



FORMOSA
LABORATORIES, INC.

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TWSE 4746

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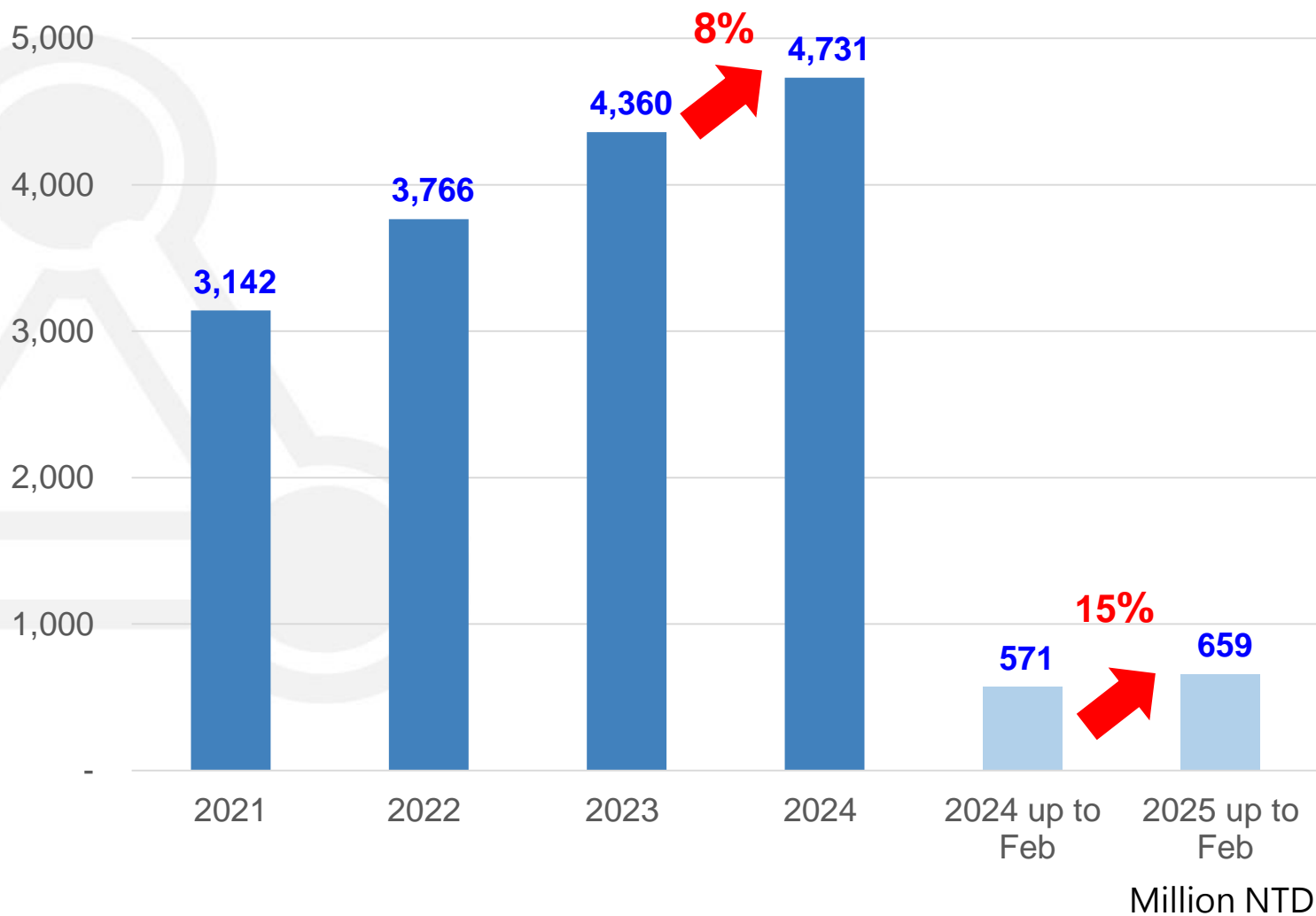
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- 2024 & 2025 up-to-Feb Operation Results
 - Business Strategy
 - Business Update

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Operation Results

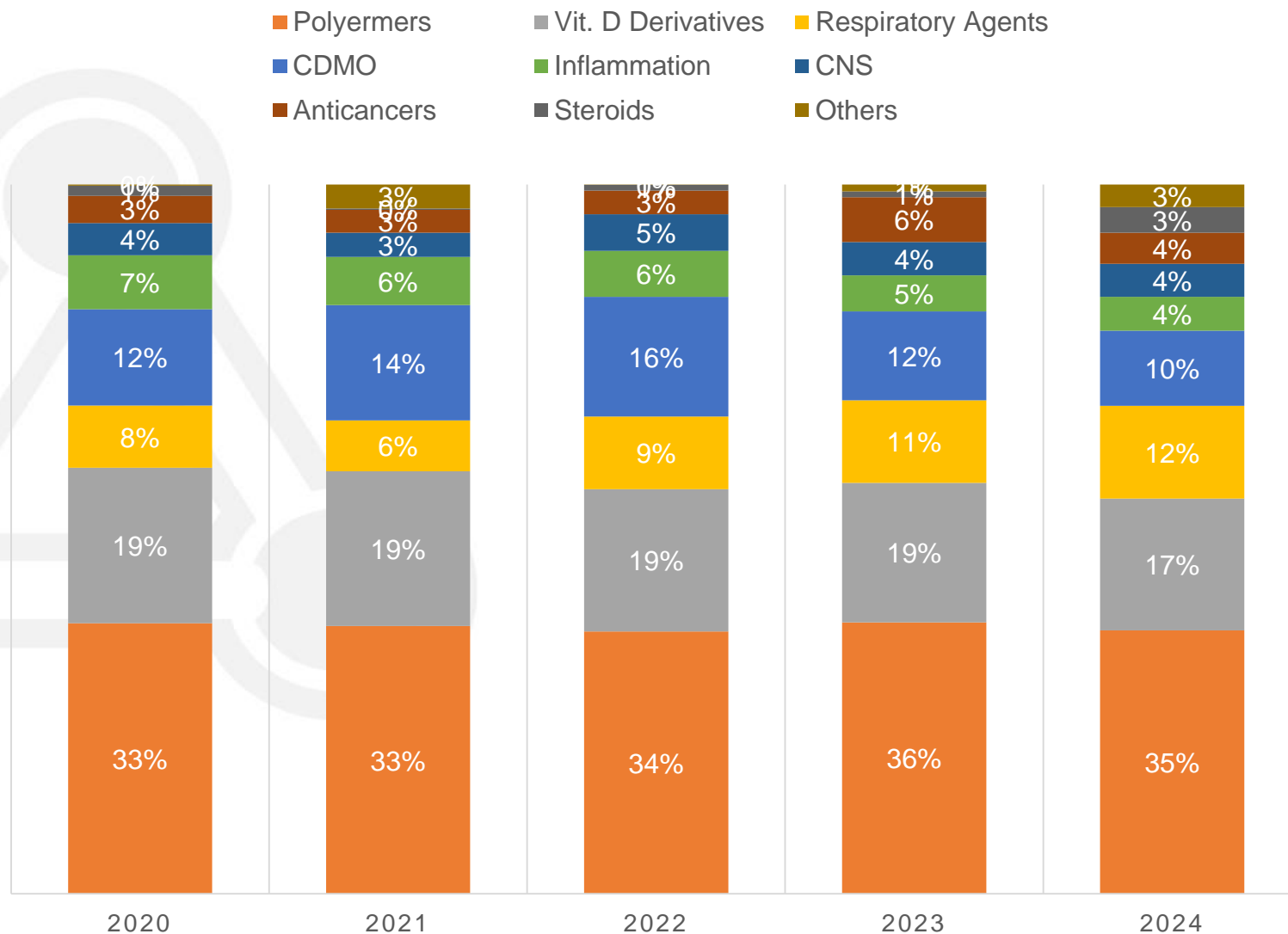
Revenues



Revenues-2024 YoY 8%

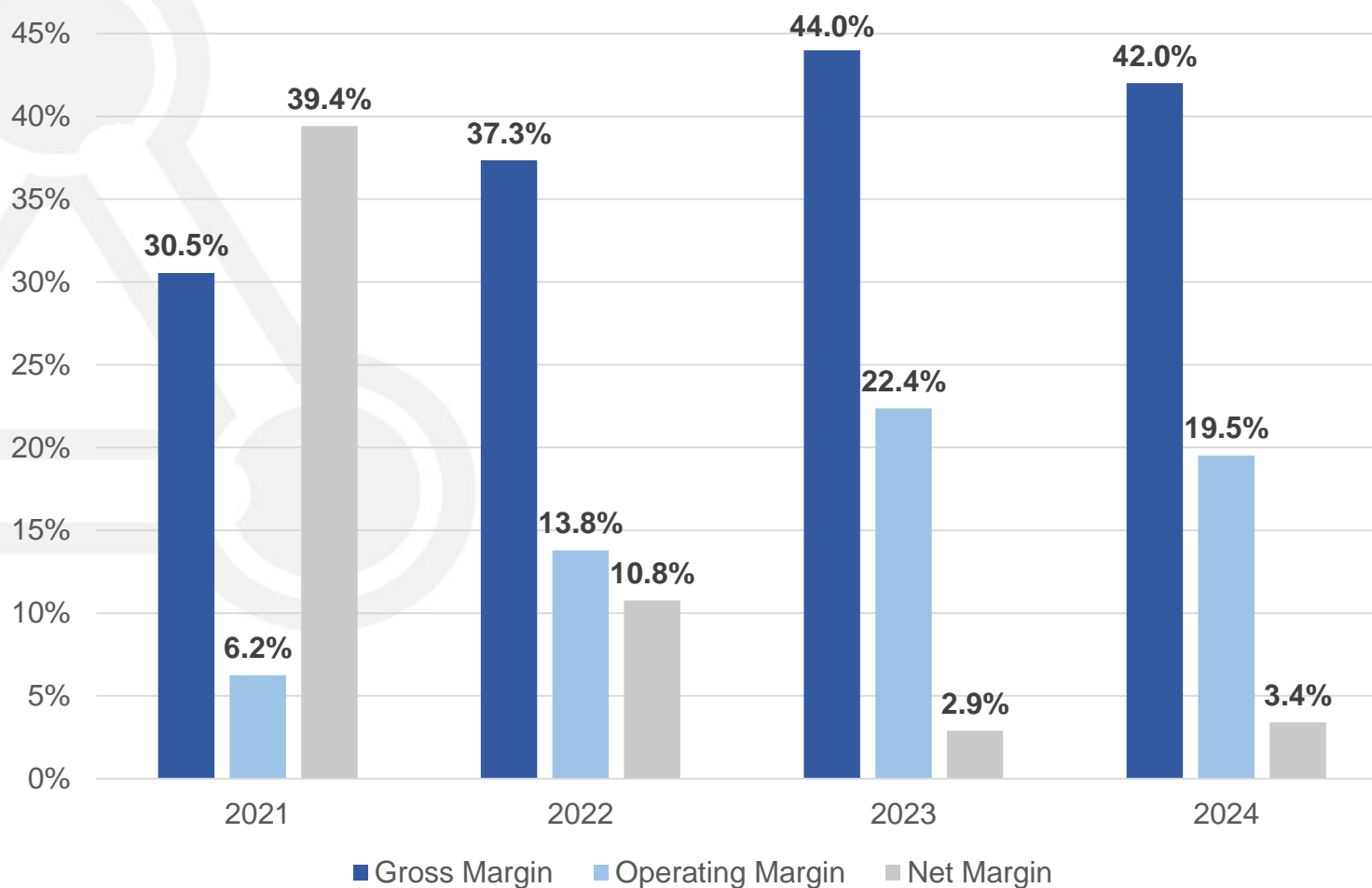
	Million NTD	YoY
Polymers	1,638	4%
Vit. D Derivatives	820	1%
Respiratory Agents	576	20%
CDMO	467	-10%
Inflammation	211	1%
CNS	206	6%
Others	813	40%
Total	4,731	8%

Product Category

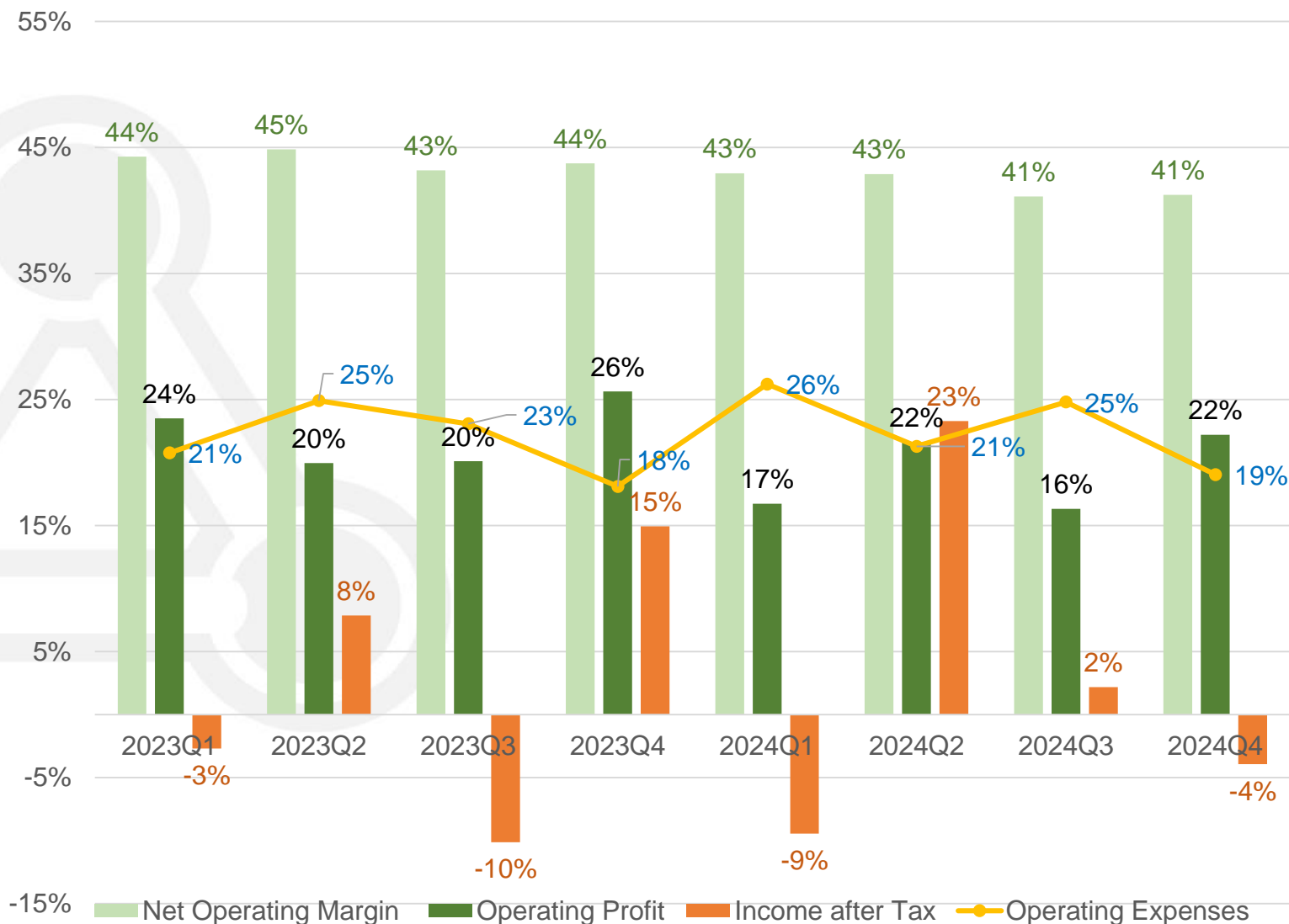


2024 Profitability

EPS NTD 1.31
BV per share NTD 63.90



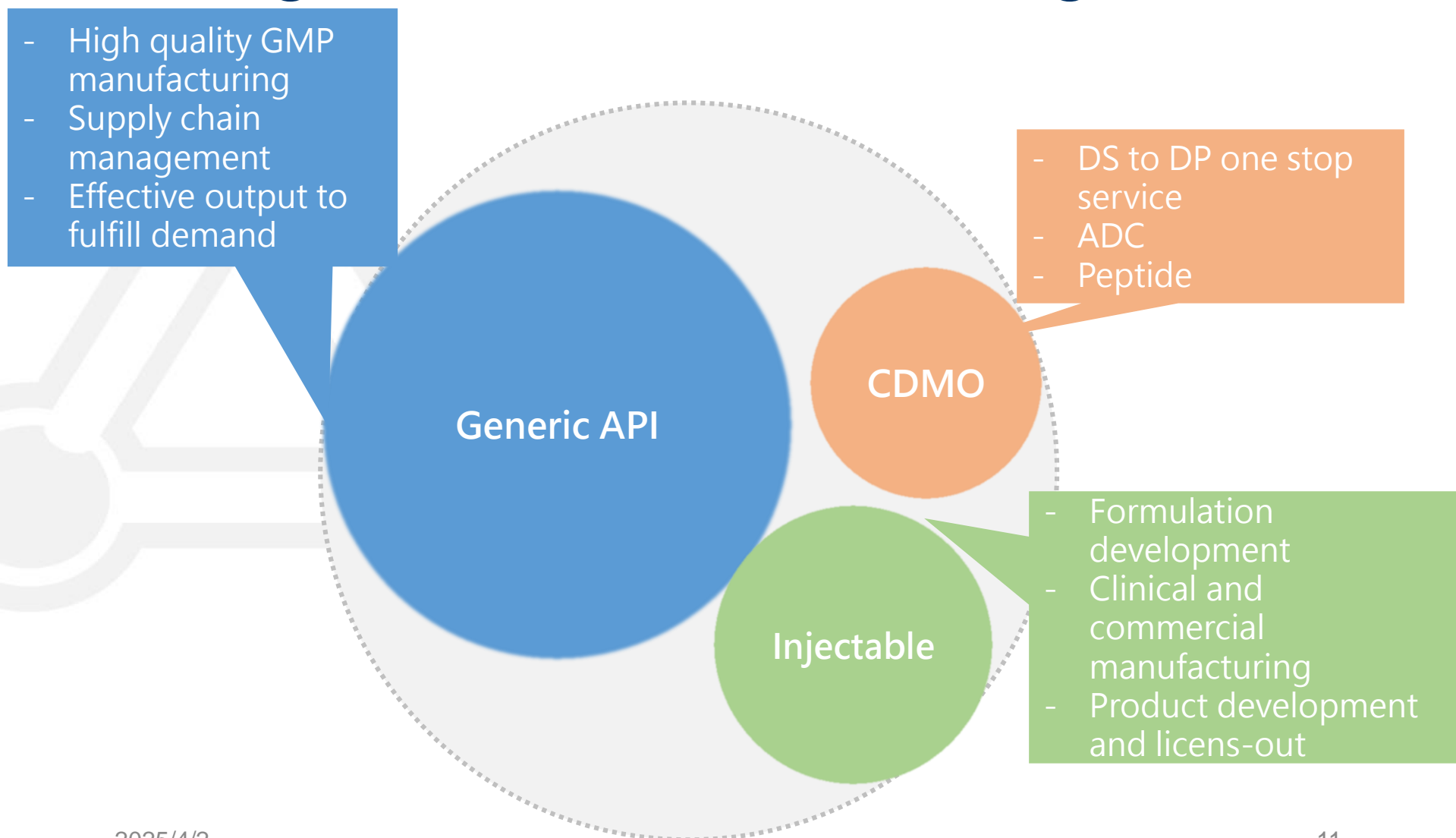
2023~2024Q4 Profitability





Business Strategy

Integrated RD and Manufacturing



Expand Overseas Footprint and Market



GLP-1 manufacturing line

- To expand the Company 's business, the manufacturing lines establishment is ongoing. The expected completion time falls in 2027.
 - Cartridge (annual capacity: 30 million)
 - Assembly line
- GLP-1 products development has started. Licensing-out partner is under discussion.
 - Take semaglutide products as example, the global market size are Ozempic 16.8 billion USD*
Wegovy 8.2 billion USD*



Key Business

Key API Products

- Improve throughput
- Expand manufacture capacity
- Stable supply chain



Highlight

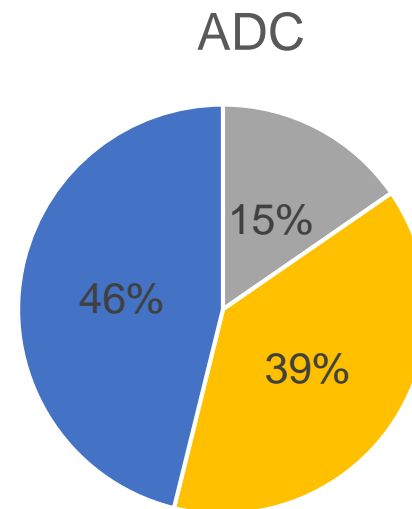
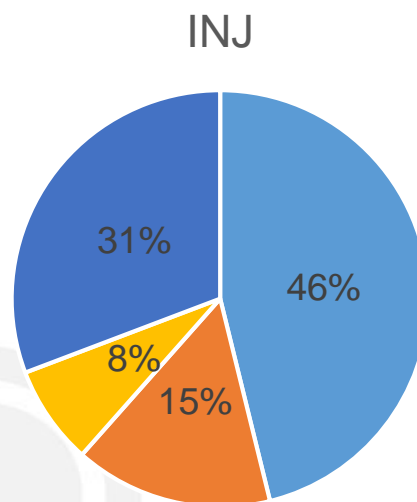
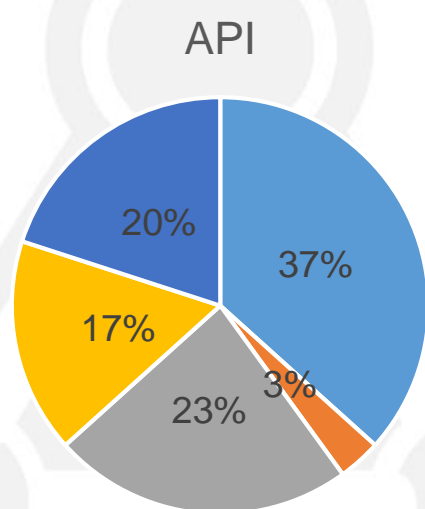
- Polymer products stay strong market share.
- Respiratory API sales keeps growing on revenue.
- Steroid products demand from originator increase and contribute to revenue.

Comprehensive CDMO Service-One Stop Shop



- One-stop service requests increase up to 30% compared to 2023
- Stable CMO supply to originator in Japan, Europe, and US. New projects discussion is completed. Two projects enters early stage development service.
- Synchem-Formosa stably expands North American CDMO business. Four projects are ongoing and few projects are under discussion.
- Multiple projects enter validation and/or commercial manufacturing
- Provide peptide synthesis services

CDMO Clients and Status



- Commercial
- Phase III
- Phase II
- Phase I
- PreIND

Once stop ADC services

Service scope

- Linker-payload, Bioconjugation, and process development
- ADC candidate screening platform
- Scale-up manufacturing. ADC-DS new line completes validation. The new line can support upto commercial scale supply
- Comprehensive analytic services

ADC CDMO highlight

- Clients include US, Europe, and Taiwan
- European client enters Phase I trial. Patient enrollment is ongoing. Scale-up manufacturing is ongoing. New ADC candidate CDMO has signed and started.
- New ADC DS and DP has completed and delivered to client for phase I global clinical trial. New scale-up production is ongoing.
- New clients in Japan, Australia, and Taiwan.
- Few ADC and peptide projects are ongoing.
- Synchem-Formosa is expanding laboratories and early stage bio-conjugation line to fulfill North American CDMO market demand.

Injectable Plant Status

- Authority inspection is ongoing
 - Cytotoxic line
 - TFDA: 2022Q2 completed
 - FDA: passed first FDA on-site GMP inspection held in July 2024.
 - EMA: 2025Q2 on-site inspection.
 - General line
 - TFDA: 2024/08 completed inspection.
Expected to receive certificate by 2025Q2.
 - Feasible for biologic and chemical drug product in liquid and lyophilization products from lab-scale, clinical trial and commercial supply.
 - Work with medical device company as alliance for product development..



Self-development Injectable Products

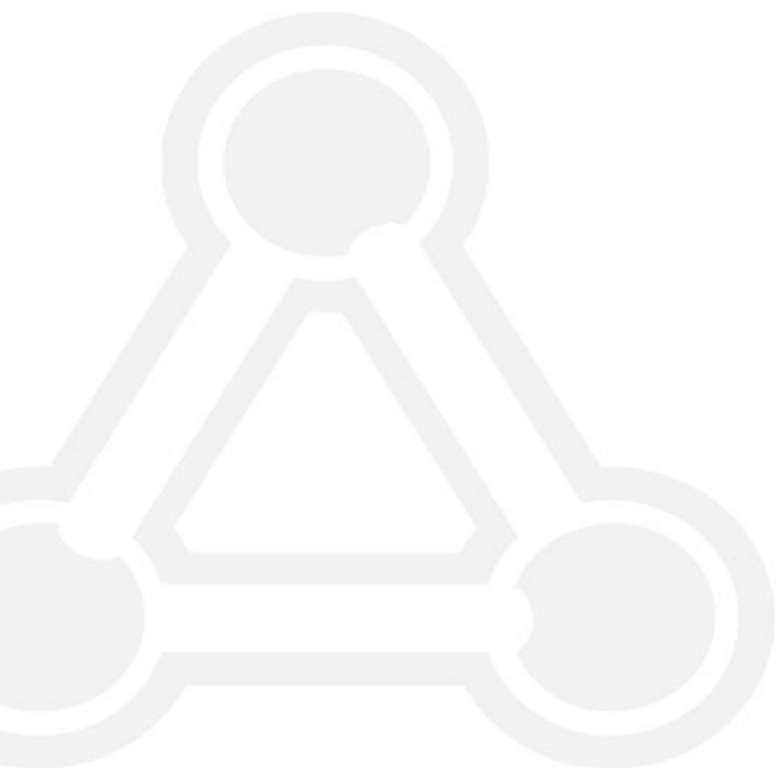
◆ Eribulin Injectable

(Global market size USD 370 million, US Market USD 130 million, TW market NTD 450 million)

Territory	Status
Taiwan	Hospital listing keeps going. Already started deliver. Expected to get first medical center invoice in 2025 1H.
Turkey	Licensing agreement signed
EU	Licensing agreement signed
US	Agreement to be signed 2024/Q4
Emerging market	- Hong Kong and Pakistan agreement signed - Discussion ongoing

◆ Other self-development injectable products are ongoing

- Anemia
- GLP-1 Products
- Chronic kidney disease related



Q & A