

1789 TT

ScinoPharm Investor Conference

— 2023 12 07



Disclaimer

This material has been prepared by ScinoPharm Taiwan, Ltd. (“ScinoPharm”).

Any opinions expressed in this material are subject to change without notice as a result of using different assumptions. ScinoPharm is under no obligation to update or keep current the information contained herein. The **information contained** in this presentation is ScinoPharm’s **confidential** information.

Any disclosure, copying, distribution or any action taken or omitted to be taken in reliance on it is prohibited and may be unlawful.

No representation or warranty, express or implied, is or **will be made** in or in relation to, and no responsibility or liability is or will be accepted by the Company **as to**, the **accuracy or completeness** of this material and any liability therefore is hereby expressly disclaimed.

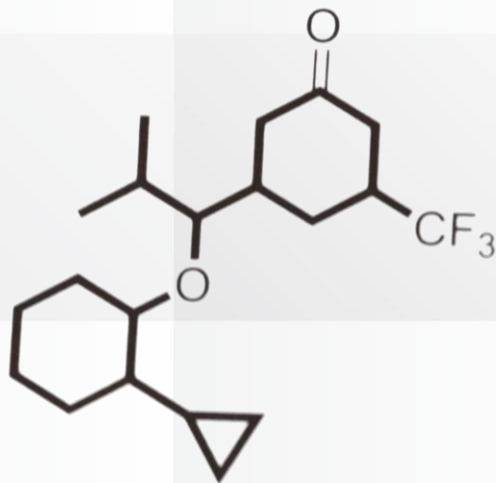
Statements made in this material include forward-looking statements, which include, without limitation, statements about the issues, plans and expectations of ScinoPharm. Without limiting the foregoing, statements including the words “believes”, “anticipates”, “plans”, “expects” and similar expressions are also forward-looking statements. Forward-looking statements reflect, among other things, management’s plans and objectives for future operations, current views with respect to future events and future economic performances and projections of various financial items. **These forward-looking statements involve known and unknown risks**, uncertainties and other factors **which may cause actual results to differ materially from those implied** by such forward-looking statements.

Agenda

01 Company Overview

02 Business Update

03 Financial Performance



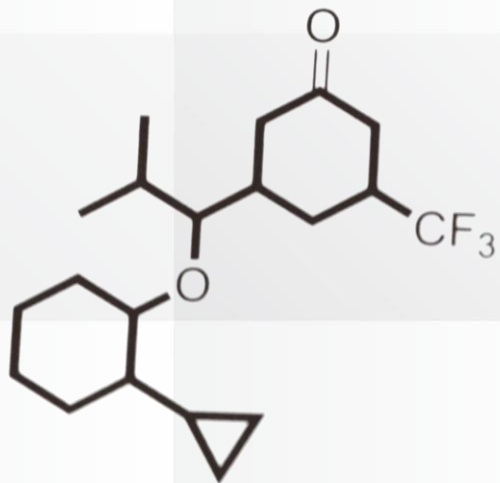
Company Overview



ScinoPharm at a Glance

- Est. 1997 in Taiwan with R&D/CGMP plants in Tainan and Changshu, China plus marketing forces in Tainan, Shanghai and Tokyo
- Specializes in providing R&D and CGMP manufacturing of APIs (cytotoxic/steroid) and injectable drug products
- 76 generic APIs in portfolio with 37 referred and approved ANDAs/NDAs*
 - 920 active DMFs worldwide with 67 US DMFs*
- 200+ contract projects with 11 approved/launched (9 NCEs) and 6 in phase 3 for NDA/MAA filing within 1-3 years*
- API plant certified by key international regulators - US FDA, EMA, EDQM, Australian TGA, Japanese PMDA, Korea KFDA, Mexico COFEPRIS and German Authority
- Injectable plant certified by US FDA and TFDA

*As of 2023/10/31



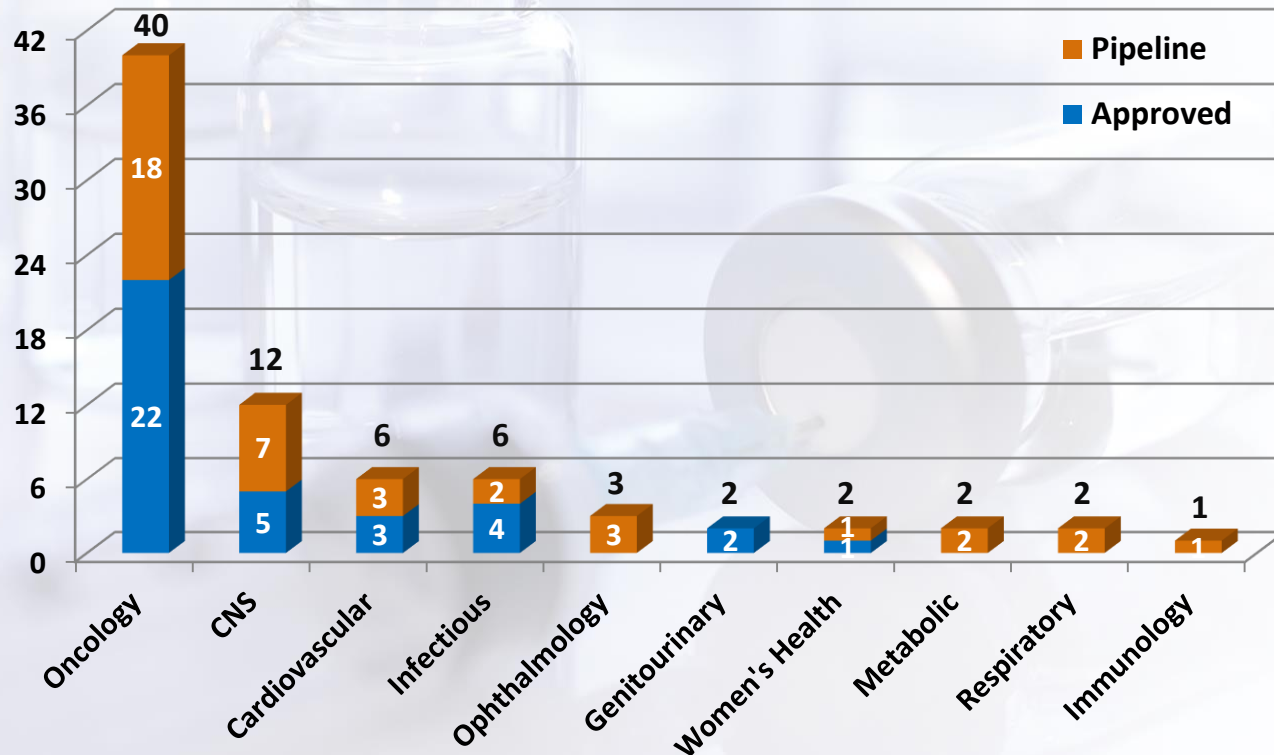
Business Updates



Optimize Generic API Portfolio

■ Generic API Portfolio

As of 2023/10/31



**Optimize
Generic API
Portfolio**

■ 2023 Generic API Product Approval Plan

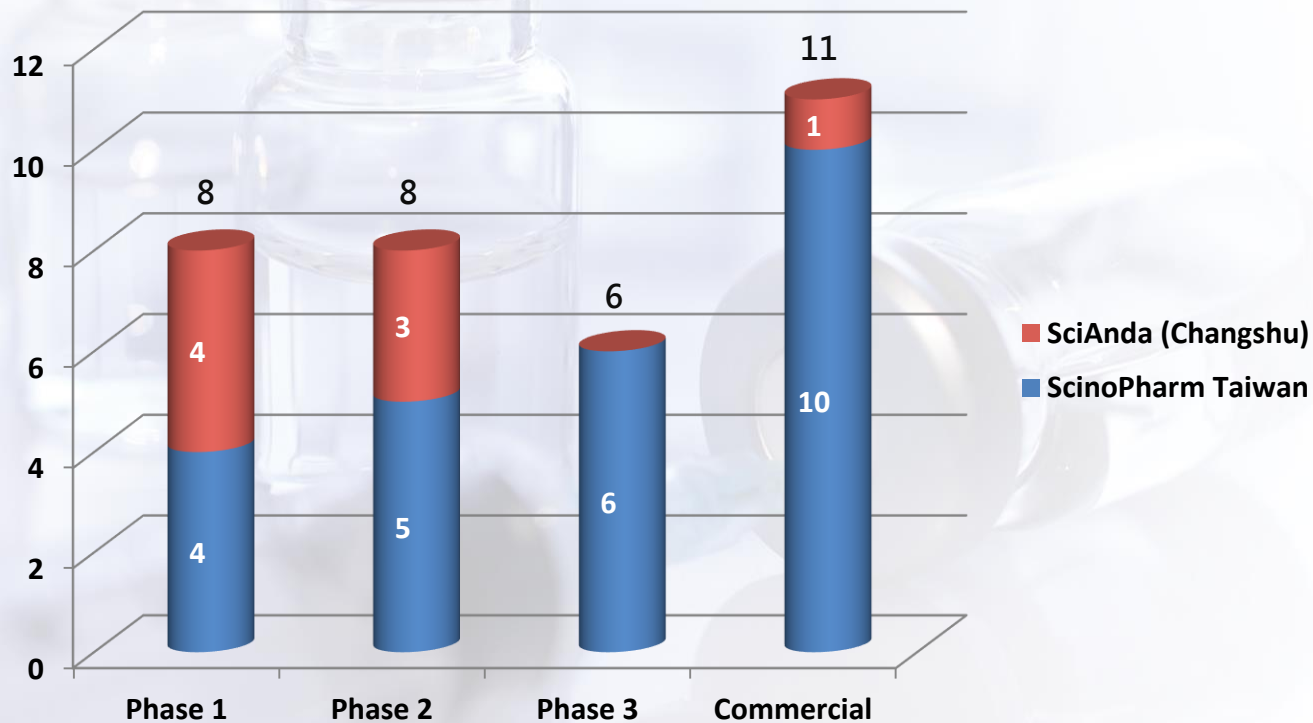
Type	Product	Region	Indication	Brand Marketer
Generic API	Bimatoprost	CN(✓)	Glaucoma	Allergan
Generic API	Cladribine	CN	Multiple sclerosis	Merck
Generic API	Galantamine HBr	CN(✓)	Alzheimer's disease	Janssen
Generic API	Azacitidine	EU	Myelodysplastic syndromes	Celgene

✓ : Approved

As of 2023/10/31

**Expand
CDMO
Business**

■ CDMO Business Status



As of 2023/10/31

**Expand
CDMO
Business**

■ 2023 CDMO Product Approval Plan

Type	Product	Region	Indication	Brand Marketer
CDMO API	Ganaxolone	EU(✓)	Genetic epilepsy	Marinus
CDMO API	Eflornithine	US/EU	FAP	Post-marketing Disclosure
CDMO API	Eflornithine	US/EU	Pediatric neuroblastoma	Post-marketing disclosure
Intermediate for CDMO API	Sotagliflozin	US(✓)	Heart failure	Lexicon

✓ : Approved

As of 2023/10/31

Advancing to Injectables

■ In-House Drug Product Submission Status

	Completed registration batches	Submitted ANDA to US FDA	US FDA's On-site Inspection	US FDA Approved ANDA
Cartridge line Prefilled-syringe				Passed US FDA Pre-Approval Inspection in May, 2022*
Vial line Liquid solution				Passed US FDA Pre-Approval Inspection in May, 2022*
Vial line Lyophilized powder				US FDA approved ANDA in Sep., 2023
Cartridge line Cartridge product				ANDA of 1 st cartridge product - Submitted to FDA in Jun., 2023

* Responded to FDA CRL

Advancing to Injectables

■ Collaborative Projects for Drug Product



**Collaboration on 505(b)(2)
for non-small cell lung cancer**

Launched in US by customer



**Collaboration on ANDA for
non-small cell lung cancer**

**Launched in US and EU by
customer**



**Collaboration on ANDA for
multiple myeloma**

**Launched in the US and EU
and submitted to other
regions by customer**



**In-house Prefilled-syringe
Product**

**Recognized by marketing
partner and ANDA review is
in the final stage**



**Contract Manufacturing of
oncology Injectable**

**Ongoing shipment of CMO
products**



In-house Cartridge Product

**ANDA submitted to FDA and
marketing partner to be
confirmed**

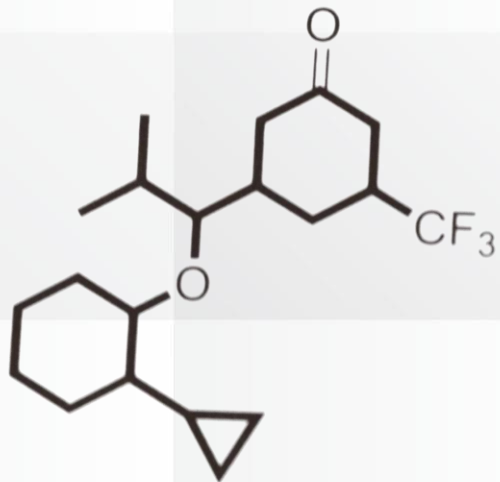
China Market

- 4 CFDI on-site inspections completed in Changshu site to facilitate China market growth

	Product	Approval	Indication	Market
2020.09	Sodium Phenylbutyrate*	2021.05	Urea cycle disorders	Orphan disease medicine
2021.02	Donafenib	2021.06	Advanced liver cancer Thyroid cancer	2023 sales projected by research report : c. RMB 680 million
2021.06	Bimatoprost	2023.02	Glaucoma	Prostaglandin drug products c. RMB 1 billion
2022.11	Azilsartan	2022.09	Hypertension	c. RMB 100 million

* Customer's clinical trial for new indication in progress

- Changshu site expects to conduct more inspections



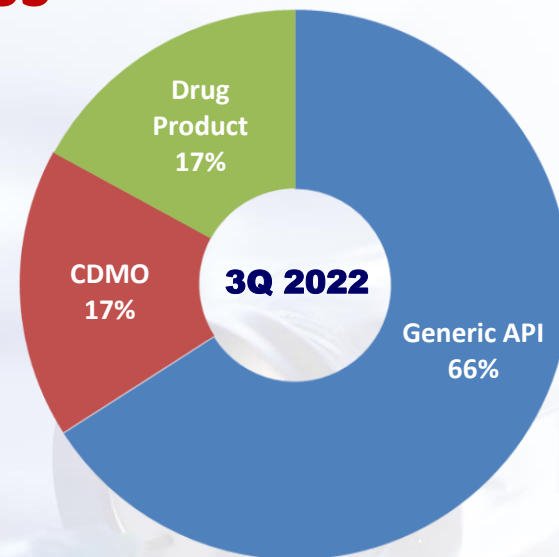
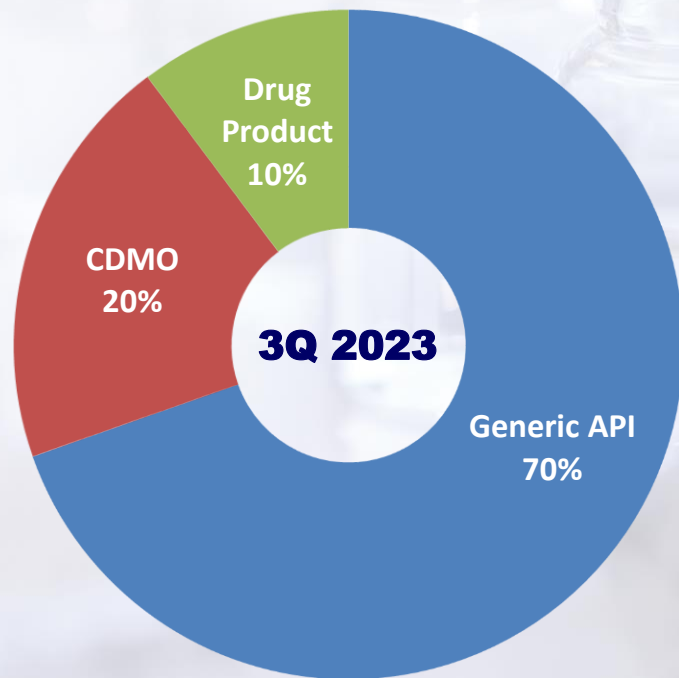
Financial Performance



Consolidated Income Statement

NTD Million except for EPS	3Q 2023		YoY	3Q 2022	
Revenue	2,063	100%	-8%	2,248	100%
Gross Profit	758	37%	-14%	884	39%
Operating Expenses	(635)	-31%	5%	(604)	-27%
Operating Profit	123	6%	-56%	280	12%
Net Profit before Tax	158	8%	-50%	314	14%
Net Profit after Tax	130	6%	-48%	252	11%
EPS (NTD)	0.16	-	-	0.32	-

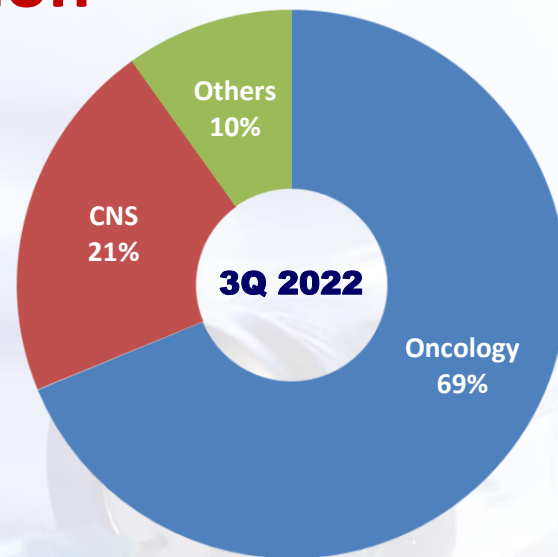
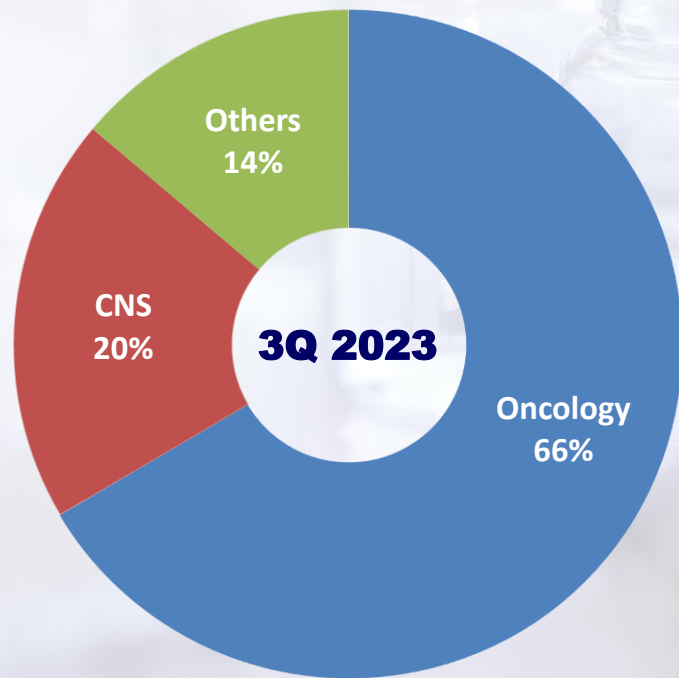
Sales Distribution – By Business



Unit: USD/M

	Generic API	CDMO	Drug Product
3Q 2023 Sales	46.4	13.4	6.8
YoY	-8.4%	2.8%	-47.9%

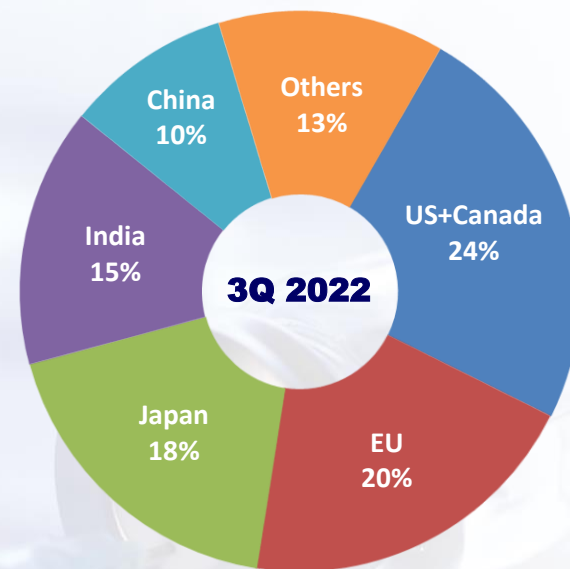
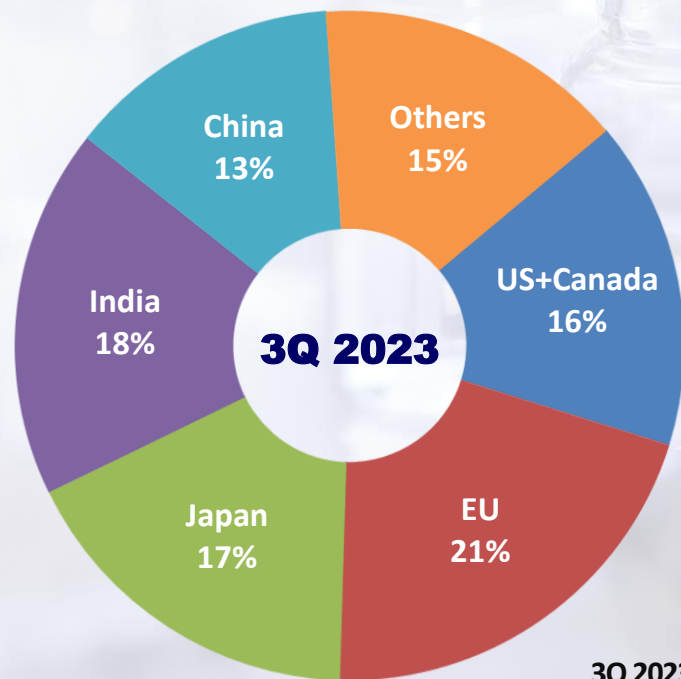
Sales Distribution – By Indication



Unit: USD/M

	Oncology	CNS	Others
3Q 2023 Sales	44.3	13.1	9.2
YoY	-16.0%	-20.3%	21.7%

Sales Distribution – By Region



Unit: USD/M

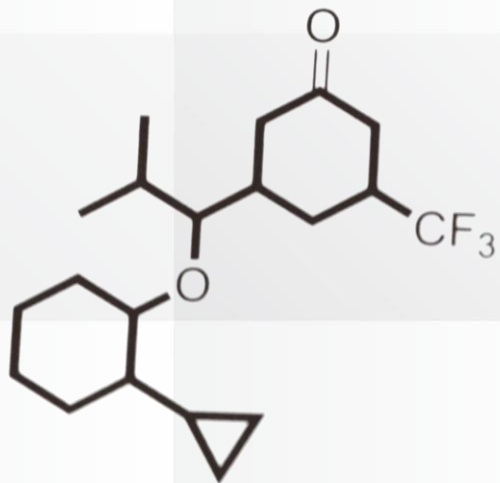
	US & Canada	EU	Japan	India	China	Others
3Q 2023 Sales	10.6	13.8	11.5	11.9	8.8	10.0
YoY	-42.4%	-11.2%	-18.0%	3.7%	20.6%	-0.1%

Consolidated Balance Sheet

NTD Million	2023/09/30		2022/09/30	
Cash and Cash Equivalents	3,995	35%	4,246	36%
Accounts Receivable	364	3%	339	3%
Inventories	1,642	14%	1,304	11%
Property, Plant & Equipment	3,653	32%	3,913	33%
Other Current/Non-Current Assets	1,913	16%	1,937	17%
Total Assets	11,567	100%	11,739	100%
Financial Debt	58	1%	49	0%
Other Current Liabilities	638	5%	629	5%
Other Non-Current Liabilities	629	5%	665	6%
Total Liabilities	1,325	11%	1,343	11%
Total Shareholders' Equities	10,242	89%	10,396	89%

Consolidated Cash Flow Statement

NTD million	3Q 2023	3Q 2022	Dif.
From Operating Activities	173	669	-496
Depreciation & Amortization	344	323	21
From Investing Activities	(157)	(172)	15
Capital Expenditure	(209)	(168)	-41
From Financing Activities	(313)	(341)	28
Effect of foreign exchange rate changes	(2)	9	-11
Net Change in Cash	(300)	165	-465
Beginning Balance	4,295	4,081	214
Ending Balance	3,995	4,246	-251



Q & A





Brand Quality with Asian Advantages

www.scinopharm.com

1789 TT