79,556	1	134,607	1
2200	Other payables	6(12)		530,299	5	407,387	4
2220	Other payables - related parties	7		7,993	-	7,732	· -
2230	Current tax liabilities			992	_	761	_
2280	Current lease liabilities			28,622	_	26,826	_
2399	Other current liabilities			2,937	_	3,104	-
21XX	Total current liabilities			707,165	6	730,892	6
	Non-current liabilities						
2540	Long-term borrowings	6(13) and 8		120,460	1	120,460	1
2570	Deferred tax liabilities	6(25)		1,380	-	874	-
2580	Non-current lease liabilities			316,085	3	311,758	3
2600	Other non-current liabilities			6	<u> </u>	294	
25XX	Total non-current liabilities			437,931	4	433,386	4
2XXX	Total liabilities			1,145,096	10	1,164,278	10
	Equity						
	Capital	6(16)					
3110	Common stock			3,060,516	28	3,043,358	26
	Capital reserve	6(17)					
3200	Capital surplus			7,830,216	70	7,734,141	65
	Accumulated deficit	6(18)					
3350	Accumulated deficit		(915,208) (8) (115,540) (1)
	Other equity interest						
3400	Other equity interest			41,939		20,919	
3XXX	Total Equity			10,017,463	90	10,682,878	90
	Significant contingent liabilities and	9					
	unrecognised contract commitments						
	Significant events after the balance	11					
	sheet date						
3X2X	Total Liabilities and Equity		\$	11,162,559	100 \$	11,847,156	100

The accompanying notes are an integral part of these consolidated financial statements.

EIRGENIX INC. AND SUBSIDIARIES CONSOLIDATED STATEMENTS OF COMPREHENSIVE INCOME YEARS ENDED DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars, except for loss per share)

			Year ended December 31				
				2023		2022	
	Items	Notes		AMOUNT	%	AMOUNT	%
4000	Operating Revenue	6(19) and 7	\$	1,022,653	100 \$	1,481,017	100
5000	Operating Costs	6(5)(10)(24) and 7	(785,912) (<u>77</u>) (724,565) (49)
5900	Gross Profit			236,741	23	756,452	51
	Operating Expenses	6(10)(24) and 7					
6100	Sales and marketing expenses		(62,232) (6) (50,844) (3)
6200	General and administrative expenses		(254,196) (25) (236,675) (16)
6300	Research and development expenses		(952,290) (93) (800,144) (54
6450	Reversal of credit impairment loss(expected credit impairment	12(2)				202	
	loss)		.—		_	392	
6000	Total operating expenses		(1,268,718) (124) (1,087,271) (73
6900	Operating Loss		(1,031,977) (<u>101</u>) (330,819) (22)
71 00	Non-operating Income and Expenses	((2) (20)		101 151	1.0	50.504	
7100	Interest income	6(3)(20)		134,471	13	59,584	4
7010	Other income	6(21)	,	5,439	-	37,644	2
7020	Other gains and losses	6(2)(9)(22)	(11,180) (1)	128,915	9
7050	Finance costs	6(9)(23) and 7	(10,403) (<u>l</u>) (9,639) (1)
7000	Total non-operating income and			110 227	1.1	216 504	1.4
5 000	expenses		.—	118,327	11	216,504	14
7900	Loss before Income Tax	((2.5)	(913,650) (90) (114,315) (8)
7950	Income tax	6(25)	(1,558)	(1,225)	
8200	Net Loss		(<u>\$</u>	915,208) (<u>90</u>) (<u>\$</u>	115,540) (<u>8</u>)
	Other Comprehensive Income						
	Components of other comprehensive						
	income that will not be reclassified to						
	profit or loss						
8316	Unrealised gains (losses) from	6(7)					
	investments in equity instruments						
	measured at fair value through other			45.000		50.004	
0210	comprehensive income		\$	45,939	<u> </u>	59,091	4
8310	Other comprehensive income(loss)						
	that will not be reclassified to			45.000	~	50, 001	
	profit or loss			45,939		59,091	4
	Components of other comprehensive						
	income that will be reclassified to						
0261	profit or loss						
8361	Exchange differences on translation			220		220	
9200	of foreign financial statements	((25)		220	-	220	-
8399	Income tax related to components of other comprehensive income that	0(23)					
	will be reclassified to profit or loss		,	41)			
8360	Other comprehensive income(loss)		(41)	<u> </u>	<u>-</u> _	
8300	that will be reclassified to profit or						
	loss			179		220	
8300	Other Comprehensive Income		<u>*</u>	46,118	5 \$	59,311	
	_		φ (¢				
8500	Total Comprehensive Loss		(<u>\$</u>	869,090) (<u>85</u>) (<u>\$</u>	56,229) (<u>4</u>)
	Logg way about	6(26)					
0750	Loss per share	6(26)	(¢		2 00 (6		0.20
9750	Loss per share		(<u>\$</u>		3.00) (<u>\$</u>		0.38)

EIRGENIX INC. AND SUBSIDIARIES CONSOLIDATED STATEMENTS OF CHANGES IN EQUITY YEARS ENDED DECEMBER 31, 2023 AND 2022

(Expressed in thousands of New Taiwan dollars)

Equity attributable to owners of the parent Other equity interest Capital Reserves Unrealised gains (losses) from financial assets Exchange measured at fair differences on value through translation of other Additional paid-in Donated assets Employee stock Capital surplus, Restricted stock to Capital surplus, foreign financial comprehensive Unearned stock options Notes Common stock capital received options employees deficit statements income compensation Total equity Year ended December 31,2022 (\$ 2,973,500) (\$ Balance at January 1, 2022 \$ 3,003,845 \$ 10,313,563 237) \$ 10,428,751 2,036 41,958 3,467 114,928 Loss for 2022 115,540) 115,540) Other comprehensive income (loss) 6(7) 220 59,091 59,311 220 59,091 Total comprehensive income (loss) 115,540 56,229) Capital surplus used to offset accumulated deficit 6(18) 2,971,464) 2.036) 2,973,500 61,651 871 155,312 Compensation costs of share-based payments 6(15) 92,790 Employee stock options exercised 8,320) 6(15)(16) 10,523 26,467 28,670 Issuance of employee restricted stocks 6(15)(16) 6,318 47,318 53,636) Redemption of employee restricted stocks 6(15)(16) 2,260) 2,260 Restricted stocks vested 59,358 59,358) Conversion of convertible bonds 6(16) 24,932 104,904 3,462) 126,374 Redemption of convertible bonds Balance at December 31, 2022 3,043,358 7,532,828 95,289 105,148 115,540) 64,922 43,986) \$ 10,682,878 Year ended December 31,2023 Balance at January 1, 2023 105,148 876 17) 43,986) \$ 10,682,878 115,540) 64,922 915,208) Loss for the 2023 915,208) Other comprehensive income (loss) 6(7) 179 45,939 46,118 Total comprehensive income (loss) 915,208 179 45,939 869,090) Capital surplus used to offset accumulated deficit 6(18) 114,664) 876) 115,540 84,285 Compensation costs of share-based payments 6(15) 96,615 180,900 Employee stock options exercised 6(15)(16) 10,264) 22,775 7,270 25,769 Employee stock options expired 6(15) 1,810) 1,810 Issuance of employee restricted stocks 6(15)(16) 11,818 109,895 121,713) Redemption of employee restricted stocks 6(15)(16) 1,930) 1,930 Restricted stocks vested 71,119 71,119)

167,500

145,854

915,208)

110,861

69,084)

\$ 10,017,463

\$ 3,060,516

\$ 7,515,052

Balance at December 31, 2023

EIRGENIX INC. AND SUBSIDIARIES CONSOLIDATED STATEMENTS OF CASH FLOWS YEARS ENDED DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars)

	Year ended December 31			er 31	
	Notes		2023		2022
CASH FLOWS FROM OPERATING ACTIVITIES					
Loss before tax		(\$	913,650)	(\$	114,315)
Adjustments		(4	713,030)	(Ψ	111,515)
Adjustments to reconcile profit (loss)					
Depreciation	6(8)(9)(24)		227,544		189,100
Amortization	6(10)(24)		11,296		16,184
Net loss (gain) on financial assets or liabilities at			,		,
fair value			1,122	(2,863)
Interest expense	6(23)		10,403	•	9,639
Interest income	6(20)	(134,471)	(59,584)
Dividend income	6(21)	Ì	475)	`	-
Compensation costs of share-based payments	6(15)(24)	`	180,900		155,312
Loss on lease modification	6(9)(22)		413		709
Loss on redemption of convertible bonds	6(22)		-		3
Reversal of credit impairment loss(expected	12(2)				
credit impairment loss)			-	(392)
Changes in operating assets and liabilities					
Changes in operating assets					
Contract assets		(59,295)	(63,802)
Notes receivable, net		(19)		1,139
Accounts receivable, net		(220,608)		46,084
Accounts receivable, net-related parties		(2,636)		546
Other receivables			6,736	(13,790)
Inventories			58,826	(324,025)
Prepayments			29,640	(17,394)
Other current assets			-		1,555
Changes in operating liabilities					
Contract liabilities		(93,709)	(93,551)
Accounts payable		(55,051)		48,151
Other payables		(5,604)		33,854
Other payables - related parties			261		2,037
Other current liabilities		(<u>167</u>)	(1,818)
Cash outflow generated from operations		(958,544)	(187,221)
Interest received			132,183		55,231
Interest paid		(10,386)	(9,316)
Dividends received			475		-
Income tax received			1,128		_
Income tax paid		(13,412)	(6,212)
Net cash flows used in operating activities		(848,556)	(147,518)

(Continued)

EIRGENIX INC. AND SUBSIDIARIES CONSOLIDATED STATEMENTS OF CASH FLOWS YEARS ENDED DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars)

	Year ended December 31			ber 31	
	Notes		2023		2022
CASH FLOWS FROM INVESTING ACTIVITIES					
Acquisition of financial assets at fair value through	6(2)				
profit or loss	,	\$	-	(\$	58,390)
Acquisition of financial assets at fair value through	6(7)				
other comprehensive income		(623)	(208,627)
Acquisition of financial assets at amortised cost		(3,700,000)	(1,032,516)
Proceeds from disposal of financial assets at					
amortised cost			4,200,403		1,636,640
Acquisition of property, plant and equipment	6(8)(27)	(575,270)	(345,792)
Acquisition of intangible assets	6(10)(27)	(15,142)		8,652)
Decrease in other financial assets			_		27,334
Decrease (increase) in refundable deposits (shown					
as other non-current assets)			56,253	(778)
Increase in prepayments for investments (shown as					
other non-current assets)		(46,270)	(20,000)
Increase in prepayments for business facilities	6(8)				
(shown as other non-current assets)		(138,453)	(433,952)
Increase in other non-current assets			84	(31,317)
Net cash flows used in investing activities		(219,018)	(476,050)
CASH FLOWS FROM FINANCING ACTIVITIES					
Repayments of bonds	6(28)		_	(200)
Proceeds from long-term borrowings	6(28)		-		120,460
Decrease (increase) in guarantee deposits	6(28)				
received(shown as other non-current liabilities)		(288)		294
Repayments of lease principal	6(28)	(29,307)	(24,435)
Employee stock options exercised			22,775		28,669
Net cash flows (used in) from financing			_		_
activities		(6,820)		124,788
Effect of exchange rate		-	692		281
Net decrease in cash and cash equivalents		(1,073,702)	(498,499)
Cash and cash equivalents at beginning of year		•	6,126,885	•	6,625,384
Cash and cash equivalents at end of year		\$	5,053,183	\$	6,126,885

EIRGENIX INC. AND SUBSIDIARIES NOTES TO THE CONSOLIDATED FINANCIAL STATEMENTS YEARS ENDED DECEMBER 31, 2023 AND 2022

(Expressed in thousands of New Taiwan dollars, except as otherwise indicated)

1. History and Organization

- (1) EirGenix, Inc. (hereinafter referred to as the "Company") was incorporated as a company limited by shares under the provisions of the Company Act of the Republic of China (R.O.C.) in December 2012. In April 2013, the Company obtained all key technologies from the biopharmaceutical pilot plant originally owned by the Development Center for Biotechnology, including its complete core competencies. The Company and its subsidiaries (hereinafter collectively referred to as the "Group") are primarily engaged in the research and development of biosimilars and new drugs, as well as biopharmaceutical contract development and manufacturing services, which included cell line construction platforms, process development platforms, analytical science and protein identification. Furthermore, the Group has two PIC/S GMP facilities certified by the Taiwan Food and Drug Administration (TFDA), one for mammalian cells and one for microbial, to provide clinical trial drug and commercial drug production.
- (2) The shares of the Company have been listed on the Taipei Exchange since June 28, 2019.
- 2. The Date of Authorisation for Issuance of the Financial Statements and Procedures for Authorisation These consolidated financial statements were authorised for issuance by the Board of Directors on March 8, 2024.
- 3. Application of New Standards, Amendments and Interpretations
 - (1) Effect of the adoption of new issuances of or amendments to International Financial Reporting Standards ("IFRS®") Accounting Standards that came into effect as endorsed by the Financial Supervisory Commission ("FSC")

New standards, interpretations and amendments endorsed by the FSC and became effective from 2023 are as follows:

	Effective date by
	International
	Accounting
New Standards, Interpretations and Amendments	Standards Board
Amendments to IAS 1, 'Disclosure of accounting policies'	January 1, 2023
Amendments to IAS 8, 'Definition of accounting estimates'	January 1, 2023
Amendments to IAS 12, 'Deferred tax related to assets and liabilities	January 1, 2023
arising from a single transaction'	
Amendments to IAS 12, 'International tax reform - pillar two model	May 23, 2023
rules'	

The above standards and interpretations have no significant impact to the Group's financial condition and financial performance based on the Group's assessment.

(2) Effect of new issuances of or amendments to IFRS Accounting Standards as endorsed by the FSC but not yet adopted by the Group

New standards, interpretations and amendments endorsed by the FSC and will become effective from 2024 are as follows:

	Effective date by
	International Accounting
New Standards, Interpretations and Amendments	Standards Board
Amendments to IFRS 16, 'Lease liability in a sale and leaseback'	January 1, 2024
Amendments to IAS 1, 'Classification of liabilities as current or non-	January 1, 2024
current'	
Amendments to IAS 1, 'Non-current liabilities with covenants'	January 1, 2024
Amendments to IAS 7 and IFRS 7, 'Supplier finance arrangements'	January 1, 2024

The above standards and interpretations have no significant impact to the Group's financial condition and financial performance based on the Group's assessment.

(3) IFRS Accounting Standards issued by IASB but not yet endorsed by the FSC

New standards, interpretations and amendments issued by IASB but not yet included in the IFRS Accounting Standards as endorsed by the FSC are as follows:

	Effective date by
	International Accounting
New Standards, Interpretations and Amendments	Standards Board
Amendments to IFRS 10 and IAS 28, 'Sale or contribution of assets	To be determined by
between an investor and its associate or joint venture'	International Accounting
	Standards Board
IFRS 17, 'Insurance contracts'	January 1, 2023
Amendments to IFRS 17, 'Insurance contracts'	January 1, 2023
Amendment to IFRS 17, 'Initial application of IFRS 17 and IFRS 9 –	January 1, 2023
comparative information'	
Amendments to IAS 21, 'Lack of exchangeability'	January 1, 2025

The above standards and interpretations have no significant impact to the Group's financial condition and financial performance based on the Group's assessment.

4. Summary of Material Accounting Policies

The principal accounting policies applied in the preparation of these consolidated financial statements are set out below. These policies have been consistently applied to all the periods presented, unless otherwise stated.

(1) Compliance statement

The consolidated financial statements of the Group have been prepared in accordance with the Regulations Governing the Preparation of Financial Reports by Securities Issuers, International Financial Reporting Standards, International Accounting Standards, IFRIC® Interpretations, and SIC® Interpretations that came into effect as endorsed by the FSC (collectively referred herein as the "IFRSs").

(2) Basis of preparation

- A. Except for the following items, the consolidated financial statements have been prepared under the historical cost convention:
 - (a) Financial assets and financial liabilities (including derivative instruments) at fair value through profit or loss.
 - (b) Financial assets at fair value through other comprehensive income.
- B. The preparation of financial statements in conformity with IFRSs requires the use of certain critical accounting estimates. It also requires management to exercise its judgement in the process of applying the Group's accounting policies. The areas involving a higher degree of judgement or complexity, or areas where assumptions and estimates are significant to the consolidated financial statements are disclosed in Note 5.

(3) Basis of consolidation

- A. Basis for preparation of consolidated financial statements:
 - (a) All subsidiaries are included in the Group's consolidated financial statements. Subsidiaries are all entities (including structured entities) controlled by the Group. The Group controls an entity when the Group is exposed, or has rights, to variable returns from its involvement with the entity and has the ability to affect those returns through its power over the entity. Consolidation of subsidiaries begins from the date the Group obtains control of the subsidiaries and ceases when the Group loses control of the subsidiaries.
 - (b) Inter-company transactions, balances and unrealised gains or losses on transactions between companies within the Group are eliminated. Accounting policies of subsidiaries have been adjusted where necessary to ensure consistency with the policies adopted by the Group.
 - (c) Profit or loss and each component of other comprehensive income are attributed to the owners of the parent and to the non-controlling interests. Total comprehensive income is attributed to the owners of the parent and to the non-controlling interests even if this results in the non-controlling interests having a deficit balance.

- (d) Changes in a parent's ownership interest in a subsidiary that do not result in the parent losing control of the subsidiary (transactions with non-controlling interests) are accounted for as equity transactions, i.e. transactions with owners in their capacity as owners. Any difference between the amount by which the non-controlling interests are adjusted and the fair value of the consideration paid or received is recognised directly in equity.
- (e) When the Group loses control of a subsidiary, the Group remeasures any investment retained in the former subsidiary at its fair value. That fair value is regarded as the fair value on initial recognition of a financial asset or the cost on initial recognition of the associate or joint venture. Any difference between fair value and carrying amount is recognised in profit or loss. All amounts previously recognised in other comprehensive income in relation to the subsidiary are reclassified to profit or loss on the same basis as would be required if the related assets or liabilities were disposed of. That is, when the Group loses control of a subsidiary, all gains or losses previously recognised in other comprehensive income in relation to the subsidiary should be reclassified from equity to profit or loss, if such gains or losses would be reclassified to profit or loss when the related assets or liabilities are disposed of.

B. Subsidiaries included in the consolidated financial statements:

			Ownersh	nip (%)
Name of	Name of		December 31,	December 31,
investor	subsidiary	Main business activities	2023	2022
The Company	EirGenix Europe GmbH	Biopharmaceutical research and development as well as business	100	100
The Company	EirGenix USA Inc.	Biopharmaceutical commissioned development, manufacturing services and consulting	100	-

- C. EirGenix USA Inc. is a subsidiary that was established in November 2023. As of December 31, 2023, no capital has been remitted.
- D. Subsidiaries not included in the consolidated financial statements: None.
- E. Adjustments for subsidiaries with different balance sheet dates: None.
- F. Significant restrictions: None.
- G. Subsidiaries that have non-controlling interests that are material to the Group: None.

(4) Foreign currency translation

Items included in the financial statements of each of the Group's entities are measured using the currency of the primary economic environment in which the entity operates (the "functional currency"). The consolidated financial statements are presented in New Taiwan dollars, which is the Company's functional and the Group's presentation currency.

A. Foreign currency transactions and balances

- (a) Foreign currency transactions are translated into the functional currency using the exchange rates prevailing at the dates of the transactions or valuation where items are remeasured. Foreign exchange gains and losses resulting from the settlement of such transactions are recognised in profit or loss in the period in which they arise.
- (b) Monetary assets and liabilities denominated in foreign currencies at the period end are retranslated at the exchange rates prevailing at the balance sheet date. Exchange differences arising upon re-translation at the balance sheet date are recognised in profit or loss.
- (c) Non-monetary assets and liabilities denominated in foreign currencies held at fair value through profit or loss are re-translated at the exchange rates prevailing at the balance sheet date; their translation differences are recognised in profit or loss. Non-monetary assets and liabilities denominated in foreign currencies held at fair value through other comprehensive income are re-translated at the exchange rates prevailing at the balance sheet date; their translation differences are recognised in other comprehensive income. However, non-monetary assets and liabilities denominated in foreign currencies that are not measured at fair value are translated using the historical exchange rates at the dates of the initial transactions.
- (d) All foreign exchange gains and losses are presented in the statement of comprehensive income within 'other gains and losses'.

B. Translation of foreign operations

The operating results and financial position of all the group entities that have a functional currency different from the presentation currency are translated into the presentation currency as follows:

- (a) Assets and liabilities for each balance sheet presented are translated at the closing exchange rate at the date of that balance sheet;
- (b) Income and expenses for each statement of comprehensive income are translated at average exchange rates of that period; and
- (c) All resulting exchange differences are recognised in other comprehensive income.

(5) Classification of current and non-current items

- A. Assets that meet one of the following criteria are classified as current assets; otherwise they are classified as non-current assets:
 - (a) Assets arising from operating activities that are expected to be realised, or are intended to be sold or consumed within the normal operating cycle;
 - (b) Assets held mainly for trading purposes;
 - (c) Assets that are expected to be realised within twelve months from the balance sheet date;
 - (d) Cash and cash equivalents, excluding restricted cash and cash equivalents and those that are to be exchanged or used to settle liabilities more than twelve months after the balance sheet date.
- B. Liabilities that meet one of the following criteria are classified as current liabilities; otherwise they are classified as non-current liabilities:
 - (a) Liabilities that are expected to be settled within the normal operating cycle;
 - (b) Liabilities arising mainly from trading activities;
 - (c) Liabilities that are to be settled within twelve months from the balance sheet date;
 - (d) Liabilities for which the repayment date cannot be extended unconditionally to more than twelve months after the balance sheet date. Terms of a liability that could, at the option of the counterparty, result in its settlement by the issue of equity instruments do not affect its classification.

(6) Cash equivalents

Cash equivalents refer to short-term, highly liquid investments that are readily convertible to known amounts of cash and which are subject to an insignificant risk of changes in value. Time deposits that meet the definition above and are held for the purpose of meeting short-term cash commitments in operations are classified as cash equivalents.

(7) Financial assets at fair value through profit or loss

- A. Financial assets at fair value through profit or loss are financial assets that are not measured at amortised cost or fair value through other comprehensive income.
- B. On a regular way purchase or sale basis, financial assets at fair value through profit or loss are recognised and derecognised using trade date accounting.
- C. At initial recognition, the Group measures the financial assets at fair value and recognises the transaction costs in profit or loss. The Group subsequently measures the financial assets at fair value, and recognises the gain or loss in profit or loss.

(8) Financial assets at fair value through other comprehensive income

- A. Financial assets at fair value through other comprehensive income comprise equity securities which are not held for trading, and for which the Group has made an irrevocable election at initial recognition to recognise changes in fair value in other comprehensive income.
- B. On a regular way purchase or sale basis, financial assets at fair value through other comprehensive income are recognised and derecognised using trade date accounting.
- C. At initial recognition, the Group measures the financial assets at fair value plus transaction costs. The Group subsequently measures the financial assets at fair value:

The changes in fair value of equity investments that were recognised in other comprehensive income are reclassified to retained earnings and are not reclassified to profit or loss following the derecognition of the investment. Dividends are recognised as revenue when the right to receive payment is established, future economic benefits associated with the dividend will flow to the Group and the amount of the dividend can be measured reliably.

(9) Financial assets at amortised cost

- A. Financial assets at amortised cost are those that meet all of the following criteria:
 - (a) The objective of the Group's business model is achieved both by collecting contractual cash flows and selling financial assets; and
 - (b) The assets' contractual cash flows represent solely payments of principal and interest.
- B. The Group's time deposits which do not fall under cash equivalents are those with a short maturity period and are measured at initial investment amount as the effect of discounting is immaterial.

(10) Accounts and notes receivable

- A. Accounts and notes receivable entitle the Group a legal right to receive consideration in exchange for transferred goods or rendered services.
- B. The short-term accounts and notes receivable without bearing interest are subsequently measured at initial invoice amount as the effect of discounting is immaterial.

(11) Impairment of financial assets

For debt instruments measured at fair value through profit or loss and financial assets at amortised cost, at each reporting date, the Group recognises the impairment provision for 12 months expected credit losses if there has not been a significant increase in credit risk since initial recognition or recognises the impairment provision for the lifetime expected credit losses (ECLs) if such credit risk has increased since initial recognition after taking into consideration all reasonable and verifiable information that includes forecasts. On the other hand, for accounts receivable or contract assets that do not contain a significant financing component, the Group recognises the impairment provision for lifetime ECLs.

(12) Derecognition of financial assets

The Group derecognises a financial asset when the contractual rights to receive the cash flows from the financial asset expire.

(13) Inventories

Inventories are stated at the lower of cost and net realisable value. Cost is determined using the first-in, first-out (FIFO) method. The cost of finished goods and work in progress comprises raw materials, direct labour, other direct costs and related production overheads. It excludes borrowing costs. The item by item approach is used in applying the lower of cost and net realisable value. Net realisable value is the estimated selling price in the ordinary course of business, less the estimated costs of completion and the estimated costs necessary to make the sale.

(14) Property, plant and equipment

- A. Property, plant and equipment are initially recorded at cost. Borrowing costs incurred during the construction period are capitalised.
- B. Subsequent costs are included in the asset's carrying amount or recognised as a separate asset, as appropriate, only when it is probable that future economic benefits associated with the item will flow to the Group and the cost of the item can be measured reliably. The carrying amount of the replaced part is derecognised. All other repairs and maintenance are charged to profit or loss during the financial period in which they are incurred.
- C. Land is not depreciated. Other property, plant and equipment apply cost model and are depreciated using the straight-line method to allocate their cost over their estimated useful lives. Each part of an item of property, plant, and equipment with a cost that is significant in relation to the total cost of the item must be depreciated separately.
- D. The assets' residual values, useful lives and depreciation methods are reviewed, and adjusted if appropriate, at each financial year-end. If expectations for the assets' residual values and useful lives differ from previous estimates or the patterns of consumption of the assets' future economic benefits embodied in the assets have changed significantly, any change is accounted for as a change in estimate under IAS 8, 'Accounting Policies, Changes in Accounting Estimates and Errors', from the date of the change. The estimated useful lives of property, plant and equipment are as follows:

Machinery and equipment	2 ~ 10 years
Office equipment	3 ~ 10 years
Buildings and structures	5 ~ 20 years
Leasehold improvements	3 ~ 20 years
Other equipment	3 ~ 10 years

(15) Leasing arrangements (lessee) — right-of-use assets/ lease liabilities

- A. Leases are recognised as a right-of-use asset and a corresponding lease liability at the date at which the leased asset is available for use by the Group. For short-term leases or leases of low-value assets, lease payments are recognised as an expense on a straight-line basis over the lease term.
- B. Lease liabilities include the net present value of the remaining lease payments at the commencement date, discounted using the incremental borrowing interest rate. Lease payments are comprised of the following:
 - (a) Fixed payments, less any lease incentives receivable; and
 - (b) Variable lease payments that depend on an index or a rate.

The Group subsequently measures the lease liability at amortised cost using the interest method and recognises interest expense over the lease term. The lease liability is remeasured and the amount of remeasurement is recognised as an adjustment to the right-of-use asset when there are changes in the lease term or lease payments and such changes do not arise from contract modifications.

- C. At the commencement date, the right-of-use asset is stated at cost comprising the following:
 - (a) The amount of the initial measurement of lease liability; and
 - (b) Any lease payments made at or before the commencement date.

The right-of-use asset is measured subsequently using the cost model and is depreciated from the commencement date to the earlier of the end of the asset's useful life or the end of the lease term. When the lease liability is remeasured, the amount of remeasurement is recognised as an adjustment to the right-of-use asset.

D. For lease modifications that decrease the scope of the lease, the lessee shall decrease the carrying amount of the right-of-use asset to reflect the partial or full termination of the lease, and recognise the difference between remeasured lease liability in profit or loss.

(16) Intangible assets

The Group's accounting policies on intangible assets are summarised below:

A. Computer software

Computer software is stated at cost and amortised on a straight-line basis over its estimated useful life of 1 to 5 years.

B. Professional expertise

Professional expertise is stated at cost and amortised on a straight-line basis over its estimated useful life of 2 to 10 years.

(17) <u>Impairment of non-financial assets</u>

The Group assesses at each balance sheet date the recoverable amounts of those assets where there is an indication that they are impaired. An impairment loss is recognised for the amount by which the asset's carrying amount exceeds its recoverable amount. The recoverable amount is the higher of an asset's fair value less costs to sell or value in use. When the circumstances or reasons for recognising impairment loss for an asset in prior years no longer exist or diminish, the impairment loss is reversed. The increased carrying amount due to reversal should not be more than what the depreciated or amortised historical cost would have been if the impairment had not been recognised.

(18) Borrowings

- A. Borrowings are recognised initially at fair value, net of transaction costs incurred. Borrowings are subsequently stated at amortised cost; any difference between the proceeds (net of transaction costs) and the redemption value is amortised over the period of the borrowings using the effective interest method.
- B. Fees paid on the establishment of loan facilities are recognised as transaction costs of the loan to the extent that it is probable that some or all of the facility will be drawn down. In this case, the fee is deferred until the draw-down occurs. To the extent there is no evidence that it is probable that some or all of the facility will be drawn down, the fee is capitalised as a pre-payment for liquidity services and amortised over the period of the facility to which it relates.

(19) Accounts payable

- A. Accounts payable are liabilities for purchases of raw materials, goods or services.
- B. The short-term accounts payable without bearing interest are subsequently measured at initial invoice amount as the effect of discounting is immaterial.

(20) Derecognition of financial liabilities

A financial liability is derecognised when the obligation specified in the contract is either discharged or cancelled or expires.

(21) Employee benefits

A. Short-term employee benefits

Short-term employee benefits are measured at the undiscounted amount of the benefits expected to be paid in respect of service rendered by employees in a period and should be recognised as expense in that period when the employees render service.

B. Pensions

For defined contribution plans, the contributions are recognised as pension expense when they are due on an accrual basis. Prepaid contributions are recognised as an asset to the extent of a cash refund or a reduction in the future payments.

C. Employees' compensation and directors' and supervisors' remuneration

Employees' compensation and directors' and supervisors' remuneration are recognised as
expense and liability, provided that such recognition is required under legal or constructive
obligation and those amounts can be reliably estimated. Any difference between the amounts
resolved by the shareholders and the actual amounts subsequently distributed is accounted
for as changes in estimates.

(22) Employee share-based payment

A. For the equity-settled share-based payment arrangements, the employee services received are measured at the fair value of the equity instruments granted at the grant date, and are recognised as compensation cost over the vesting period, with a corresponding adjustment to equity. The fair value of the equity instruments granted shall reflect the impact of market vesting conditions and non-vesting conditions. Compensation cost is subject to adjustment based on the service conditions that are expected to be satisfied and the estimates of the number of equity instruments that are expected to vest under the non-market vesting conditions at each balance sheet date. Ultimately, the amount of compensation cost recognised is based on the number of equity instruments that eventually vest.

B. Restricted stocks:

- (a) Restricted stocks issued to employees are measured at the fair value of the equity instruments granted at the grant date, and are recognised as compensation cost over the vesting period.
- (b) The restricted stocks issued by the Group cannot be transferred during the vesting period, but voting right and dividend right are not restricted on these stocks. Employees are not required to return the dividends received if they resign during the vesting period.
- (c) For restricted stocks where employees do not need to pay to acquire those stocks, if employees resign during the vesting period, they are considered not meeting the vesting condition from the date of resignation and the Group will redeem and retire those stocks at the initial issuance price.

(23) Income tax

A. The tax expense for the period comprises current and deferred tax. Tax is recognised in profit or loss, except to the extent that it relates to items recognised in other comprehensive income or items recognised directly in equity, in which cases the tax is recognised in other comprehensive income or equity

- B. The current income tax expense is calculated on the basis of the tax laws enacted or substantively enacted at the balance sheet date. Management periodically evaluates positions taken in tax returns with respect to situations in accordance with applicable tax regulations. It establishes provisions where appropriate based on the amounts expected to be paid to the tax authorities. An additional tax is levied on the unappropriated retained earnings and is recorded as income tax expense in the year the stockholders resolve to retain the earnings.
- C. Deferred tax is recognised, using the balance sheet liability method, on temporary differences arising between the tax bases of assets and liabilities and their carrying amounts in the consolidated balance sheet. However, the deferred tax is not accounted for if it arises from initial recognition of goodwill or of an asset or liability in a transaction other than a business combination that at the time of the transaction affects neither accounting nor taxable profit or loss and does not give rise to equal taxable and deductible temporary differences. Deferred tax is provided on temporary differences arising on investments in subsidiaries, except where the timing of the reversal of the temporary difference is controlled by the Group and it is probable that the temporary difference will not reverse in the foreseeable future. Deferred tax is determined using tax rates (and laws) that have been enacted or substantially enacted by the balance sheet date and are expected to apply when the related deferred tax asset is realised or the deferred tax liability is settled.
- D. Deferred tax assets are recognised only to the extent that it is probable that future taxable profit will be available against which the temporary differences can be utilised. At each balance sheet date, unrecognised and recognised deferred tax assets are reassessed.
- E. A deferred tax asset shall be recognised for the carryforward of unused tax credits resulting from research and development expenditures to the extent that it is possible that future taxable profit will be available against which the unused tax credits can be utilised.

(24) Share capital

Ordinary shares are classified as equity. Incremental costs directly attributable to the issue of new shares or stock options are shown in equity as a deduction, net of tax, from the proceeds.

(25) Revenue recognition

A. Service revenue

(a) The Group provides biopharmaceutical contract testing and development services. Revenue from providing services is recognised in the accounting period in which the services are rendered. For fixed-price contracts, revenue is recognised based on the actual service provided to the end of the reporting period as a proportion of the total services to be provided. This is determined based on the actual cost relative to the total expected cost. The customer pays at the time specified in the payment schedule. If the services rendered exceed the payment, a contract asset is recognised. If the payments

exceed the services rendered, a contract liability is recognised.

(b) The Group's estimate about revenue, costs and progress towards complete satisfaction of a performance obligation is subject to a revision whenever there is a change in circumstances. Any increase or decrease in revenue or costs due to an estimate revision is reflected in profit or loss during the period when the management become aware of the changes in circumstances.

B. Sales revenue

The Group sells self-developed products. Sales are recognised when control of the products has transferred, being when the products are delivered to the customer, the customer has full discretion over the channel and price to sell the products, and there is no unfulfilled obligation that could affect the customer's acceptance of the products. Delivery occurs when the products have been shipped to the specific location, the risks of obsolescence and loss have been transferred to the customer, and either the customer has accepted the products in accordance with the sales contract, or the Group has objective evidence that all criteria for acceptance have been satisfied.

C. Authorisation and cooperative development revenue

(a) The Group's authorisation and cooperative development transactions mainly arise from authorising intellectual property rights of pharmaceutical products to pharmaceutical factories. Although the Group will continuously provide research and development services on the pharmaceutical products, pharmaceutical factories can access the research and development outcome at any time. Based on the Group's assessment, the Group uses its special technologies in manufacturing pharmaceutical cell lines, which are unique so that pharmaceutical factories would have difficulty finding another similar service provider who offers the same services in terms of the subsequent research and development on the authorised pharmaceutical products. The authorisation and subsequent research and development services provided by the Group are bonded and highly interrelated, which does not meet the criteria of being distinct, and hence are accounted for as a single performance obligation to be delivered over time. Pharmaceutical factories pay a non-refundable up-front payment upon signing of the contracts, and make milestone payments upon each milestone achieved. The transaction prices, net of variable considerations that are not highly probable to be realised, are recognised as revenue based on the progress of performance obligations that are satisfied over time. The aforementioned stage of completion is determined based on the ratio of the actual research and development costs incurred at the end of the reporting period to the estimated total research and development costs for the authorisation contracts. The Group uses input method to measure progress towards the satisfaction of a performance obligation as there is a direct relationship between the transfer of control of services to

customers and the Group's inputs, including costs of contract research and development services, contract manufacturing services and medicines. Revenue is only recognised when it is highly probable that a significant reversal will not occur. The customer pays at the time specified in the payment schedule. If the services rendered exceed the payment, a contract asset is recognised. If the payments exceed the services rendered, a contract liability is recognised. A contract liability recognised as revenue through the performance obligation is satisfied over time.

(b) The Group also entered into contracts with pharmaceutical factories, whereby the Group is entitled to a sales-based royalty in exchange for a license of manufacturing and the right to sell pharmaceutical products. In accordance with the contracts, the Group will not undertake any activities that will significantly affect the intellectual property to which the customer has rights. The Group recognises revenue at the later of when the performance obligation has been satisfied and the subsequent transfer of control or sale occurs.

(26) Government grants

Government grants are recognised at their fair value only when there is reasonable assurance that the Group will comply with any conditions attached to the grants and the grants will be received. Government grants are recognised in profit or loss on a systematic basis over the periods in which the Group recognises expenses for the related costs for which the grants are intended to compensate.

(27) Operating segments

Operating segments are reported in a manner consistent with the internal reporting provided to the chief operating decision-maker. The Group's chief operating decision-maker, who is responsible for allocating resources and assessing performance of the operating segments.

5. Critical Accounting Judgements, Estimates and Key Sources of Assumption Uncertainty

The preparation of these consolidated financial statements requires management to make critical judgements in applying the Group's accounting policies and make critical assumptions and estimates concerning future events. Assumptions and estimates may differ from the actual results and are continually evaluated and adjusted based on historical experience and other factors. Such assumptions and estimates have a significant risk of causing a material adjustment to the carrying amounts of assets and liabilities within the next financial year; and the related information is addressed below:

- (1) <u>Critical judgements in applying the Group's accounting policies</u> None.
- (2) Critical accounting estimates and assumptions
 - A. Impairment on property, plant and equipment
 - (a) The Group assesses impairment based on its internal and external information and industry

characteristics and determines the separate cash flows of a specific group of assets, useful lives of assets and the future possible income and expenses arising from the assets depending on how assets are utilised and industrial characteristics. Any changes of economic circumstances or estimates due to the change of Group strategy might cause material impairment on assets in the future.

- (b) As of December 31, 2023, the carrying amount of property, plant and equipment was \$3,337,685.
- B. Recognition of service revenue and authorisation and cooperative development revenue
 - (a) Service revenue and authorisation and cooperative development revenue are recognised based on the stage of completion. The Group sets the key assumption factors for estimating total future cost based on the past operating experience, and regularly reviews and assesses the reasonableness of the basis for relevant assumptions.
 - (b) For the year ended December 31, 2023, the service revenue and authorisation and cooperative development revenue amounted to \$605,990 and \$141,472, respectively.

6. Details of Significant Accounts

(1) Cash and cash equivalents

	Dece	mber 31, 2023	December 31, 2022		
Cash on hand and petty cash	\$	61	\$	61	
Demand deposits		448,160		756,773	
Time deposits	-	4,604,962		5,370,051	
	\$	5,053,183	\$	6,126,885	

The Group transacts with a variety of financial institutions all with high credit quality to disperse credit risk, so it expects that the probability of counterparty default is remote.

(2) Financial assets at fair value through profit or loss

Items	Decei	mber 31, 2023	Dec	cember 31, 2022
Non-current items:				
Financial assets mandatorily measured				
at fair value through profit or loss				
Profit-sharing investment in new				
drug development	\$	58,390	\$	58,390
Limited partnership				
venture capital		20,000		
		78,390		58,390
Valuation adjustment		1,908		3,030
· · · · · · · · · · · · · · · · · · ·	\$	80,298	\$	61,420

- A. The Group recognised net (losses) gains amounting to (\$1,122) and \$2,863 on financial assets at fair value through profit or loss for the years ended December 31, 2023 and 2022, respectively.
- B. On April 18, 2022, the Group entered into a new drug development profit-sharing agreement for TSY-0110 (EG12043) (the "Product") with FORMOSA PHARMACEUTICALS, INC. to replace the original development and manufacturing related cooperation agreement. Raw materials for the product development stage were provided by the Group at a reasonable market price, and FORMOSA PHARMACEUTICALS, INC. was responsible for the research and development of the product, and the implementation of the product in and manufacturing of the product after completing the development of the product Either party may commercialize the product in the global market, and each party is entitled to receive 50% licensing interest in any future revenue or interest derived from the development and commercialization of the product. Under the agreement, the Group paid a consideration amounting to US\$30,000 thousand for the licensing interest, which will be paid in accordance with the agreement and the development schedule. As of December 31, 2023, the Group has paid US\$2,000 thousand.

(3) Financial assets at amortised cost

Items	Decen	nber 31, 2023	December 31, 2022		
Current items:					
Time deposits (Note)	\$	500,000	\$	1,000,000	
Non-current items:					
Government bonds	\$	31,930	\$	32,452	
Pledged time deposits		8,790		8,671	
	\$	40,720	\$	41,123	

Note: The deposit period for time deposits ranges between three months and a year.

A. Amounts recognised in profit or loss in relation to financial assets at amortised cost are listed below:

	Year ended December 31				
		2023	2022		
Interest income	\$	28,235	\$	1,722	

- B. Details of the Group's financial assets at amortised cost pledged to others as collateral are provided in Note 8.
- C. Information relating to credit risk of financial assets at amortised cost is provided in Note 12(2). The counterparties of the Group's investments in certificates of deposits and government bonds are financial institutions and governments with high credit quality, so the Group expects that the probability of counterparty default is remote.

(4) Notes and accounts receivable

	Decer	nber 31, 2023	December 31, 2022		
Notes receivable	\$	19	\$	<u> </u>	
Accounts receivable Less: Allowance for uncollectible	\$	253,687	\$	33,079	
accounts	(297)	(297)	
	\$	253,390	\$	32,782	

A. The ageing analysis of notes receivable and accounts receivable that were past due but not impaired is as follows:

	Decemb	er 31	, 2023	December 31, 2022			
	Notes receivable	Acc	counts receivable	Notes receivable	Accoun	nts receivable	
Not past due	\$ 19	\$	176,990	\$ -	\$	32,782	
Up to 30 days past due	-		76,400	-		-	
31 to 90 days past due	-		-	-		-	
91 to 180 days past due	-		-	-		-	
Over 181 days past due			297			297	
	\$ 19	\$	253,687	\$ -	\$	33,079	

The above ageing analysis was based on past due date.

- B. As of December 31, 2023 and 2022, notes receivable and accounts receivable(including related parties) were all from contracts with customers. Also, as of January 1, 2022, the balance of receivables from contracts with customers amounted to \$80,159.
- C. As at December 31, 2023 and 2022, without taking into account any collateral held or other credit enhancements, the maximum exposure to credit risk in respect of the amount that best represents the notes and accounts receivable (including related parties) held by the Group was \$256,045 and \$32,782, respectively.
- D. The Group did not hold any collateral.
- E. Information relating to credit risk of accounts receivable and notes receivable is provided in Note 12(2).

(5) <u>Inventories</u>

	December 31, 2023						
		Cost		Allowance for valuation loss		Book value	
Raw materials	\$	426,217	(\$	51,483)	\$	374,734	
Work in progress		127,143		-		127,143	
Finished goods		178,690		(165)		178,525	
Merchandise inventory		235		_		235	
	\$	732,285	(\$	51,648)	\$	680,637	
	December 31, 2022						
	Allowance for						
		Cost		valuation loss		Book value	
Raw materials	\$	377,424	(\$	18,327)	\$	359,097	
Work in progress		281,739		-		281,739	
Finished goods		98,150		-		98,150	
Merchandise inventory		477		-		477	
	\$	757,790	(\$	18,327)	\$	739,463	

The cost of inventories recognized as expense for the year:

	Year ended December 31				
		2023		2022	
Cost of goods used	\$	327,739	\$	199,628	
Cost of goods sold		149,041		138,672	
Loss on decline in market value		33,321		1,012	
Loss on disposal inventory		1,014		-	
(Gain) loss on physical inventory	(4)		34	
	\$	511,111	\$	339,346	

(6) Prepayments

	Decen	nber 31, 2023	December 31, 2022		
Office supplies	\$	-	\$	9,009	
Prepayments for contracted research expense		17,151		11,310	
Excess business tax paid (or Net Input VAT)		24,454		6,267	
Prepayments to suppliers		26,187		50,100	
Other prepaid expenses		26,010		46,756	
	\$	93,802	\$	123,442	

(7) Financial assets at fair value through other comprehensive income

Items	 December 31, 2023	ecember 31, 2023 December 31, 2		
Non-current items:				
Equity instruments				
Emerging and unlisted stocks	\$ 215,026	\$	214,403	
Valuation adjustment	 110,861		64,922	
	\$ 325,887	\$	279,325	

- A. The Group has elected to classify shares that are considered to be strategic investments as financial assets at fair value through other comprehensive income. The fair value of such investments amounted to \$325,887 and \$279,325 as at December 31, 2023 and 2022, respectively.
- B. The Group acquired equity instruments amounting to \$623 and \$208,627 for the year ended December 31, 2023 and 2022.
- C. Amounts recognised in profit or loss and other comprehensive income in relation to the financial assets at fair value through other comprehensive income are listed below:

	Year ended December 31								
		2023	2022						
Equity instruments at fair value									
through other comprehensive									
income									
Fair value change recognised									
in other comprehensive income	\$	45,939	\$	59,091					
Dividend income recognised									
in profit or loss Held at end									
of period	\$	475	\$	_					

(8) Property, plant and equipment

$^{\circ}$	^	1	1
7	u	12	1

		achinery and equipment	_	Office equipment	E	Buildings and structures	in	Leasehold nprovements		Other equipment	equ	Unfinished astruction and aipment under acceptance		Total	busi (sh	epayments for iness facilities own as other non-current assets)
At January 1 Cost Accumulated depreciation	\$ (978,923 317,142)	\$	75,921 30,726)	\$	1,434,479 229,062)	\$	45,596 12,142)	\$	32,925 12,788)	\$	642,864	\$	3,210,708 601,860)	\$	98,273
•	\$	661,781	\$	45,195	\$	1,205,417	\$	33,454	\$	20,137	\$	642,864	\$	2,608,848	\$	98,273
Opening net book amount as at January 1 Additions Reclassifications	\$	661,781 116,776 147,987	\$	45,195 6,447	\$	1,205,417 15,263 528,357	\$	33,454 1,665	\$	20,137 5,768		642,864 557,083 676,344)	\$	2,608,848 703,002	\$	98,273 138,453
Transfers from other non- current assets		62,723		32		-		59		1,294		158,129		222,237	(222,237)
Depreciation expense	(98,594)	(9,399)	(78,412)	(4,809)	(5,219)		-	(196,433)		-
Net exchange differences Closing net book amount		-		31				<u> </u>	_	<u> </u>		<u> </u>		31		<u>-</u>
as at December 31	\$	890,673	\$	42,306	\$	1,670,625	\$	30,369	\$	21,980	\$	681,732	\$	3,337,685	\$	14,489
At December 31 Cost Accumulated depreciation	\$ (<u> </u>	1,301,038 410,365)	\$ (<u> </u>	80,678 38,372)	\$ (<u> </u>	1,978,099 307,474)	\$ (<u></u>	47,320 16,951)	\$ (<u> </u>	37,142 15,162)	\$	681,732	\$ (<u> </u>	4,126,009 788,324)	\$	14,489
	\$	890,673	\$	42,306	\$	1,670,625	\$	30,369	\$	21,980	\$	681,732	\$	3,337,685	\$	14,489

At January 1 Cost Accumulated depreciation		chinery and quipment 813,793 239,109) 574,684		Office equipment 68,349 24,341) 44,008	\$ (Buildings and structures 1,295,911 164,219) 1,131,692	in	Leasehold nprovements 24,495 8,974) 15,521	- \$ (Other equipment 26,524 8,870) 17,654	eq	Unfinished nstruction and uipment under acceptance 103,265 103,265	 \$ (\$	Total 2,332,337 445,513) 1,886,824	bus (sł	epayments for iness facilities nown as other non-current assets) 65,456
									_	<u> </u>			_			
Opening net book amount as at January 1 Additions Reclassifications	\$	574,684 92,578 7,701	\$	44,008 8,863	\$	1,131,692 76,679 61,890	\$	15,521 21,101	\$	17,654 6,516		103,265 278,775 69,591)	\$	1,886,824 484,512	\$	65,456 433,952
Transfers from other non- current assets		69,453		862		-		-		405	(330,415		401,135	(401,135)
Depreciation expense	(80,909)	(8,566)	(64,844)	(3,168)	(4,438)		-	(161,925)		-
Reclassified to inventories	(1,726)		-		-		-		-		-	(1,726)		-
Net exchange differences				28							_			28	_	
Closing net book amount as at December 31	\$	661,781	\$	45,195	\$	1,205,417	\$	33,454	\$	20,137	\$	642,864	\$	2,608,848	\$	98,273
At December 31 Cost Accumulated depreciation	\$ (978,923 317,142)	\$ (<u></u>	75,921 30,726)	\$ (<u> </u>	1,434,479 229,062)	\$ (<u></u>	45,596 12,142)	\$ (<u> </u>	32,925 12,788)	\$	642,864	\$ (<u></u>	3,210,708 601,860)	\$	98,273
	\$	661,781	\$	45,195	\$	1,205,417	\$	33,454	\$	20,137	\$	642,864	\$	2,608,848	\$	98,273

Information about the property, plant and equipment that were pledged to others as collateral is provided in Note 8.

(9) <u>Leasing arrangements - lessee</u>

- A. The Group leases various assets including land, buildings, machinery and equipment, multifunction printers and business vehicles. Rental contracts are typically made for periods of 1 to 20 years. Lease terms are negotiated on an individual basis and contain a wide range of different terms and conditions. The lease agreements do not impose covenants, but leased assets may not be used as security for borrowing purposes.
- B. Short-term leases with a lease term of 12 months or less comprise certain offices, dormitories, business vehicles and warehouses. Low-value assets comprise multifunction printers.
- C. The carrying amount of right-of-use assets and the depreciation charge are as follows:

		December 31, 2023	December 31, 2022						
		Carrying amount		Carrying amount					
Land	\$	187,939	\$	202,394					
Buildings		73,893		84,031					
Machinery and equipment		65,921		35,305					
Transportation equipment		967		2,584					
(Business vehicles)									
Office equipment (Multifunction printers)		516	_	1,016					
running,	\$	329,236	\$	325,330					
	Year ended December 31								
		2023	2022						
		Depreciation expense		Depreciation expense					
Land	\$	14,648	\$	14,543					
Buildings		10,292		8,288					
Machinery and equipment		4,055		2,310					
Transportation equipment (Business vehicles)		1,616		1,640					
Office equipment (Multifunction printers)	-	500	_	396					
	\$	31,111	\$	27,177					

- D. For the years ended December 31, 2023 and 2022, the additions to right-of-use assets were \$35,017 and \$54,767, respectively.
- E. The information on profit and loss accounts relating to lease contracts is as follows:

Van	andad	December	21
I cai	enaea	December	21

	2023	2022		
Items affecting profit or loss				
Interest expense on lease liabilities	\$ 8,096	\$ 8,208		
Expense on short-term lease contracts	27,123	13,877		
Expense on leases of low-value assets	307	376		
Loss on lease modification	413	709		

F. For the years ended December 31, 2023 and 2022, the Group's total cash outflow for leases were \$64,833 and \$46,896, respectively.

(10) Intangible assets

	2023									
		Software	Prof	essional expertise		Total				
At January 1										
Cost	\$	45,851	\$	107,953	\$	153,804				
Accumulated amortisation	(21,678)	(104,059)	(125,737)				
	\$	24,173	\$	3,894	\$	28,067				
Opening net book amount as										
at January 1	\$	24,173	\$	3,894	\$	28,067				
Additions		3,339		8,159		11,498				
Amortisation expense	(7,129)	(4,167)	(11,296)				
Closing net book amount as at December 31	\$	20,383	\$	7,886	\$	28,269				
At December 31										
Cost	\$	49,190	\$	116,112	\$	165,302				
Accumulated amortisation	(28,807)	(108,226)	(137,033)				
	\$	20,383	\$	7,886	\$	28,269				

			2022		
	Software	Profe	ssional expertise		Total
\$	21,153	\$	107,953	\$	129,106
(16,438)	(93,115)	(109,553)
\$	4,715	\$	14,838	\$	19,553
\$	4,715	\$	14,838	\$	19,553
	8,652		-		8,652
	16,046		-		16,046
(5,240)	(10,944)	(16,184)
\$	24,173	\$	3,894	\$	28,067
\$	45,851	\$	107,953	\$	153,804
(21,678)	(104,059)	(125,737)
\$	24,173	\$	3,894	\$	28,067
	\$ ((16,438) \$ 4,715 \$ 4,715 \$ 8,652 16,046 (5,240) \$ 24,173 \$ 45,851 (21,678)	\$ 21,153 \$ (16,438) (\$ 4,715 \$ \$ \$ 4,715 \$ \$ \$ 8,652 \$ 16,046 \$ 5,240) (\$ 24,173 \$ \$ \$ 45,851 \$ \$ (21,678) (Software Professional expertise \$ 21,153 \$ 107,953 (16,438) (93,115) \$ 4,715 \$ 14,838 \$ 8,652 - 16,046 - (5,240) (10,944) \$ 24,173 \$ 3,894 \$ 45,851 \$ 107,953 (21,678) (104,059)	Software Professional expertise \$ 21,153 \$ 107,953 \$ (16,438) (93,115) (93,115) (

2022

Note: Transfers pertain to assets transferred from prepaid intangible assets (shown as "other non-current assets").

A. Details of amortisation on intangible assets are as follows:

		per 31		
		2023		2022
Operating costs	\$	5,840	\$	10,456
General and administrative expenses		1,193		1,005
Research and development expenses		4,156		4,705
Sales and marketing expenses		107		18
	\$	11,296	\$	16,184

- B. The basic information of the professional expertise that is material to the Group is as follows:
 - (a) In April 2013, the Group acquired professional expertise, including cell line establishment, process development, process optimisation, analytical method development and validation, product qualification, GMP manufacturing and stability test, etc., amounting to \$92,483 from the Development Center for Biotechnology cGMP biopharmaceutical pilot plant facility.
 - (b) In July 2013, the Group acquired professional expertise of Herceptin from FORMOSA PHARMACEUTICALS, INC. amounting to \$7,143.
 - (c) In July 2013, the Group acquired commercial authorisation of recombinant protein cell line from Life Technologies Corporation amounting to \$7,485.

(d) In September 2023, the Group obtained an authorisation from American Type Culture Collection for the detection of cancer cell lines with a total price of \$8,159, which can be applied on the commercial implementation of the marketing and manufacturing of subsequent cancer drug products.

(11) Other non-current assets

	Dece	ember 31, 2023	December 31, 2022			
Non-current prepayments for investments	\$	46,270	\$	20,000		
Long-term prepayments to suppliers		30,000		30,000		
Prepayments for business facilities		14,489		98,273		
Guarantee deposits paid		8,795		65,048		
Other assets		5,404		1,844		
	\$	104,958	\$	215,165		

(12) Other payables

	Decen	nber 31, 2023	December 31, 2022			
Payable on equipment	\$	285,960	\$	158,228		
and intangible assets						
Salary and bonus payable		99,260		95,239		
Service expense payable		44,882		52,083		
Payable on consumables		18,604		25,012		
Payable on repairs and		28,856		19,732		
maintenance expense						
Others		52,737	-	57,093		
	\$	530,299	\$	407,387		

(13) Long-term borrowings

Type of borrowings	Borrowing period and repayment term	Interest rate range	Collateral	December 31, 2023	
Long-term bank borrowings	S				
Credit borrowing	Borrowing period is from February 15, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from March 2025	1.7250%~ 1.9500%	None	\$	39,560
п	Borrowing period is from June 30, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from March 2025.	1.7250%~ 1.9500%	"		
				-	80,900
				\$	120,460

Type of borrowings	Borrowing period and repayment term	Interest rate range	Collateral	Decem	December 31, 2022	
Long-term bank borrowings						
Credit borrowing	Borrowing period is from February 15, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from March 2025	1.3500%~ 1.8250%	None	\$	39,560	
n	Borrowing period is from June 30, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from March 2025.	1.4750%~ 1.8250%	n.			
	,				80,900	
				\$	120,460	

- A. Information on the Group's undrawn borrowing facilities is provided in Note 12(2) C.
- B. On December 23, 2021, the Company entered into a \$714,000 syndicated loan agreement with Hua Nan Commercial Bank Ltd. and the government will subsidize 0.5% handling fee of the bank for the Company's compliance with the "Action Plan for Accelerated Investment by Domestic Corporations".
- C. Information about assets pledged as collateral for long-term borrowings is provided in Note 8.

(14) Pensions

- A. The Company has established a defined contribution pension plan (the "New Plan") under the Labor Pension Act (the "Act"), covering all regular employees with R.O.C. nationality. Under the New Plan, the Company contributes monthly an amount not lower than 6% of the employees' monthly salaries and wages to the employees' individual pension accounts at the Bureau of Labor Insurance. The benefits accrued are paid monthly or in lump sum upon termination of employment.
- B. EirGenix Europe GmbH contributed pension under local regulations.
- C. The pension costs under defined contribution pension plans of the Group for the years ended December 31, 2023 and 2022, were \$20,915 and \$18,290, respectively.

(15) Share based payment

A. For the years ended December 31, 2023 and 2022, the Group's share-based payment arrangements were as follows:

		Quantity granted		
Type of		(shares in		
arrangement	Grant date	thousands)	Contract period	Vesting conditions
Employee stock	2015. 07. 01	1,270	10 years	1 to 4 years'
options - B			·	service
"	2015. 07. 01	130	"	"
"	2015. 07. 06	250	"	"
"	2016. 01. 01	270	"	"
Employee stock	2016. 05. 05	100	10 years	2 to 4 years'
options - C				service
Employee stock	2016. 10. 12	515	10 years	2 to 4 years'
options - D				service
"	2016. 12. 29	85	"	"
Employee stock	2017. 08. 08	395	10 years	2 to 4 years'
options - E				service
"	2017. 12. 27	570	"	"
"	2018. 03. 23	175	"	"
Employee stock	2019. 01. 25	520	10 years	2 to 4 years'
options - F				service
"	2019. 05. 13	285	"	"
Restricted stocks	2016. 11. 18	1,660	N/A	Conditions of
to employees - A				service years and
				performance
"	2017. 08. 08	257	"	"
Employee stock	2019. 11. 12	960	10 years	2 to 4 years'
options - G			•	service
"	2020. 04. 15	775	"	"
11	2020. 08. 12	205	"	"
Restricted stocks	2020. 05. 13	455	N/A	0.25 to 3 years'
to employees - B				service
"	2020. 12. 10	144	"	"
Restricted stocks	2020. 08. 14	905	N/A	Performance
to employees - D				conditions
"	2020. 12. 10	94	"	"
Employee stock	2020. 12. 23	830	10 years	2 to 4 years'
options - H				service
"	2021. 05. 12	315	"	"
"	2021. 08. 12	505	"	"
"	2021. 10. 01	1,185	"	"

		Quantity granted		
Type of		(shares in		
arrangement	Grant date	thousands)	Contract period	Vesting conditions
Restricted stocks	2021. 10. 15	613	N/A	Performance
to employees - E				conditions
"	2022. 01. 10	184	"	"
"	2022. 09. 08	190	"	"
Restricted stocks	2021. 10. 15	340	N/A	Performance
to employees - F				conditions
Employee stock	2022. 03. 22	160	10 years	2 to 4 years'
options - I				service
"	2022. 05. 12	225	"	"
"	2022. 08. 11	685	"	"
"	2022. 09. 08	510	"	"
Restricted stocks	2022. 09. 08	63	N/A	Performance
to employees - G				conditions
"	2022. 11. 08	195	"	"
"	2023. 03. 10	6	"	"
"	2023. 11. 09	325	"	"
Employee stock	2022. 11. 08	615	10 years	2 to 4 years'
options - J				service
"	2023. 03. 10	1,105	"	"
"	2023. 05. 10	255	"	"
"	2023. 08. 08	225	"	"
"	2023. 12. 22	270	"	"
Restricted stocks	2023. 11. 09	826	N/A	Performance
to employees - H				conditions
Restricted stocks	2023. 12. 22	26	N/A	Performance
to employees - I				conditions

- (a) The restricted stocks issued by the Group cannot be transferred during the vesting period, but voting right and dividend right are not restricted on these stocks. If employees resign during the vesting period, they are considered not meeting the vesting condition from the date of resignation and the Group will redeem and retire those stocks at the initial issuance price, but employees are not required to return the dividends received.
- (b) The above-mentioned share-based payment arrangements are equity-settled.

B. Details of the share-based payment arrangements are as follows:

(a) Employee stock options

		2023	20	2022			
	No. of	Weighted-	No. of	Weighted-			
	options	average	options	average			
	(shares in	exercise price	(shares in	exercise price			
	thousands)	(in dollars)	thousands)	(in dollars)			
Options outstanding at January 1	5,666	\$15~146.4	5,282	\$15~146.4			
Options granted	1,855	100.5~120	2,195	71.6~118.5			
Options forfeited	(894	25.2~146.4	(759)	25.2~146.4			
Options exercised	(727	<u>/</u>) 15~51.2	(1,052)	15~51.2			
Options outstanding at December 31	5,900	15~146.4 <u>)</u>	5,666	15~146.4			
Options exercisable at December 31	1,608	3=	1,238				

(b) Restricted stocks to employees

		2023	2022
		(shares in thousands)	(shares in thousands)
Stocks outstanding at January 1		2,571	2,869
Stocks granted		1,182	632
Stocks vested	(1,167) (704)
Stocks retired	(193) (226)
Stocks outstanding		2,393	2,571
at December 31			

C. The weighted-average stock prices of stock options at exercise dates for the years ended December 31, 2023 and 2022 were \$105.6 (in dollars) and \$99.8 (in dollars), respectively.

D. The expiry date and exercise price of stock options outstanding at the balance sheet dates are as follows:

			December 31, 2023		December 31, 2022	
				Exercise		Exercise
			No. of shares	price	No. of shares	price
Type of	Issue date		(shares in		(shares in	
arrangement	approved	Expiry date	thousands)	(in dollars)	thousands)	(in dollars)
Employee		2025. 06. 30	50	\$ 15	140	\$ 15
stock				,		,
options - B						
"	2015 07 01	2025. 06. 30	5	20	20	20
"		2025. 00. 30	15	20	25	20
"		2025. 12. 31	25	20	25	20
Employee		2026. 05. 04	10	29.2	10	29.2
stock						
options - C						
Employee	2016 10 12	2026. 10. 11	150	29.2	180	29.2
stock	2010. 10. 12	2020. 10. 11	130	27.2	100	27.2
options - D						
"	2016 12 29	2026. 12. 28	15	37.5	15	37.5
Employee		2027. 08. 07	4	29.2	18	29.2
stock						
options - E						
"	2017 12 27	2027. 12. 26	79	25	112	25
"		2028. 03. 22	48	23.5	52	23.5
Employee		2029. 01. 24	34	28.7	103	28.7
stock						
options - F						
"	2019. 05. 13	2029. 05. 12	94	34.3	141	34.3
Employee	2019. 11. 12	2029. 11. 11	207	25.2	325	25.2
stock						
options - G						
"	2020. 04. 15	2030. 04. 14	89	28.8	175	28.8
"	2020. 08. 12	2030. 08. 11	79	51.2	140	51.2
Employee	2020. 12. 23	2030. 12. 22	341	42.1	515	42.1
stock						
options - H	2024 27 27	2024 27 1:				
"		2031. 05. 11	215	146.4	235	146.4
"		2031. 08. 11	250	128.4	305	128.4
••	2021. 10. 01	2031. 09. 30	835	117.5	990	117.5

			December 31, 2023		December 31, 2022		2022	
				Exercis	se		E	xercise
			No. of shares	price		No. of shares		price
Type of	Issue date		(shares in			(shares in		
arrangement	approved	Expiry date	_thousands)	(in dolla	rs)	_thousands)	(in	dollars)
Employee	2022 .03. 22	2032. 03. 21	80	\$ 9	3.5	145	\$	93.5
stock								
options - I								
"	2022. 05. 12	2032. 05. 11	195	7	1.6	225		71.6
"	2022. 08. 11	2032. 08. 10	440	8	5.9	645		85.9
"	2022. 09. 08	2032. 09. 07	345	11	8.5	510		118.5
Employee	2022. 11. 08	2032. 11. 07	510	10	3.5	615		103.5
stock								
options - J								
"	2023. 03 .10	2033. 03. 09	1,035	11	1.5	-		-
"	2023. 05. 10	2033. 05. 09	255	12	0.0	-		-
"	2023. 08. 08	2033. 08. 07	225	10	1.5	-		_
"	2023. 12. 22	2033. 12. 21	270	10	0.5	-		-

E. The fair value of stock options granted is measured using the Black-Scholes option-pricing model to estimate the fair value of employee stock options, cash capital increase reserved for employee preemption and restricted stocks to employees. Relevant information is as follows:

Type of		Quantity granted (shares in	Stock price	Exercise price	Expected price	Expected option	Risk-free	Fair value per
arrangement	Grant date	thousands)	(in dollars)	(in dollars)	volatility	life	interest rate	unit (in dollars)
Employee stock options - B	2015. 07. 01	1,270	\$ 14.88	\$ 15	36.58~ 37.13%	5.5 ~ 7 years	1.15~ 1.35%	\$5.22 ~ 6.01
"	2015. 07. 01	130	14.88	20	36.58~ 37.13%	5.5 ~ 7 years	1.15~ 1.35%	3.83~ 4.69
"	2015. 07. 06	250	14.60	20	37.09~ 37.64%	5.5 ~ 7 years	1.15~ 1.35%	3.75~ 4.6
"	2016. 01. 01	270	16.03	20	40.11~ 40.30%	5.5 ~ 7 years	0.79~ 0.90%	4.91~ 5.76
Employee stock options - C	2016. 05. 05	100	13.27	29.2	40.75~ 40.91%	6 ~ 7 years	0.70~ 0.77%	1.86 ~ 2.30
Employee stock options - D	2016. 10. 12	515	21.42	29.2	39.82~ 39.91%	6 ~ 7 years	0.71~ 0.75%	5.19~ 5.93
"	2016. 12. 29	85	20.40	37.5	39.39~ 39.48%	6 ~ 7 years	1.16~ 1.20%	3.49~ 4.18

Type of arrangement	Grant date	Quantity granted (shares in thousands)	Stock price (in dollars)	Exercise price (in dollars)	Expected price volatility	Expected option life	Risk-free	Fair value per unit (in dollars)
Employee stock options - E	2017. 08. 08	395	\$ 18.75	\$ 29.2	38.13~ 38.22%	6 ~ 7 years	0.82~ 0.88%	\$3.64~ 4.23
"	2017. 12. 27	570	18.07	25	36.97~ 37.23%	6 ~ 7 years	0.74~ 0.80%	3.81~ 4.41
"	2018. 03. 23	175	19.16	23.5	36.87~ 37.17%	6 ~ 7 years	0.79~ 0.84%	4.71 ~ 5.38
Employee stock options - F	2019. 01. 25	520	21.96	28.7	36.03~ 36.90%	6 ~ 7 years	0.72~ 0.78%	4.85~ 5.74
"	2019. 05. 13	285	25.75	34.3	35.50~ 36.35%	6 ~ 7 years	0.64~ 0.67%	5.39 ~ 6.40
Restricted stocks to employees - A	2016. 11. 18	1,660	22.88	-	-	-	-	22.88
"	2017. 08. 08	257	19.61	-	-	-	-	19.61
Employee stock options - G	2019. 11. 12	960	29.05	25.2	26.38%	6 ~ 7 years	0.63~ 0.66%	7.77 ~ 8.42
"	2020. 04. 15	775	33.10	28.8	50.33%	6 ~ 7 years	0.47~ 0.49%	15.56 ~ 16.65
"	2020. 08. 12	205	57.80	51.2	64.08%	6 ~ 7 years	0.36~ 0.38%	33.07 ~ 35.18
Restricted stocks to employees - B	2020. 05. 13	455	46.85	-	-	-	-	46.85
"	2020. 12. 10	144	48.60	-	-	-	-	48.60
Restricted stocks to employees - D	2020. 08. 14	905	55.70	-	-	-	-	55.70
"	2020. 12. 10	94	48.60	-	-	-	-	48.60
Employee stock options - H	2020. 12. 23	830	47.55	42.1	61.28%	6 ~ 7 years	0.22~ 0.26%	26.15~ 27.88
"	2021. 05. 12	315	154.5	146.4	65.02%	6 ~ 7 years	0.31~ 0.35%	89.32~ 95.02
"	2021. 08. 12	505	135.5	128.4	67.02%	6 ~ 7 years	0.32~ 0.34%	80.24~ 85.25
"	2021. 10. 01	1,185	124.0	117.5	65.78%	6 ~ 7 years	0.34% 0.34~ 0.38%	72.39~ 76.99

Type of arrangement	Grant date	Quantity granted (shares in thousands)	x price	Exercise price (in dollars)	Expected price volatility	Expected option life	Risk-free	Fair value per
Restricted stocks to employees - E	2021. 10. 15	613	\$ 106.5	\$ -	-	-	-	\$106.5
- <u>L</u>	2022. 01. 10	184	108.5	-	-	-	-	108.5
"	2022. 09. 08	190	118.5	-	-	-	-	118.5
Restricted stocks to employees - F	2021. 10. 15	340	106.5	-	-	-	-	106.5
Employee stock options - I	2022. 03. 22	160	93.5	93.5	62.20%	6 ~ 7 years	0.86~ 0.87%	52.85~ 56.27
"	2022. 05. 12	225	71.6	71.6	61.32%	6 ~ 7 years	1.22~ 1.27%	40.37~ 43.04
"	2022. 08. 11	685	85.9	85.9	60.04%	6 ~ 7 years	1.10~ 1.14%	47.51~ 50.67
"	2022. 09. 08	510	118.5	118.5	60.29%	6 ~ 7 years	1.19~ 1.23%	65.9~ 70.28
Restricted stocks to employees - G	2022. 09. 08	63	118.5	-	-	-	-	118.5
"	2022. 11. 08	195	103.5	-	-	-	-	103.5
"	2023. 03. 10	6	111.5	-	-	-	-	111.5
"	2023. 11. 09	325	103.0	-	-	-	-	103.0
Employee stock options - J	2022. 11. 08	615	103.5	103.5	60.00%	6 ~ 7 years	1.63~ 1.70%	57.97~ 61.88
"	2023. 03. 10	1,150	111.5	111.5	59.15%	6 ~ 7 years	1.12~ 1.14%	60.98~ 65.04
"	2023. 05. 10	255	120.0	120.0	58.70%	6 ~ 7 years	1.07~ 1.09%	65.15~ 69.50
"	2023. 08. 08	225	101.5	101.5	57.40%	6 ~ 7 years	1.10~ 1.12%	54.18~ 57.84
"	2023. 12. 22	270	100.5	100.5	55.38%	6 ~ 7 years	1.18~ 1.19%	52.26~ 55.82
Restricted stocks to employees - H	2023. 11. 09	826	103.0	-	-	-	-	103.0
Restricted stocks to employees - I	2023. 12. 22	26	100.5	-	-	-	-	100.5

F. Expenses incurred on share-based payment transactions are shown below:

	Year ended December 31				
		2023	2022		
Employee stock options	\$	84,285	\$	62,522	
Restricted stocks to employees		96,615		92,790	
	\$	180,900	\$	155,312	

(16) Share capital

A. As of December 31, 2023, the Company's authorised capital was \$4,000,000, consisting of 400,000 thousand shares of ordinary share (including 12 million shares reserved for employee stock options, preferred shares with warrants or convertible bonds issued by the Company), and the paid-in capital was \$3,060,516 with a par value of \$10 (in dollars) per share, consisting of 306,052 thousand shares. All proceeds from shares issued have been collected.

Movements in the number of the Company's ordinary shares outstanding are as follows (unit: shares in thousands):

		2023	2022
At January 1		304,336	300,385
Employee stock options			
exercised		727	1,052
Issuance of employee restricted			
stocks		1,182	632
Redemption of employee restricted			
stocks	(193) (226)
Conversions of convertible bonds			2,493
At December 31	\$	306,052	\$ 304,336

- B. For the years ended December 31, 2023 and 2022, the Company issued 727 thousand and 1,052 thousand ordinary shares related to the exercise of employee share options in accordance with the employee share options plan with a par value of \$10 (in dollars) per share, totalling \$7,270 and \$10,523, respectively.
- C. For the years ended December 31, 2023 and 2022, as employee restricted stocks distributed to certain employees did not meet the vesting conditions in accordance with the terms of restricted shares, the Company's Board of Directors resolved to repurchase and retire the employee restricted stocks amounting to 193 thousand and 226 thousand shares, respectively.

- D. The shareholders during their meeting on August 3, 2021 resolved to issue the 1st and 2nd restricted stocks to employees amounting to 1,000 thousand and 340 thousand shares with no subscription price, respectively. For the year ended December 31, 2021, the Board of Directors of the Company resolved to issue the 1st and 2nd restricted stocks to employees amounting to 797 thousand and 340 thousand shares in 2021, respectively. The Board of Directors of the Company resolved to issue the 1st restricted stocks to employees amounting to 190 thousand shares in 2021.
- E. The shareholders during their stockholders' meeting on August 3, 2021 resolved to issue 55,000 thousand ordinary shares through the private placement. The Board of Directors of the Company resolved the issuance price of \$91.5 (in dollars) and the total consideration of issuing common stock was \$5,032,500 on October 1, 2021, and the effective date was set on October 15, 2021. The registration has been completed on December 13, 2021. Pursuant to the Securities and Exchange Act, the ordinary shares raised through the private placement are subject to certain transfer restrictions and cannot be listed on the stock exchange until three years after they have been issued and have been offered publicly. Other than these restrictions, the rights and obligations of the ordinary shares raised through the private placement are the same as other issued ordinary shares.
- F. The shareholders during their meeting on June 10, 2022, resolved to issue the 1st restricted stocks to employees amounting to 850 thousand shares with no subscription price. On September 8, 2022, the Board of Directors of the Company resolved to issue restricted stocks to employees amounting to 63 thousand shares with the effective date set on September 8, 2022. On November 8, 2022, the Board of Directors of the Company resolved to issue restricted stocks to employees amounting to 195 thousand shares with the effective date set on November 8, 2022. On March 10, 2023, the Board of Directors resolved to issue restricted stocks to employees amounting to 6 thousand shares with the effective date set on March 10, 2023. On November 9, 2023, the Board of Directors of the Company resolved to issue restricted stocks to employees amounting to 325 thousand shares with the effective date set on November 9, 2023.
- G. The shareholders during their meeting on May 31, 2023 resolved to issue the 1st and 2nd restricted stocks to employees amounting to 805 thousand and 870 thousand shares with no subscription price, respectively. On November 9, 2023, the Board of Directors of the Company resolved to issue the 2nd restricted stocks to employees amounting to 826 thousand shares in 2023, with the effective date set on November 9, 2023. On December 22, 2023, the Board of Directors of the Company resolved to. issue the 1st restricted stocks to employees amounting to 26 thousand shares in 2023, with the effective date set on December 22, 2023.
- H. The shareholders during their meeting on May 31, 2023 adopted a resolution to raise cash capital through private placement. The maximum number of shares to be issued through the

private placement is 30,000 thousand shares and the private placement may be made in three installments as authorised by the shareholders during their meeting. The private placement was in accordance with the Securities and Exchange Act and the Directions for Public Companies Conducting Private Placements of Securities. The Company's Board of Directors resolved not to execute the private placement on March 8, 2024.

(17) Capital surplus

Pursuant to the R.O.C. Company Act, capital surplus arising from paid-in capital in excess of par value on issuance of common stocks and donations can be used to cover accumulated deficit or to issue new stocks or cash to shareholders in proportion to their share ownership, provided that the Company has no accumulated deficit. Further, the R.O.C. Securities and Exchange Act requires that the amount of capital surplus to be capitalised mentioned above should not exceed 10% of the paid-in capital each year. However, capital surplus should not be used to cover accumulated deficit unless the legal reserve is insufficient.

(18) Accumulated deficit

- A. Under the Company's Articles of Incorporation, the current year's earnings, if any, shall first be used to pay all taxes and offset prior years' operating losses and then 10% of the remaining amount shall be set aside as legal reserve. After the provision or reversal of special reserve in accordance with laws or regulations, the appropriation of the remaining earnings along with the unappropriated earnings of prior years shall be proposed by the Board of Directors and resolved at shareholders' meetings.
- B. The Company's dividend policy is summarised below: The Board of Directors would consider the earnings situation of current year, capital and financial structure, future operating needs, retained earnings and legal reserve, as well as the market competition to propose the appropriation of earnings to the shareholders during their meetings for resolution, and cash dividends shall account for at least 10% of the total dividends distributed.
- C. On June 10, 2022, the shareholders at their meeting resolved the deficit compensation for the year ended December 31, 2021. The Company offset the accumulated deficit by capital surplus. Refer to the website of "Market Observation Post System" for information about earnings appropriation to offset deficit as proposed by the Board of Directors and resolved by the shareholders.
- D. On May 31, 2023, the shareholders at their meetings resolved the deficit compensation for the year ended December 31, 2022. The Company offset the accumulated deficit against the capital surplus. Refer to the website of "Market Observation Post System" for information about earnings appropriation to offset deficit as proposed by the Board of Directors and resolved by the shareholders.

- E. On March 8, 2024, the Board of Directors proposed the deficit compensation for the year ended December 31, 2023. The Company offset the accumulated deficit against the capital surplus. Refer to the website of "Market Observation Post System" for information about earnings appropriation to offset deficit as proposed by the Board of Directors and resolved by the shareholders.
- F. As of December 31, 2023 and 2022, there was no earnings to be distributed.

(19) Operating revenue

	Year ended December 31			
		2023	2022	
Revenue from contracts with				
customers	\$	1,022,653	\$	1,481,017

A. Disaggregation of revenue

The Group derives revenue from the transfer of services, authorisation and goods over time and at a point in time in the following major categories:

	Year ended December 31, 2023			
		Sales of authorisation and cooperative		
	Sales of services	development	Sales of goods	Total
Timing of revenue recognition		-		
At a point in time	\$ -	\$ -	\$ 161,594	\$ 161,594
Over time	605,990	141,472	113,597	861,059
	\$ 605,990	<u>\$ 141,472</u>	\$ 275,191	\$ 1,022,653
	Year ended December 31, 2022			
		Sales of authorisation and cooperative		
	Sales of services	authorisation	Sales of goods	Total
Timing of revenue recognition	Sales of services	authorisation and cooperative	Sales of goods	Total
=	Sales of services \$ -	authorisation and cooperative	Sales of goods \$ 417,774	Total \$ 417,774
recognition		authorisation and cooperative development		

B. Contract assets and liabilities

(a) The Group has recognised the following revenue-related contract assets and liabilities:

	Decer	December 31, 2023		December 31, 2022		January 1, 2022	
Contract assets:							
Services	\$	240,564	\$	213,981	\$	144,831	
Sales		53,130		20,418		25,766	
	\$	293,694	\$	234,399	\$	170,597	
Current contract liabilities		_		_			
Services	\$	41,739	\$	104,384	\$	102,289	
Authorisation and cooperative		15,027		46,091		121,678	
Non-current contract liabili	ties						
Authorisation and cooperative						20,059	
	\$	56,766	\$	150,475	\$	244,026	

(b) Revenue recognised that was included in the contract liability balance at the beginning of the year

Revenue recognised that was included in the contract liability balance at the beginning of the year

Services

Authorisation and cooperative

 Year ended	Deceml	per 31
2023		2022
\$ 100,624	\$	92,362
 32,211		101,380
\$ 132,835	\$	193,742

(c). Unfulfilled long-term contracts

development

Aggregate amount of the transaction price allocated to long-term technology service contracts, authorisation and cooperative development contracts that are partially or fully unsatisfied, and all of the milestone payment as at December 31, 2023 amounted to \$1,341,954. The management expects to recognise the amount in the future.

C. Details on authorisation and cooperative development revenue arising from providing drug development, commercialization service and authorising intellectual property rights of pharmaceutical products to the pharmaceutical factory are as follows:

In April 2019, the Group entered into an authorisation and cooperative development contract of EG12014 with Sandoz AG. The contract includes up-front payment, milestone payment at each stage and profit-sharing royalty on sales of products in the authorised markets in proportion to the ratios specified in the contract. The contract is mainly for providing the biosimilars development and commercialisation services and authorising intellectual

property rights to the customer in the authorised regions. As of December 31, 2023, the Group has received the aforementioned up-front payment and part of the milestone payment in accordance with the contract terms. The revenue of up-front payment and milestone payment achieved is recognised based on the satisfaction percentage during research and development period. If the drug was successfully launched, the supply price based on the supply terms and quantities, and the profit-sharing royalty calculated based on sales could also be collected. For the years ended December 31, 2023 and 2022, the Group recognised the revenue from authorisation and cooperative development contract amounting to \$141,472 and \$261,876, respectively.

The European Medicines Agency and the US Food and Drug Administration accepted the Sandoz AG's application for marketing review in January 2022 and February 2022, respectively. Sandoz AG received a complete response letter from the US Food and Drug Administration in December 2022. Within the complete response letter (CRL):

- (a) There were no clinical or safety or biosimilarity deficiencies cited in the CRL.
- (b) The CRL cites certain drug product deficiencies related to the manufacturing facility identified by the agency during a pre-license inspection of the site.

In January 2023, the Company received an EIR (Establishment Inspection Report) from the US Food and Drug Administration, which indicated that the Company's Zhubei plant had passed the US FDA's pre-marketing drug inspection. Sandoz is in close contact with the FDA to meet the satisfactory resolution of the FDA observations in a timely manner and plans a BLA resubmission in due course.

- D. In April 2023, the Company received a letter from the Taiwan Food and Drug Administration (TFDA) to which indicated that the Company had obtained the domestic active pharmaceutical ingredients "EG12014 Trastuzumab" license and a drug master file number. In September 2023, the Company received the approval by the National Health Insurance Administration with respect to its enrollment in the reimbursement system which became effective from October 1, 2023.
- E. On November 16, 2023, Sandoz AG received the marketing authorisation from Committee for Medicinal Products for Human Use (CHMP) for the trastuzumab biosimilar, EG12014, which was licensed by the Company for sale.

(20) Interest income

Interest income from bank deposits
Interest income from financial
assets measured at amortised cost

Y ear ended	Decemb	per 31	
 2023		2022	
\$ 106,236	\$		57,862
 28,235			1,722
\$ 134,471	\$		59,584

(21) Other income

	 Year ended l	December 31		
	 2023		2022	
Government grant revenues	\$ 4,712	\$	37,214	
Dividend income	475		-	
Other income	 252		430	
	\$ 5,439	\$	37,644	

The Company received a grant for the 'Breast Cancer Targeted Antibody similar to EG12014 Trastuzumab Biosimilar phase III clinical trial program' from Ministry of Economic Affairs (MOEA). The program execution period is from November 1, 2019 to June 30, 2023 and the limit on total grant amounted to \$80,000. For the years ended December 31, 2023 and 2022, the Company recognised government grants revenue of \$4,591 and \$36,994, respectively.

(22) Other gains and losses

	Year ended December 31				
		2023	2022		
Foreign exchange (losses) gains	(\$	9,431) \$	126,788		
(Losses) gains on financial assets at					
fair value through profit or loss	(1,122)	2,863		
Loss on lease modification	(413) (709)		
Miscellaneous disbursements	(214) (24)		
Loss on redemption of convertible					
bonds		<u> </u>	3)		
	(\$	11,180) \$	128,915		

(23) Finance costs

		per 31		
		2023		2022
Interest expense on lease liabilities Interest expense on bank	\$	8,096	\$	8,208
borrowings		2,300		1,205
Other interest expense		7		226
	\$	10,403	\$	9,639

(24) Employee benefits, depreciation and amortisation expenses

Function	Year ended December 31, 2023			Year ended December 31, 2022				
	Classified as	Classified as		Classified as				
	Operating	Operating		Operating	Operating			
Nature	Costs	Expenses	Total	Costs	Expenses	Total		
Employee								
benefit								
Wages and	\$ 121,946	\$ 271,494	\$ 393,440	\$ 147,873	\$ 220,847	\$ 368,720		
salaries								
Share based	70,628	110,272	180,900	60,275	95,037	155,312		
payment								
Labour and	14,408	23,704	38,112	13,771	19,354	33,125		
health								
insurance								
Pension	9,257	11,658	20,915	7,427	10,863	18,290		
costs								
Directors'	-	4,125	4,125	-	3,948	3,948		
remuneration								
Other	6,358	15,092	21,450	5,557	13,245	18,802		
personnel								
expenses								
Depreciation	114,746	112,799	227,545	99,536	89,564	189,100		
expense								
Amortisation	5,840	5,456	11,296	10,456	5,728	16,184		
expense								

- A. In accordance with to the Articles of Incorporation of the Company, a ratio of distributable profit of the current year, after covering accumulated losses, shall be distributed as employees' compensation and directors' and supervisors' remuneration. The ratio shall be 1% to 5% for employees' compensation and shall not be higher than 3% for directors' remuneration.
- B. No employees' compensation and directors' remuneration was accrued due to the net loss incurred for the years ended December 31, 2023 and 2022.
- C. Information about employees' compensation and directors' and supervisors' remuneration of the Company as resolved at the meeting of Board of Directors and resolved at the shareholders' meeting will be posted in the "Market Observation Post System" at the website of the Taiwan Stock Exchange.

(25) Income taxes

A. Income tax expense

(a) Components of income tax expense:

	Year ended December 31				
		2023		2022	
Current tax: Current tax on profits for the year	\$	1,093	\$	1,023	
Prior year income tax over estimation			(136)	
Total current tax		1,093		887	
Deferred tax: Origination and reversal of temporary differences		465		338	
Income tax expense	\$	1,558	\$	1,225	

(b) The income tax (charge)/credit relating to components of other comprehensive income is as follows:

	Year ended December 31				
		2023		2022	
Currency translation differences	\$	41	\$		

B. Reconciliation between income tax expense and accounting profit

	Year ended December 31						
		2023		2022			
Tax calculated based on loss before							
tax and statutory tax rate	(\$	181,856)	(\$	22,863)			
Expenses disallowed by tax regulation		25		49			
Tax exempt income by tax regulation		-	(3,099)			
Taxable losses not recognised as deferred							
tax assets		172,451		27,274			
Prior year income tax under							
(over)estimation		-	(136)			
Temporary differences not							
recognised as deferred tax assets		10,938		<u>-</u>			
Income tax expenses	\$	1,558	\$	1,225			

C. Amounts of deferred tax assets or liabilities as a result of temporary differences:

						2023			
		Ian	uary 1		ognised in	comp	ised in other rehensive	Dec	ember 31
-Deferred tax assets: Share of profit (loss) of associates and subsidiaries accounted		Jan	uary 1	_ pror	it or ioss		come	<u>Dec</u>	ember 31
for using the equity method, net differen	ces	\$	874	\$	465	\$	-	\$	1,339
Currency translation differences			_		_		41	\$	41
		\$	874	\$	465	\$	41	\$	1,380
						2022			
				Reco	ognised in	•	ised in other rehensive		
-Deferred tax assets: Share of profit (loss associates and subsidiaries account for using the equity method, net differen	ed	\$	536	\$	338	\$	_	\$	874
D. Details of the amount	t the (Compa	ny is en	titled a	s investme	ent tax cre	edit and unre	ecogn	ised
deferred tax assets are	e as fo	llows:							
			Decem	ber 31,	2023				
					Unrecogn	nised			
Qualifying items	•	nused t	ax credit		eferred tax		Expiry		
Research and development	\$		960,90	00 \$	Ģ	960,900	Note	e	
Machinery and equipment			8,84	44		8,844	Note	е	
			Decem	ber 31,	2022				
					Unrecogn				
Qualifying items		nused t	ax credit		eferred tax		Expiry		
Research and development	\$		887,10	50 \$	8	387,160	Note	e	

Note: The Company was entitled to the incentives conferred under the Biotech and New Pharmaceutical Development Act following the Company's incorporation as a biotech pharmaceutical company pursuant to the Letter No. Jing-Shou-Gong-Zi-10920401340 issued by the MOEA on February 3, 2020. Subsequently, the MOEA approved the Company's additional items pursuant to the Letter No. Jing-Shou-Gong-Zi-11120426560 on August 29, 2022. The incentive measures are valid for five years beginning on the next date of the issuance of MOEA's Letter. The investment tax credit can be first used to offset expenditure on research and development and staff training when there is taxable business income. Any unused tax credit can be first used to offset expenditure on machinery, equipment, or systems when there is taxable business income. Any unused tax credit is available for the following two years. As of December 31, 2023, the Company has no profit-seeking enterprise income tax.

E. Expiration dates of unused tax losses and amounts of unrecognised deferred tax assets are as follows:

December 31, 2023

Amount filed/ deferred tax								
Year incurred	assessed	Un	used amount		assets	Expiry year		
2014	Amount assessed	\$	131,762	\$	131,762	2024		
2015	Amount assessed		133,257		133,257	2025		
2016	Amount assessed		109,737		109,737	2026		
2017	Amount assessed		163,949		163,949	2027		
2018	Amount assessed		371,827		371,827	2028		
2019	Amount assessed		858,819		858,819	2029		
2020	Amount assessed		1,009,168		1,009,168	2030		
2021	Amount assessed		56,144		56,144	2031		
2022	Amount filed		135,927		135,927	2032		
2023	Amount expected		862,256		862,256	2033		
		\$	3,832,846	\$	3,832,846			

December 31, 2022

	Amount filed/			d	eferred tax	
Year incurred	assessed	Un	used amount		assets	Expiry year
2013	Amount assessed	\$	104,540	\$	104,540	2023
2014	Amount assessed		131,762		131,762	2024
2015	Amount assessed		133,257		133,257	2025
2016	Amount assessed		109,737		109,737	2026
2017	Amount assessed		163,949		163,949	2027
2018	Amount assessed		371,827		371,827	2028
2019	Amount assessed		858,819		858,819	2029
2020	Amount assessed		1,009,168		1,009,168	2030
2021	Amount filed		56,144		56,144	2031
2022	Amount expected		132,140		132,140	2032
		\$	3,071,343	\$	3,071,343	

F. The amounts of deductible temporary differences that ware not recognised as deferred tax assets are as follows:

	I	December 31, 2023	 December 31, 2022
Deductible temporary differences	\$	64,721	\$ 10,031

G. The Company's income tax returns through 2021 have been assessed and approved by the Tax Authority.

(26) Loss per share

	Year ended December 31, 2023				
			Weighted average		
			number of ordinary		
			shares outstanding		Loss per share
	Amoun	t after tax	(share in thousands)		(in dollars)
Basic loss per share					
Loss for the year	(<u>\$</u>	915,208)	304,888	<u>(\$</u>	3.00)
		Year	ended December 31,	20	22
			Weighted average		
			number of ordinary		
			shares outstanding		Loss per share
	Amoun	t after tax	(share in thousands)		(in dollars)
Basic loss per share					
Loss for the year	(\$	115,540)	303,258	(<u>\$</u>	0.38)

Diluted loss per share would not be calculated as the Company had loss for the years ended December 31, 2023 and 2022.

(27) Supplemental cash flow information

A. Investing activities with partial cash payments:

	Year ended December 31				
	2023		2022		
Purchase of property, plant and equipment	\$	703,002	\$	484,512	
Add: Opening balance of other payables		158,228		19,508	
Less: Ending balance of other payables	(285,960)	(158,228)	
Cash paid during the year	\$	575,270	\$	345,792	
	Year ended December 31 2023 202			per 31 2022	
Purchase of intangible assets Add: Ending balance of	\$	11,498	\$	8,562	
prepayment for intangible assets (Note) Less: Opening balance of		5,209		1,565	
prepayment for intangible assets (Note)	(1,565)	(1,565)	
Cash paid during the period	\$	15,142	\$	8,562	

Note: Shown as "other non-current assets".

B. Financing activities with no cash flow effects

	Year ended December 31				
		2023		2022	
Conversion of convertible bonds	\$		- \$	126,375	

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		ong-term rrowings	Leas	e liabilitie		arantee		abilities from financing tivities-gross
At January 1 Changes in cash flow from financing	\$	120,460	\$	338,58	4 \$	294	\$	459,338
activities Impact of changes in		-	(29,30	7) (288)	(29,595)
foreign exchange rate Changes in other		-		35,01	7	-		35,017
non-cash items				41	3			413
At December 31	\$	120,460	\$	344,70	<u>7</u> \$	6	\$	465,173
	2022							
		g-term	ase liab	I (ii	Bonds payable ncluding current portion)	Guarant deposit receive	S	Liabilities from financing activities-
At January 1	\$	owings Lea - \$				\$	u	gross \$ 434,612
At January 1 Changes in cash flow from financing activities Changes in right-of-		20,460 (,542 \$,435) (127,070 200)		- 294	\$ 434,612 96,119
use assets		-	54	,767	-		-	54,767
Impact of changes in foreign exchange rate Changes in other		-		1	-		-	1
non-cash items		<u> </u>		709 (126,870)			(126,161)
At December 31	\$ 1	20,460 \$	338	,584 \$		\$	294	\$ 459,338

7. Related Party Transactions

(1) Parent and ultimate controlling party

The Group has no ultimate parent company and ultimate controlling party.

(2) Names of related parties and relationship

Names of related parties	Relationship with the Group
FORMOSA LABORATORIES, INC.	Other related party
Development Center for Biotechnology (DCB)(Note 1)	<i>"</i>
FORMOSA PHARMACEUTICALS, INC.	<i>"</i>
TFBS Bioscience Inc. (Note 2)	<i>"</i>
Forward BioT Venture Capital	<i>"</i>

- Note 1: DCB's term as a director expired after re-election of directors at the Company's shareholders' meeting on June 10, 2022. Accordingly, it became a non-related party. (The transaction amounts for the period from January 1, 2022 to June 10, 2022 are disclosed in the financial statements.)
- Note 2: The Company was elected as one of the directors of TFBS Bioscience, Inc. on June 8, 2022. Accordingly, the Company became a related party. (The transaction amount for the year ended December 31, 2023 and period from June 8, 2022 to December 31, 2022 are disclosed in the financial statements.)

(3) Significant related party transactions

A. Operating revenue

	Year ended December 31				
		2023		2022	
Sales of goods:					
Other related parties	\$	972	\$	12,850	
Sales of services:					
Other related parties		9,477		5,622	
	\$	10,449	\$	18,472	

- (a) No similar transaction can be compared with for the sales of service. Prices and terms are determined based on mutual agreements.
- (b) On December 31, 2023 and 2022, the Group has recognised the revenue-related contract assets amounting to \$1,994 and \$744, and contract liabilities amounting to \$372 and \$620, respectively.
- B. Service expense (shown as 'research and development expense')

	Year ended December 31				
		2023		2022	
Other related parties	\$	12,377	\$	17,651	

It refers to service expense of contracted Biopharmaceutical research and development with other related parties. Prices and terms are determined based on mutual agreements.

C. Testing expense (shown as 'operating costs')

	Year ended December 31				
		2023		2022	
Other related parties-TFBS					
Bioscience Inc.	\$	7,517	\$	15,152	
Other related parties		2,627		2,460	
	\$	10,144	\$	17,612	

D. Other expenses (shown as 'administrative expenses')

	Year ended December 31					
	2023		2022			
Other related parties-DCB	\$	- \$	2,463			

It refers to repair and maintenance fees, based on the price specified in the contract as mutually agreed, allocated from leasing plant and lab from DCB, and the expense shall be paid before the 25th day of the first month of each quarter as specified in the contract.

E. Receivables from related parties

	Decemb	per 31, 2023	Decemb	per 31, 2022
Other receivables:				
Other related parties	\$	2,636	\$	_

F. Payables to related parties

	Dec	cember 31, 2022	D	ecember 31, 2022
Other payables:				
Other related parties	\$	7,993	\$	7,732

G. Property transactions

(a) Acquisition of property, plant and equipment:

	 Year ended December 31					
	 2023	20)22			
Other related parties	\$ 645	\$	_			

(b) Acquisition of financial assets:

		Year ended December 31				
		2	023		2022	
	Accounts	Consi	deration	Con	sideration	
Other related party-FORMOSA PHARMACEUTIC	Non-current financial assets at fair value through profit or loss					
ALS, INC.	loss	\$		\$	58,390	
Other related party-TFBS	Non-current financial assets at fair value through other					
Bioscience	comprehensive income	\$	-	\$	40,627	
Other related party-Forward BioT Venture Capital	Non-current prepayments for investments	\$	15,000	\$		

Refer to Note 6(2) B. and Note 6(7) B. for details of the transactions relating to the Company's acquisition of assets from related parties.

H. Lease transactions - lessee

(a) The Group leases plant, laboratory, instrument and equipment from DCB. Rental contract period is expected to be 20 years with initial rental period of 5 years plus the extension options. Rents are paid before the 25th day of the first month of each quarter.

(b) Right-of-use assets

As of December 31, 2023 and 2022, DCB was no longer a related party, and therefore the carrying amount of its related right-of-use assets was not disclosed.

	Year ended December 31		
	2022		
	Deprecia	ation expense	
Land	\$	3,061	
Buildings		2,279	
Machinery and equipment		1,022	
	\$	6,362	

(c) Lease liabilities

i. Outstanding balance

As of December 31, 2023 and 2022, DCB was no longer a related party, and therefore the carrying amount of its related lease liabilities was not disclosed.

ii. Interest expense

	Year ended	December 31
	2	022
Other related party -		
DCB	\$	2,185

(d) Rent expense (shown as 'operating cost' and 'operating expenses')

	Year ende	d December 31
		2022
Other related party -		
DCB	\$	505

(4) Key management compensation

	Year ended December 31					
		2023		2022		
Salaries and other short-term employee benefits	\$	31,942	\$	30,390		
Post-employment benefits		491		556		
Share based payment		30,145		31,043		
	\$	62,578	\$	61,989		

8. Pledged Assets

The Group's assets pledged as collateral are as follows:

	Book value		E	Book value		
Pledged asset	Decemb	per 31, 2023	Dece	mber 31, 2022	Purpose	
Pledged time deposits	\$	8,790	\$	8,671	Note 1	
(shown as non-current						
financial assets at amortised cost)						
Guarantee deposits paid	\$	8,795	\$	65,048	Note 2	
(shown as other non-current assets, others)						
Property, plant and equipment	\$	1,551,633	\$	1,158,399	Note 3	
Pledged government bonds						
(shown as non-current						
financial assets at amortised cost)	\$	31,390	\$		Note 4	

- Note 1: It refers to guarantee for lease of land.
- Note 2: It refers to deposits for research commissioned contract, equipment and office, guarantee for gas meter as well as certificates of deposit for customs post-release duty payment.
- Note 3: In April 2022, the Company terminated the syndicated loan agreement with 6 financial institutions including Taiwan Business Bank. However, the guarantee for the pledged buildings has not yet been released.
- Note 4: It refers to guarantee for investment.

9. Significant Contingent Liabilities and Unrecognised Contract Commitments

(1) Contingencies

None.

(2) Commitments

- A. As of December 31, 2023 and 2022, the remaining payments contracted for research commissioned contracts at the balance sheet date but not yet incurred amounted to \$59,156 and \$105,637, respectively.
- B. As of December 31, 2023 and 2022, the remaining payments contracted for equipment purchase and plant design at the balance sheet date but not yet incurred amounted to \$876,590 and \$815,285, respectively.
- C. The Group entered into a long-term consignment contract with a supplier to ensure the future supply of goods and pay the guarantee amounting to \$30,000. As of December 31, 2023, the aforementioned amount was shown as other non-current assets, others of \$30,000.

10. Significant Disaster Loss

None.

11. Significant Events after the Balance Sheet Date

- A. In January 2024, in response to the expansion of Zhubei production line, the Company entered into a construction contract with an engineering company for a total amount of \$1,373,643 (including tax).
- B. The Board of Directors on March 8, 2024 resolved to issue the 1st restricted stocks to employees amounting to 1,400 thousand shares with no subscription price, which has not yet been resolved by the shareholders as of March 8, 2024.
- C. The Board of Directors on March 8, 2024 resolved to raise additional cash through private placement. The maximum number of shares to be issued through the private placement is 30,000 thousand, and the private placement can be completed in three instalments after the authorisation by shareholders. However, the issuance has not yet been resolved at the shareholders' meeting as of March 8, 2024.
- D. As the Company's self-developed product, EG1206A, will begin phase III clinical trial according to the research and development schedule. On March 8, 2024, the Board of

Directors resolved to authorise the chairman to enter into a commissioned research project for the phase III clinical trial with a CRO and other companies.

12. Others

(1) Capital management

The Group's objectives when managing capital are to safeguard the Group's ability to continue as a going concern in order to provide returns for shareholders and to maintain an optimal capital structure to reduce the cost of capital. In order to maintain or adjust the capital structure, the Group may adjust the amount of dividends paid to shareholders, return capital to shareholders, issue new shares or sell assets to reduce debt.

(2) Financial instruments

A. Financial instruments by category

		December 31, 2023		December 31, 2022
Financial assets				
Financial assets at fair value				
through profit or loss				
Financial assets mandatorily measured				
at fair value through profit or loss	\$	80,298	\$	61,420
Financial assets at fair value				
through other comprehensive				
income				
Designation of equity	\$	325,887	\$	270 225
instrument	Ф	323,887	Ф	279,325
Financial assets at amortised cost				
Cash and cash equivalents	\$	5,053,183	\$	6,126,885
Financial assets at amortised cost		540,720		1,041,123
Notes receivable		19		-
Accounts receivable		253,390		32,782
Accounts receivable - related parties		2,636		-
Other receivables		20,497		24,944
Guarantee deposits paid (shown				
as other non-current assets)		8,795		65,048
	\$	5,879,240	\$	7,290,782

	Decen	mber 31, 2023	Dece	mber 31, 2022
Financial liabilities				
Financial liabilities at amortised				
cost				
Accounts payable	\$	79,556	\$	134,607
Other payables		530,299		407,387
Other payables-related parties		7,993		7,732
Long-term borrowings		120,460		120,460
Guarantee deposits received				
(shown as other non-current				
liabilities)		6		294
	\$	738,314	\$	670,480
Lease liability (current and non-	\$	344,707	\$	338,584
current)				

B. Financial risk management policies

- (a) The Group's activities expose it to a variety of financial risks: market risk (including foreign exchange risk, interest rate risk and price risk), credit risk and liquidity risk. The Group's overall risk management programme focuses on the unpredictability of financial markets and seeks to minimise potential adverse effects on the Group's financial position and financial performance.
- (b) Risk management is carried out by a central treasury department (Group treasury) under policies approved by the Board of Directors. Group treasury identifies, evaluates and hedges financial risks in close co-operation with the Group's operating units. The Board provides written principles for overall risk management, as well as written policies covering specific areas and matters, such as foreign exchange risk, interest rate risk, credit risk, use of derivative financial instruments and non-derivative financial instruments, and investment of excess liquidity.

C. Significant financial risks and degrees of financial risks

(a) Market risk

i. Exchange rate risk

- (i) The Group operates internationally and is exposed to exchange risk arising from various currency exposures, primarily with respect to the USD, EUR, GBP and JPY. Foreign exchange rate risk arises from future commercial transactions and recognised assets and liabilities.
- (ii) Management has set up a policy to require group companies to manage their foreign exchange risk against their functional currency. The companies are required to hedge their entire foreign exchange risk exposure with the Group treasury.

(iii) The Group's businesses involve some non-functional currency operations (the Company's functional currency: NTD; subsidiaries' functional currency: EUR). The information on assets and liabilities denominated in foreign currencies whose values would be materially affected by the exchange rate fluctuations is as follows:

]	December 31, 2023		
	_	n currency ount (In			
	tho	usands)	Exchange rate	Book	value (NTD)
Financial assets					
Monetary items					
USD:NTD	\$	53,756	30.71	\$	1,650,847
EUR:NTD		363	33.98		12,335
GBP:NTD		67	39.15		2,623
JPY:NTD		10,751	0.22		2,365
Financial liabilities					
Monetary items					
USD:NTD	\$	932	30.71	\$	28,622
EUR:NTD		546	33.98		18,553
GBP:NTD		17	39.15		666
JPY:NTD		57,505	0.22		12,651
]	December 31, 2022	,	
	Foreig	n currency			
	_	ount (In			
		usands)	Exchange rate	Rook	value (NTD)
Financial assets	1110	usanus)	Exchange rate	DOOK	value (IVID)
Monetary items					
USD:NTD	\$	44,053	30.71	\$	1,352,868
EUR:NTD	φ	191	32.72	Ф	6,250
GBP:NTD		110	37.09		4,080
JPY:NTD		8,476	0.23		1,949
Financial liabilities		0,470	0.23		1,949
·					
Monetary items USD:NTD	\$	708	30.71	\$	21,743
EUR:NTD	ψ	1,048	32.72	Ψ	34,291
GBP:NTD		30	37.09		1,113
ODF.NID		30	37.09		1,113

⁽iv) Analysis of foreign currency market risk arising from significant foreign exchange variation:

Year ended December 31, 2023

	Sensitivity analysis					
				Effec	ct on other	
	Degree of	Effec	et on profit or	comp	orehensive	
	variation		loss	iı	ncome	
Financial assets						
Monetary items						
USD:NTD	1%	\$	16,508	\$	-	
EUR:NTD	1%		60		63	
GBP:NTD	1%		26		-	
JPY:NTD	1%		24		-	
Financial liabilities						
Monetary items						
USD:NTD	1%	\$	286	\$	-	
EUR:NTD	1%		186		-	
GBP:NTD	1%		7		-	
JPY:NTD	1%		127		-	

Y ear	ende	d L	ece	ember	31,	2022
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		Sensitivity analysis										
				Effect on other								
	Degree of	Effec	et on profit or	comprehensive								
	variation		loss	income								
Financial assets												
Monetary items												
USD:NTD	1%	\$	13,529	\$	-							
EUR:NTD	1%		8		55							
GBP:NTD	1%		41		-							
JPY:NTD	1%		19		-							
Financial liabilities												
Monetary items												
USD:NTD	1%	\$	217	\$	-							
EUR:NTD	1%		343		-							
GBP:NTD	1%		11		-							

(v) The total exchange (losses) gains, including realised and unrealised, arising from significant foreign exchange variation on the monetary items held by the Group for the years ended December 31, 2023 and 2022, amounted to (\$9,431) and \$126,788, respectively.

ii. Price risk

- (i.) The Group's equity securities, which are exposed to price risk, are the held financial assets at fair value through other comprehensive income. To manage its price risk arising from investments in equity securities, the Group diversifies its portfolio. Diversification of the portfolio is done in accordance with the limits set by the Group.
- (ii.) The Group's investments in equity securities comprise. The prices of equity securities would change due to the change of the future value of investee companies. If the prices of these equity securities had increased/decreased by 1% with all other variables held constant, post-tax profit for the years ended December 31, 2023 and 2022 would have increased/decreased by \$189 and \$0, respectively, as a result of gains/losses on equity securities classified as at fair value through profit or loss. Other comprehensive income for the years ended December 31, 2023 and 2022 would have increased/decreased by \$3,259 and \$2,793, respectively, as a result of other comprehensive income classified as equity investment at fair value through other comprehensive income.

iii. Cash flow and fair value interest rate risk

The Group's main interest rate risk arises from long-term borrowings with variable rates. Borrowings issued at variable rates expose the Group to cash flow interest rate risk. During 2023 and 2022, the Group's borrowings at variable rate were mainly denominated in New Taiwan dollars.

(b) Credit risk

- i. Credit risk refers to the risk of financial loss to the Group arising from default by the clients or counterparties of financial instruments on the contract obligations. The main factor is that counterparties could not repay in full the accounts receivable based on the agreed terms.
- ii. The Group manages their credit risk taking into consideration the entire group's concern. According to the Group's credit policy, each local entity in the Group is responsible for managing and analysing the credit risk for each of their new clients before standard payment and delivery terms and conditions are offered. Internal risk control assesses the credit quality of the customers, taking into account their financial position, past experience and other factors. Individual risk limits are set based on internal or external ratings in accordance with limits set by the Board of Directors. The utilisation of credit limits is regularly monitored.
- iii. The Group adopts the assumptions under IFRS 9, that is, the default occurs when the contract payments are past due over 90 days.

- iv. The Group adopts the following assumption under IFRS 9 to assess whether there has been a significant increase in credit risk on that instrument since initial recognition: If the contract payments were past due over 30 days based on the terms, there has been a significant increase in credit risk on that instrument since initial recognition.
- v. The following indicators are used to determine whether the credit impairment of debt instruments has occurred:
 - (i) It becomes probable that the issuer will enter bankruptcy or other financial reorganisation due to their financial difficulties;
 - (ii) The disappearance of an active market for that financial asset because of financial difficulties;
 - (iii) Default or delinquency in interest or principal repayments;
 - (iv) Adverse changes in national or regional economic conditions that are expected to cause a default.
- vi. The Group classifies customers' accounts receivable, and contract assets in accordance with customer types. The Group applies the modified approach using individual provision to estimate expected credit loss.
- vii. The Group's notes and accounts receivable were generated from the customers who have optimal credit rating, and the expected credit loss rate is 0.03% after using the forecastability of future boom. As of December 31, 2023 and 2022, the total carrying amount of notes and accounts receivable (including related parties) amounted to \$256,342 and \$33,079, respectively. Although some accounts receivable were past due over 90 days, the expected credit risk is insignificant based on individual assessment, thus, loss allowance was recognised amounting to \$297 and \$297, respectively. The counterparties of time deposits over 3 months are financial institutions all with high credit quality and the expected credit risk is insignificant based on the assessment, thus, no loss allowance was recognised.

viii. Movements in loss allowance for accounts receivable are as follows:

	Year ended December 31							
		2023	2022					
At January 1	\$	297 \$	689					
Reversal of impairment loss		- (392)					
At December 31	\$	297 \$	297					

(c) Liquidity risk

- i. Cash flow forecasting is performed in the operating entities of the Group and aggregated by Group treasury. Group treasury monitors rolling forecasts of the Company's liquidity requirements to ensure it has sufficient cash to meet operational needs.
- ii. Surplus cash held by the operating entities over and above balance required for working capital management are transferred to the Group treasury. Group treasury invests surplus cash in interest bearing current accounts and time deposits, choosing instruments with appropriate maturities or sufficient liquidity to provide sufficient head-room as determined by the above-mentioned forecasts.
- iii. The Group has the following undrawn borrowing facilities:

	D	December 31, 2023	 December 31, 2022			
Floating rate:						
Expiring within one year	\$	1,410,000	\$ 1,020,000			
Expiring beyond one year		593,540	 593,540			
	\$	2,003,540	\$ 1,613,540			

iv. The table below analyses the Group's non-derivative financial liabilities into relevant maturity groupings based on the remaining period at the balance sheet date to the contractual maturity date. The amounts disclosed in the table are the contractual undiscounted cash flows.

	L	Less than		etween 1			
December 31, 2023	1 year		an	d 5 years	Ov	er 5 years	 Total
Non-derivative financial							
<u>liabilities</u>							
Accounts payable	\$	79,556	\$	-	\$	-	\$ 79,556
Other payables		530,299		-		-	530,299
Other payables- related parties		7,993		-		-	7,993
Lease liabilities		36,273		118,543		245,982	400,798
Long-term borrowings		2,376		123,322		-	125,698
Guarantee deposit received (show as other non-current liabilities)		6		-		-	6

	Less than Between 1						
December 31, 2022		1 year		nd 5 years	Over 5 years		 Total
Non-derivative financial							
<u>liabilities</u>							
Accounts payable	\$	134,607	\$	-	\$	-	\$ 134,607
Other payables		407,387		-		-	407,387
Other payables- related parties		7,732		-		-	7,732
Lease liabilities		34,828		115,926		247,968	398,722
Long-term borrowings		2,216		125,265		-	127,481
Guarantee deposit received (show as other non-current liabilities)		294		-		-	294

v. The Group does not expect the timing of occurrence of the cash flows estimated through the maturity date analysis will be significantly earlier, nor expect the actual cash flow amount will be significantly different.

(3) Fair value information

- A. The different levels that the inputs to valuation techniques are used to measure fair value of financial and non-financial instruments have been defined as follows:
 - Level 1: Quoted prices (unadjusted) in active markets for identical assets or liabilities that the entity can access at the measurement date. An active market refers to a market in which transactions for an asset or liability take place with sufficient frequency and volume to provide pricing information on an ongoing basis.
 - Level 2: Inputs other than quoted prices included within Level 1 that are observable for the asset or liability, either directly or indirectly.
 - Level 3: Unobservable inputs for the asset or liability. The fair value of the Group's investment in equity investment without active market and the call options and put options embedded in convertible bonds issued by the Group are included in Level 3.

B. Financial instruments not measured at fair value

Except for financial assets at fair value through profit or loss and financial assets measured at fair value through other comprehensive income, the carrying amounts of cash and cash equivalents, financial assets at amortised cost, notes receivable, accounts receivable (including related parties), other receivables, guarantee deposits paid (shown as other non-current assets), accounts payable, other payables (including related parties), long-term borrowings, guarantee deposits received (shown as other non-current liabilities) and lease liabilities are approximate to their fair values.

- C. The related information on financial and non-financial instruments measured at fair value by level on the basis of the nature, characteristics and risks of the assets and liabilities are as follows:
 - (a) The related information on the nature of the assets and liabilities is as follows:

December 31, 2023	Level 1	_	Level 2		Level 3	Total	
Assets							
Recurring fair value							
measurements							
Financial assets at fair							
value through profit or							
loss							
Profit-sharing investments	\$	-	\$ -	\$	61,410	\$	61,410
in new drug							
development							
Limited partnership		-	-		18,888		18,888
venture capital							
Financial assets at fair value							
through other							
comprehensive income							
Equity securities		_			325,887		325,887
	\$	_	\$ -	\$	406,185	\$	406,185
December 31, 2022	Level 1		Level 2	_	Level 3		Total
Assets							
Recurring fair value							
<u>measurements</u>							
Financial assets at fair							
value through profit or							
loss							
Profit-sharing investments	\$	-	\$ -	\$	61,420	\$	61,420
in new drug							
development							
Financial assets at fair value							
through other							
comprehensive income							
Equity securities		_			279,325		279,325
	\$	_	<u>\$</u>	\$	340,745	\$	340,745

(b). The fair value of financial instruments measured by using valuation techniques can be referred to current fair value of instruments with similar terms and characteristics in substance, discounted cash flow method or other valuation methods.

D. The following chart is the movement of Level 3 for the years ended December 31, 2023 and 2022:

	2023									
				Profit-						
				sharing						
				estment in		Limited				
	Е	quity	n	ew drug	partnership					
	instr			development		venture capital		Total		
At January 1	\$ 2	79,325	\$	61,420	\$	_	\$	340,745		
Additions		623		_		20,000		20,623		
Gains or losses recognised in profit or loss shown as other gains and losses								·		
Gains (losses) on valuation		-	(10)	(1,112)	(1,122)		
Gains and losses recognised in other comprehensive income										
Gains (losses) on valuation		45,939						45,939		
At December 31	\$ 3	25,887	\$	61,410	\$	18,888	\$	406,185		
						2022				
						Profit-				
			Equity		sharing investment in new					
	Der	rivative								
	instr	uments	ins	struments	drug	development		Total		
At January 1	\$	891	\$	11,607	\$	-	\$	12,498		
Additions		-		208,627		58,390		267,017		
Conversions of convertible bonds	(723)		-		-	(723)		
Gains or losses recognised in profit or loss shown as other gains and losses										
Gains (losses) on valuation	(167)		-		3,030		2,863		
Gains and losses recognised in other comprehensive income										
Gains (losses) on valuation		-		59,091		-		59,091		
Settled during the year	(1)					(1)		
At December 31	\$	-	\$	279,325	\$	61,420	\$	340,745		

- E. For the years ended December 31, 2023 and 2022, there was no transfer into or out from Level 3.
- F. Appointed external appraiser is in charge of valuation procedures for fair value measurements being categorised within Level 3, and frequently calibrating valuation model, performing back-testing, updating inputs used to the valuation model and making any other necessary adjustments to the fair value.

G. The following is the qualitative information of significant unobservable inputs and sensitivity analysis of changes in significant unobservable inputs to valuation model used in Level 3 fair value measurement:

Fair value at Signific		Significant	Range			
	Decen	nber 31,	Valuation	unobservable	(weighted	Relationship of inputs
	20)23	technique	input	average)	to fair value
Non-derivative equity instrument:						
Unlisted shares	\$	8,236	Price-Book Ratio	Price-to -book ratio Discount for lack of marketability	2.66~3.75 (3.67) 30% (30%)	The higher the multiple, the higher the fair value; The higher the discount for lack of marketability, the lower the fair value
Unlisted shares	3	17,651	Price-Book Ratio	Price-to-book ratio	1.24~2.54 (1.97)	The higher the multiple, the higher the fair value; The higher the discount for lack of marketability,
				Discount for lack of marketability	7.25% (7.25%)	the lower the fair value
Profit-sharing investments in new drug development		61,410	Royalty relief method of income approach	Discount rate	24.69%	The higher the discount rate, the lower the fair value
•				Market share	2.0%~5.9%	The higher the market share, the higher the fair value
Limited partnership venture capital		18,888	Net asset value	N/A	N/A	N/A

	Fair value at				Range				
	December	31, Val	luation	unobservable	(weighted	Relationship of inputs			
	2022	tecl	nnique	input	average)	to fair value			
Non-derivative equity instrument:									
Unlisted shares	\$ 6,2		e-Book atio	Price-to -book ratio Discount for lack of marketability	1.54~8.46 (3.05) 30% (30%)	The higher the multiple, the higher the fair value; The higher the discount for lack of marketability, the lower the fair value			
Unlisted shares	273,1		e-Book atio	Price-to-book ratio	2.01~2.54 (2.19)	The higher the multiple, the higher the fair value; The higher the discount for lack of marketability,			
				Discount for lack of marketability	30% (30%)	the lower the fair value			
Profit-sharing investments in new drug development	61,4	met inc	Ity relief hod of come broach	Discount rate	24.58%	The higher the discount rate, the lower the fair value			
-		••		Market share	1.0%~5.4%	The higher the market share, the higher the fair value			

H. The Group has carefully assessed the valuation models and assumptions used to measure fair value. However, use of different valuation models or assumptions may result in different measurement. The following is the effect of profit or loss or of other comprehensive income from financial assets and liabilities categorised within Level 3 if the inputs used to valuation models have changed:

			December 31, 2023								
			Rec	cognised i	n pr	ofit or loss		Recognis			
			Fa	vourable	Uı	nfavourable	F	Favourable	Ur	nfavourable	
	Input	Change		change		change		change	change		
Financial assets											
Profit-sharing	Discount	±5%	\$	3,071	(\$	3,071)	\$	-	\$	-	
investments	Rate										
in new drug development	Market Share										
Limited	NA	±5%		944	(944)		_		_	
partnership venture capital	1421	±3 70		711	•	744)					
Unlisted shares	Price-Book Ratio	±5%		-		-		16,294	(16,294)	
	Lack of	±5%									
	marketability	_2 70						16,294	(16,294)	
	J		\$	4,015	(\$	4,015)	\$	32,588	(\$	32,588)	
						December	31	, 2022			
								Recognis	ed i	n other	
			Rec	cognised i	n pr	ofit or loss		comprehen	sive	e income	
			Fa	vourable	Uı	nfavourable	F	Favourable	Uı	nfavourable	
	Input	Change		hange		change		change		change	
Financial assets											
Profit-sharing	Discount	±5%	\$	3,071	(\$	3,071)	\$	-	\$	-	
investments	Rate										
in new drug	Market										
development	Share	. 50/						12.066	,	12.0(()	
Unlisted shares	Price-Book Ratio	±5%		-		-		13,966	(13,966)	
	Lack of	±5%									
	marketability	_5/0				_		13,966	(_	13,966)	
	,		\$	3,071	(\$	3,071)	\$	27,932	(\$	27,932)	

13. <u>Supplementary Disclosures</u>

(1) Significant transactions information

- A. Loans to others: None.
- B. Provision of endorsements and guarantees to others: None.
- C. Holding of marketable securities at the end of the period (not including subsidiaries, associates and joint ventures): Please refer to table 1.

- D. Acquisition or sale of the same security with the accumulated cost exceeding \$300 million or 20% of the Company's paid-in capital: None.
- E. Acquisition of real estate reaching NT\$300 million or 20% of paid-in capital or more: Please refer to table 2.
- F. Disposal of real estate reaching NT\$300 million or 20% of paid-in capital or more: None.
- G. Purchases or sales of goods from or to related parties reaching NT\$100 million or 20% of paid-in capital or more: None.
- H. Receivables from related parties reaching NT\$100 million or 20% of paid-in capital or more: None.
- I. Trading in derivative instruments undertaken during the reporting periods: None.
- J. Significant inter-company transactions during the reporting periods: Please refer to table 3.

(2) Information on investees

Names, locations and other information of investee companies (not including investees in Mainland China): Please refer to table 4.

(3) Information on investments in Mainland China

None.

(4) Major shareholders information

Major shareholders information: Please refer to table 5.

14. Segment Information

(1) General information

The Group is primarily engaged in the biosimilar and new drug research and development as well as biopharmaceutical contract development and manufacturing services, including cell line construction platforms, process development platforms, analytical science and protein characterisation, as well as PIC/S GMP facilities to provide clinical trial drug and listed drug production, etc. The Group operates business only in a single industry. The Chief Operating Decision-maker who allocates resources and assesses performance of the Group as a whole, has identified that the Group has only one reportable operating segment.

(2) Segment Information

The accounting policies of the operating segments are in agreement with the significant accounting policies summarised in Note 4. The Group's segment profit (loss) is measured with the loss before tax, which is used as a basis for the Group in assessing the performance of the operating segments.

(3) Information about segment profit or loss, assets and liabilities

The Group has only one reportable operating segment, thus, the reportable information is in agreement with those in the consolidated financial statements.

(4) Reconciliation for segment income (loss)

The amounts provided to the Chief Operating Decision-maker with respect to segment assets, liabilities and loss before tax from continuing operations are measured in a manner consistent with that in the financial statements. Thus, no reconciliation is needed.

(5) Information on products and services

The Group's revenue is mainly from biopharmaceutical contract development and manufacturing services, authorisation and cooperative development and sales. Details of revenue are as follows:

	Year ended December 31							
Service revenue		2023	2022					
	\$	605,990	\$	757,680				
Sales revenue Authorisation and cooperative		141,472		261,876				
development revenue		275,191		461,461				
	\$	1,022,653	\$	1,481,017				

(6) Geographical information

Geographical information for the years ended December 31, 2023 and 2022 is as follows:

	Year ended December 31											
		20			20)22						
			Ion-current			N	Von-current					
	Revenue			assets		Revenue		assets				
Taiwan	\$	474,123	\$	3,799,337	\$	746,845	\$	3,176,241				
Japan		186,584		-		133,023		-				
American & Canada		119,293		-		304,969		-				
Europe		242,186		811		283,327		1,169				
Others		467				12,853	_	_				
	\$	1,022,653	\$	3,800,148	\$	1,481,017	\$	3,177,410				

(7) Major customer information

Major customers which contributed more than 10% of the Group's total operating revenues for the years ended December 31, 2023 and 2022 are listed below:

		Year ended December 31								
		202	23	2022						
	_ F	Revenue	Segment	<u>F</u>	Revenue	Segment				
A	\$	174,644	Note	\$	123,081	Note				
В		144,479	"		261,876	"				
C		125,354	"		514,208	"				
D		113,042	"		136,272	"				
E		109,280	"		-	"				

Note: The Group has only one reportable operating segment.

Holding of marketable securities at the end of the period (not including subsidiaries, associates and joint ventures)

Non-current financial assets a

fair value through profit or loss

Non-current financial assets at

amortised cost

None

December 31, 2023

Table 1

Forward BioT Venture Capital

VI government bonds

93 Central Government Bonds A

equity

Expressed in thousands of NTD (Except as otherwise indicated)

18,888

31,930

As of December 31, 2023

18,888

31,930

5.69%

Securities held by	Marketable securities	Relationship with the securities issuer	General ledger account	Number of shares	Book value	Ownership	Fair value	Footnote
EirGenix Inc.	Oncomatryx Biopharma S.L. common stock	None	Non-current financial assets at fair value through other comprehensive	31,801	8,236	0.37% \$	8,236	
"	TFBS Bioscience, Inc. common stock	The Company's other related party	"	4,942,455	317,651	14.20%	317,651	

Acquisition of real estate reaching NT\$300 million or 20% of paid-in capital or more

Year ended December 31, 2023

Table 2

Expressed in thousands of NTD (Except as otherwise indicated)

If the counterparty is a related party, information as to the

last transaction of the real estate is disclosed below:

							Original owner					Reason for	
Real estate	Real		Transaction	Status of		Relationship with the	who sold the real estate to the		Date of the original		Basis or reference used in setting	acquisition of real estate and	Other
acquired by	estate	Date of the event	 Amount	payment	Counterparty	counterparty	counterparty	Relationships	Transfer Date	Amount	the price	status of the real estate	commitments
EirGenix Inc.	Factory building	2021/9/30 (Note 4)	\$	Based on the terms in the purchase order	Lee Ming Construction Co., Ltd. China Ecotek Corporation	None	NA	NA	NA	NA	Price comparison and price negotiation	Manufacturing purpose	None
					Getinge Group Taiwan Co., Ltd. Jian-Yi Biotech Co., Ltd.								
					Min-Pin,Chen Architects & Associates								

Note 1: The appraisal result should be presented in the 'Basis or reference used in setting the price' column if the real estate acquired should be appraised pursuant to the regulations.

Note 2: Paid-in capital referred to herein is the paid-in capital of parent company. In the case that shares were issued with no par value or a par value other than NT\$10 per share, the 20 % of paid-in capital shall be replaced by 10% of equity attributable to owners of the parent in the calculation.

Note 3: Date of the event referred to herein is the date of contract signing date, date of payment, date of execution of a trading order, date of title transfer, date of board resolution, or other date that can confirm the counterparty and the monetary amount of the transaction, whichever is earlier.

Note 4: This is the signing date of the first transaction, which is mainly arising from the construction of the factory. The Company continuously signed contracts with relevant suppliers, of which the contract amount has reached \$300,000.

Significant inter-company transactions during the reporting period

Year ended December 31, 2023

Table 3

Expressed in thousands of NTD (Except as otherwise indicated)

Transaction

				Transaction					
								Percentage of consolidated total	
Number (Note 1)	Company name	Counterparty	Relationship	General ledger account		Amount	Transaction terms	operating revenues or total assets (Note 3)	
0	EirGenix Inc.	EirGenix Europe GmbH	(1)	Operating expense	\$	72,270	Note 4	7.07%	
0	EirGenix Inc.	EirGenix Europe GmbH	(1)	Other payables		12,758	"	0.11%	

Note 1: The numbers filled in for the transaction company in respect of inter-company transactions are as follows:

- (1) Parent company is '0'.
- (2) The subsidiaries are numbered in order starting from '1'.
- Note 2: Relationship between transaction company and counterparty is classified into the following three categories; fill in the number of category each case belongs to (If transactions between parent company and subsidiaries or refer to the same transaction, it is not required to disclose twice. For example, if the parent company has already disclosed its transaction with a subsidiary, then the subsidiary is not required to disclose the transaction; for transactions between two subsidiaries, if one of the subsidiaries has disclosed the transaction, then the other is not required to disclose the transaction.):
 - (1) Parent company to subsidiary.
 - (2) Subsidiary to parent company.
 - (3) Subsidiary to subsidiary.
- Note 3: Regarding percentage of transaction amount to consolidated total operating revenues or total assets, it is computed based on period-end balance of transaction to consolidated total assets for balance sheet accounts and based on accumulated transaction amount for the period to consolidated total operating revenues for income statement accounts.
- Note 4: Prices and terms for services are based on the mutual agreement and payments are collected quarterly in advance.
- Note 5: Transactions between the parent company and subsidiaries are eliminated.
- Note 6: Individual amounts less than \$1,000 are not disclosed.

Information on investees

Year ended December 31, 2023

Table 4

Expressed in thousands of NTD (Except as otherwise indicated)

				Initial i	nvestn	men	nt amount	Shares held as at December 31, 2023		Net profit (loss) of		Investment income (loss)			
Investor		Location	Main business activities	Balance as December 2023			Balance as at December 31, 2022	Number of shares	Ownership (%)	Во	ok value		investee for the year ended ember 31, 2023	recognised by the Company for the year ended December 31, 2023	Footnote
EirGenix Inc.	EirGenix Europe GmbH	Germany	Biopharmaceutical research and development as well as business development	\$	845	\$	845	-	100.00	\$	7,743	\$	2,324	\$ 2,324	None
EirGenix Inc.	EirGenix USA Inc.	USA	Biopharmaceutical commissioned development, manufacturing services and consulting	\$	-	\$	-	-	100.00	\$	-	\$	-	\$ -	None

Major shareholders information

December 31, 2023

Table 5

	Shares						
Name of major shareholders	Number of shares held	Ownership (%)					
Foxconn Technology Co., Ltd.	27,500,000	8.98					
Yonglin Capital Holding Co., Ltd.	26,500,000	8.65					
Formosa Laboratories, Inc.	18,582,818	6.06					

EIRGENIX INC.

PARENT COMPANY ONLY FINANCIAL

STATEMENTS AND INDEPENDENT AUDITORS'

REPORT

DECEMBER 31, 2023 AND 2022

For the convenience of readers and for information purpose only, the auditors' report and the accompanying financial statements have been translated into English from the original Chinese version prepared and used in the Republic of China. In the event of any discrepancy between the English version and the original Chinese version or any differences in the interpretation of the two versions, the Chinese-language auditors' report and financial statements shall prevail.

INDEPENDENT AUDITORS' REPORT TRANSLATED FROM CHINESE

To the Board of Directors and Shareholders of EirGenix Inc.

Opinion

We have audited the accompanying parent company only balance sheets of EirGenix Inc. (the "Company") as at December 31, 2023 and 2022, and the related parent company only statements of comprehensive income, of changes in equity and of cash flows for the years then ended, and notes to the parent company only financial statements, including a summary of material accounting policies.

In our opinion, the accompanying parent company only financial statements present fairly, in all material respects, the financial position of the Company as at December 31, 2023 and 2022, and its financial performance and its cash flows for the years then ended in accordance with the Regulations Governing the Preparation of Financial Reports by Securities Issuers.

Basis for opinion

We conducted our audits in accordance with the Regulations Governing Financial Statement Audit and Attestation Engagements of Certified Public Accountants and Standards on Auditing of the Republic of China. Our responsibilities under those standards are further described in the Auditors' Responsibilities for the Audit of the Parent Company Only Financial Statements section of our report. We are independent of the Company in accordance with the Norm of Professional Ethics for Certified Public Accountants of the Republic of China, and we have fulfilled our other ethical responsibilities in accordance with these requirements. We believe that the audit evidence we have obtained is sufficient and appropriate to provide a basis for our opinion.

Key audit matters

Key audit matters are those matters that, in our professional judgment, were of most significance in our audit of the Company's 2023 parent company only financial statements. These matters were addressed in the context of our audit of the parent company only financial statements as a whole and, in forming our opinion thereon, we do not provide a separate opinion on these matters.

The key audit matters for the Company's 2023 parent company only financial statements are stated as follows:

Accuracy of service revenue and authorisation and cooperative development revenue

Description

Refer to Note 4(25) for accounting policy on service revenue and authorisation and cooperative development revenue recognition, Note 5(2) for significant accounting estimates and assumptions, and Note 6(20) for details of operating revenue. The amount of service revenue and authorisation and cooperative development revenue for the year ended December 31, 2023 were NTD 605,990 thousand and NTD 141,472 thousand, respectively.

The Company's service revenue and authorisation and cooperative development revenue primarily arise from offering biopharmaceutical contract development and manufacturing services and authorising intellectual property rights of medicine development to pharmaceutical factory. Revenue is recognised based on the stage of completion at the balance sheet date provided that such transaction amounts can be reliably estimated. Since the information process, recording and maintenance are partially performed manually and the recognition of service revenue and authorisation and cooperative development revenue contains a high degree of uncertainty resulting in a complex calculation process, and revenue recognition is significant to the financial statements, we considered the accuracy of service revenue recognition a key audit matter.

How our audit addressed the matter

We performed the following audit procedures on the above key audit matter:

- 1. Obtained management's accounting policies on the service revenue and authorisation and cooperative development revenue recognition and confirmed that they are reasonable.
- 2. Selected samples and examined the contract in order to confirm whether the judgement made by the management was in line with the contract and generally accepted accounting principles..
- 3. For the performance obligation which was satisfied over time, selected samples and examined each data of contract costs and assessed whether the method and parameters used to measure the completion of performance obligation are reasonable.
- 4. Recalculated the accuracy of amount recognised as revenue and respective timing of recognition.

Impairment assessment of property, plant and equipment

Description

Refer to Note 4(17) for accounting policy on impairment of non-financial assets, Note 5(2) for uncertainty of accounting estimates and assumptions in relation to property, plant and equipment and Note 6(9) for description of property, plant and equipment.

On December 31, 2023, property, plant and equipment amounted to NTD 3,337,069 thousand, which were constructed to extend the production capacity of GMP. The Company assesses at each balance sheet date the fair value or recoverable value of those assets whether there is any indication that they may be impaired based on internal and external information. Since the impairment indication assessment and information and assumptions used to assess recoverable amount of assets have significant impact to property, plant and equipment, we considered impairment assessment of property, plant and equipment a key audit matter.

How our audit addressed the matter

We performed the following audit procedures on the above key audit matter:

- 1. Reviewed and assessed the reasonableness of each data in the impairment assessment.
- 2. Assessed the estimation procedure of future cash flows, and checked whether the cash flows listed in the assessment is consistent with operating plans.
- 3. Interviewed management to discuss the Company's operations and reviewed the actual performance of prior years' operating plans in order to understand the Company's intention and ability and ascertained whether there was any significant postponement on research and development.
- 4. Assessed the reasonableness of the significant assumptions adopted in estimating cash flows.

Responsibilities of management and those charged with governance for the parent company only financial statements

Management is responsible for the preparation and fair presentation of the parent company only financial statements in accordance with the Regulations Governing the Preparation of Financial Reports by Securities Issuers, and for such internal control as management determines is necessary to enable the preparation of parent company only financial statements that are free from material misstatement, whether due to fraud or error.

In preparing the parent company only financial statements, management is responsible for assessing the Company's ability to continue as a going concern, disclosing, as applicable, matters related to going concern and using the going concern basis of accounting unless management either intends to liquidate the Company or to cease operations, or has no realistic alternative but to do so.

Those charged with governance, including the audit committee, are responsible for overseeing the Company's financial reporting process.

Auditors' responsibilities for the audit of the parent company only financial statements

Our objectives are to obtain reasonable assurance about whether the parent company only financial statements as a whole are free from material misstatement, whether due to fraud or error, and to issue an auditors' report that includes our opinion. Reasonable assurance is a high level of assurance, but is not a guarantee that an audit conducted in accordance with the Standards on Auditing of the Republic of China will always detect a material misstatement when it exists. Misstatements can arise from fraud or error and are considered material if, individually or in the aggregate, they could reasonably be expected to influence the economic decisions of users taken on the basis of these parent company only financial statements.

As part of an audit in accordance with the Standards on Auditing of the Republic of China, we exercise professional judgment and professional skepticism throughout the audit. We also:

- 1. Identify and assess the risks of material misstatement of the parent company only financial statements, whether due to fraud or error, design and perform audit procedures responsive to those risks, and obtain audit evidence that is sufficient and appropriate to provide a basis for our opinion. The risk of not detecting a material misstatement resulting from fraud is higher than for one resulting from error, as fraud may involve collusion, forgery, intentional omissions, misrepresentations, or the override of internal control.
- 2. Obtain an understanding of internal control relevant to the audit in order to design audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the Company's internal control.
- 3. Evaluate the appropriateness of accounting policies used and the reasonableness of accounting estimates and related disclosures made by management.

- 4. Conclude on the appropriateness of management's use of the going concern basis of accounting and, based on the audit evidence obtained, whether a material uncertainty exists related to events or conditions that may cast significant doubt on the Company's ability to continue as a going concern. If we conclude that a material uncertainty exists, we are required to draw attention in our auditors' report to the related disclosures in the parent company only financial statements or, if such disclosures are inadequate, to modify our opinion. Our conclusions are based on the audit evidence obtained up to the date of our auditors' report. However, future events or conditions may cause the Company to cease to continue as a going concern.
- 5. Evaluate the overall presentation, structure and content of the parent company only financial statements, including the disclosures, and whether the parent company only financial statements represent the underlying transactions and events in a manner that achieves fair presentation.
- 6. Obtain sufficient appropriate audit evidence regarding the financial information of the entities or business activities within the Company and its subsidiaries to express an opinion on the parent company only financial statements. We are responsible for the direction, supervision and performance of the audit. We remain solely responsible for our audit opinion.

We communicate with those charged with governance regarding, among other matters, the planned scope and timing of the audit and significant audit findings, including any significant deficiencies in internal control that we identify during our audit.

We also provide those charged with governance with a statement that we have complied with relevant ethical requirements regarding independence, and to communicate with them all relationships and other matters that may reasonably be thought to bear on our independence, and where applicable, related safeguards.

From the matters communicated with those charged with governance, we determine those matters that were of most significance in the audit of the parent company only financial statements of the current period and are therefore the key audit matters. We describe these matters in our auditors' report unless law or regulation precludes public disclosure about the matter or when, in extremely rare circumstances, we determine that a matter should not be communicated in our report because the adverse consequences of doing so would reasonably be expected to outweigh the public interest benefits of such communication.

Teng, Sheng-Wei

Yen, Yu-Fang

For and on behalf of PricewaterhouseCoopers, Taiwan March 8, 2024

The accompanying parent company only financial statements are not intended to present the financial position and results of operations and cash flows in accordance with accounting principles generally accepted in countries and jurisdictions other than the Republic of China. The standards, procedures and practices in the Republic of China governing the audit of such financial statements may differ from those generally accepted in countries and jurisdictions other than the Republic of China. Accordingly, the accompanying parent company only financial statements and independent auditors' report are not intended for use by those who are not informed about the accounting principles or auditing standards generally accepted in the Republic of China, and their applications in practice.

As the financial statements are the responsibility of the management, PricewaterhouseCoopers cannot accept any liability for the use of, or reliance on, the English translation or for any errors or misunderstandings that may derive from the translation.

EIRGENIX INC. PARENT COMPANY ONLY BALANCE SHEETS DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars)

			December 31, 2023	December 31, 2022			
	Assets	Notes	 AMOUNT	%	 AMOUNT	%	
	Current assets						
1100	Cash and cash equivalents	6(1)	\$ 5,048,604	45	\$ 6,108,994	52	
1136	Current financial assets at amortised	6(3)					
	cost		500,000	5	1,000,000	9	
1140	Current contract assets	6(20) and 7	293,694	3	234,399	2	
1150	Notes receivable, net	6(4)	19	-	-	-	
1170	Accounts receivable, net	6(4)	253,390	2	32,782	-	
1180	Accounts receivable, net-related	7					
	parties		2,636	-	-	-	
1200	Other receivables		20,497	-	24,944	-	
1220	Current income tax assets		17,648	-	5,963	-	
130X	Inventories	6(5)	680,637	6	739,463	6	
1410	Prepayments	6(6)	 92,677	1	 122,502	1	
11XX	Total current assets		 6,909,802	62	 8,269,047	70	
	Non-current assets						
1510	Non-current financial assets at fair	6(2) and 7					
	value through profit or loss		80,298	1	61,420	1	
1517	Non-current financial assets at fair	6(7)					
	value through other comprehensive						
	income		325,887	3	279,325	2	
1535	Non-current financial assets at	6(3) and 8					
	amortised cost		40,720	-	41,123	-	
1550	Investments accounted for using	6(8)					
	equity method		7,743	-	5,200	-	
1600	Property, plant and equipment, net	6(9), 7 and 8	3,337,069	30	2,607,958	22	
1755	Right-of-use assets	6(10)	329,236	3	325,330	3	
1780	Intangible assets	6(11)	28,269	-	28,067	-	
1990	Other non-current assets	6(9)(12) and 8	 104,764	1	214,887	2	
15XX	Total non-current assets		 4,253,986	38	 3,563,310	30	
1XXX	Total assets		\$ 11,163,788	100	\$ 11,832,357	100	

(Continued)

EIRGENIX INC. PARENT COMPANY ONLY BALANCE SHEETS DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars)

	Liabilities and Equity	Notes		December 31, 2023 AMOUNT	<u>%</u>	December 31, 2022 AMOUNT		
	Current liabilities			INTO CITY		TIMOGRA	%	
2130	Current contract liabilities	6(20) and 7	\$	56,766	- \$	150,475	2	
2170	Accounts payable			79,556	1	134,607	1	
2200	Other payables	6(13)		519,762	5	384,682	3	
2220	Other payables - related parties	7		20,751	-	16,397	-	
2280	Current lease liabilities			28,622	-	26,826	-	
2399	Other current liabilities			2,937	<u>-</u> _	3,106		
21XX	Total current liabilities			708,394	6	716,093	6	
	Non-current liabilities							
2540	Long-term borrowings	6(14) and 8		120,460	1	120,460	1	
2570	Deferred tax liabilities	6(26)		1,380	-	874	-	
2580	Non-current lease liabilities			316,085	3	311,758	3	
2600	Other non-current liabilities			6	<u> </u>	294		
25XX	Total non-current liabilities			437,931	4	433,386	4	
2XXX	Total liabilities			1,146,325	10	1,149,479	10	
	Equity							
	Capital	6(17)						
3110	Common stock			3,060,516	28	3,043,358	26	
	Capital reserve	6(18)						
3200	Capital surplus			7,830,216	70	7,734,141	65	
	Accumulated deficit	6(19)						
3350	Accumulated deficit		(915,208) (8) (115,540) (1)	
	Other equity interest							
3400	Other equity interest			41,939		20,919		
3XXX	Total equity			10,017,463	90	10,682,878	90	
	Significant contingent liabilities and	9						
	unrecognised contract commitments							
	Significant events after the balance	11						
	sheet date							
3X2X	Total liabilities and equity		\$	11,163,788	100 \$	11,832,357	100	

The accompanying notes are an integral part of these parent company only financial statements.

EIRGENIX INC. PARENT COMPANY ONLY STATEMENTS OF COMPREHENSIVE INCOME YEARS ENDED DECEMBER 31, 2023 AND 2022

(Expressed in thousands of New Taiwan dollars, except for loss per share)

			Year ended December 31				
				2023		2022	
	Items	Notes		AMOUNT	%	AMOUNT	%
4000	Operating Revenue	6(20) and 7	\$	1,022,653	100 \$	1,481,017	100
5000	Operating Costs	6(5)(11)(25) and 7	(785,912) (<u>77</u>) (724,565) (49)
5900	Gross Profit			236,741	23	756,452	51
	Operating Expenses	6(11)(25) and 7					
6100	Sales and marketing expenses		(62,593) (6) (51,130) (4)
6200	General and administrative expenses		(254,196) (25) (236,675) (16)
6300	Research and development expenses		(955,346) (94) (802,439) (54)
6450	Reversal of credit impairment loss(expected credit impairment	12(2)					
	loss)			<u>-</u>	<u> </u>	392	
6000	Total operating expenses		(1,272,135) (125) (1,089,852) (74)
6900	Operating Loss		(1,035,394) (102) (333,400) (23)
	Non-operating Income and Expenses		`			, ·	
7100	Interest income	6(3)(21)		134,471	13	59,584	4
7010	Other income	6(22)		5,439	1	37,644	3
7020	Other gains and losses	6(2)(23)	(11,180) (1)	128,915	9
7050	Finance costs	6(10)(24) and 7	(10,403) (1) (9,635) (1)
7070	Share of profit of subsidiaries \	6(8)	(10,103)(1)(7,033)(1)
7070	associates and joint ventures	0(0)					
	accounted for using equity method			2,324	_	1,690	
7000	Total non-operating income and			2,324	 -	1,090	<u>-</u>
7000				120 651	10	210 100	1.5
7000	expenses		,—	120,651	12	218,198	15
7900	Loss before Income Tax	((2))	(914,743) (90) (115,202) (8)
7950	Income tax	6(26)	(465)	(338)	
8200	Net Loss		(\$	915,208) (90) (\$	115,540) (8)
8316	Other Comprehensive Income Components of other comprehensive income that will not be reclassified to profit or loss Unrealised gains (losses) from investments in equity instruments measured at fair value through other	6(7)					
	comprehensive income		\$	45,939	5 \$	59,091	1
8310	Other comprehensive income(loss) that will not be reclassified to		φ	45,939	<u> </u>	39,091	<u>4</u>
	profit or loss			45,939	5	59,091	4
	Components of other comprehensive income that will be reclassified to profit or loss						
8361	Exchange differences on translation of foreign financial statements			220		220	
8399	Income tax relating to components of other comprehensive income(loss) that will be reclassified to profit or	6(26)			-	220	-
	loss		(<u>41</u>)	<u> </u>	<u> </u>	<u>-</u>
8360	Other comprehensive income that will be reclassified to profit or loss			179	<u> </u>	220	
8300	Other Comprehensive Income		\$	46,118	5 \$	59,311	4
8500	Total Comprehensive Loss		(\$	869,090) (<u>85</u>) (<u>\$</u>	56,229) (4)
0750	Loss per share	6(27)	<i>(</i>		2.00\ / #		0.200
9750	Loss per share		(<u>\$</u>		3.00) (<u>\$</u>		0.38)

The accompanying notes are an integral part of these parent company only financial statements.

EIRGENIX INC. PARENT COMPANY ONLY STATEMENTS OF CHANGES IN EQUITY YEARS ENDED DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars)

			Capital Reserves					_					
	Notes	Common stock	Additional paid-in capital	Donated assets received	Employee stock options	Capital surplus, stock options	Restricted stock to employees	Capital surplus, others	Accumulated deficit	Exchange differences on translation of foreign financial statements	Unrealised gains (losses) from financial assets measured at fair value through other comprehensive income	Unearned compensation	Total equity
Year ended December 31, 2022													
Balance at January 1, 2022		\$ 3,003,845	\$ 10,313,563	\$ 2,036	\$ 41,958	\$ 3,467	\$ 114,928	\$ -	(\$ 2,973,500)	(\$ 237)	\$ 5,831	(\$ 83,140)	\$ 10,428,751
Loss for 2022		-						-	(115,540)				(115,540)
Other comprehensive income (loss)	6(7)	-	-	-	-	-	-	-	-	220	59,091	-	59,311
Total comprehensive income (loss)		-						-	(115,540)	220	59,091	-	(56,229)
Capital surplus used to offset accumulated deficit	6(19)		(2,971,464)	(2,036)					2,973,500				-
Compensation costs of share-based payments	6(16)	-	-	-	61,651	-	-	871	-	-	-	92,790	155,312
Employee stock options exercised	6(16)(17)	10,523	26,467	-	(8,320)	-	-	-	-	-	-	-	28,670
Issuance of employee restricted stocks	6(16)(17)	6,318	-	-	-	-	47,318	-	-	-	-	(53,636)	-
Redemption of employee restricted stocks	6(16)(17)	(2,260)	-	-	-	-	2,260	-	-	-	-	-	-
Restricted stocks vested		-	59,358	-	-	-	(59,358)	-	-	-	-	-	-
Conversion of convertible bonds	6(17)	24,932	104,904	-	-	(3,462)	-	-	-	-	-	-	126,374
Redemption of convertible bonds		<u>-</u>	<u>-</u>			(5)		5					<u>-</u>
Balance at December 31, 2022		\$ 3,043,358	\$ 7,532,828	\$ -	\$ 95,289	\$ -	\$ 105,148	\$ 876	(\$ 115,540)	(\$ 17)	\$ 64,922	(\$ 43,986)	\$ 10,682,878
Year ended December 31, 2023													
Balance at January 1, 2023		\$ 3,043,358	\$ 7,532,828	\$ -	\$ 95,289	\$ -	\$ 105,148	\$ 876	(\$ 115,540)	(\$ 17)	\$ 64,922	(\$ 43,986)	\$ 10,682,878
Loss for 2023		-	-	-	-	-	-	-	(915,208)	-	-	-	(915,208)
Other comprehensive income (loss)	6(7)									179	45,939		46,118
Total comprehensive income (loss)									(915,208)	179	45,939		(869,090)
Capital surplus used to offset accumulated deficit	6(19)	-	(114,664)	-	-		-	(876)	115,540	-	-	-	-
Compensation costs of share-based payments	6(16)	-	-	-	84,285	-	-	-	-	-	-	96,615	180,900
Employee stock options exercised	6(16)(17)	7,270	25,769	-	(10,264)	-	-	-	-	-	-	-	22,775
Employee stock options expired	6(16)	-	-	-	(1,810)	-	-	1,810	-	-	-	-	-
Issuance of employee restricted stocks	6(16)(17)	11,818	-	-	-	-	109,895	-	-	-	-	(121,713)	-
Redemption of employee restricted stocks	6(16)(17)	(1,930)	-	-	-	-	1,930	-	-	-	-	-	-
Restricted stocks vested			71,119				(71,119)						
Balance at December 31, 2023		\$ 3,060,516	\$ 7,515,052	\$ -	\$ 167,500	\$ -	\$ 145,854	\$ 1,810	(\$ 915,208)	\$ 162	\$ 110,861	(\$ 69,084)	\$ 10,017,463

EIRGENIX INC. PARENT COMPANY ONLY STATEMENTS OF CASH FLOWS YEARS ENDED DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars)

	Year ended December 31			per 31	
	Notes		2023		2022
CASH ELOWS EDOM ODED ATING A CTIVITIES					
CASH FLOWS FROM OPERATING ACTIVITIES Loss before tax		(\$	914,743)	(©	115,202)
Adjustments		(4)	914,743)	(4)	113,202)
Adjustments to reconcile profit (loss)					
Depreciation	6(9)(10)(25)		227,208		187,987
Amortization	6(11)(25)		11,296		16,184
Net loss (gain) on financial assets or liabilities at			11,290		10,104
fair value	0(2)(23)		1,122	(2,863)
Interest expense	6(24)		10,403	(9,635
Interest income	6(21)	(134,471)	(59,584)
Dividend income	6(22)	(475)	(57,504)
Compensation costs of share-based payments	6(16)(25)	(180,900		155,312
Share of profit of subsidiaries \(\) associates and	6(8)		100,700		155,512
joint ventures accounted for using equity	0(0)				
method		(2,324)	(1,690)
Loss on lease modification	6(10)(23)	(413	(709
Reversal of credit impairment loss (expected	12(2)		113		10)
credit impairment loss)	12(2)		_	(392)
Loss on redemption of convertible bonds	6(23)		_	(3
Changes in operating assets and liabilities	0(20)				J
Changes in operating assets Changes in operating assets					
Contract assets		(59,295)	(63,802)
Notes receivable, net		(19)	(1,139
Accounts receivable, net		(220,608)		46,084
Accounts receivable, net-related parties		(2,636)		546
Other receivables		`	6,736	(13,793)
Inventories			58,826	ì	324,025)
Prepayments			29,825	ì	16,719)
Other current assets					1,555
Changes in operating liabilities					-,
Contract liabilities		(93,709)	(93,551)
Accounts payable		Ì	55,051)	`	48,151
Other payables		`	7,338		19,212
Other payables - related parties			4,354		5,601
Other current liabilities		(169)	(1,816)
Cash outflow generated from operations		(945,079)	(201,319)
Interest received		`	132,183	`	55,232
Interest paid		(10,393)	(9,314)
Dividends received			475		-
Income tax received			1,128		-
Income tax paid		(12,813)	(4,835)
Net cash flows used in operating activities		(834,499)	(160,236)

(Continued)

EIRGENIX INC. PARENT COMPANY ONLY STATEMENTS OF CASH FLOWS YEARS ENDED DECEMBER 31, 2023 AND 2022 (Expressed in thousands of New Taiwan dollars)

	Year ended D			December 31			
	Notes		2023		2022		
CASH FLOWS FROM INVESTING ACTIVITIES							
Acquisition of financial assets at fair value through	6(2) and 7						
profit or loss		\$	-	(\$	58,390)		
Acquisition of financial assets at fair value through	6(7)						
other comprehensive income		(623)	(208,627)		
Acquisition of financial assets at amortised cost		(3,700,000)	(1,032,516)		
Proceeds from disposal of financial assets at							
amortised cost			4,200,403		1,636,640		
Acquisition of property, plant and equipment	6(28)	(575,239)	(345,548)		
Acquisition of intangible assets	6(28)	(15,142)	(8,652)		
Decrease in other financial assets			-		27,334		
Decrease (increase) in refundable deposits (shown							
as other non-current assets)			56,253	(778)		
Increase in prepayments for investments (shown as							
other non-current assets)		(46,270)	(20,000)		
Increase in prepayments for business facilities	6(9)						
(shown as other non-current assets)		(138,453)	(433,952)		
Increase in other non-current assets				(31,274)		
Net cash flows used in investing activities		(219,071)	(475,763)		
CASH FLOWS FROM FINANCING ACTIVITIES							
Repayments of bonds	6(29)		-	(200)		
Proceeds from long-term borrowings	6(29)		-		120,460		
Decrease (increase) in guarantee deposits received	6(29)						
(shown as other non-current liabilities)		(288)		294		
Repayments of lease principal	6(29)	(29,307)	(23,657)		
Employee stock options exercised			22,775		28,669		
Net cash flows (used in) from financing							
activities		(6,820)		125,566		
Net decrease in cash and cash equivalents		(1,060,390)	(510,433)		
Cash and cash equivalents at beginning of year			6,108,994		6,619,427		
Cash and cash equivalents at end of year		\$	5,048,604	\$	6,108,994		

EIRGENIX INC.

NOTES TO THE PARENT COMPANY ONLY FINANCIAL STATEMENTS YEARS ENDED DECEMBER 31, 2023 AND 2022

(Expressed in thousands of New Taiwan dollars, except as otherwise indicated)

1. History and Organization

- (1) EirGenix, Inc. (hereinafter referred to as the "Company") was incorporated as a company limited by shares under the provisions of the Company Act of the Republic of China (R.O.C.) in December 2012. In April 2013, the Company obtained all key technologies from the biopharmaceutical pilot plant originally owned by the Development Center for Biotechnology, including its complete core competencies. The Company is primarily engaged in the research and development of biosimilars and new drugs, as well as biopharmaceutical contract development and manufacturing services, which included cell line construction platforms, process development platforms, analytical science and protein identification. Furthermore, the Company has two PIC/S GMP facilities certified by the Taiwan Food and Drug Administration (TFDA), one for mammalian cells and one for microbial, to provide clinical trial drug and commercial drug production.
- (2) The shares of the Company have been listed on the Taipei Exchange since June 28, 2019.
- 2. The Date of Authorisation for Issuance of the Financial Statements and Procedures for Authorisation

These parent company only financial statements were authorised for issuance by the Board of Directors on March 8, 2024.

3. Application of New Standards, Amendments and Interpretations

(1) Effect of the adoption of new issuances of or amendments to International Financial Reporting

Standards ("IFRS®") Accounting Standards that came into effect as endorsed by the Financial

Supervisory Commission ("FSC")

New standards, interpretations and amendments endorsed by the FSC and became effective from 2023 are as follows:

	Effective date by
	International
	Accounting
New Standards, Interpretations and Amendments	Standards Board
Amendments to IAS 1, 'Disclosure of accounting policies'	January 1, 2023
Amendments to IAS 8, 'Definition of accounting estimates'	January 1, 2023
Amendments to IAS 12, 'Deferred tax related to assets and liabilities	January 1, 2023
arising from a single transaction'	
Amendments to IAS 12, 'International tax reform - pillar two model	May 23, 2023
rules'	

The above standards and interpretations have no significant impact to the Company's financial condition and financial performance based on the Company's assessment.

(2) Effect of new issuances of or amendments to IFRS Accounting Standards as endorsed by the FSC but not yet adopted by the Company

New standards, interpretations and amendments endorsed by the FSC and will become effective from 2024 are as follows:

	Effective date by International Accounting
New Standards, Interpretations and Amendments	Standards Board
Amendments to IFRS 16, 'Lease liability in a sale and leaseback' Amendments to IAS 1, 'Classification of liabilities as current or non-current'	January 1, 2024 January 1, 2024
Amendments to IAS 1, 'Non-current liabilities with covenants' Amendments to IAS 7 and IFRS 7, 'Supplier finance arrangements'	January 1, 2024 January 1, 2024

The above standards and interpretations have no significant impact to the Company's financial condition and financial performance based on the Company's assessment.

(3) IFRS Accounting Standards issued by IASB but not yet endorsed by the FSC

New standards, interpretations and amendments issued by IASB but not yet included in the IFRS Accounting Standards as endorsed by the FSC are as follows:

	Effective date by
	International Accounting
New Standards, Interpretations and Amendments	Standards Board
Amendments to IFRS 10 and IAS 28, 'Sale or contribution of assets	To be determined by
between an investor and its associate or joint venture'	International Accounting
	Standards Board
IFRS 17, 'Insurance contracts'	January 1, 2023
Amendments to IFRS 17, 'Insurance contracts'	January 1, 2023
Amendment to IFRS 17, 'Initial application of IFRS 17 and IFRS 9 –	January 1, 2023
comparative information'	
Amendments to IAS 21, 'Lack of exchangeability'	January 1, 2025

The above standards and interpretations have no significant impact to the Company's financial condition and financial performance based on the Company's assessment.

4. Summary of Material Accounting Policies

The principal accounting policies applied in the preparation of these parent company only financial statements are set out below. These policies have been consistently applied to all the periods presented, unless otherwise stated.

(1) Compliance statement

The parent company only financial statements of the Company have been prepared in accordance with the Regulations Governing the Preparation of Financial Reports by Securities Issuers.

(2) Basis of preparation

- A. Except for the following items, the parent company only financial statements have been prepared under the historical cost convention:
 - (a) Financial assets and financial liabilities (including derivative instruments) at fair value through profit or loss.
 - (b) Financial assets at fair value through other comprehensive income.
- B. The preparation of parent company only financial statements in conformity with the Regulations Governing the Preparation of Financial Reports by Securities Issuers requires the use of certain critical accounting estimates. It also requires management to exercise its judgement in the process of applying the Company's accounting policies. The areas involving a higher degree of judgement or complexity, or areas where assumptions and estimates are significant to the parent company only financial statements are disclosed in Note 5.

(3) Foreign currency translation

Items included in the parent company only financial statements of each of the Company's entities are measured using the currency of the primary economic environment in which the entity operates (the "functional currency"). The parent company only financial statements are presented in New Taiwan dollars, which is the Company's functional and currency.

A. Foreign currency transactions and balances

- (a) Foreign currency transactions are translated into the functional currency using the exchange rates prevailing at the dates of the transactions or valuation where items are remeasured. Foreign exchange gains and losses resulting from the settlement of such transactions are recognised in profit or loss in the period in which they arise.
- (b) Monetary assets and liabilities denominated in foreign currencies at the period end are retranslated at the exchange rates prevailing at the balance sheet date. Exchange differences arising upon re-translation at the balance sheet date are recognised in profit or loss.
- (c) Non-monetary assets and liabilities denominated in foreign currencies held at fair value through profit or loss are re-translated at the exchange rates prevailing at the balance sheet date; their translation differences are recognised in profit or loss. Non-monetary assets and liabilities denominated in foreign currencies held at fair value through other comprehensive income are re-translated at the exchange rates prevailing at the balance sheet date; their translation differences are recognised in other comprehensive income. However, non-monetary assets and liabilities denominated in foreign currencies that are not measured at fair value are translated using the historical exchange rates at the dates of the initial transactions.
- (d) All foreign exchange gains and losses are presented in the statement of comprehensive income within 'other gains and losses'.

B. Translation of foreign operations

The operating results and financial position of all the subsidiaries, associates and joint arrangements that have a functional currency different from the presentation currency are translated into the presentation currency as follows:

- (a) Assets and liabilities for each balance sheet presented are translated at the closing exchange rate at the date of that balance sheet;
- (b) Income and expenses for each statement of comprehensive income are translated at average exchange rates of that period; and

(c) All resulting exchange differences are recognised in other comprehensive income.

(4) Classification of current and non-current items

- A. Assets that meet one of the following criteria are classified as current assets; otherwise they are classified as non-current assets:
 - (a) Assets arising from operating activities that are expected to be realised, or are intended to be sold or consumed within the normal operating cycle;
 - (b) Assets held mainly for trading purposes;
 - (c) Assets that are expected to be realised within twelve months from the balance sheet date;
 - (d) Cash and cash equivalents, excluding restricted cash and cash equivalents and those that are to be exchanged or used to settle liabilities more than twelve months after the balance sheet date.
- B. Liabilities that meet one of the following criteria are classified as current liabilities; otherwise they are classified as non-current liabilities:
 - (a) Liabilities that are expected to be settled within the normal operating cycle;
 - (b) Liabilities arising mainly from trading activities;
 - (c) Liabilities that are to be settled within twelve months from the balance sheet date;
 - (d) Liabilities for which the repayment date cannot be extended unconditionally to more than twelve months after the balance sheet date. Terms of a liability that could, at the option of the counterparty, result in its settlement by the issue of equity instruments do not affect its classification.

(5) Cash equivalents

Cash equivalents refer to short-term, highly liquid investments that are readily convertible to known amounts of cash and which are subject to an insignificant risk of changes in value. Time deposits that meet the definition above and are held for the purpose of meeting short-term cash commitments in operations are classified as cash equivalents.

(6) Financial assets at fair value through profit or loss

A. Financial assets at fair value through profit or loss are financial assets that are not measured at amortised cost or fair value through other comprehensive income.

- B. On a regular way purchase or sale basis, financial assets at fair value through profit or loss are recognised and derecognised using trade date accounting.
- C. At initial recognition, the Company measures the financial assets at fair value and recognises the transaction costs in profit or loss. The Company subsequently measures the financial assets at fair value, and recognises the gain or loss in profit or loss.

(7) Financial assets at fair value through other comprehensive income

- A. Financial assets at fair value through other comprehensive income comprise equity securities which are not held for trading, and for which the Company has made an irrevocable election at initial recognition to recognise changes in fair value in other comprehensive income.
- B. On a regular way purchase or sale basis, financial assets at fair value through other comprehensive income are recognised and derecognised using trade date accounting.
- C. At initial recognition, the Company measures the financial assets at fair value plus transaction costs. The Company subsequently measures the financial assets at fair value:

The changes in fair value of equity investments that were recognised in other comprehensive income are reclassified to retained earnings and are not reclassified to profit or loss following the derecognition of the investment. Dividends are recognised as revenue when the right to receive payment is established, future economic benefits associated with the dividend will flow to the Company and the amount of the dividend can be measured reliably.

(8) Financial assets at amortised cost

- A. Financial assets at amortised cost are those that meet all of the following criteria:
 - (a) The objective of the Company's business model is achieved both by collecting contractual cash flows and selling financial assets; and
 - (b) The assets' contractual cash flows represent solely payments of principal and interest.
- B. The Company's time deposits which do not fall under cash equivalents are those with a short maturity period and are measured at initial investment amount as the effect of discounting is immaterial.

(9) Accounts and notes receivable

A. Accounts and notes receivable entitle the Company a legal right to receive consideration in exchange for transferred goods or rendered services.

B. The short-term accounts and notes receivable without bearing interest are subsequently measured at initial invoice amount as the effect of discounting is immaterial.

(10) Impairment of financial assets

For debt instruments measured at fair value through profit or loss and financial assets at amortised cost, at each reporting date, the Company recognises the impairment provision for 12 months expected credit losses if there has not been a significant increase in credit risk since initial recognition or recognises the impairment provision for the lifetime expected credit losses (ECLs) if such credit risk has increased since initial recognition after taking into consideration all reasonable and verifiable information that includes forecasts. On the other hand, for accounts receivable or contract assets that do not contain a significant financing component, the Company recognises the impairment provision for lifetime ECLs.

(11) Derecognition of financial assets

The Company derecognises a financial asset when the contractual rights to receive the cash flows from the financial asset expire.

(12) Inventories

Inventories are stated at the lower of cost and net realisable value. Cost is determined using the first-in, first-out (FIFO) method. The cost of finished goods and work in progress comprises raw materials, direct labour, other direct costs and related production overheads. It excludes borrowing costs. The item by item approach is used in applying the lower of cost and net realisable value. Net realisable value is the estimated selling price in the ordinary course of business, less the estimated costs of completion and the estimated costs necessary to make the sale.

(13) Investments accounted for using equity method - subsidiaries

- A. Subsidiaries are all entities (including structured entities) controlled by the Company. The Company controls an entity when the Company is exposed, or has rights, to variable returns from its involvement with the entity and has the ability to affect those returns through its power over the entity.
- B. Inter-company transactions, balances and unrealised gains or losses on transactions between companies are eliminated. Accounting policies of subsidiaries have been adjusted where necessary to ensure consistency with the policies adopted by the Company.
- C. The Company's share of its subsidiaries' post-acquisition profits or losses is recognised in profit or loss, and its share of post-acquisition movements in other comprehensive income is recognised in other comprehensive income. When the Company's share of losses in a subsidiary equals or

exceeds its interest in the subsidiary, the Company recognise loss continuously in proportion to its ownership.

D. In accordance with "Regulations Governing the Preparation of Financial Reports by Securities Issuers", the profit or loss and other comprehensive income or loss presented on the parent company only financial statements are consistent with those presented on the parent company only financial statements. In addition, owner's equity presented on the parent company only financial statements is consistent with equity attributable to owners of parent presented on the consolidated financial statements.

(14) Property, plant and equipment

- A. Property, plant and equipment are initially recorded at cost. Borrowing costs incurred during the construction period are capitalised.
- B. Subsequent costs are included in the asset's carrying amount or recognised as a separate asset, as appropriate, only when it is probable that future economic benefits associated with the item will flow to the Company and the cost of the item can be measured reliably. The carrying amount of the replaced part is derecognised. All other repairs and maintenance are charged to profit or loss during the financial period in which they are incurred.
- C. Land is not depreciated. Other property, plant and equipment apply cost model and are depreciated using the straight-line method to allocate their cost over their estimated useful lives. Each part of an item of property, plant, and equipment with a cost that is significant in relation to the total cost of the item must be depreciated separately.
- D. The assets' residual values, useful lives and depreciation methods are reviewed, and adjusted if appropriate, at each financial year-end. If expectations for the assets' residual values and useful lives differ from previous estimates or the patterns of consumption of the assets' future economic benefits embodied in the assets have changed significantly, any change is accounted for as a change in estimate under IAS 8, 'Accounting Policies, Changes in Accounting Estimates and Errors', from the date of the change. The estimated useful lives of property, plant and equipment are as follows:

Machinery and equipment	2 ~ 10 years
Office equipment	3 ~ 10 years
Buildings and structures	5 ~ 20 years
Leasehold improvements	3 ~ 20 years
Other equipment	$3 \sim 10 \text{ years}$

(15) Leasing arrangements (lessee) — right-of-use assets/ lease liabilities

- A. Leases are recognised as a right-of-use asset and a corresponding lease liability at the date at which the leased asset is available for use by the Company. For short-term leases or leases of low-value assets, lease payments are recognised as an expense on a straight-line basis over the lease term.
- B. Lease liabilities include the net present value of the remaining lease payments at the commencement date, discounted using the incremental borrowing interest rate. Lease payments are comprised of the following:
 - (a) Fixed payments, less any lease incentives receivable; and
 - (b) Variable lease payments that depend on an index or a rate.

The Company subsequently measures the lease liability at amortised cost using the interest method and recognises interest expense over the lease term. The lease liability is remeasured and the amount of remeasurement is recognised as an adjustment to the right-of-use asset when there are changes in the lease term or lease payments and such changes do not arise from contract modifications.

- C. At the commencement date, the right-of-use asset is stated at cost comprising the following:
 - (a) The amount of the initial measurement of lease liability; and
 - (b) Any lease payments made at or before the commencement date.

The right-of-use asset is measured subsequently using the cost model and is depreciated from the commencement date to the earlier of the end of the asset's useful life or the end of the lease term. When the lease liability is remeasured, the amount of remeasurement is recognised as an adjustment to the right-of-use asset.

D. For lease modifications that decrease the scope of the lease, the lessee shall decrease the carrying amount of the right-of-use asset to reflect the partial or full termination of the lease, and recognise the difference between remeasured lease liability in profit or loss.

(16) <u>Intangible assets</u>

The Company's accounting policies on intangible assets are summarised below:

A. Computer software

Computer software is stated at cost and amortised on a straight-line basis over its estimated useful life of 1 to 5 years.

B. Professional expertise

Professional expertise is stated at cost and amortised on a straight-line basis over its estimated useful life of 2 to 10 years.

(17) <u>Impairment of non-financial assets</u>

The Company assesses at each balance sheet date the recoverable amounts of those assets where there is an indication that they are impaired. An impairment loss is recognised for the amount by which the asset's carrying amount exceeds its recoverable amount. The recoverable amount is the higher of an asset's fair value less costs to sell or value in use. When the circumstances or reasons for recognising impairment loss for an asset in prior years no longer exist or diminish, the impairment loss is reversed. The increased carrying amount due to reversal should not be more than what the depreciated or amortised historical cost would have been if the impairment had not been recognised.

(18) Borrowings

- A. Borrowings are recognised initially at fair value, net of transaction costs incurred. Borrowings are subsequently stated at amortised cost; any difference between the proceeds (net of transaction costs) and the redemption value is amortised over the period of the borrowings using the effective interest method.
- B. Fees paid on the establishment of loan facilities are recognised as transaction costs of the loan to the extent that it is probable that some or all of the facility will be drawn down. In this case, the fee is deferred until the draw-down occurs. To the extent there is no evidence that it is probable that some or all of the facility will be drawn down, the fee is capitalised as a pre-payment for liquidity services and amortised over the period of the facility to which it relates.

(19) Accounts payable

- A. Accounts payable are liabilities for purchases of raw materials, goods or services.
- B. The short-term accounts payable without bearing interest are subsequently measured at initial invoice amount as the effect of discounting is immaterial.

(20) <u>Derecognition of financial liabilities</u>

A financial liability is derecognised when the obligation specified in the contract is either discharged or cancelled or expires.

(21) Employee benefits

A. Short-term employee benefits

Short-term employee benefits are measured at the undiscounted amount of the benefits expected to be paid in respect of service rendered by employees in a period and should be recognised as expense in that period when the employees render service.

B. Pensions

For defined contribution plans, the contributions are recognised as pension expense when they are due on an accrual basis. Prepaid contributions are recognised as an asset to the extent of a cash refund or a reduction in the future payments.

C. Employees' compensation and directors' and supervisors' remuneration

Employees' compensation and directors' and supervisors' remuneration are recognised as expense and liability, provided that such recognition is required under legal or constructive obligation and those amounts can be reliably estimated. Any difference between the amounts resolved by the shareholders and the actual amounts subsequently distributed is accounted for as changes in estimates.

(22) Employee share-based payment

A. For the equity-settled share-based payment arrangements, the employee services received are measured at the fair value of the equity instruments granted at the grant date, and are recognised as compensation cost over the vesting period, with a corresponding adjustment to equity. The fair value of the equity instruments granted shall reflect the impact of market vesting conditions and non-vesting conditions. Compensation cost is subject to adjustment based on the service conditions that are expected to be satisfied and the estimates of the number of equity instruments that are expected to vest under the non-market vesting conditions at each balance sheet date. Ultimately, the amount of compensation cost recognised is based on the number of equity instruments that eventually vest.

B. Restricted stocks:

- (a) Restricted stocks issued to employees are measured at the fair value of the equity instruments granted at the grant date, and are recognised as compensation cost over the vesting period.
- (b) The restricted stocks issued by the Company cannot be transferred during the vesting period, but voting right and dividend right are not restricted on these stocks. Employees are not required to return the dividends received if they resign during the vesting period.

(c) For restricted stocks where employees do not need to pay to acquire those stocks, if employees resign during the vesting period, they are considered not meeting the vesting condition from the date of resignation and the Company will redeem and retire those stocks at the initial issuance price.

(23) Income tax

- A. The tax expense for the period comprises current and deferred tax. Tax is recognised in profit or loss, except to the extent that it relates to items recognised in other comprehensive income or items recognised directly in equity, in which cases the tax is recognised in other comprehensive income or equity.
- B. The current income tax expense is calculated on the basis of the tax laws enacted or substantively enacted at the balance sheet date. Management periodically evaluates positions taken in tax returns with respect to situations in accordance with applicable tax regulations. It establishes provisions where appropriate based on the amounts expected to be paid to the tax authorities. An additional tax is levied on the unappropriated retained earnings and is recorded as income tax expense in the year the stockholders resolve to retain the earnings.
- C. Deferred tax is recognised, using the balance sheet liability method, on temporary differences arising between the tax bases of assets and liabilities and their carrying amounts in the parent company only balance sheet. However, the deferred tax is not accounted for if it arises from initial recognition of goodwill or of an asset or liability in a transaction other than a business combination that at the time of the transaction affects neither accounting nor taxable profit or loss and does not give rise to equal taxable and deductible temporary differences. Deferred tax is provided on temporary differences arising on investments in subsidiaries, except where the timing of the reversal of the temporary difference is controlled by the Company and it is probable that the temporary difference will not reverse in the foreseeable future. Deferred tax is determined using tax rates (and laws) that have been enacted or substantially enacted by the balance sheet date and are expected to apply when the related deferred tax asset is realised or the deferred tax liability is settled.
- D. Deferred tax assets are recognised only to the extent that it is probable that future taxable profit will be available against which the temporary differences can be utilised. At each balance sheet date, unrecognised and recognised deferred tax assets are reassessed.
- E. A deferred tax asset shall be recognised for the carryforward of unused tax credits resulting from research and development expenditures to the extent that it is possible that future taxable profit will be available against which the unused tax credits can be utilised.

(24) Share capital

Ordinary shares are classified as equity. Incremental costs directly attributable to the issue of new shares or stock options are shown in equity as a deduction, net of tax, from the proceeds.

(25) Revenue recognition

A. Service revenue

- (a) The Company provides biopharmaceutical contract testing and development services. Revenue from providing services is recognised in the accounting period in which the services are rendered. For fixed-price contracts, revenue is recognised based on the actual service provided to the end of the reporting period as a proportion of the total services to be provided. This is determined based on the actual cost relative to the total expected cost. The customer pays at the time specified in the payment schedule. If the services rendered exceed the payment, a contract asset is recognised. If the payments exceed the services rendered, a contract liability is recognised.
- (b) The Company's estimate about revenue, costs and progress towards complete satisfaction of a performance obligation is subject to a revision whenever there is a change in circumstances. Any increase or decrease in revenue or costs due to an estimate revision is reflected in profit or loss during the period when the management becomes aware of the changes in circumstances.

B. Sales revenue

The Company sells self-developed products. Sales are recognised when control of the products has been transferred, being when the products are delivered to the customer, the customer has full discretion over the channel and price to sell the products, and there is no unfulfilled obligation that could affect the customer's acceptance of the products. Delivery occurs when the products have been shipped to the specific location, the risks of obsolescence and loss have been transferred to the customer, and either the customer has accepted the products in accordance with the sales contract, or the Company has objective evidence that all criteria for acceptance have been satisfied.

C. Authorisation and cooperative development revenue

(a) The Company's authorisation and cooperative development transactions mainly arise from authorising intellectual property rights of pharmaceutical products to pharmaceutical factories. Although the Company will continuously provide research and development services on the pharmaceutical products, pharmaceutical factories can access the research and

development outcome at any time. Based on the Company's assessment, the Company uses its special technologies in manufacturing pharmaceutical cell lines, which are unique so that pharmaceutical factories would have difficulty finding another similar service provider who offers the same services in terms of the subsequent research and development on the authorised pharmaceutical products. The authorisation and subsequent research and development services provided by the Company are bonded and highly interrelated, which does not meet the criteria of being distinct, and hence are accounted for as a single performance obligation to be delivered over time. Pharmaceutical factories pay a nonrefundable up-front payment upon signing of the contracts, and make milestone payments upon each milestone achieved. The transaction prices, net of variable considerations that are not highly probable to be realised, are recognised as revenue based on the progress of performance obligations that are satisfied over time. The aforementioned stage of completion is determined based on the ratio of the actual research and development costs incurred at the end of the reporting period to the estimated total research and development costs for the authorisation contracts. The Company uses input method to measure progress towards the satisfaction of a performance obligation as there is a direct relationship between the transfer of control of services to customers and the Company's inputs, including costs of contract research and development services, contract manufacturing services and medicines. Revenue is only recognised when it is highly probable that a significant reversal will not occur. The customer pays at the time specified in the payment schedule. If the services rendered exceed the payment, a contract asset is recognised. If the payments exceed the services rendered, a contract liability is recognised. A contract liability recognised as revenue through the performance obligation is satisfied over time.

(b) The Company also entered into contracts with pharmaceutical factories, whereby the Company is entitled to a sales-based royalty in exchange for a license of manufacturing and the right to sell pharmaceutical products. In accordance with the contracts, the Company will not undertake any activities that will significantly affect the intellectual property to which the customer has rights. The Company recognises revenue at the later of when the performance obligation has been satisfied and the subsequent transfer of control or sale occurs.

(26) Government grants

Government grants are recognised at their fair value only when there is reasonable assurance that the Company will comply with any conditions attached to the grants and the grants will be received. Government grants are recognised in profit or loss on a systematic basis over the periods in which the Company recognises expenses for the related costs for which the grants are intended to compensate.

5. Critical Accounting Judgements, Estimates and Key Sources of Assumption Uncertainty

The preparation of these parent company only financial statements requires management to make critical judgements in applying the Company's accounting policies and make critical assumptions and estimates concerning future events. Assumptions and estimates may differ from the actual results and are continually evaluated and adjusted based on historical experience and other factors. Such assumptions and estimates have a significant risk of causing a material adjustment to the carrying amounts of assets and liabilities within the next financial year; and the related information is addressed below:

(1) Critical judgements in applying the Company's accounting policies

None.

(2) Critical accounting estimates and assumptions

A. Impairment on property, plant and equipment

- (a) The Company assesses impairment based on its internal and external information and industry characteristics and determines the separate cash flows of a specific group of assets, useful lives of assets and the future possible income and expenses arising from the assets depending on how assets are utilised and industrial characteristics. Any changes of economic circumstances or estimates due to the change of Company strategy might cause material impairment on assets in the future.
- (b) As of December 31, 2023, the carrying amount of property, plant and equipment was \$3,337,069.

B. Recognition of service revenue and authorisation and cooperative development revenue

- (a) Service revenue and authorisation and cooperative development revenue are recognised based on the stage of completion. The Company sets the key assumption factors for estimating total future cost based on the past operating experience, and regularly reviews and assesses the reasonableness of the basis for relevant assumptions.
- (b) For the year ended December 31, 2023, the service revenue and authorisation and cooperative development revenue amounted to \$605,990 and \$141,472, respectively.

6. Details of Significant Accounts

(1) Cash and cash equivalents

	Dece	ember 31, 2023	 December 31, 2022
Cash on hand and petty cash	\$	61	\$ 61
Demand deposits		443,581	738,882
Time deposits		4,604,962	 5,370,051
	\$	5,048,604	\$ 6,108,994

The Company transacts with a variety of financial institutions all with high credit quality to disperse credit risk, so it expects that the probability of counterparty default is remote.

(2) Financial assets at fair value through profit or loss

Items		December 31, 2023	December 31, 2022		
Non-current items:					
Financial assets mandatorily					
measured at fair value					
through profit or loss					
Profit-sharing investment in new drug	\$	58,390	\$	58,390	
development	Ψ	30,370	Ψ	30,370	
Limited partnership		20,000		_	
venture capital		20,000			
		78,390		58,390	
Valuation adjustment		1,908		3,030	
	\$	80,298	\$	61,420	

- A. The Company recognised net (losses) gains amounting to (\$1,122) and \$2,863 on financial assets at fair value through profit or loss for the years ended December 31, 2023 and 2022, respectively.
- B. On April 18, 2022, the Company entered into a new drug development profit-sharing agreement for TSY-0110 (EG12043) (the "Product") with FORMOSA PHARMACEUTICALS, INC. to replace the original development and manufacturing related cooperation agreement. Raw materials for the product development stage were provided by the Company at a reasonable market price, and FORMOSA PHARMACEUTICALS, INC. was responsible for the research and development of the product, and the implementation of the production and manufacturing of the product after completing the development of the product. Either party may commercialize this product in the global market, and each party is entitled to receive 50% licensing interest in any future revenue or interest derived from the development and commercialization of the product. Under the aforementioned agreement, the Company paid a consideration amounting to US\$30,000 thousand for the licensing interest, which will be paid in accordance with the agreement and the development schedule. As of December 31, 2023, the Company had paid US\$2,000 thousand.

(3) Financial assets at amortised cost

Items	Decer	mber 31, 2023	December 31, 2022				
Current items:							
Time deposits (Note)	\$	500,000	\$	1,000,000			
Non-current items:							
Government bonds	\$	31,930	\$	32,452			
Pledged time deposits		8,790		8,671			
	\$	40,720	\$	41,123			

Note: The deposit period for time deposits ranges between three months and a year.

A. Amounts recognised in profit or loss in relation to financial assets at amortised cost are listed below:

	 Year ended December 31									
Interest income	 2023	2022								
	\$ 28,235	\$	1,722							

- B. Details of the Company's financial assets at amortised cost pledged to others as collateral are provided in Note 8.
- C. Information relating to credit risk of financial assets at amortised cost is provided in Note 12(2). The counterparties of the Company's investments in certificates of deposits and government bonds are financial institutions and governments with high credit quality, so the Company expects that the probability of counterparty default is remote.

(4) Notes and accounts receivable

	Decen	nber 31, 2023	Decemb	per 31, 2022
Notes receivable	\$	19	\$	<u> </u>
Accounts receivable	\$	253,687	\$	33,079
Less: Allowance for uncollectible accounts	(297)	(297)
	\$	253,390	\$	32,782

A. The ageing analysis of notes receivable and accounts receivable that were past due but not impaired is as follows:

	Decemb	er 3	1, 2023	December 31, 2022							
	Notes receivable	Acc	counts receivable	Notes	receivable	Acc	ounts receivable				
Not past due	\$ 19	\$	176,990	\$	-	\$	32,782				
Up to 30 days past due	-		76,400		-		-				
31 to 90 days past due	-		-		-		-				
91 to 180 days past due	-		-		-		-				
Over 181 days past due			297	-			297				
	<u>\$ 19</u>	<u>\$</u>	253,687	\$		\$	33,079				

The above ageing analysis was based on past due date.

- B. As of December 31, 2023 and 2022, notes receivable and accounts receivable (including related parties) were all from contracts with customers. Also, as of January 1, 2022, the balance of receivables from contracts with customers amounted to \$80,159.
- C. As at December 31, 2023 and 2022, without taking into account any collateral held or other credit enhancements, the maximum exposure to credit risk in respect of the amount that best represents the notes and accounts receivable (including related parties) held by the Company was \$256,045 and \$32,782, respectively.
- D. The Company did not hold any collateral.
- E. Information relating to credit risk of accounts receivable and notes receivable is provided in Note 12(2).

(5) <u>Inventories</u>

	December 31, 2023											
		Cost	Book value									
Raw materials	\$	426,217	(\$	51,483)	\$	374,734						
Work in progress		127,143		-		127,143						
Finished goods		178,690	(165)		178,525						
Merchandise inventory		235				235						
	\$	732,285	(\$_	51,648)	\$	680,637						
	December 31, 2022											
				Allowance for								
		Cost		valuation loss		Book value						
Raw materials	\$	377,424	(\$	18,327)	\$	359,097						
Work in progress		281,739		-		281,739						
Finished goods		98,150		-		98,150						
Merchandise inventory		477		<u> </u>		477						
	\$	757,790	(<u>\$</u>	18,327)	\$	739,463						

The cost of inventories recognised as expense for the year:

	Year ended December 31										
		2023		2022							
Cost of goods used	\$	327,739	\$	199,628							
Cost of goods sold		149,041		138,672							
Loss on decline in market value		33,321		1,012							
Loss on disposal inventory		1,014		-							
(Gain) loss on physical inventory	(4)		34							
	\$	511,111	\$	339,346							

(6) Prepayments

	Decen	nber 31, 2023	Decen	nber 31, 2022
Office supplies	\$	-	\$	9,009
Prepayments for contracted research expense		17,151		11,310
Excess business tax paid (or Net Input VAT)		24,504		6,106
Prepayments to suppliers		26,187		50,100
Other prepaid expenses		24,835		45,977
	\$	92,677	\$	122,502

(7) Financial assets at fair value through other comprehensive income

Items	Dec	cember 31, 2023	December 31, 2022				
Non-current items:							
Equity instruments							
Emerging and unlisted stocks	\$	215,026	\$	214,403			
Valuation adjustment		110,861		64,922			
	\$	325,887	\$	279,325			

- A. The Company has elected to classify shares that are considered to be strategic investments as financial assets at fair value through other comprehensive income. The fair value of such investments amounted to \$325,887 and \$279,325 as at December 31, 2023 and 2022, respectively.
- B. The Company acquired equity instruments amounting to \$623 and \$208,627 for the years ended December 31, 2023 and 2022, respectively.
- C. Amounts recognised in profit or loss and other comprehensive income in relation to the financial assets at fair value through other comprehensive income are listed below:

		Year ended	Dec	ember 31			
		2023	2022				
Equity instruments at fair value through other comprehensive income							
Fair value change recognised in other comprehensive income	<u>\$</u>	45,939	<u>\$</u>	59,091			
Dividend income recognised in profit or loss Held at end of period	<u>\$</u>	475	<u>\$</u>				
(8) Investments accounted for using equity	<u>meth</u>						
		December 31, 2023		December 31, 2022			

A. Please refer to Note 4(3) in the consolidated financial statements for the year ended December 31, 2023 for the information regarding the Company's subsidiaries.

7,743

5,200

Subsidiary:

EirGenix Europe GmbH

B. EirGenix USA Inc. is a subsidiary that was established in November 2023. As of December 31, 2023, no capital has been remitted.

C. Share of profit of subsidiaries, associates and joint ventures accounted for using equity method is as follows:

	Year ended December 31								
		2023	2022						
Subsidiary:									
EirGenix Europe GmbH	\$	2,324	\$	1,690					

(9) Property, plant and equipment

2023

								20	25							
		achinery and equipment		Office equipment	I	Buildings and structures		Leasehold provements		Other equipment	con	Unfinished astruction and ipment under acceptance		Total	bus (sł	epayments for siness facilities nown as other non-current assets)
At January 1	_		_		_		_		_		_		_		_	
Cost	\$	978,923		74,309	\$	1,434,479	\$	45,596	\$	- ,	\$	642,864	\$	3,209,096		98,273
Accumulated depreciation	(317,142)	\$	30,004)	(_	229,062)	\$	12,142)	<u>_</u>	12,788)	Φ	612.961	<u></u>	601,138)	\$	- 09 272
	<u> </u>	661,781	Φ	44,305	ф	1,205,417	Ф	33,454	Ф	20,137	Ф	642,864	<u> </u>	2,607,958	Φ_	98,273
Opening net book amount as at January 1	\$	661,781	\$	44,305	\$	1,205,417	\$	33,454	\$	20,137	\$	642,864	\$	2,607,958	\$	98,273
Additions	Ψ	116,776	Ψ	6,416	φ	15,263	Ψ	1,665	φ	5,768	Ψ	557,083	Ψ	702,971	φ	138,453
Reclassifications		147,987		-,		528,357		-,		-	(676,344)		-		-
Transfers from other non- current assets		62,723		32		-		59		1,294		158,129		222,237	(222,237)
Depreciation expense	(98,594)	(9,063)	(78,412)	(4,809)	(5,219)		-	(196,097)		_
Closing net book amount as at December 31	\$	890,673	\$	41,690	\$	1,670,625	\$	30,369	\$	21,980	\$	681,732	\$	3,337,069	\$	14,489
At December 31 Cost	\$	1,301,038	\$	78,972	\$	1,978,099	\$	47,320	\$	37,142	\$	681,732	\$	4,124,303	\$	14,489
Accumulated depreciation	(410,365)		37,282)	(307,474)		16,951)		15,162)	Ψ	-	(787,234)		- 1,107
•	\$	890,673	\$	41,690	\$	1,670,625	\$	30,369	\$	21,980	\$	681,732	\$	3,337,069	\$	14,489

	M	Iachinery and equipment	Office equipment]	Buildings and structures		Leasehold nprovements		Other equipment	con	Unfinished struction and ipment under acceptance		Total	bus (sł	epayments for iness facilities nown as other non-current assets)
At January 1 Cost Accumulated depreciation	\$	813,793 239,109)	\$ 67,037 23,995)	\$	1,295,911 164,219)	\$	24,495 8,974)		26,524 8,870)	\$	103,265	\$ (2,331,025 445,167)	\$	65,456
. room.maneed doprocum.	\$	574,684	\$ 43,042	\$	1,131,692	\$	15,521	\$	17,654	\$	103,265	\$	1,885,858	\$	65,456
Opening net book amount as at January 1 Additions	\$	574,684 92,578	\$ 43,042 8,619	\$	1,131,692 76,679	\$	15,521 21,101	\$	17,654 6,516	\$	103,265 278,775	\$	1,885,858 484,268	\$	65,456 433,952
Reclassifications Transfers from other non- current assets		7,701 69,453	862		61,890		-		405	(69,591) 330,415		401,135	(401,135)
Depreciation expense Reclassified to inventories Closing net book amount	(80,909) 1,726)	8,218)	(64,844)	(3,168)	(4,438)		- -	(161,577) 1,726)		- -
as at December 31	\$	661,781	\$ 44,305	\$	1,205,417	\$	33,454	\$	20,137	\$	642,864	\$	2,607,958	\$	98,273
At December 31 Cost Accumulated depreciation	\$ (978,923 317,142)	74,309 30,004)	\$ (_	1,434,479 229,062)	\$ (45,596 12,142)	\$ (_	32,925 12,788)	\$	642,864	\$ (3,209,096 601,138)		98,273
•	\$	661,781	\$ 44,305	\$	1,205,417	\$	33,454	\$	20,137	\$	642,864	\$	2,607,958	\$	98,273

Information about the property, plant and equipment that were pledged to others as collateral is provided in Note 8.

(10) <u>Leasing arrangements - lessee</u>

- A. The Company leases various assets including land, buildings, machinery and equipment, multifunction printers and business vehicles. Rental contracts are typically made for periods of 1 to 20 years. Lease terms are negotiated on an individual basis and contain a wide range of different terms and conditions. The lease agreements do not impose covenants, but leased assets may not be used as security for borrowing purposes.
- B. Short-term leases with a lease term of 12 months or less comprise of certain offices, dormitories, business vehicles and warehouses. Low-value assets comprise multifunction printers.
- C. The carrying amount of right-of-use assets and the depreciation expense are as follows:

	<u></u>	December 31, 2023		December 31, 2022		
		Carrying amount		Carrying amount		
Land	\$	187,939	\$	202,394		
Buildings		73,893		84,031		
Machinery and equipment		65,921		35,305		
Transportation equipment (Business vehicles)		967		2,584		
Office equipment						
(Multifunction printers)		516	_	1,016		
	\$	329,236	\$	325,330		
	Year ended December 31					
		2023		2022		
		Depreciation expense		Depreciation expense		
Land	\$	14,648	\$	14,543		
Buildings		10,292		7,521		
Machinery and equipment		4,055		2,310		
Transportation equipment (Business vehicles)		1,616		1,640		
Office equipment						
(Multifunction printers)		500		396		
	\$	31,111	\$	26,410		

D. For the years ended December 31, 2023 and 2022, the additions to right-of-use assets were \$35,017 and \$54,767, respectively.

The information on profit and loss accounts relating to lease contracts is as follows:

	Year ended December 31				
		2023		2022	
Items affecting profit or loss					
Interest expense on lease	\$	8,096	\$	8,204	
liabilities					
Expense on short-term lease contracts		22,508		11,889	
Expense on leases of low-value assets		307		376	
Loss on lease modification		413		709	

E. For the years ended December 31, 2023 and 2022, the Company's total cash outflow for leases were \$60,218 and \$44,126, respectively.

(11) Intangible assets

		2023				
		Software	Pro	fessional expertise		Total
At January 1						
Cost	\$	45,851	\$	107,953	\$	153,804
Accumulated amortisation	(21,678)	(104,059)	(125,737)
	\$	24,173	\$	3,894	\$	28,067
Opening net book amount as at January 1	\$	24,173	\$	3,894	\$	28,067
Additions		3,339		8,159		11,498
Amortisation expense	(7,129)	(4,167)	(11,296)
Closing net book amount as at December 31	\$	20,383	\$	7,886	\$	28,269
At December 31						
Cost	\$	49,190	\$	116,112	\$	165,302
Accumulated amortisation	(28,807)	(108,226)	(137,033)
	\$	20,383	\$	7,886	\$	28,269

				2022		
		Software	Profess	sional expertise		Total
At January 1						
Cost	\$	21,153	\$	107,953	\$	129,106
Accumulated amortisation	(16,438)	(93,115)	(109,553)
	<u>\$</u>	4,715	\$	14,838	\$	19,553
Opening net book amount as at January 1	\$	4,715	\$	14,838	\$	19,553
Additions		8,652		-		8,652
Transfers(Note)		16,046		-		16,046
Amortisation expense	(5,240)	(10,944)	(16,184)
Closing net book amount as at December 31	\$	24,173	\$	3,894	\$	28,067
At December 31						
Cost	\$	45,851	\$	107,953	\$	153,804
Accumulated amortisation	(21,678)	(104,059)	(125,737)
	\$	24,173	\$	3,894	\$	28,067

Note:Transfers pertain to the assets transferred from prepaid intangible assets (shown as 'other non-current assets').

A. Details of amortisation on intangible assets are as follows:

	Year ended December 31				
		2023		2022	
Operating costs	\$	5,840	\$	10,456	
General and administrative expenses		1,193		1,005	
Research and development expenses		4,156		4,705	
Sales and marketing expenses		107		18	
	\$	11,296	\$	16,184	

- B. The basic information of the professional expertise that is material to the Company is as follows:
 - (a) In April 2013, the Company acquired professional expertise, including cell line establishment, process development, process optimisation, analytical method development and validation, product qualification, GMP manufacturing and stability test, etc., amounting to \$92,483 from the Development Center for Biotechnology cGMP biopharmaceutical pilot plant facility.

- (b) In July 2013, the Company acquired professional expertise of Herceptin from FORMOSA PHARMACEUTICALS, INC. amounting to \$7,143.
- (c) In July 2013, the Company acquired commercial authorisation of recombinant protein cell line from Life Technologies Corporation amounting to \$7,485.
- (d) In September 2023, the Company obtained an authorisation from American Type Culture Collection for the detection of cancer cell lines with a total price of \$8,159, which can be applied on the commercial implementation of the marketing and manufacturing of subsequent cancer drug products.

(12) Other non-current assets

	Dece	mber 31, 2023	 December 31, 2022
Non-current prepayments for investments	\$	46,270	\$ 20,000
Long-term prepayments to suppliers		30,000	30,000
Prepayments for business facilities		14,489	98,273
Guarantee deposits paid		8,795	65,048
Other assets		5,210	 1,566
	\$	104,764	\$ 214,887

(13) Other payables

	Decen	nber 31, 2023	Decer	mber 31, 2022
Payable on construction and equipment	\$	285,960	\$	158,228
Salary and bonus payable		88,898		87,593
Service expense payable		44,968		51,108
Payable on consumables		18,604		25,012
Payable on repairs and maintenance expense		28,856		19,732
Others		52,476		43,009
	\$	519,762	\$	384,682

(14) Long-term borrowings

,	Borrowing period and			
Type of borrowings	repayment term	Interest rate range	Collateral	December 31, 2023
Long-term bank borrowi	ngs			
Credit borrowings	Borrowing period is from February 15, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from March 2025	1.7250% ~1.9500%	None	\$ 39,560
"	Borrowing period is from June 30, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from March 2025	1.7250% ~1.9500%	"	80,900
				\$ 120,460
	Borrowing period and			
Type of borrowings	Borrowing period and repayment term	Interest rate range	Collateral	December 31, 2022
Long-term bank borrowi	repayment term			December 31, 2022
	repayment term	1.3500% ~1.8250%	None None	
Long-term bank borrowi	repayment term ngs Borrowing period is from February 15, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from March 2025 Borrowing period is from June 30, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from	1.3500%		December 31, 2022
Long-term bank borrowings Credit borrowings	repayment term ngs Borrowing period is from February 15, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of every month from March 2025 Borrowing period is from June 30, 2022 to February 15, 2027; interest is payable monthly; principal is payable on the 15th of	1.3500% ~1.8250% 1.4750%	None	December 31, 2022

- A. Information on the Company's undrawn borrowing facilities is provided in Note 12(2) C.
- B. On December 23, 2021, the Company entered into a \$714,000 syndicated loan agreement with Hua Nan Commercial Bank Ltd. and the government will subsidize 0.5% handling fee of the bank for the Company's compliance with the "Action Plan for Accelerated Investment by Domestic Corporations".
- C. Information about assets pledged as collateral for long-term borrowings is provided in Note 8.

(15) Pensions

- A. The Company has established a defined contribution pension plan (the "New Plan") under the Labor Pension Act (the "Act"), covering all regular employees with R.O.C. nationality. Under the New Plan, the Company contributes monthly an amount not lower than 6% of the employees' monthly salaries and wages to the employees' individual pension accounts at the Bureau of Labor Insurance. The benefits accrued are paid monthly or in lump sum upon termination of employment.
- B. The pension costs under the defined contribution pension plans of the Company for the years ended December 31, 2023 and 2022, were \$17,928 and \$16,051, respectively.

(16) Share-based payment

A. For the years ended December 31, 2023 and 2022, the Company's share-based payment arrangements were as follows:

Type of		Quantity granted		
arrangement	Grant date	(shares in thousands)	Contract period	Vesting conditions
Employee stock options - B	2015. 07. 01	1,270	10 years	1 to 4 years' service
"	2015. 07. 01	130	"	"
"	2015. 07. 06	250	"	"
"	2016. 01. 01	270	"	"
Employee stock options - C	2016. 05. 05	100	10 years	2 to 4 years' service
Employee stock options - D	2016. 10. 12	515	10 years	2 to 4 years' service
"	2016. 12. 29	85	"	"

Type of		Quantity granted		
arrangement	Grant date	(shares in thousands)	Contract period	Vesting conditions
Employee stock	2017. 08. 08	395	10 years	2 to 4 years'
options - E				service
"	2017. 12. 27	570	"	"
"	2018. 03. 23	175	"	"
Employee stock options - F	2019. 01. 25	520	10 years	2 to 4 years' service
"	2019. 05. 13	285	"	"
Restricted stocks	2016. 11. 18	1,660	NA	Conditions of
to employees - A				service years and performance
"	2017. 08. 08	257	"	"
Employee stock	2019. 11. 12	960	10 years	2 to 4 years'
options - G		, , ,	3	service
"	2020. 04. 15	775	"	"
"	2020. 08. 12	205	"	11
Restricted stocks	2020. 05. 13	455	NA	0.25 to 3 years'
to employees - B				service
"	2020. 12. 10	144	"	"
Restricted stocks	2020. 08. 14	905	NA	Performance
to employees - D				conditions
"	2020 12 10	0.4	"	"
	2020. 12. 10	94		
Employee stock options - H	2020. 12. 23	830	10 years	2 to 4 years' service
"	2021. 05. 12	315	"	"
"	2021. 08. 12	505	"	"
"	2021. 10. 01	1,185	"	"
Restricted stocks	2021. 10. 15	613	NA	Performance
to employees - E				conditions
"	2022. 01. 10	184	"	"
"	2022. 09. 08	190	"	"
Restricted stocks to employees - F	2021. 10. 15	340	NA	Performance conditions
Employee stock options - I	2022. 03. 22	160	10 years	2 to 4 years' service
"	2022. 05. 12	225	"	"
"	2022. 08. 11	685	"	"
"	2022. 09. 08	510	"	"

Type of		Quantity granted		
arrangement	Grant date	(shares in thousands)	Contract period	Vesting conditions
Restricted stocks to employees - G	2022. 09. 08	63	NA	Performance conditions
"	2022. 11. 08	195	"	"
"	2023. 03. 10	6	"	"
"	2023. 11. 09	325	"	"
Employee stock	2022. 11. 08	615	10 years	2 to 4 years'
options - J				service
"	2023. 03. 10	1,105	"	"
"	2023. 05. 10	255	"	"
"	2023. 08. 08	225	"	"
"	2023. 12. 22	270	"	"
Restricted stocks to employees - H	2023. 11. 09	826	NA	Performance conditions
Restricted stocks to employees - I	2023. 12. 22	26	NA	Performance conditions

- (a) The restricted stocks issued by the Company cannot be transferred during the vesting period, but voting right and dividend right are not restricted on these stocks. If employees resign during the vesting period, they are considered not meeting the vesting condition from the date of resignation and the Company will redeem and retire those stocks at the initial issuance price, but employees are not required to return the dividends received.
- (b) The above-mentioned share-based payment arrangements are equity-settled.
- B. Details of the share-based payment arrangements are as follows:
 - (a) Employee stock options

	2023		2022		
		Weighted-		Weighted-	
		average		average	
	No. of options	exercise price	No. of options	exercise price	
	(shares in thousands)	(in dollars)	(shares in thousands)	(in dollars)	
Options outstanding at January 1	5,666	\$15~146.4	5,282	\$15~146.4	
Options granted	1,855	100.5~120	2,195	71.6~118.5	
Options forfeited	(894)	25.2~146.4	(759)	25.2~146.4	
Options exercised	(15~51.2	(1,052)	15~51.2	
Options outstanding at December 31	5,900	15~146.4	5,666	15~146.4	
Options exercisable at December 31	1,608		1,238		

(b) Restricted stocks to employees

		2023	2022		
		(shares in thousands)	(shares in thousands)		
Stocks outstanding at January 1		2,571	2,869		
Stocks granted		1,182	632		
Stocks vested	(1,167) (704)		
Stocks retired	(_	193) (226)		
Stocks outstanding at December 31	_	2,393	2,571		

- C. The weighted-average stock prices of stock options at exercise dates for the years ended December 31, 2023 and 2022 were \$105.6 (in dollars) and \$99.8 (in dollars), respectively.
- D. The expiry date and exercise price of stock options outstanding at the balance sheet dates are as follows:

			December	31, 2023	December 31, 2022			
Type of	Issue date		No. of shares (shares in	Exercise price	No. of shares (shares in	Exercise price		
		F 1 .	`	-	•	-		
arrangement	approved	Expiry date	thousands)	(in dollars)	thousands)	(in dollars)		
Employee	2015. 07. 01	2025. 06. 30	50	\$ 15	140	\$ 15		
stock								
options - B								
"	2015. 07. 01	2025. 06. 30	5	20	20	20		
"	2015. 07. 06	2025. 07. 05	15	20	25	20		
"	2016. 01. 01	2025. 12. 31	25	20	25	20		
Employee	2016. 05. 05	2026. 05. 04	10	29.2	10	29.2		
stock								
options - C								
Employee	2016. 10. 12	2026. 10. 11	150	29.2	180	29.2		
stock								
options - D								
"	2016. 12. 29	2026. 12. 28	15	37.5	15	37.5		
Employee	2017. 08. 08	2027. 08. 07	4	29.2	18	29.2		
stock								
options - E								
"	2017. 12. 27	2027. 12. 26	79	25	112	25		
"	2018. 03. 23	2028. 03. 22	48	23.5	52	23.5		

			December	31, 2023	December 31, 2022		
			No. of shares	Exercise	No. of shares	Exercise	
Type of	Issue date		(shares in	price	(shares in	price	
arrangement	approved	Expiry date	thousands)	(in dollars)	thousands)	(in dollars)	
Employee	2019. 01. 25	2029. 01. 24	34	\$ 28.7	103	\$ 28.7	
stock							
options - F							
"	2019. 05. 13	2029. 05. 12	94	34.3	141	34.3	
Employee	2019. 11. 12	2029. 11. 11	207	25.2	325	25.2	
stock							
options - G							
"	2020. 04. 15	2030. 04. 14	89	28.8	175	28.8	
"	2020. 08. 12	2030. 08. 11	79	51.2	140	51.2	
Employee	2020. 12. 23	2030. 12. 22	341	42.1	515	42.1	
stock							
options - H							
"	2021. 05. 12	2031. 05. 11	215	146.4	235	146.4	
"	2021. 08. 12	2031. 08. 11	250	128.4	305	128.4	
"	2021. 10. 01	2031. 09. 30	835	117.5	990	117.5	
Employee	2022. 03. 22	2032 .03. 21	80	93.5	145	93.5	
stock							
options - I							
"	2022. 05. 12	2032. 05. 11	195	71.6	225	71.6	
"	2022. 08. 11	2032. 08. 10	440	85.9	645	85.9	
"	2022. 09. 08	2032. 09. 07	345	118.5	510	118.5	
Employee	2022. 11. 08	2032. 11. 07	510	103.5	615	103.5	
stock							
options - J							
"	2023. 03. 10	2033. 03. 09	1,035	111.5	-	-	
"	2023. 05. 10	2033. 05. 09	255	120	-	-	
"	2023. 08. 08	2033. 08. 07	225	101.5	-	-	
"	2023. 12 .22	2033. 12. 21	270	100.5	-	-	

E. The fair value of stock options granted is measured using the Black-Scholes option-pricing model to estimate the fair value of employee stock options, cash capital increase reserved for employee preemption and restricted stocks to employees. Relevant information is as follows:

				Exercise	Expected	Expected		
Type of	a .	Quantity granted	Stock price	price	price	option	Risk-free	Fair value per
arrangement		(shares in thousands)	(in dollars)	(in dollars)	volatility	life	interest rate	unit (in dollars)
Employee	2015. 07. 01	1,270	\$ 14.88	\$ 15	36.58~	5.5 ~ 7	1.15~	\$5.22 ~
stock options - B					37.13%	years	1.35%	6.01
"	2015. 07. 01	130	14.88	20	36.58~	5.5 ~ 7	1.15~	3.83~
					37.13%	years	1.35%	4.69
"	2015. 07. 06	250	14.60	20	37.09~	5.5 ~ 7	1.15~	3.75~
					37.64%	years	1.35%	4.60
"	2016. 01. 01	270	16.03	20	40.11~	5.5 ~ 7	0.79~	4.91~
					40.30%	years	0.90%	5.76
Employee	2016. 05. 05	100	13.27	29.2	40.75~	6 ~ 7 years	0.70~	1.86 ~
stock options - C					40.91%		0.77%	2.30
Employee	2016. 10. 12	515	21.42	29.2	39.82~	$6 \sim 7 \text{ years}$	0.71~	5.19~
stock options - D					39.91%		0.75%	5.93
"	2016. 12. 29	85	20.40	37.5	39.39~	6 ~ 7 years	1.16~	3.49~
					39.48%	,	1.20%	4.18
Employee	2017. 08. 08	395	18.75	29.2	38.13~	6 ~ 7 years	0.82~	3.64~
stock options - E					38.22%		0.88%	4.23
_ "	2017. 12. 27	570	18.07	25	36.97~	6 ~ 7 years	0.74~	3.81~
					37.23%		0.80%	4.41
"	2018. 03. 23	175	19.16	23.5	36.87~	6 ~ 7 years	0.79~	4.71 ~
					37.17%		0.84%	5.38
Employee	2019. 01. 25	520	21.96	28.7	36.03~	6 ~ 7 years	0.72~	4.85~
stock options - F					36.90%		0.78%	5.74
"	2019. 05. 13	285	25.75	34.3	35.50%~ 36.35%	6 ~ 7 years	0.64~ 0.67%	5.39 ~ 6.40
Restricted stocks to employees -	2016. 11. 18	1,660	22.88	-	-	-	-	22.88
A								
"	2017. 08. 08	257	19.61	_	_	_	_	19.61
Employee	2019. 11. 12	960	29.05	25.2	26.38%	6 ~ 7 years	0.63~	7.77 ~
stock options - G						J	0.66%	8.42
"	2020. 04. 15	775	33.10	28.8	50.33%	6 ~ 7 years	0.47~ 0.49%	15.56 ~ 16.65
"	2020. 08. 12	205	57.80	51.2	64.08%	6 ~ 7 years	0.45%	33.07 ~
	2020. 00. 12	203	37.00	31.2	04.0070	o 7 years	0.38%	35.18
Restricted	2020. 05. 13	455	46.85	_	_	-	-	46.85
stocks to								
employees - B								
"	2020. 12. 10	144	48.60	_	_	_	_	48.60
Restricted	2020. 12. 10	905	55.70	_	<u>-</u>	_	<u>-</u>	55.70
stocks to	_0_0, 00, 14	703	33.70	_				33.70
employees -								
D								
"	2020. 12. 10	94	48.60					48.60
	2020. 12. 10	94	48.00	-	-	-	-	40.00

Type of arrangement	Grant date	Quantity granted (shares in thousands)	Stock price (in dollars)	Exercise price (in dollars)	Expected price volatility	Expected option life	Risk-free interest rate	Fair value per unit (in dollars)
Employee stock options - H	2020. 12. 23	830	\$ 47.55	\$ 42.1	61.28%	6 ~ 7 years	0.22~ 0.26%	\$26.15~ 27.88
"	2021. 05. 12	315	154.5	146.4	65.02%	6 ~ 7 years	0.31~ 0.35%	89.32~ 95.02
"	2021. 08. 12	505	135.5	128.4	67.02%	6 ~ 7 years	0.32~ 0.34%	80.24~ 85.25
"	2021. 10. 01	1,185	124.0	117.5	65.78%	6 ~ 7 years	0.34~ 0.38%	72.39~ 76.99
Restricted stocks to employees - E	2021. 10. 15	613	106.5	-	-	-	-	106.5
"	2022. 01. 10	184	108.5	_	_	_	_	108.5
"	2022. 09. 08	190	118.5	_	_	_	_	118.5
Restricted stocks to employees - F	2021. 10. 15	340	106.5	-	-	-	-	106.5
Employee stock options - I	2022. 03. 22	160	93.5	93.5	62.20%	6 ~ 7 years	0.86~ 0.87%	52.85~ 56.27
"	2022. 05. 12	225	71.6	71.6	61.32%	6 ~ 7 years	1.22~ 1.27%	40.37~ 43.04
"	2022. 08. 11	685	85.9	85.9	60.04%	6 ~ 7 years	1.10~ 1.14%	47.51~ 50.67
"	2022. 09. 08	510	118.5	118.5	60.29%	6 ~ 7 years	1.19~ 1.23%	65.9~ 70.28
Restricted stocks to employees - G	2022. 09. 08	63	118.5	-	-	-	-	118.5
"	2022. 11. 08	195	103.5	-	-	-	-	103.5
"	2023. 03. 10	6	111.5	-	-	-	-	111.5
"	2023. 11. 09	325	103.0	-	-	-	-	103.0
Employee stock options - J	2022. 11. 08	615	103.5	103.5	60.00%	6 ~ 7 years	1.63~ 1.70%	57.97~ 61.88
	2023. 03. 10	1,150	111.5	111.5	59.15%	6 ~ 7 years	1.12~ 1.14%	60.98~ 65.04
"	2023. 05. 10	255	120.0	120.0	58.70%	6 ~ 7 years	1.07~ 1.09%	65.15~ 69.50
"	2023. 08. 08	225	101.5	101.5	57.40%	6 ~ 7 years	1.10~ 1.12%	54.18~ 57.84
"	2023. 12. 22	270	100.5	100.5	55.38%	6 ~ 7 years	1.12% 1.18~ 1.19%	52.26~ 55.82

Type of arrangement	Grant date	Quantity granted (shares in thousands)	ock price dollars)	p	ercise orice dollars)	Expected price volatility	Expected option life	Risk-free interest rate	Fair value per unit (in dollars)
Restricted stocks to	2023. 11. 09	826	\$ 103.0	\$	-	-	-	-	\$103.0
employees - H Restricted stocks to employees - I	2023. 12. 22	26	100.5		-	-	-	-	100.5

F. Expenses incurred on share-based payment transactions are shown below:

	Year ended December 31						
		2023	2022				
Employee stock options	\$	84,285	\$	62,522			
Restricted stocks to employees		96,615		92,790			
	\$	180,900	\$	155,312			

(17) Share capital

A. As of December 31, 2023, the Company's authorised capital was \$4,000,000, consisting of 400,000 thousand shares of ordinary share (including 12 million shares reserved for employee stock options, preferred shares with warrants or convertible bonds issued by the Company), and the paid-in capital was \$3,060,516 with a par value of \$10 (in dollars) per share, consisting of 306,052 thousand shares. All proceeds from shares issued have been collected.

Movements in the number of the Company's ordinary shares outstanding are as follows (unit: shares in thousands):

	2023	2022
At January 1	304,336	300,385
Employee stock options		
exercised	727	1,052
Issuance of employee restricted		
stocks	1,182	632
Redemption of employee restricted		
stocks (193)	(226)
Conversion of convertible bonds	<u> </u>	2,493
At December 31	306,052	304,336

B. For the years ended December 31, 2023 and 2022, the Company issued 727 thousand and 1,052 thousand ordinary shares related to the exercise of employee share options in accordance with

- the employee share options plan with a par value of \$10 (in dollars) per share, totalling \$7,270 and \$10,523, respectively.
- C. For the years ended December 31, 2023 and 2022, the Company's Board of Directors resolved to repurchase and retire the employee restricted stocks because employee restricted stocks distributed to certain employees amounting to 193 thousand shares and 226 thousand shares, respectively, did not meet the vesting conditions in accordance with the terms of restricted shares.
- D. The shareholders during their meeting on August 3, 2021 resolved to issue the 1st and 2nd restricted stocks to employees amounting to 1,000 thousand and 340 thousand shares with no subscription price, respectively. The Board of Directors of the Company resolved to issue the 1st and 2nd restricted stocks to employees amounting to 797 thousand and 340 thousand shares in 2021, respectively. The Board of Directors of the Company resolved to issue the 1st restricted stocks to employees amounting to 190 thousand shares in 2021.
- E. The shareholders during their stockholders' meeting on August 3, 2021 resolved to issue 55,000 thousand ordinary shares through the private placement. The Board of Directors of the Company resolved the issuance price of \$91.5 (in dollars) and the total consideration of issuing common stock was \$5,032,500 on October 1, 2021, and the effective date was set on October 15, 2021. The registration has been completed on December 13, 2021. Pursuant to the Securities and Exchange Act, the ordinary shares raised through the private placement are subject to certain transfer restrictions and cannot be listed on the stock exchange until three years after they have been issued and have been offered publicly. Other than these restrictions, the rights and obligations of the ordinary shares raised through the private placement are the same as other issued ordinary shares.
- F. The shareholders during their meeting on June 10, 2022, resolved to issue the 1st restricted stocks to employees amounting to 850 thousand shares with no subscription price. On September 8, 2022, the Board of Directors of the Company resolved to issue restricted stocks to employees amounting to 63 thousand shares with the effective date set on September 8, 2022. On November 8, 2022, the Board of Directors of the Company resolved to issue restricted stocks to employees amounting to 195 thousand shares with the effective date set on November 8, 2022. On March 10, 2023, the Board of Directors resolved to issue restricted stocks to employees amounting to 6 thousand shares with the effective date set on March 10, 2023. On November 9, 2023, the Board of Directors of the Company resolved to issue restricted stocks to employees amounting to 325 thousand shares with the effective date set on November 9, 2023.
- G. The shareholders during their meeting on May 31, 2023 resolved to issue the 1st and 2nd restricted stocks to employees amounting to 805 thousand and 870 thousand shares with no

subscription price, respectively. On November 9, 2023, the Board of Directors of the Company resolved to issue the 2nd restricted stocks to employees amounting to 826 thousand shares in 2023, with the effective date set on November 9, 2023. On December 22, 2023, the Board of Directors of the Company resolved to issue the 1st restricted stocks to employees amounting to 26 thousand shares in 2023, with the effective date set on December 22, 2023.

H. The shareholders during their meeting on May 31, 2023 adopted a resolution to raise cash capital through private placement. The maximum number of shares to be issued through the private placement is 30,000 thousand shares and the private placement may be made in three installments as authorised by the shareholders during their meeting. The private placement was in accordance with the Securities and Exchange Act and the Regulations for Public Companies Conducting Private Placements of Securities. The Company's Board of Directors resolved not to execute the private placement on March 8, 2024.

(18) Capital surplus

Pursuant to the R.O.C. Company Act, capital surplus arising from paid-in capital in excess of par value on issuance of common stocks and donations can be used to cover accumulated deficit or to issue new stocks or cash to shareholders in proportion to their share ownership, provided that the Company has no accumulated deficit. Further, the R.O.C. Securities and Exchange Act requires that the amount of capital surplus to be capitalised mentioned above should not exceed 10% of the paid-in capital each year. However, capital surplus should not be used to cover accumulated deficit unless the legal reserve is insufficient.

(19) Accumulated deficit

- A. Under the Company's Articles of Incorporation, the current year's earnings, if any, shall first be used to pay all taxes and offset prior years' operating losses and then 10% of the remaining amount shall be set aside as legal reserve. After the provision or reversal of special reserve in accordance with laws or regulations, the appropriation of the remaining earnings along with the unappropriated earnings of prior years shall be proposed by the Board of Directors and resolved at shareholders' meetings.
- B. The Company's dividend policy is summarised below: The Board of Directors would consider the earnings situation of current year, capital and financial structure, future operating needs, retained earnings and legal reserve, as well as the market competition to propose the appropriation of earnings to the shareholders during their meetings for resolution, and cash dividends shall account for at least 10% of the total dividends distributed.

- C. On June 10, 2022, the shareholders at their meetings resolved the deficit compensation for the year ended December 31, 2021. The Company offset the accumulated deficit by capital surplus. Refer to the website of "Market Observation Post System" for information about earnings appropriation to offset deficit as proposed by the Board of Directors and resolved by the shareholders.
- D. On May 31, 2023, the shareholders resolved the deficit compensation for the year ended December 31, 2022. The Company offset the accumulated deficit against the capital surplus. Refer to the website of "Market Observation Post System" for information about earnings appropriation to offset deficit as proposed by the Board of Directors and resolved by the shareholders.
- E. On March 8, 2024, the Board of Directors proposed the deficit compensation for the year ended December 31, 2023. The Company offset the accumulated deficit against the capital surplus. Refer to the website of "Market Observation Post System" for information about earnings appropriation to offset deficit as proposed by the Board of Directors and resolved by the shareholders.
- F. As of December 31, 2023 and 2022, there was no earnings to be distributed.

(20) Operating revenue

	Year ended December 31						
		2023	2022				
Revenue from contracts with							
customers	\$	1,022,653	\$	1,481,017			

A. Disaggregation of revenue

The Company derives revenue from the transfer of services, authorisation and goods over time and at a point in time in the following major categories:

		Year ended December 31, 2023								
			Sales of authorisation and cooperative							
	Sales of services		dev	velopment	Sale	es of goods	ls Total			
Timing of revenue recognition										
At a point in time	\$	-	\$	-	\$	161,594	\$	161,594		
Over time		605,990		141,472		113,597		861,059		
	\$	605,990	\$	141,472	\$	275,191	\$	1,022,653		

		Year ended December 31, 2022										
				Sales of								
			aut	horisation								
			and	cooperative								
	Sales	of services	de	velopment	Sales of goo			Total				
Timing of revenue recognition												
At a point in time	\$	-	\$	-	\$	417,774	\$	417,774				
Over time		757,680		261,876		43,687		1,063,243				
	\$	757,680	\$	261,876	\$	461,461	\$	1,481,017				

B. Contract assets and liabilities

(a) The Company has recognised the following revenue-related contract assets and liabilities:

	December 31, 20			nber 31, 2022	January 1, 2022	
Contract assets:						
Services	\$	240,564	\$	213,981	\$	144,831
Sales		53,130		20,418		25,766
	\$	293,694	\$	234,399	\$	170,597
Current contract liabilities		_				
Services	\$	41,739	\$	104,384	\$	102,289
Authorisation and cooperative		15,027		46,091		121,678
Non-current contract liabilities						
Authorisation and						
cooperative development						
			-			20,059
	\$	56,766	\$	150,475	\$	244,026

(b) Revenue recognised that was included in the contract liability balance at the beginning of the year.

included in the contract liabilit	5
balance at the beginning of the	,
year	
Services	
Authorisation and cooperative	
development	

Revenue recognised that was

2023	2022
\$ 100,624	\$ 92,362
32,211	101,380
\$ 132,835	\$ 193,742

(C) Unfulfilled long-term contracts

Aggregate amount of the transaction price allocated to long-term technology service contracts, authorisation and cooperative development contracts that are partially or fully unsatisfied, and all of the milestone payment as at December 31, 2023 amounted to \$ 1,341,954. The management expects to recognise the amount in the future.

C. Details of authorisation and cooperative development revenue arising from providing drug development, commercialization service and authorising intellectual property rights of pharmaceutical products to the pharmaceutical factory are as follows:

In April 2019, the Company entered into an authorisation and cooperative development contract of EG12014 with Sandoz AG. The contract includes up-front payment, milestone payment at each stage and profit-sharing royalty on sales of products in the authorised markets in proportion to the ratios specified in the contract. The contract is mainly for providing the biosimilars development and commercialisation services and authorising intellectual property rights to the customer in the authorised regions. As of December 31, 2023, the Company has received the aforementioned up-front payment and part of the milestone payment in accordance with the contract terms. The revenue of up-front payment and milestone payment achieved is recognised based on the satisfaction percentage during research and development period. If the drug was successfully launched, the supply price based on the supply terms and quantities, and the profit-sharing royalty calculated based on sales could also be collected. For the years ended December 31, 2023 and 2022, the Company recognised the revenue from authorisation and cooperative development contract amounting to \$141,472 and \$261,876, respectively.

The European Medicines Agency and the US Food and Drug Administration accepted the Sandoz AG's application for marketing review in January 2022 and February 2022, respectively. Sandoz AG received a complete response letter from the US Food and Drug Administration in December 2022. Within the complete response letter (CRL):

- A. There were no clinical or safety or biosimilarity deficiencies cited in the CRL.
- B. The CRL cites certain drug product deficiencies related to the manufacturing facility identified by the agency during a pre-license inspection of the site.

In January 2023, the Company received an EIR (Establishment Inspection Report) from the US Food and Drug Administration, which indicated that the Company's Zhubei plant had passed the US FDA's pre-marketing drug inspection. Sandoz is in close contact with the FDA to meet the satisfactory resolution of the FDA observations in a timely manner and plans a BLA resubmission in due course.

- D.In April 2023, the Company received a letter from the Taiwan Food and Drug Administration (TFDA) to which indicated that the Company had obtained the domestic active pharmaceutical ingredients "EG12014 Trastuzumab" license and a drug master file number. In September 2023, the Company received the approval by the National Health Insurance Administration with respect to its enrollment in the reimbursement system which became effective from October 1, 2023.
- E.On November 16, 2023, Sandoz AG received the marketing authorisation from Committee for Medicinal Products for Human Use (CHMP) for the trastuzumab biosimilar, EG12014, which was licensed by the Company for sale.

(21) Interest income

Interest income from bank deposits
Interest income from financial
assets measured at amortised cost

Year ended	Decemb	per 31
2023		2022
\$ 106,236	\$	57,862
28,235		1,722
\$ 134,471	\$	59,584

(22) Other income

Year ended December 31 2023 2022 \$ 4,712 \$ 37,214 Government grant revenues Dividend income 475 \$ Other income 252 430 \$ 5,439 37,644

The Company received a grant for the 'Breast Cancer Targeted Antibody similar to EG12014 Trastuzumab Biosimilar phase III clinical trial program' from Ministry of Economic Affairs (MOEA). The program execution period is from November 1, 2019 to June 30, 2023 and the limit on total grant amounted to \$80,000. For the years ended December 31, 2023 and 2022, the Company recognised government grants revenue of \$4,591 and \$36,994, respectively.

(23) Other gains and losses

	Year ended December 31				
		2023	2022		
Foreign exchange (losses) gains	(\$	9,431) \$	126,788		
(Losses) gains on financial assets at fair value through profit or loss	(1,122)	2,863		
Loss on lease modification	(413) (709)		
Miscellaneous disbursements	(214) (24)		
Loss on redemption of convertible					
bonds		- (3)		
	(\$	11,180) \$	128,915		

(24) Finance costs

	Year ended December 31					
		2023		2022		
Interest expense on lease liabilities	\$	8,096	\$	8,204		
Interest expense on bank		2,300		1,205		
borrowings						
Other interest expense		7		226		
	\$	10,403	\$	9,635		

(25) Employee benefits, depreciation and amortisation expenses

Function	Year end	ed December	31, 2023	Year ended December 31, 2022			
	Classified as	Classified as		Classified as Classified as			
	Operating	Operating		Operating	Operating		
Nature	Costs	Expenses	Total	Costs	Expenses	Total	
Employee benefit expense							
Wages and salaries	\$ 121,946	\$ 224,178	\$ 346,124	\$ 147,873	\$ 185,966	\$ 333,839	
Share based payment	70,628	110,272	180,900	60,275	95,037	155,312	
Labour and health insurance fees	14,408	20,005	34,413	13,771	16,809	30,580	
Pension costs	9,257	8,671	17,928	7,427	8,624	16,051	
Directors' remuneration	-	4,125	4,125	-	3,948	3,948	
Other personnel expenses	6,358	15,092	21,450	5,557	13,245	18,802	
Depreciation expense	114,746	112,462	227,208	99,536	88,451	187,987	
Amortisation expense	5,840	5,456	11,296	10,456	5,728	16,184	

- A. In accordance with to the Articles of Incorporation of the Company, a ratio of distributable profit of the current year, after covering accumulated losses, shall be distributed as employees' compensation and directors' and supervisors' remuneration. The ratio shall be 1% to 5% for employees' compensation and shall not be higher than 3% for directors' remuneration.
- B. No employees' compensation and directors' remuneration was accrued due to the net loss incurred for the years ended December 31, 2023 and 2022.
- C. Information about employees' compensation and directors' and supervisors' remuneration of the Company as resolved at the meeting of Board of Directors and resolved at the shareholders' meeting will be posted in the "Market Observation Post System" at the website of the Taiwan Stock Exchange.

(26) Income taxes

A. Income tax expense

(a) Components of income tax expense:

	Year ended December 31					
		2023		2022		
Current tax:						
Current tax on profits for the year	\$		\$			
Deferred tax:						
Origination and reversal						
of temporary differences		465		338		
Income tax expense	\$	465	\$	338		

(b) The income tax (charge)/credit relating to components of other comprehensive income is as follows:

	Year ended December 31					
		2023	2022			
Currency translation differences	\$	41	\$ -			

B. Reconciliation between income tax expense and accounting profit:

	Year ended December 31					
		2023	2022			
Tax calculated based on loss before tax and statutory tax rate	(\$	182,949) (\$	23,040)			
Expenses disallowed by tax						
regulation		25	49			
Tax exempt income by tax						
regulation		- (3,099)			
Taxable losses not recognised as						
deferred tax assets		172,451	26,428			
Temporary differences not						
recognised as deferred tax assets		10,938				
Income tax expenses	\$	465 \$	338			

C. Amounts of deferred tax assets or liabilities as a result of temporary differences:

	2023							
			Recog	nised in	Recognise	ed in other		
	Jan	uary 1	y 1 profit or loss		comprehensive income		December 31	
 Deferred tax assets: Share of profit (loss) of associates and subsidiaries accounted for using the equity method, 								
net differences	\$	874	\$	465	\$	-	\$	1,339
Currency translation differences		-		-		41		41
Income tax expense	\$	874	\$	465	\$	41	\$	1,380
					2022			
			Recog	nised in	Recognise	ed in other		
	Jan	uary 1	profit	or loss	comprehens	sive income	Dece	ember 31
-Deferred tax assets: Share of profit (loss) of associates and subsidiaries accounted for using the equity method,	ď	527	¢	220	¢.		¢.	974
net differences	\$	536	\$	338	\$		\$	874

D. Details of the amount the Company is entitled as investment tax credit and unrecognised deferred tax assets are as follows:

			Unrec	cognised deferred	
Qualifying items	Unus	sed tax credits		tax assets	Expiry year
Research and development	\$	960,900	\$	960,900	Note
Machinery and equipment		8,844		8,844	Note

December 31, 2023

December 31, 2022

Unrecognised deferred

Qualifying items
Unused tax credits
tax assets
Expiry year

Research and development
\$887,160 \$887,160 Note

Note: The Company was entitled to the incentives conferred under the Biotech and New Pharmaceutical Development Act following the Company's incorporation as a biotech pharmaceutical company pursuant to the Letter No. Jing-Shou-Gong-Zi-10920401340 issued by the MOEA on February 3, 2020. Subsequently, the MOEA approved the Company's additional items pursuant to the Letter No. Jing-Shou-Gong-Zi-11120426560 on August 29, 2022. The incentive measures are valid for five years beginning on the next date of the issuance of MOEA's Letter. The investment tax credit can be first used to offset expenditure on research and development and staff training when there is taxable business income. Any unused tax credit is available for the following four years. Additionally, the investment tax credit can be first used to offset expenditure on machinery, equipment, or systems when there is taxable business income. Any unused tax credit is available for the following two years. As of December 31, 2023, the Company has no profit-seeking enterprise income tax.

E. Expiration dates of unused tax losses and amounts of unrecognised deferred tax assets are as follows:

December 31, 2023

December 31, 2023							
	Amount filed/			U	nrecognised		
Year incurred	assessed	_ Un	used amount	defe	rred tax assets	Expiry year	
2014	Amount assessed	\$	131,762	\$	131,762	2024	
2015	Amount assessed		133,257		133,257	2025	
2016	Amount assessed		109,737		109,737	2026	
2017	Amount assessed		163,949		163,949	2027	
2018	Amount assessed		371,827		371,827	2028	
2019	Amount assessed		858,819		858,819	2029	
2020	Amount assessed		1,009,168		1,009,168	2030	
2021	Amount assessed		56,144		56,144	2031	
2022	Amount filed		135,927		135,927	2032	
2023	Amount expected		862,256		862,256	2033	
		\$	3,832,846	\$	3,832,846		

December 31, 2022

	Amount filed/ Unrecognised					
Year incurred	assessed	Un	used amount	defe	erred tax assets	Expiry year
2013	Amount assessed	\$	104,540	\$	104,540	2023
2014	Amount assessed		131,762		131,762	2024
2015	Amount assessed		133,257		133,257	2025
2016	Amount assessed		109,737		109,737	2026
2017	Amount assessed		163,949		163,949	2027
2018	Amount assessed		371,827		371,827	2028
2019	Amount assessed		858,819		858,819	2029
2020	Amount assessed		1,009,168		1,009,168	2030
2021	Amount filed		56,144		56,144	2031
2022	Amount expected		132,140		132,140	2032
		\$	3,071,343	\$	3,071,343	

F. The amounts of deductible temporary differences that were not recognised as deferred tax assets are as follows:

December 31, 2023 December 31, 2022

Deductible temporary differences \$ 64,721 \$ 10,031

G. The Company's income tax returns through 2021 have been assessed and approved by the Tax Authority.

(27) Loss per share

		Yea	r ended December 31,	202	23			
	Weighted average							
			number of ordinary					
			shares outstanding		Loss per share			
	Amoun	t after tax	(shares in thousands)		(in dollars)			
Basic loss per share								
Loss for the year	(<u>\$</u>	915,208)	304,888	<u>(\$</u>	3.00)			
		Year	r ended December 31,	202	22			
			Weighted average					
			number of ordinary					
			shares outstanding		Loss per share			
	Amoun	t after tax	(shares in thousands)		(in dollars)			
Basic loss per share								
Loss for the year	(\$	115,540)	303,258	<u>(\$</u>	0.38)			

Diluted loss per share would not be calculated as the Company had loss for the years ended December 31, 2023 and 2022.

(28) Supplemental cash flow information

A. <u>Investing activities with partial cash payments:</u>

	Year ended December 31				
		2023	2022		
Purchase of property, plant and equipment	\$	702,971	\$	484,268	
Add: Opening balance of other					
payables		158,228		19,508	
Less: Ending balance of other					
payables	(285,960)	(158,228)	
Cash paid during the year	\$	575,239	\$	345,548	
		Year ended 3	December 31 202	22	
Purchase of intangible assets Add: Ending balance of prepayment for intangible	\$	11,498	\$	8,562	
assets (Note) Less: Opening balance of prepayment for intangible		5,209		1,565	
assets (Note)	(1,565)	(1,565)	
Cash paid during the period	\$	15,142	\$	8,562	

Note: Shown as "other non-current assets".

B. Financing activities with no cash flow effects:

	Year ended December 31						
	2023		2022				
Conversion of convertible bonds	\$	<u>-</u> \$	126,375				

(29) Changes in liabilities from financing activities

2022	
70173	
2023	

		Long-term orrowings	Le	ase lia	ıbilitie		arantee ts received		fina	ities from ancing ies-gross
At January 1	\$	120,460	\$	3	38,58	34 \$	294	\$		459,338
Changes in cash flow from financing activities		-	(29,30	07) (288)	(29,595)
Changes in right-of- use assets		-			35,01	17	-			35,017
Changes in other non-cash items		-			41	13				413
At December 31	\$	120,460	\$	3	344,70	<u>)7</u> \$	6	\$		465,173
						2022				
						Bonds				
					ŗ	oayable			Ι	Liabilities
					(iı	ncluding	Guarant	ee	fror	m financing
	Lo	ng-term			(current	deposit	S	a	ctivities-
	bor	owings Le	ase lia	bilitie	s p	ortion)	receive	d		gross
At January 1	\$	- \$	300	5,765	\$	127,070	\$	-	\$	433,835
Changes in cash flow from financing activities		120,460 (23	3,657)) (200)	2	294		96,897
Changes in right-of- use assets		_	5,	1,767		_		_		54,767
Changes in other		-	ر.	+,/0/		_		-		J 4 ,/0/
non-cash items		<u> </u>		709	(126,870)		_	(126,161)
At December 31	\$	120,460 \$	338	3,584	\$	-	\$ 2	294	\$	459,338

7. Related Party Transactions

(1) Parent and ultimate controlling party

The Company has no ultimate parent company and ultimate controlling party.

(2) Names of related parties and relationship

Names of related parties	Relationship with the Group
EirGenix Europe GmbH	Subsidiary
FORMOSA LABORATORIES, INC.	Other related party
Development Center for Biotechnology (DCB)(Note 1)	<i>"</i>
FORMOSA PHARMACEUTICALS, INC.	<i>"</i>
TFBS Bioscience Inc.(Note 2)	<i>"</i>
Forward BioT Venture Capital	//

Note 1: DCB's term as a director expired after re-election of directors at the Company's shareholders' meeting on June 10, 2022. Accordingly, it became a non-related party. (The transaction amounts for the period from January 1, 2022 to June 10, 2022 are disclosed in the financial statements.)

Note 2: The Company was elected as one of the directors of TFBS Bioscience, Inc. on June 8, 2022. Accordingly, the Company became a related party. (The transaction amount for the year ended December 31, 2023 and period from June 8, 2022 to December 31, 2022 are disclosed in the financial statements.)

(3) Significant related party transactions

A. Operating revenue

	Year ended December 31					
		2023		2022		
Sales of goods:						
Other related parties	\$	972	\$	12,850		
Sales of services:						
Other related parties		9,477		5,622		
	\$	10,449	\$	18,472		

- (a) No similar transaction can be compared with for the sales of service. Prices and terms are determined based on mutual agreements.
- (b) On December 31, 2023 and 2022, the Company has recognised the revenue-related contract assets amounting to \$1,994 and \$744, and contract liabilities amounting to \$372 and \$620, respectively.

B. Service expense (shown as 'sales and marketing expense' and 'research and development expense')

	Year ended December 31					
		2023	2022			
Subsidiary-EirGenix Europe GmbH	\$	72,270	\$	66,663		
Other related parties		12,377		17,651		
	\$	84,647	\$	84,314		

It refers to service expense that the Company commissioned its subsidiaries and other related parties to perform biopharmaceutical research and development as well as business development. Prices and terms are determined based on mutual agreements.

C. <u>Testing expense</u> (shown as 'operating costs')

	Year ended December 31					
		2023		2022		
Other related parties-TFBS Bioscience Inc.	\$	7,517	\$	15,152		
Other related parties		2,627		2,460		
-	\$	10,144	\$	17,612		

D. Other expenses (shown as 'administrative expenses')

	Year ended December 31						
	2023		2022				
Other related parties-DCB	\$	- \$	2,463				

It refers to repair and maintenance fees, based on the price specified in the contract as mutually agreed, allocated from leasing plant and lab from DCB, and the expense shall be paid before the 25th day of the first month of each quarter as specified in the contract.

E. Receivables from related parties:

	Decemb	er 31, 2023	December 31, 2022			
Accounts receivable:						
Other related parties	\$	2,636	\$	_		

F. Payables to related parties

	December 31, 2023		December 31, 2022		
Other payables:					
Subsidiary	\$	12,758	\$	8,665	
Other related parties		7,993		7,732	
	\$	20,751	\$	16,397	

G. Property transactions

(a) Acquisition of property, plant and equipment:

	 Year ended December 31				
	 2023	2022			
Other related parties	\$ 645	\$			

(b) Acquisition of financial assets:

		Year ended December 31				
			2023		2022	
	Accounts	Co	onsiderat	ion	(Consideration
Other related party- FORMOSA LABORATORIES, INC.	Non-current financial assets at fair value through profit or loss	\$		_	\$	58,390
TFBS Bioscience Inc.	Non-current financial assets at fair value through other comprehensive income	\$			\$	40,627
Other related party- Forward BioT Venture Capital	Non-current prepayments for investments	\$	1	5,000	\$	<u>-</u>

Refer to Note 6(2) B. and Note 6(7) B. for details of the transactions relating to the Company's acquisition of assets from related parties.

H. <u>Lease transactions - lessee</u>

(a) The Company leases plant, laboratory, instrument and equipment from DCB. Rental contract period is expected to be 20 years with initial rental period of 5 years plus the extension options. Rents are paid before the 25th day of the first month of each quarter.

(b) Right-of-use assets

As of December 31, 2023 and 2022, DCB was no longer a related party, and therefore the carrying amount of its related right-of-use assets was not disclosed.

	Year ended	Year ended December 31		
	2	022		
	Depreciat	ion expense		
Land	\$	3,061		
Buildings		2,279		
Machinery and equipment		1,022		
	\$	6,362		

(c) Lease liabilities

i. Outstanding balance

As of December 31, 2023 and 2022, DCB was no longer a related party, and therefore the carrying amount of its related lease liabilities was not disclosed.

ii.Interest expense

	Year ende	d December 31
		2022
Other related party -		
DCB	\$	2,185

(d) Rent expense (shown as 'operating cost' and 'operating expenses')

	Year ended D	ecember 31
	202	2
Other related party -		
DCB	\$	505

(4) Key management compensation

		er 31		
		2023		2022
Salaries and other short-term employee benefits	\$	25,092	\$	24,790
Post-employment benefits		216		309
Share based payment		30,125		30,244
	\$	55,433	\$	55,343

8. Pledged Assets

The Company's assets pledged as collateral are as follows:

	Bo		
Pledged asset	December 31, 2023	December 31, 2022	Purpose
Pledged time deposits			
(shown as non-current			
financial assets at			
amortised cost)	\$ 8,79	0 \$ 8,671	Note 1
Guarantee deposits paid			
(shown as other non-current			
assets)	\$ 8,79	5 \$ 65,048	Note 2
Property, plant and			
equipment	\$ 1,551,63	3 \$ 1,158,399	Note 3
Pledged government bonds			
(shown as non-current financial			
assets at amortised cost)	\$ 31,93	0 \$ -	Note 4

- Note 1: It refers to guarantee for lease of land.
- Note 2: It refers to deposits for research commissioned contract, equipment and office, guarantee for gas meter as well as certificates of deposit for customs post-release duty payment.
- Note 3: In April 2022, the Company terminated the syndicated loan agreement with 6 financial institutions including Taiwan Business Bank. However, the guarantee for the pledged buildings has not yet been released.

Note 4: It refers to guarantee for investment.

9. Significant Contingent Liabilities and Unrecognised Contract Commitments

(1) Contingencies

None.

(2) Commitments

- A. As of December 31, 2023 and 2022, the remaining payments contracted for research commissioned contracts at the balance sheet date but not yet incurred amounted to \$59,156 and \$105,637, respectively.
- B. As of December 31, 2023 and 2022, the remaining payments contracted for equipment purchase and plant design at the balance sheet date but not yet incurred amounted to \$876,590 and \$815,285, respectively.

C. The Company entered into a long-term consignment contract with a supplier to ensure the future supply of goods and pay the guarantee amounting to \$30,000. As of December 31, 2023, the aforementioned amount was shown as other non-current assets, others of \$30,000.

10. Significant Disaster Loss

None.

11. Significant Events after the Balance Sheet Date

- A. In January 2024, in response to the expansion of Zhubei production line, the Company entered into a construction contract with an engineering company for a total amount of \$1,373,643 (including tax).
- B. The Board of Directors during its meeting on March 8, 2024 resolved to issue the 1st restricted stocks to employees amounting to 1,400 thousand shares with no subscription price, which has not yet been resolved by the shareholders as of March 8, 2024.
- C. The Board of Directors during its meeting on March 8, 2024 resolved to raise additional cash through private placement. The maximum number of shares to be issued through the private placement is 30,000 thousand, and the private placement can be completed in three installments after the authorisation by shareholders. However, the issuance has not yet been resolved at the shareholders' meeting as of March 8, 2024.
- D. The Company's self-developed product, EG1206A, will begin phase III clinical trial according to the research and development schedule. On March 8, 2024, the Board of Directors resolved to authorise the chairman to enter into a commissioned research project for the phase III clinical trial with a CRO and other companies.

12. Others

(1) Capital management

The Company's objectives when managing capital are to safeguard the Company's ability to continue as a going concern in order to provide returns for shareholders and to maintain an optimal capital structure to reduce the cost of capital. In order to maintain or adjust the capital structure, the Company may adjust the amount of dividends paid to shareholders, return capital to shareholders, issue new shares or sell assets to reduce debt.

(2) Financial instruments

A. Financial instruments by category

	 December 31, 2023	December 31, 2022
Financial assets		
Financial assets at fair value through		
profit or loss		
Financial assets mandatorily measured		
at fair value through profit or loss		
on initial recognition	\$ 80,298	\$ 61,420
Financial assets at fair value through other comprehensive income		
Designation of equity instrument	\$ 325,887	\$ 279,325
Financial assets at amortised cost		
Cash and cash equivalents	\$ 5,048,604	\$ 6,108,994
Financial assets at amortised cost	540,720	1,041,123
Notes receivable	19	-
Accounts receivable	253,390	32,782
Accounts receivable - related parties	2,636	-
Other receivables	20,497	24,944
Guarantee deposits paid (shown as other non-current assets)	8,795	 65,048
,	\$ 5,874,661	\$ 7,272,891
	December 31, 2023	December 31, 2022
Financial liabilities		
Financial liabilities at amortised cost		
Accounts payable	\$ 79,556	\$ 134,607
Other payables	519,762	384,682
Other payables-related parties	20,751	16,397
Long-term borrowings	120,460	120,460
Guarantee deposits received		
(shown as other non-current		
liabilities)	 6	 294
	\$ 740,535	\$ 656,440
Lease liability	\$ 344,707	\$ 338,584

B. Financial risk management policies

(a) The Company's activities expose it to a variety of financial risks: market risk (including foreign exchange risk, interest rate risk and price risk), credit risk and liquidity risk. The Company's overall risk management programme focuses on the unpredictability of financial markets and seeks to minimise potential adverse effects on the Company's financial position and financial performance.

- (b) Risk management is carried out by a central treasury department (Company treasury) under policies approved by the Board of Directors. Company treasury identifies, evaluates and hedges financial risks in close cooperation with the Company's operating units. The Board provides written principles for overall risk management, as well as written policies covering specific areas and matters, such as foreign exchange risk, interest rate risk, credit risk, use of derivative financial instruments and non-derivative financial instruments, and investment of excess liquidity.
- C. Significant financial risks and degrees of financial risks
 - (a) Market risk

i. Exchange rate risk

- (i) The Company operates internationally and is exposed to exchange risk arising from various currency exposures, primarily with respect to the USD, GBP, EUR, and JPY. Foreign exchange rate risk arises from future commercial transactions and recognised assets and liabilities.
- (ii) Management has set up a policy to require Company units to manage their foreign exchange risk against their functional currency. The units are required to hedge their entire foreign exchange risk exposure with the Company treasury.
- (iii) The Company's businesses involve some non-functional currency operations (the Company's functional currency: NTD). The information on assets and liabilities denominated in foreign currencies whose values would be materially affected by the exchange rate fluctuations is as follows:

		December 31, 2023					
	a	eign currency mount (In housands)	Exchange rate		Book value (NTD)		
Financial assets							
Monetary items							
USD:NTD	\$	53,756	30.71	\$	1,650,847		
EUR:NTD		363	33.98		12,335		
GBP:NTD		67	39.15		2,623		
JPY:NTD		10,751	0.22		2,365		
Financial liabilities							
Monetary items							
USD:NTD	\$	932	30.71	\$	28,622		
EUR:NTD		546	33.98		18,553		
GBP:NTD		17	39.15		666		
JPY:NTD		57,505	0.22		12,651		

December	$^{\circ}$	2022
I Jecember	- K I	7077
	21	. 4044

	ar	ign currency nount (In ousands)	Exchange rate	Book value (NTD)
Financial assets				
Monetary items				
USD:NTD	\$	44,053	30.71	\$ 1,352,868
EUR:NTD		191	32.72	6,250
GBP:NTD		110	37.09	4,080
JPY:NTD		8,476	0.23	1,949
Financial liabilities				
Monetary items				
USD:NTD	\$	708	30.71	\$ 21,743
EUR:NTD		1,048	32.72	34,291
GBP:NTD		30	37.09	1,113

(iv) Analysis of foreign currency market risk arising from significant foreign exchange variation:

	Year ended December 31, 2023				
	Sensitivity analysis				
				Effect	on other
	Degree of	Effect on profit or loss		comprehensive	
	variation			income	
Financial assets					
Monetary items					
USD:NTD	1%	\$	16,508	\$	-
EUR:NTD	1%		60		63
GBP:NTD	1%		26		-
JPY:NTD	1%		24		-
Financial liabilities					
Monetary items					
USD:NTD	1%	\$	286	\$	-
EUR:NTD	1%		186		-
GBP:NTD	1%		7		-
JPY:NTD	1%		127		-

Year ended December 31, 2022 Sensitivity analysis Effect on other Degree of Effect on profit or comprehensive variation loss income Financial assets Monetary items \$ **USD:NTD** 1% 13,529 \$ 55 **EUR:NTD** 1% **GBP:NTD** 1% 41 19 JPY:NTD 1% Financial liabilities Monetary items 1% \$ \$ **USD:NTD** 217 **EUR:NTD** 1% 343 **GBP:NTD** 1% 11

(v) The total exchange gains (losses), including realised and unrealised, arising from significant foreign exchange variation on the monetary items held by the Company for the years ended December 31, 2023 and 2022, amounted to (\$9,431) and \$126,788, respectively.

ii. Price risk

- (i) The Company's equity securities, which are exposed to price risk, are the held financial assets at fair value through other comprehensive income. To manage its price risk arising from investments in equity securities, the Company diversifies its portfolio. Diversification of the portfolio is done in accordance with the limits set by the Company.
- (ii)The Company's investments comprise equity securities. The prices of equity securities would change due to the change of the future value of investee companies. If the prices of these equity securities had increased/decreased by 1% with all other variables held constant, post-tax profit for the years ended December 31, 2023 and 2022 would have increased/decreased by \$189 and \$0, respectively, as a result of gains/losses on equity securities classified as at fair value through profit or loss. Other components of equity would have increased/decreased by \$3,259 and \$2,793, respectively, as a result of other comprehensive income classified as equity investment at fair value through other comprehensive income.

iii. Cash flow and fair value interest rate risk

The Company's main interest rate risk arises from long-term borrowings with variable rates. Borrowings issued at variable rates expose the Company to cash flow interest rate risk. During 2023 and 2022, the Company's borrowings at variable rate were mainly denominated in New Taiwan dollars.

(b) Credit risk

- i. Credit risk refers to the risk of financial loss to the Company arising from default by the clients or counterparties of financial instruments on the contract obligations. The main factor is that counterparties could not repay in full the accounts receivable based on the agreed terms.
- ii. According to the Company's credit policy, each local entity in the Company is responsible for managing and analysing the credit risk for each of their new clients before standard payment and delivery terms and conditions are offered. Internal risk control assesses the credit quality of the customers, taking into account their financial position, past experience and other factors. Individual risk limits are set based on internal or external ratings in accordance with limits set by the Board of Directors. The utilisation of credit limits is regularly monitored.
- iii. The Company adopts the assumption under IFRS 9, that is, the default occurs when the contract payments are past due over 90 days.
- iv. The Company adopts the following assumption under IFRS 9 to assess whether there has been a significant increase in credit risk on that instrument since initial recognition:

 If the contract payments were past due over 30 days based on the terms, there has been a significant increase in credit risk on that instrument since initial recognition.
- v. The following indicators are used to determine whether the credit impairment of debt instruments has occurred:
 - (i) It becomes probable that the issuer will enter bankruptcy or other financial reorganisation due to their financial difficulties;
 - (ii) The disappearance of an active market for that financial asset because of financial difficulties;
 - (iii) Default or delinquency in interest or principal repayments;
 - (iv) Adverse changes in national or regional economic conditions that are expected to cause a default.
- vi. The Company classifies customers' accounts receivable, and contract assets in accordance with customer types. The Company applies the modified approach using individual provision to estimate expected credit loss.

vii. The Company's notes and accounts receivable were generated from the customers who have optimal credit rating, and the expected credit loss rate is 0.03% after using the forecastability of future boom. As of December 31, 2023 and 2022, the carrying amount of notes and accounts receivable (including related parties) amounted to \$256,342 and \$33,079, respectively. Although some accounts receivable were past due over 90 days, the expected credit risk is insignificant based on individual assessment, thus, loss allowance amounted to \$297 and \$297, respectively. The counterparties of time deposits over 3 months are financial institutions all with high credit quality and the expected credit risk is insignificant based on the assessment, thus, no loss allowance was recognised.

viii. Movements in loss allowance for accounts receivable are as follows:

	Year ended December 31							
		2023		2022				
At January 1	\$	297	\$		689			
Reversal of impairment loss			(392)			
At December 31	\$	297	\$		297			

(c) Liquidity risk

- i. Cash flow forecasting is performed in the operating units of the Company and aggregated by Company treasury. Company treasury monitors rolling forecasts of the Company's liquidity requirements to ensure it has sufficient cash to meet operational needs.
- ii. Surplus cash held by the operating entities over and above balance required for working capital management are transferred to the Company treasury. Company treasury invests surplus cash in interest bearing current accounts and time deposits, choosing instruments with appropriate maturities or sufficient liquidity to provide sufficient headroom as determined by the abovementioned forecasts.
- iii. The Company has the following undrawn borrowing facilities:

	Dece	ember 31, 2023	Dece	ember 31, 2022
Floating rate:				
Expiring within one year	\$	1,410,000	\$	1,020,000
Expiring beyond one year		593,540		593,540
	\$	2,003,540	\$	1,613,540

iv. The table below analyses the Company's non-derivative financial liabilities into relevant maturity groupings based on the remaining period at the balance sheet date to the contractual maturity date. The amounts disclosed in the table are the contractual undiscounted cash flows.

	Ι	Less than	В	etween 1			
December 31, 2023		1 year	and	d 5 years	Ov	er 5 years	Total
Non-derivative							
financial liabilities							
Accounts payable	\$	79,556	\$	-	\$	-	\$ 79,556
Other payables		519,762		-		-	519,762
Other payables-		20,751		-		-	20,751
related parties							
Lease liabilities		36,273		118,543		245,982	400,798
Long-term borrowings		2,376		123,322		-	125,698
Guarantee deposits received							
(shown as other non- current liabilities)		6		-		-	6
current nuclinies)							
	Ι	Less than	В	etween 1			
December 31, 2022	I	ess than 1 year		etween 1 d 5 years	Ov	ver 5 years	Total
Non-derivative					Ov	ver 5 years	 Total
•					Ov	ver 5 years	Total
Non-derivative	\$				<u>Ov</u>	ver 5 years	\$ Total 134,607
Non-derivative financial liabilities		1 year	and			ver 5 years - -	\$
Non-derivative financial liabilities Accounts payable Other payables Other payables-		1 year 134,607	and			ver 5 years - - -	\$ 134,607
Non-derivative financial liabilities Accounts payable Other payables		1 year 134,607 384,682	and			ver 5 years	\$ 134,607 384,682
Non-derivative financial liabilities Accounts payable Other payables Other payables-		1 year 134,607 384,682	and			ver 5 years 247,968	\$ 134,607 384,682
Non-derivative financial liabilities Accounts payable Other payables Other payables- related parties		1 year 134,607 384,682 16,397	and	d 5 years		- - -	\$ 134,607 384,682 16,397
Non-derivative financial liabilities Accounts payable Other payables Other payables- related parties Lease liabilities		1 year 134,607 384,682 16,397 34,828	and	- - - 115,926		- - -	\$ 134,607 384,682 16,397 398,722
Non-derivative financial liabilities Accounts payable Other payables Other payables- related parties Lease liabilities Long-term borrowings		1 year 134,607 384,682 16,397 34,828	and	- - - 115,926		- - -	\$ 134,607 384,682 16,397 398,722

v. The Company does not expect the timing of occurrence of the cash flows estimated through the maturity date analysis will be significantly earlier, nor expect the actual cash flow amount will be significantly different.

(3) Fair value information

- A. The different levels that the inputs to valuation techniques are used to measure fair value of financial and non-financial instruments have been defined as follows:
 - Level 1: Quoted prices (unadjusted) in active markets for identical assets or liabilities that the entity can access at the measurement date. An active market refers to a market in which transactions for an asset or liability take place with sufficient frequency and volume to provide pricing information on an ongoing basis.
 - Level 2: Inputs other than quoted prices included within Level 1 that are observable for the asset or liability, either directly or indirectly.
 - Level 3: Unobservable inputs for the asset or liability. The fair value of the Company's investment in equity investment without active market and the call options and put options embedded in convertible bonds issued by the Company are included in Level 3.
- B. Financial instruments not measured at fair value
 - Except for financial assets at fair value through profit or loss and financial assets at fair value through other comprehensive income, the carrying amounts of cash and cash equivalents, financial assets at amortised cost, notes receivable, accounts receivable (including related parties), other receivables, guarantee deposits paid (shown as other non-current assets), accounts payable, other payables (including related parties), long-term borrowings, guarantee deposits received (shown as other non-current liabilities) and lease liabilities are approximate to their fair values.
- C. The related information on financial and non-financial instruments measured at fair value by level on the basis of the nature, characteristics and risks of the assets and liabilities are as follows:
 - (a) The related information on the nature of the assets and liabilities is as follows:

December 31, 2023	Level 1		Level	2	 Level 3	 Total
Assets						
Recurring fair value measurements						
Financial assets at fair value through profit or loss						
Profit-sharing investments in						
new drug development	\$	-	\$	-	\$ 61,410	\$ 61,410
Limited partnership						
venture capital		-		-	18,888	18,888
Financial assets at fair value						
through other comprehensive						
income						
Equity securities		_		_	325,887	 325,887
	\$	_	\$		\$ 406,185	\$ 406,185

December 31, 2022	Level 1		Level 2		 Level 3	Total	
Assets							
Recurring fair value measurements							
Financial assets at fair value							
through profit or loss							
Profit-sharing investments in							
new drug development	\$	-	\$	-	\$ 61,420	\$	61,420
Financial assets at fair value							
through other comprehensive							
income							
Equity securities		_			 279,325		279,325
	\$		\$	_	\$ 340,745	<u>\$</u>	340,745

- (b) The fair value of financial instruments measured by using valuation techniques can be referred to current fair value of instruments with similar terms and characteristics in substance, discounted cash flow method or other valuation methods.
- D. The following chart is the movement of Level 3 for the years ended December 31, 2023 and 2022:

				2	023			
			5	sharing				
			inv	estments		Limited		
		Equity	in 1	new drug	pa	rtnership		
	in	struments	dev	elopment	vent	ture capital		Total
At January 1	\$	279,325	\$	61,420	\$	-	\$	340,745
Additions		623		-		20,000		20,623
Gains or losses recognised in profit or loss shown as other gains and losses								
Gains (losses) on valuation		-	(10)	(1,112)	(1,122)
Gains and losses recognised in other comprehensive income								
Gains (losses) on valuation		45,939				_		45,939
At December 31	\$	325,887	\$	61,410	\$	18,888	\$	406,185

	De	rivative		Equity	ne	ew drug		
	inst	ruments	ins	struments	dev	elopment		Total
At January 1	\$	891	\$	11,607	\$	-	\$	12,498
Additions		-		208,627		58,390		267,017
Conversions of convertible bonds	(723)		-		-	(723)
Gains or losses recognised in profit or loss shown as other gains and losses								
Gains (losses) on valuation	(167)		-		3,030		2,863
Gains and losses recognised in other comprehensive income								
Gains (losses) on valuation		-		59,091		-		59,091
Settled during the year	(1)					(1)
At December 31	\$		\$	279,325	\$	61,420	\$	340,745

2022

- E. For the years ended December 31, 2023 and 2022, there was no transfer into or out from Level 3.
- F. Appointed external appraiser is in charge of valuation procedures for fair value measurements being categorised within Level 3, and frequently calibrating valuation model, performing backtesting, updating inputs used to the valuation model and making any other necessary adjustments to the fair value.
- G. The following is the qualitative information of significant unobservable inputs and sensitivity analysis of changes in significant unobservable inputs to valuation model used in Level 3 fair value measurement:

	Fair value at December 31, 2023	Valuation technique	Significant unobservable input	Range (weighted average)	Relationship of inputs to fair value
Non-derivative equity instrument:					
Unlisted shares	\$ 8,236	Price-Book Ratio	Price-to-book ratio	2.66~3.75 (3.67)	The higher the multiple, the higher the fair value
			Discount for lack of marketability	30% (30%)	The higher the discount for lack of marketability, the lower the fair value
Unlisted shares	317,651	Price-Book Ratio	Price-to-book ratio	1.24~2.54 (1.97)	The higher the multiple, the higher the fair value
			Discount for lack of marketability	7.25% (7.25%)	The higher the discount for lack of marketability, the lower the fair value
Profit-sharing investments in new drug development	61,410	Royalty relief method of income approach	Discount rate	24.69%	The higher the discount rate, the lower the fair value
•			Market share	2.0%~5.9%	The higher the market share, the higher the fair
Limited partnership venture capital	18,888	Net asset value	N/A	N/A	N/A
	Fair value at December 31, 2022	Valuation technique	Significant unobservable input	Range (weighted average)	Relationship of inputs to fair value
Non-derivative equity instrument:		teermique		uvorugo	Tur value
Unlisted shares	\$ 6,207	Price-Book Ratio	Price-to-book ratio	1.54-8.46 (3.05)	The higher the multiple, the higher the fair value
			Discount for lack of marketability	30% (30%)	The higher the discount for lack of marketability, the lower the fair value
Unlisted shares	273,118	Price-Book Ratio	Price-to-book ratio	2.01-2.54 (2.19)	The higher the multiple, the higher the fair value
			Discount for lack of marketability	30% (30%)	The higher the discount for lack of marketability, the lower the fair value
Profit-sharing investments in new drug development	61,420	Royalty relief method of income approach	Discount rate	24.58%	The higher the discount rate, the lower the fair value
			Market share	1.0%~5.4%	The higher the market share, the higher the fair

H. The Company has carefully assessed the valuation models and assumptions used to measure fair value. However, use of different valuation models or assumptions may result in different measurement. The following is the effect on profit or loss or on other comprehensive income from financial assets and liabilities categorised within Level 3 if the inputs used to valuation models have

						December	31.	, 2023		onangea.
			Re	cognised	in p	rofit or loss		Recogni comprehe		
			Fa	vourable	U	nfavourable	Fa	avourable	Uı	nfavourable
	Input	Change	C	hange		change	change			change
Financial assets Profit-sharing investments in new drug	Discount rate Market	±5%	\$	3,071	(\$	3,071)	\$	-	\$	-
development Limited partnership venture capital	share N/A	±5%		944	(944)		-		-
Unlisted shares	Price-Book Ratio	±5%		-		-		16,294	(16,294)
	Lack of marketability	±5%	\$	4,015	(\$	4,015)	\$	16,294 32,588	(<u></u> (<u>\$</u>	16,294) 32,588)
						December	31,	, 2022		
			Re	cognised	in p	rofit or loss		Recogni comprehe		
			Fa	vourable	U	nfavourable	Fa	avourable	Uı	nfavourable
	Input	Change		hange		change		change		change
Financial assets Profit-sharing investments in new drug development	Discount rate Market share	±5%	\$	3,071	(\$	3,071)	\$	-	\$	-
Unlisted shares	Price-Book Ratio	±5%		-		-		13,966	(13,966)
	Lack of marketability	±5%		_		_		13,966	(13,966)
	marketaomity		\$	3,071	(\$	3,071)	\$	27,932	(\$	27,932)
					_		_			

13. Supplementary Disclosures

(1) Significant transactions information

- A. Loans to others: None.
- B. Provision of endorsements and guarantees to others: None.
- C. Holding of marketable securities at the end of the period (not including subsidiaries, associates and joint ventures): Please refer to table 1.
- D. Acquisition or sale of the same security with the accumulated cost exceeding NT\$300 million or 20% of the Company's paid-in capital: None.
- E. Acquisition of real estate reaching NT\$300 million or 20% of paid-in capital or more: Please refer to table 2.
- F. Disposal of real estate reaching NT\$300 million or 20% of paid-in capital or more: None.
- G. Purchases or sales of goods from or to related parties reaching NT\$100 million or 20% of paid-in capital or more: None.
- H. Receivables from related parties reaching NT\$100 million or 20% of paid-in capital or more: None.
- I. Trading in derivative instruments undertaken during the reporting periods: None.
- J. Significant inter-company transactions during the reporting periods: Please refer to table 3.

(2) <u>Information on investees</u>

Names, locations and other information of investee companies (not including investees in Mainland China): Please refer to table 4.

(3) <u>Information on investments in Mainland China</u>

None.

(4) Major shareholders information

Major shareholders information: Please refer to table 5.

14. Segment Information

None.

EirGenix Inc. Holding of marketable securities at the end of the period (not including subsidiaries, associates and joint ventures)

December 31, 2023

Table 1

Expressed in thousands of NTD (Except as otherwise indicated)

Ac of	December	21	2023

Securities held by	Marketable securities	Relationship with the securities issuer	General ledger account	Number of shares	Book value	Ownership	Fair value	Footnote
EirGenix Inc.	Oncomatryx Biopharma S.L. common stock	None	Non-current financial assets at fair value through other comprehensive	31,801	\$ 8,236	0.37% \$	8,236	
"	TFBS Bioscience, Inc. common stock	The Company's other related party	п	4,942,455	317,651	14.20%	317,651	
11	Forward BioT Venture Capital equity	"	Non-current financial assets at fair value through profit or loss	-	18,888	5.69%	18,888	
"	93 Central Government Bonds A VI government bonds	None	Non-current financial assets at amortised cost	-	31,930	-	31,930	

EirGenix Inc. and its subsidiaries

Acquisition of real estate reaching NT\$300 million or 20% of paid-in capital or more

Year ended December 31, 2023

Table 2

Expressed in thousands of NTD (Except as otherwise indicated)

If the counterparty is a related party, information as to the last transaction of the real estate is disclosed below:

Real estate acquired by	Real estate	Date of the event	Transaction Amount	Status of payment	Counterparty	Relationship with the counterparty	Original owner who sold the real estate to the counterparty	Relationships	Date of the original Transfer Date	Amount	Basis or reference used in setting the price	Reason for acquisition of real estate and status of the real	Other commitmen
EirGenix Inc.	Factory building	2021/9/30 (Note 4)	\$ 666,130	Based on the terms in the purchase order	China Ecotek Corporation Getinge Group Taiwan Co., Ltd. Jian-Yi Biotech	None	NA	NA	NA	NA	Price comparison and price negotiation	Manufacturing purpose	None
					Co., Ltd. Min-Pin,Chen Architects & Associates								

Note 1: The appraisal result should be presented in the 'Basis or reference used in setting the price' column if the real estate acquired should be appraised pursuant to the regulations.

Note 2: Paid-in capital referred to herein is the paid-in capital of parent company. In the case that shares were issued with no par value or a par value other than NT\$10 per share, the 20 % of paid-in capital shall be replaced by 10% of equity attributable to owners of the parent in the calculation.

Note 3: Date of the event referred to herein is the date of contract signing date, date of payment, date of execution of a trading order, date of title transfer, date of board resolution, or other date that can confirm the counterparty and the monetary amount of the transaction, whichever is earlier.

Note 4: This is the signing date of the first transaction, which is mainly arising from the construction of the factory. The Company continuously signed contracts with relevant suppliers, of which the contract amount has reached \$300,000.

EirGenix Inc.

Significant inter-company transactions during the reporting period Year ended December 31, 2023

Table 3

Expressed in thousands of NTD (Except as otherwise indicated)

							Transaction
Number			Relationship			Transaction	
(Note 1)	Company name	Counterparty	(Note 2)	General ledger account	Amount	terms	Percentage of consolidated total operating revenues or total assets (Note 3)
0	EirGenix Inc.	EirGenix Europe GmbH	(1)	Operating expense	\$ 72,270	Note 4	7.07%
0	EirGenix Inc.	EirGenix Europe GmbH	(1)	Other payables	12,758	Note 4	0.11%

Transaction

- Note 1: The numbers filled in for the transaction company in respect of inter-company transactions are as follows:
 - (1) Parent company is '0'.
 - (2) The subsidiaries are numbered in order starting from '1'.
- Note 2: Relationship between transaction company and counterparty is classified into the following three categories; fill in the number of category each case belongs to (If transactions between parent company and subsidiaries
 - refer to the same transaction, it is not required to disclose twice. For example, if the parent company has already disclosed its transaction with a subsidiary, then the subsidiary is not required to disclose the transaction; for transactions between two subsidiaries, if one of the subsidiaries has disclosed the transaction, then the other is not required to disclose the transaction.):
 - (1) Parent company to subsidiary.
 - (2) Subsidiary to parent company.
 - (3) Subsidiary to subsidiary.
- Note 3: Regarding percentage of transaction amount to consolidated total operating revenues or total assets, it is computed based on period-end balance of transaction to consolidated total assets for balance sheet accounts and based on accumulated transaction amount for the period to consolidated total operating revenues for income statement accounts.
- Note 4: Prices and terms for services are based on the mutual agreement and payments are collected quarterly in advance.
- Note 5: Transactions between the parent company and subsidiaries are eliminated.
- Note 6: Individual amounts less than \$1,000 are not disclosed.

EirGenix Inc. Information on investees Year ended December 31, 2023

Table 4

Expressed in thousands of NTD (Except as otherwise indicated)

												Ne	t profit (loss)			
								Shares he	eld as at Dec	emb	er 31,	of	the investee	Ir	vestment income	
]	Initial invest	mer	nt amount		2023			f	or the year	(lo	oss) recognised by	
				Ва	alance as at	Ва	alance as at						ended	the	e Company for the	
				De	ecember 31,	De	ecember 31,	Number of	Ownership			De	ecember 31,	yea	ar ended December	
Investor	Investee	Location	Main business activities		2023		2022	shares	(%)	Boo	ok value		2023		31, 2023	Footnote
EirGenix Inc.	EirGenix Europe GmbH	Germany	Biopharmaceutical research and development as well as business development	\$	845	\$	845	-	100.00	\$	7,743	\$	2,324	\$	2,324	None
EirGenix Inc.	EirGenix USA Inc.	U.S.A	Biopharmaceutical commissioned development, manufacturing services and consulting		-		-	-	100.00	\$	-	\$	-	\$	-	None

EirGenix Inc.

Major shareholders information

December 31, 2023

Table 5

	Shares		
Name of major shareholders	Number of shares held	Ownership (%)	
Foxconn Technology Co., Ltd	27,500,000	8.98	
Yonglin Capital Holding Co., Ltd.	26,500,000	8.65	
Formosa Laboratories, Inc.	18,552,818	6.06	

EIRGENIX INC. STATEMENT OF CASH AND CASH EQUIVALENTS DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Statement 1

Item	Description	Amount	
Cash on hand and petty cash		\$	61
Demand deposit-			347,726
Deposit of NTD			
Demand deposit-			
Deposit of foreign currency	USD 2,765 thousand Exchange rate 30.71		84,887
	GBP 67 thousand Exchange rate 39.15		2,621
	JPY 10,751 thousand Exchange rate 0.22		2,335
	ERU 177 thousand Exchange rate 33.98		6,009
	Others		3
Time deposits-			3,300,000
Deposit of NTD			
Time deposits-			
Deposit of foreign currency	USD 42,500 thousand Exchange rate 30.71		1,304,962
		\$	5,048,604

The above mentioned time deposits are all maturing in three months with interest rate of 1.38%~6.20%.

<u>EIRGENIX INC.</u> <u>STATEMENT OF FINANCIAL ASSETS MEASURED AT AMORTIZED COST - CURRENT DECEMBER 31, 2023</u>

(Expressed in thousands of New Taiwan dollars)

		Amount of			
		certificate of		Carrying	
Name	Description	deposit	Rates	Amount	Note
Time deposits at Lu Chou Branch-Chang Hwa Commercial Bank, Ltd.	2023/11/06~2024/04/06	1	1.50%	\$ 500,000	

EIRGENIX INC. STATEMENT OF INVENTORIES DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

		Amo				
Item		Cost	Net R	ealizable Value	Note	
Raw material Work in progress	\$	426,217	\$	434,464	Replacement cost is based on the market value Net realisable value is based on	
Finished goods Merchandise inventory		127,143 178,690 235 732,285	\$	542,431 696,837 237 1,673,969	the market value	
Less: Allowance for inventory Valuation losses	(51,648) 680,637				

EIRGENIX INC. STATEMENT OF OPERATING COSTS FOR THE YEAR ENDED DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Item	Description	Total
Raw materials at January 1	\$	377,424
Add: Raw materials purchased during the year		422,982
Surplus of raw materials		4
Less: Raw materials at December 31	(426,217)
Raw materials deficit	(1,014)
Raw materials reclassified to merchandise inventory	(48,966)
Reclassified to research and development expenses	(198,101)
Labor cost used for the year	(66,006)
Raw materials during the year		60,106
Direct labor		2,030
Manufacturing expense		17,963
Manufacturing cost		80,099
Add: Beginning work in progress		281,739
Processing cost		8,984
Less: Ending work in progress	(127,143)
Reclassified to research and development expenses	(40,442)
Cost of finished goods		203,237
Add: Beginning finished goods		98,150
Less: Ending finished goods	(178,690)
Reclassified to research and development expenses	(22,767)
Cost of goods manufactured and sold		99,930
Merchandise inventory at January 1		477
Add: Purchase during the year		326
Raw materials reclassified to merchandise inventory		48,966
Less: Merchandise inventory at December 31	(235)
Reclassified to research and development expenses	(423)
Cost of goods sold for the merchandise inventory		49,111
Other operating costs		602,540
Gain on physical inventory	(4)
Loss on disposal inventory		1,014
Loss on decline in market value of inventory		33,321
Operating costs	\$	785,912

EIRGENIX INC. STATEMENT OF MANUFACTURING EXPENSE FOR THE YEAR ENDED DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Item	Amount	Note
Inspection fees	\$ 5,001	
Depreciation expense	2,961	
Consumables	2,263	
Rent expense	1,744	
Wages and salaries	1,294	
Repairs and maintenance expense	1,004	
Service fees	1,072	
		Each of the account
		was less than 5% of the
Other expenses	2,624	total account balance.
	\$ 17,963	

EIRGENIX INC. STATEMENT OF OTHER OPERATING COSTS FOR THE YEAR ENDED DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Amount		Note
\$	118,622	
	111,785	
	70,628	
	66,006	
	49,630	
	46,279	
	40,426	
	36,391	
		Each of the account
		was less than 5% of the
	62,773	total account balance.
\$	602,540	
		\$ 118,622 111,785 70,628 66,006 49,630 46,279 40,426 36,391

EIRGENIX INC. STATEMENT OF SALES AND MARKETING EXPENSES FOR THE YEAR ENDED DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Item	 Amount	Note
Wages and salaries	\$ 23,521	
Share-based payments	12,555	
Service fees	10,984	
Advertisement expense	6,001	
		Each of the account
		was less than 5% of the
Other expenses	 9,532	total account balance.
	\$ 62,593	

EIRGENIX INC. STATEMENT OF GENERAL AND ADMINISTRATIVE EXPENSES FOR THE YEAR ENDED DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Item	 Amount	Note
Wages and salaries	\$ 82,165	
Share-based payments	55,877	
Depreciation expense	24,063	
Utilities expense	11,975	
Insurance expense	11,665	
		Each of the account
		was less than 5% of the
Other expenses	 68,451	total account balance.
	\$ 254,196	

EIRGENIX INC. STATEMENT OF RESEARCH AND DEVELOPMENT EXPENSES FOR THE YEAR ENDED DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Item	_	Amount	Note	
Consumables	\$	340,477		
Service fees		242,647		
Wages and salaries		118,492		
Depreciation expense		86,999		
			Each of the account	
			was less than 5% of the	
Other expenses		166,731	total account balance.	
	\$	955,346		

EIRGENIX INC.

SUMMARY STATEMENT OF CURRENT PERIOD EMPLOYEE BENEFITS, DEPRECIATION, AND AMORTIZATION EXPENSES BY FUNCTION

FOR THE YEAR ENDED DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Statement 10

Function	Year ended December 31, 2023			Year ended December 31, 2022		
Nature	Classified as Operating Costs	Classified as Operating Expenses	Total	Classified as Operating Costs	Classified as Operating Expenses	Total
Employee Benefit Expense						
Wages and salaries	\$ 121,946	\$ 224,178	\$ 346,124	\$ 147,873	\$ 185,966	\$ 333,839
Share based payment	70,628	110,272	180,900	60,275	95,037	155,312
Labour and health insurance fees	14,408	20,005	34,413	13,771	16,809	30,580
Pension costs	9,257	8,671	17,928	7,427	8,624	16,051
Directors' remuneration	-	4,125	4,125	-	3,948	3,948
Other personnel expenses	6,358	15,092	21,450	5,557	13,245	18,802
Depreciation Expense	114,746	112,462	227,208	99,536	88,451	187,987
Amortisation Expense	5,840	5,456	11,296	10,456	5,728	16,184

Note:

- A. As at December 31, 2023 and 2022, the Company had 400 and 369 employees, both including 9 non-employee directors, respectively.
- B. The Company's stock is listed for trading on the over-the-counter securities exchange and shall additionally disclose the following information:
- (a) Average employee benefit expense were \$1,537 and \$1,541 for the years ended December 31, 2023 and 2022, respectively.
- (b) Average employee salaries were \$1,348 and \$1,359 for the years ended December 31, 2023 and 2022, respectively.
- (c) Adjustment of average employee salaries was (0.81%).
- (d) The Company has no supervisors' remuneration as it set up the audit committees.
- C. The Company' salary and compensation policy:
- (a) Directors

In accordance with the Articles of Incorporation of the Company, the Board of Directors shall propose a ratio not higher than 3% of distributable profit of directors' remuneration, and the distribution shall be approved by a resolution adopted by a majority vote at a meeting of Board of Directors attended by two-thirds of the total number of directors, and shall be reported to the shareholders during their meeting. If the Company has accumulated deficit, earnings should be reserved to cover losses and then be appropriated based on the abovementioned ratios.

The Company did not distribute directors' remuneration in the previous two years. The directors only receive the transportation allowance for the professional practice execution, while the independent directors' additionally receive the fixed remuneration for the professional practice execution. The rest of the director's emolument is the salary expenses of the director who concurrently acts as an employee. The above directors' transportation allowance and the independent directors' remuneration for the professional practice execution have been reviewed by the Company's remuneration committee and approved by the Board of Directors.

EIRGENIX INC.

SUMMARY STATEMENT OF CURRENT PERIOD EMPLOYEE BENEFITS, DEPRECIATION, AND AMORTIZATION EXPENSES BY FUNCTION (Cont.) FOR THE YEAR ENDED DECEMBER 31, 2023

(Expressed in thousands of New Taiwan dollars)

Statement 10

(b) The president and vice presidents

The remuneration for the president and vice presidents is the employeess' compensation distributed from retained earnings. In accordance with the Articles of Incorporation of the Company, a ratio of 1% to 5% of distributable profit of the current year shall be distributed as employees' compensation, which shall be distributed in the form of shares or in cash. Qualification requirements of employees include the employees of subsidiaries of the company meeting certain specific requirements. Distribution of employees' compensation shall be approved by a resolution adopted by a majority vote at a meeting of Board of Directors attended by two-thirds of the total number of directors and shall be reported to the shareholders during their meeting. If the Company has accumulated deficit, earnings should be reserved to cover losses and then be appropriated based on the abovementioned ratios.

The emolument of the president and vice presidents of the Company shall be determined according to the position, contribution to the Company and by reference to the general pay levels of the industry, and shall be reviewed by the remuneration committee and approved by the Board of Directors. The Company did not distribute employee compensation in the previous two years.

(c) Employees:

The salary and compensation package of the Company's employees consists of three parts: basic fixed salary, bonus and welfare; the payment standard: the basic fixed salary is determined based on the time devoted in and the responsibilities undertaken for the position, and by reference to the salary situation in the industry; bonus is awarded based on the achievement of employee and department goals as well as the Company's operation performance; the welfare system stipulates the benefits that employees can enjoy according to law and regulation and takes into account the needs of employees. Employee reward system is based on the individual performance, contribution to the Company and the market value of the position, which are positively associated with the operation performance with the Articles of Incorporation of the Company, a ratio of 1% to 5% of distributable profit of the current year shall be distributed as employees' compensation, which shall be distributed in the form of shares or in cash. Qualification requirements of employees include the employees of subsidiaries of the company meeting certain specific requirements.





EirGenix, Inc.

President: Lee-Cheng Liu

Address: No.101, Lane 169, Kangning St., Xizhi Dist., New Taiepi City

Tel: +886-2-7708-0123 Fax: +886-2-7708-1666