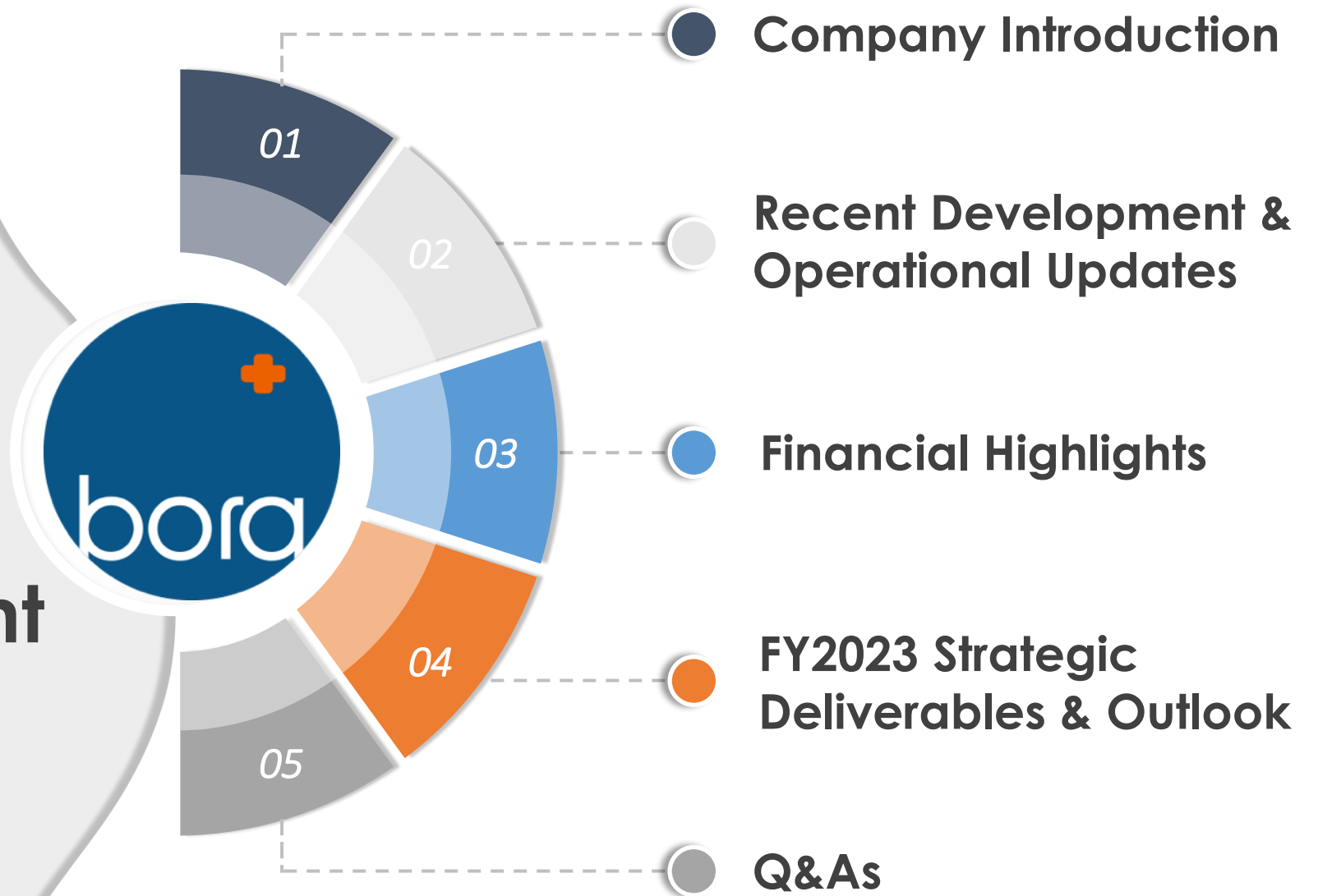




Bora Pharmaceuticals Investor Conference

March 2023

Today to Present



Bora – Largest Pharma CDMO in Taiwan with Global Reach



1972

Hoan Pharmaceuticals is established by John Sheng.



2007

Bora Pharmaceuticals is established as a development scale CDMO with headquarters in Taipei, Taiwan.



2017

Bora's Initial Public Offering (IPO) on the Taipei Stock Exchange (TICKER 6472.TWO).



2020

Bora acquires GSK Mississauga facility and establishes a North American Headquarters.



2022

Bora acquires TWi assets, adding two manufacturing facilities to the network



2002

Bobby Sheng becomes Chairman and CEO of Hoan Pharmaceuticals.



2013

Bora acquires Eisai PICS Tainan facility.



2018

Bora acquires Impax/Amneal's USFDA Zhunan facility.



2022

Bora acquires Eden Biologics Zhubei facility.
Bora Biologics is established.

Well-structured Portfolio Companies with Dedicated Function



Bora
Pharmaceuticals
(Tainan Facility)

Bora
Pharmaceutical
Laboratories
(Zhunan)

Bora
Pharmaceuticals
USA (USA)

Bora
Pharmaceutical
Services
(Canada)

Bora Biologics
(Zhubei)

TWi Pharmaceuticals

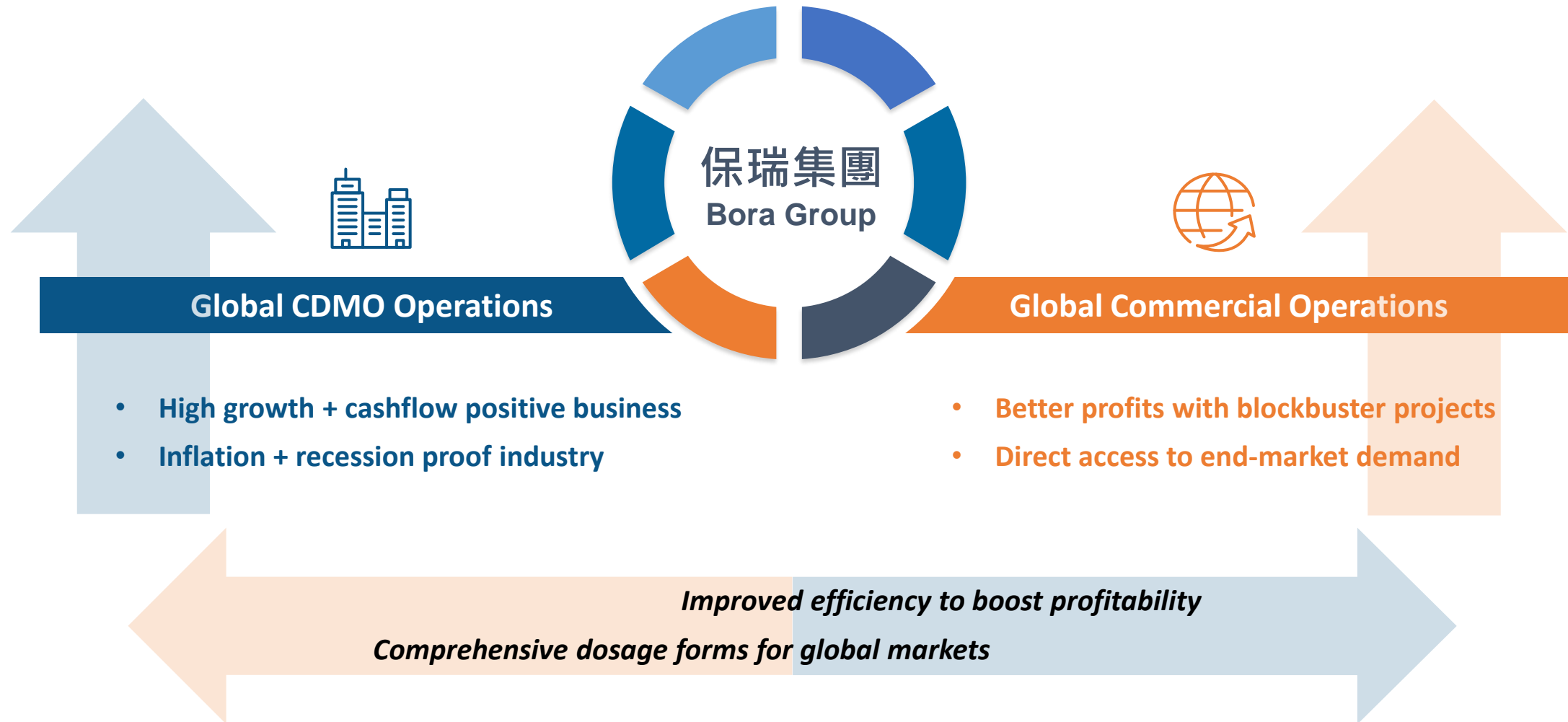
聯邦化學製藥股份
有限公司

保瑞聯邦股份
有限公司

Global CDMO Operations

Global Commercial Operations

Dual-Engine Growth Strategy to “Race to the Top”



Experienced Leadership Team with Proven Track Record



Bobby Sheng
Group CEO & Chairman
> 25 years of experiences in pharma industry, including M&A, strategy planning and operational management



Simon Chen
R&D VP, Spokesman



Alice Wang
Group CFO & Deputy Spokesperson



Sally Langa
Senior VP, S&M CDMO



Don Liscombe
GM, Bora Pharma Services



Tom Chang
GM, Bora Pharma Laboratories Inc



Jennifer Kuan
TW Site of Operations VP, Bora Biologics



Goff Baker
Quality VP



Nick Liu
GM, TWi Pharma



Henry Kuo
GM, Bora Health

20+ years of pharmaceutical experience in every function with global exposure

Bora by the Numbers

50+ Billion NTD Market Cap*

1250+ Employees

100+ Export Countries

7 Manufacturing Sites

10 Billion NTD+ FY2022 Revenue

95% Revenues outside of Taiwan

#1 Pharma Manufacturer in Taiwan (by volume) *

Recent Developments & Operational Highlights

Best Year Achieved with Annual Revenue over NT\$10bn



Robust Business Growth

- **FY2022 revenue of NT\$10,494 million, 2nd consecutive year delivering over 100% growth with dual-engine strategy being successfully executed**
 - **Global CDMO Operations continue to be the cornerstone** with total revenue of NT\$4,788 million
 - ✓ 17 new customers awarded + 26 new molecules added
 - ✓ Completed US FDA audit for SK, the first Taiwan ophthalmic site to be approved by the US FDA
 - ✓ Bora Biologics set-up to build a presence in the fast-growing biological macromolecules market globally – 11 Projects closed with growing contribution from global CDMO clients and 13 change orders/work orders signed during H2'22 to boost future growth
 - **Global Commercial Sale Operations serve as a catalyst to accelerate the growth with the completion of TWi acquisition**
 - ✓ 5 ANDA submission made + 4 US ANDA approvals received
 - ✓ 5 product launches in the US, including the first-to-market Dexamethasone AG
 - ✓ 1 licensing-in & 2 licensing-out deals signed to further boost the potentials

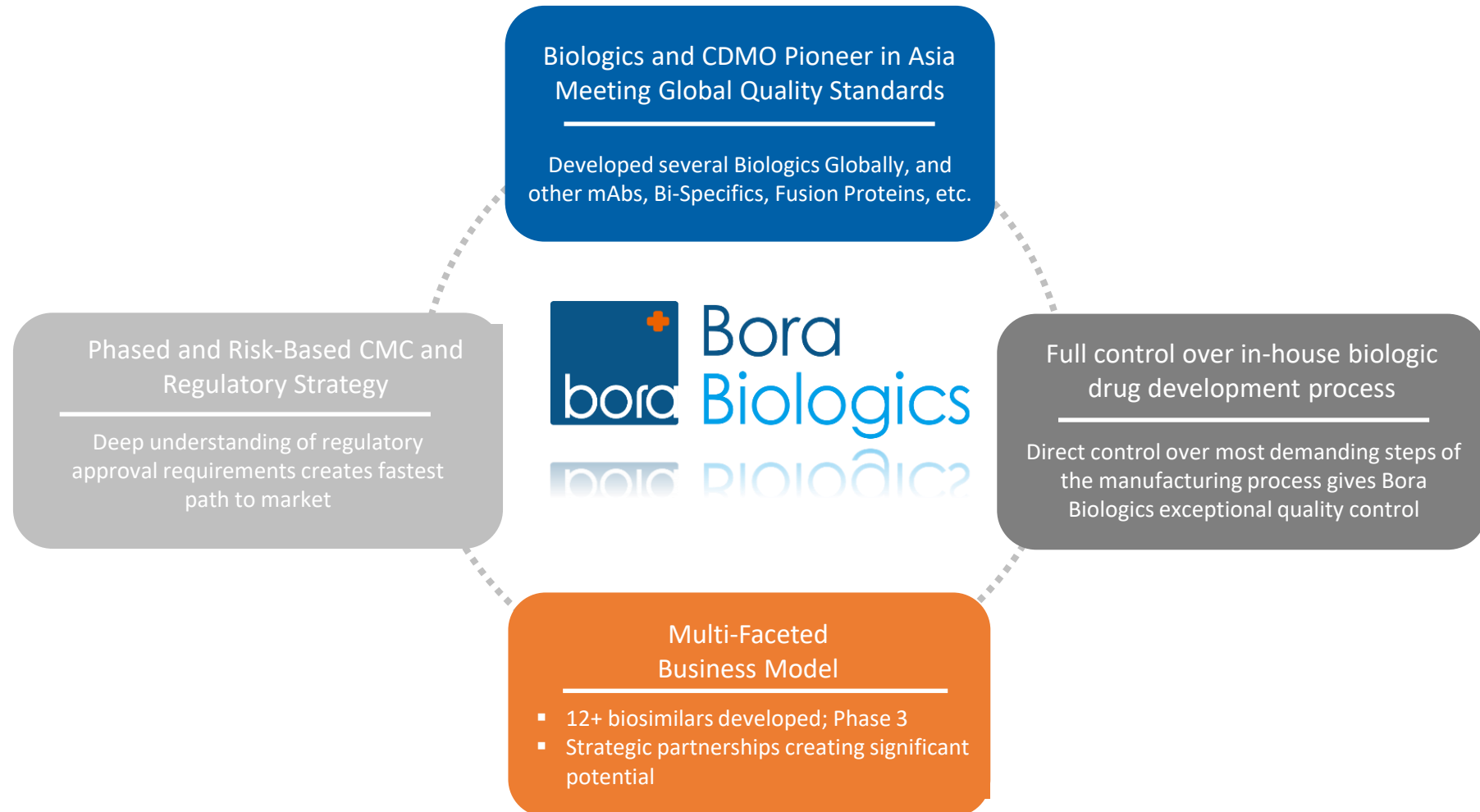
Long-term Value Creation

- **Net income of NT\$1,401 million, increasing by over 30% for 3 years in a row to achieve the highest EPS of NT\$18.52**
 - Dividends of NT\$11 per share, which includes cash dividends of NT\$8 plus stock dividends of NT\$3 and represents approximately 60% payout, set another historic record to maximize shareholders' return
 - Margin profile is expected to be further enhanced with the launch of TWi's self-manufactured Dexamethasone in 2023
- ✓ **Proven track record of delivering sustainable growth being recognized by global professionals**
 - The Company is recognized as one of the **High-Growth Companies Asia Pacific 2023** by Financial Times (FT) and the only Taiwan-based pharmaceutical company among the Top 500 from all Asian countries
 - Continuous value creation through robust business growth together with the improved profitability leads the Company being part of **MSCI Global Small Cap Index** effective November 30, 2022 and **FTSE Global Equity Index Series Small Cap Index** effective March 17, 2023

2 Transformative Transactions in 2022 – Bora Biologic



Value creation by gaining complex technology in the most cost efficient way



Biologic Platform – Making Bora #1 & the Only CDMO Offering Both Small Molecules & Biologics



High Titer Platform starting from Cell Line Development



One of the most comprehensive onsite Analytical Labs in the region w/ experience in early protein characterization and meeting innovator ranges



Manufacturability and Quality Policies that are in line with Global Regulatory Standards and meets all regulations



Minimizing Time to IND and BLA & Platform processes are proven



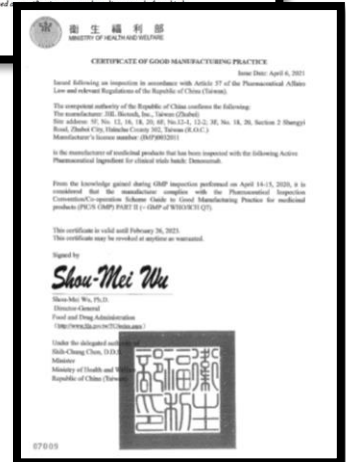
Innovative global clinical trials strategy creates fastest path to market

EU QP audit (2021)

PPD		CONFIDENTIAL
AUDIT CERTIFICATE		Global Quality and Compliance
PR #	13040	
AUDITEE	Eden Biologics	
AUDIT ADDRESS	5F, No.18, Sec.2, Shengye Rd. Zhubei City Taiwan 302	
DATES OF AUDIT	09-Mar-2021 to 11-Mar-2021	
AUDIT TYPE	Vendor Audit: Qualification	
AUDIT STRATEGY	Remote	
ENGAGEMENT	IMP Drug Substance: Manufacture and Quality Control Testing of Drug Substance intended use in EU and non-EU Clinical Trials	
PURPOSE	To determine if appropriate internal processes and documentation are in place at Eden Biologics to provide IMP Drug Substance manufacture, Quality Control Analyses and Stability Testing	
AREAS AUDITED	Quality Management System Personnel Production Premises & Equipment Documentation Outsourced Activities Complaints & Results Self Inspection Data Integrity	
LEAD AUDITOR	O'Donohue, Darragh M	
OTHER AUDITORS	Hardy K Farmer Holley P McCloskey	
APPROVED BY	Irene Connell	
APPROVED ON	08-Apr-2021 9:12 pm	
ISSUED BY	Darragh M O'Donohue	
ISSUED ON	09-Apr-2021 10:41 am	

Disclaimer: This Audit Certificate signifies an audit was conducted by PPD Global Quality and Compliance. This Audit Certificate should not be viewed as a guarantee of quality.

Report run by: Darragh M O'Donohue



TFDA GMP (2021)

2 Transformative Transactions in 2022 – TWi



Value creation by adding another strong catalyst to fuel the growth



Approved ANDAs – Majority are PIVs

13

PIV

Product	Indication	Form	Technology	Approval Time	Launch Time
Donepezil hydrochloride (Aricept)	Dementia	ER tablet	Matrix	10/2014	Q1 2015
Guanfacine hydrochloride (INTUNIV)	ADHD	ER tablet	Matrix	06/2015	Q2 2015
Megestrol acetate (Megace ES)	Anorexia, Cachexia	Oral suspension	Nanoparticle	08/2014	Q3 2015
Bupropion HCl (WELLBUTRIN XL)	Depressive Disorder	ER tablet	Membrane-controlled	11/2017	Q4 2018
Cyclobenzaprine hydrochloride (AMRIX)	Muscle relaxants	ER capsule	Pellets	01/2013	Q2 2019
Testosterone (AndroGel)	Hypogonadism	Gel 1.62%	Topical gel (pump)	09/2019	Q2 2020
Choline fenofibrate (TriLipix)	Hypertriglyceridemia	DR capsule	Mini-tablets	07/2019	Q3 2020
Terbutaline Sulfate (BRETHINE)	Asthma	Tablet	IR tablets	06/2020	Q3 2020
Dimethyl Fumarate (Tecfidera)	Multiple sclerosis	DR capsule	Mini-tablets	01/2019	Q4 2020
Metformin HCl (Fortamet)	Diabetes	ER tablet	OROS	09/2021	Q4 2021
Testosterone (Axiron)	Hypogonadism	Solution	Topical solution	10/2021	Q1 2023
Dexlansoprazole (Dexilant)	Proton-pump inhibitor	DR capsule	Pellets	09/2022	Q1 2023
Oxcarbazepine ER (Oxtellar XR)*	Seizures	ER tablet	Matrix	11/2018	

8

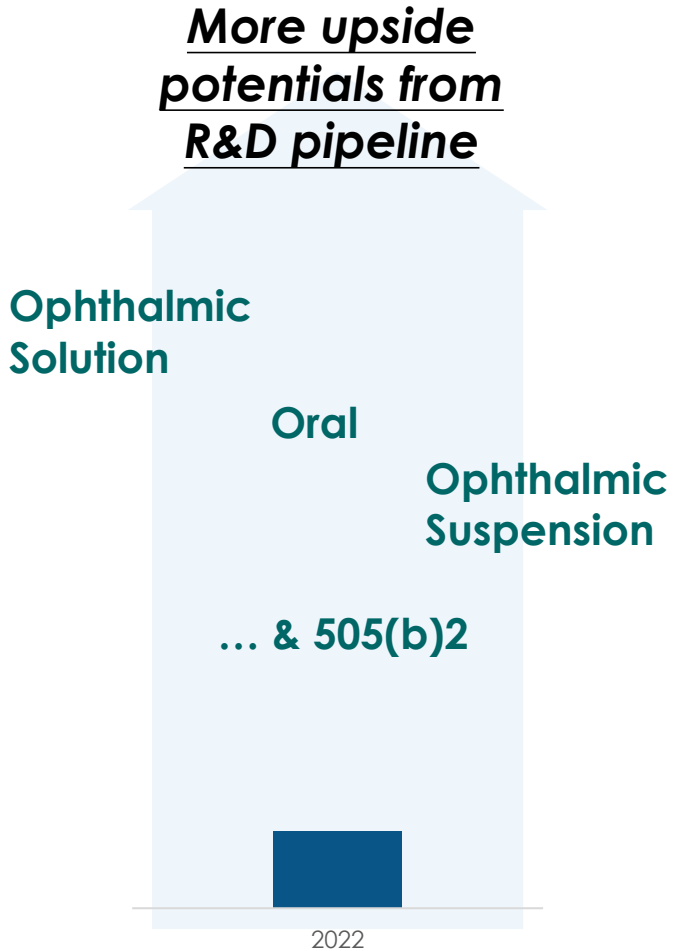
High-entry-barrier

Product	Indication	Form	Technology	Approval Time	Launch Time
Nifedipine (PROCARDIA XL)	Hypertension	ER tablet	OROS	04/2014	Q1 2015
Diltiazem hydrochloride (Cardizem CD)	Hypertension; Angina	ER capsule	Pellets	08/2018	Q2 2019
Propafenone HCl (Rythmol SR)	Anti-arrhythmic	ER capsule	Mini-tablets	06/2020	Q3 2020
Mycophenolic acid (MYFORTIC)	Immuno-suppressant	DR tablet	DR coating	11/2021	Q1 2022
Fluphenazine HCl	Antipsychotic	Tablet	IR tablets	04/2022	Q4 2022
Guanfacine HCl	Hypertension	Tablet	IR tablets	05/2022	Q1 2023
Dicyclomine HCl (BENTYL)	Irritable bowel syndrome	Capsule	Capsules	12/2022	Q1 2023
Diltiazem HCL SR (Cardizem SR)	Hypertension	SR Capsule	Capsules	03/2023	

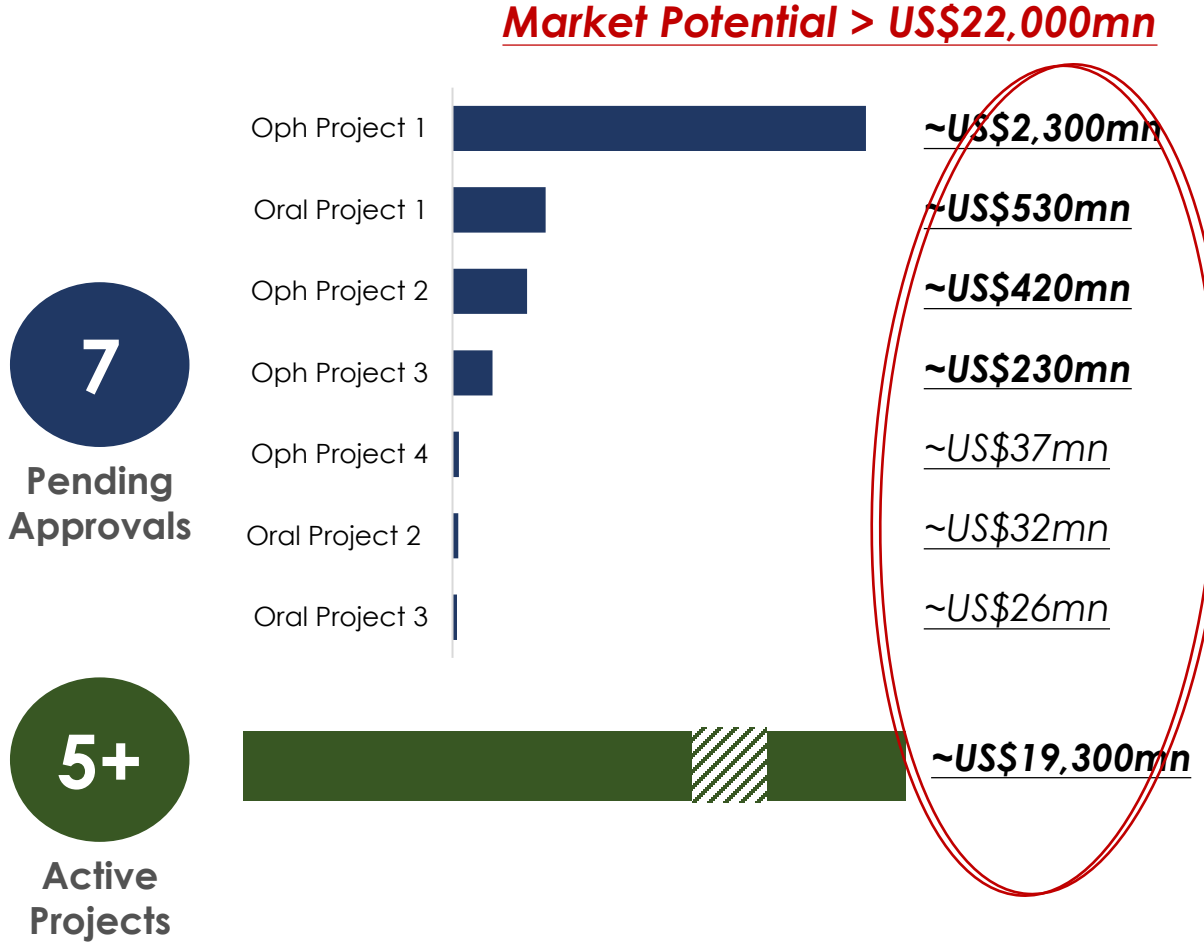
More to Come from Robust R&D Pipeline



TWi Sales



TTL Addressable Market Opportunities*

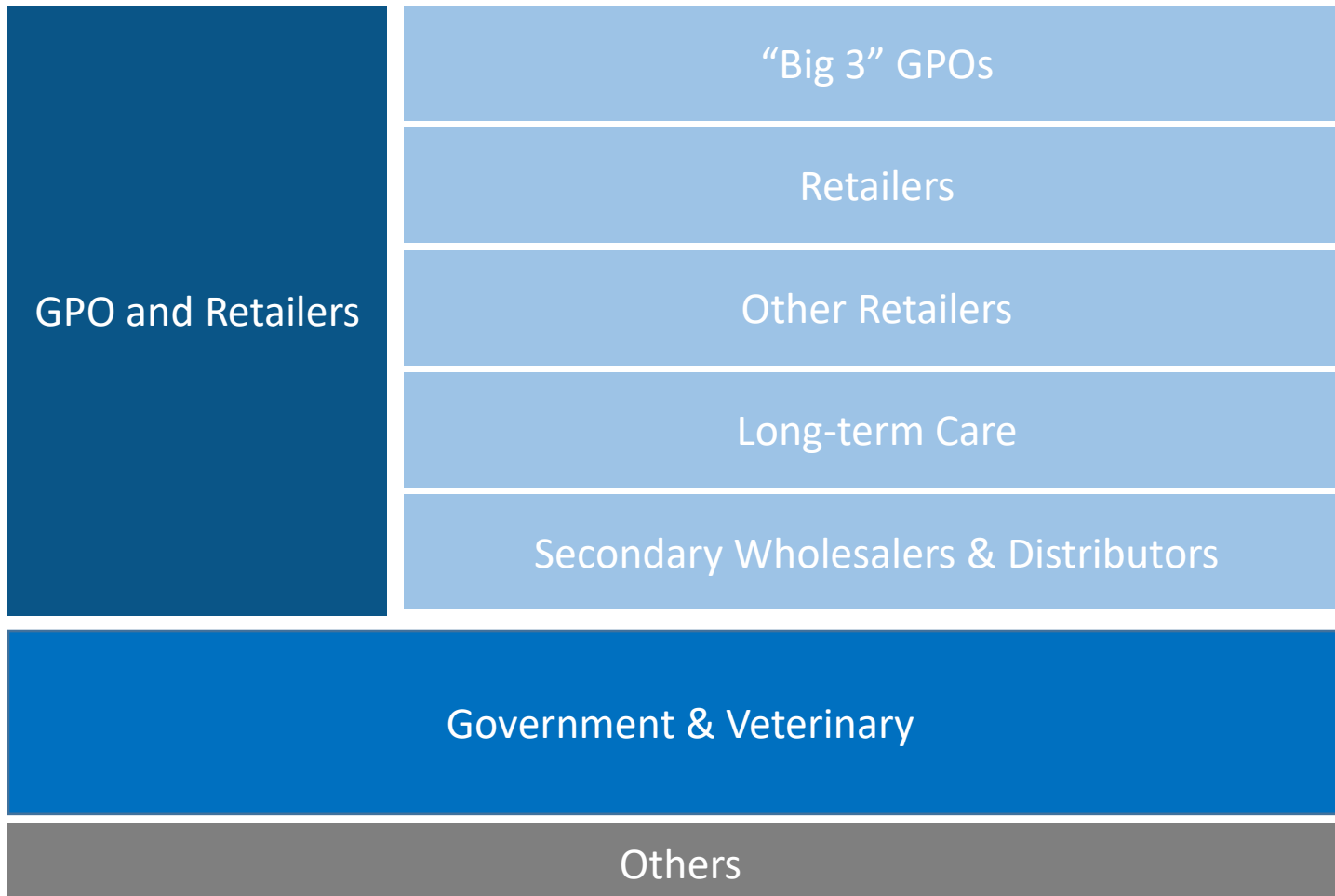


*Based on IQVIA data as of December 2022

Superior S&M Recognized by Major Channels in the US



- Accessible to ~90% of the demand in the US through well-connected customers



Ranked **#1** in the US for ***New Generic Launches*** November 2022

Top 50 Corporations		Contribution Rate
1	TWi Pharma US	12.21%
2	Prasco Labs	8.82%
3	Quallent Pharma	7.10%
4	Dr Reddy Inc	6.73%
5	Viartis Inc.	5.08%
6	Apotex Corp	4.78%
7	Teva Pharm USA	4.71%
8	Cipla USA, Inc.	4.05%
9	Johnson & Johnson	3.42%
10	Sun Pharma	3.32%
11	Heteropharmaceutic	3.27%
12	Accord Hlthcnc Inc	2.21%
13	Novartis	2.06%
14	Lupin	1.66%
15	Zydus Pharm	1.53%
16	Amneal Inc	1.39%
17	Strides Pharma	1.36%
18	Northstar Rx	1.22%
19	Hikma Pharm	1.16%
20	Ascend Labs	1.13%
21	Auromedics Pharma	1.12%
22	KvK-Tech	1.10%
23	Alvogen	1.06%
24	Braintree Labs Inc	0.92%
25	Alembic Pharm	0.91%

Source: Raymond James, U.S. FDA, Wolters Kluwer, IQVIA

More Synergies to be Realized through Integration

Efficiency Improvement for Synergies of Operations

Short Term

- ✓ Move products to the most efficient site -> 6 TWi products are under TT study at Zhunan site
- ✓ Reduce redundant operating costs -> Centralizing packaging of TWi products in Zhunan Site
- ✓ Remove low margin contributing (working capital consuming) products -> PAC to be terminated by the end of Mar'23
- ✓ Optimize R&D pipeline -> Pipeline selection meeting kicking off across different functions with support from sites
- ✓ Get SK pass USFDA audit + fine tune SK org to support CMO business -> Successful USFDA inspection and EIR received in Dec'22

Mid Term

- Accelerate portfolio expansion via M&A or partnership for Branded, OTC, and Biologics
- Diversify product portfolio
- Synergize CDMO business by leveraging commercial sale networks
- Expand sales in Taiwan and China
- Invest in business systems to further enhance the operating/managerial efficiency

Long Term

- Create sustainable growth with well-diversified portfolio
- Expand into more overseas markets with meaningful market presence
- Become a truly global pharmaceutical company

Revenue Growth with More Diversified Portfolio

Best in Class



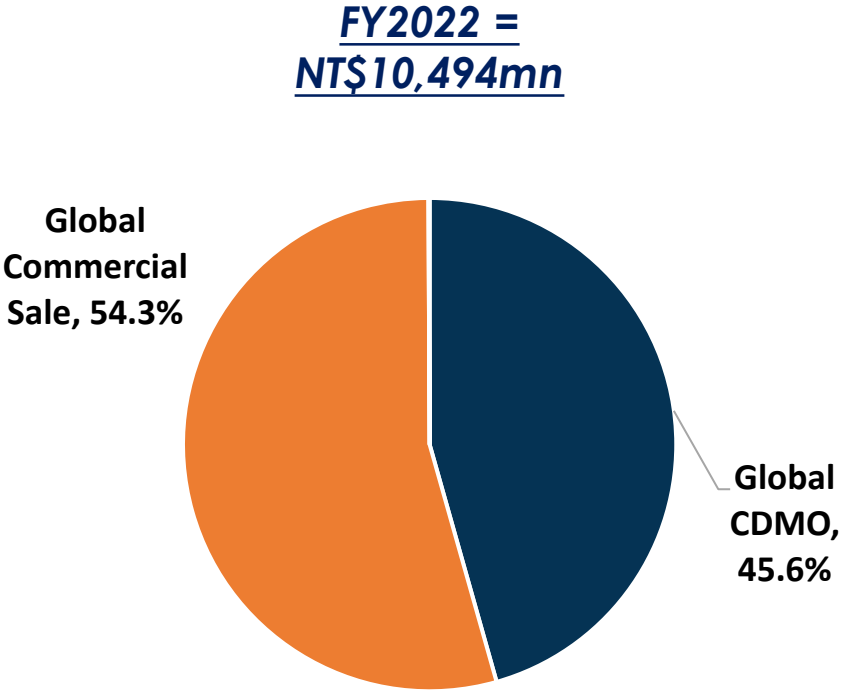
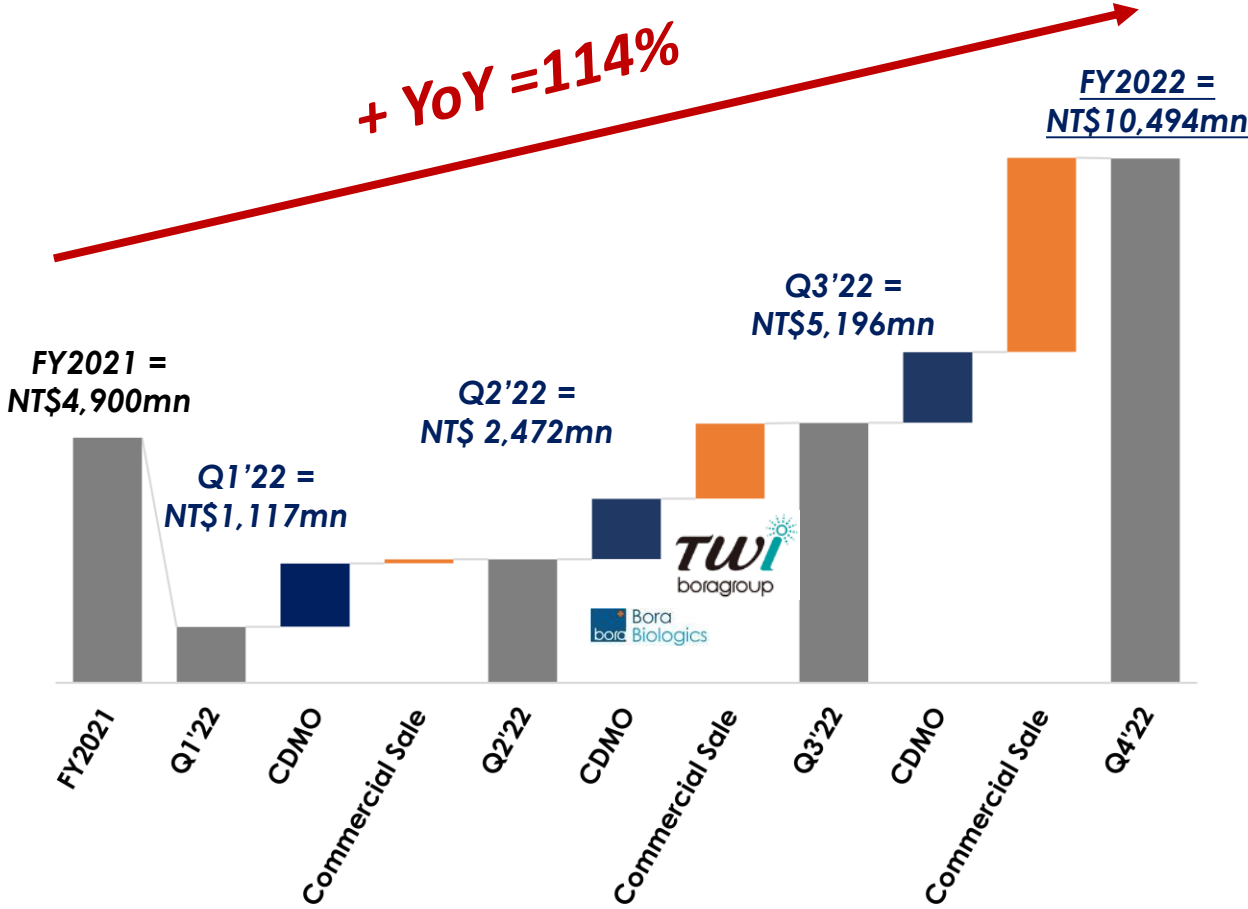
Financial Highlights



Strong Growth Catalysts Continue to Set New Record



FY2022 Quarterly Revenue Breakdown and Key Growth Drivers



Financial Highlights

Key Financials (in NTD millions, except for EPS)	FY2021	FY2022	YoY %
Revenue	4,900	10,494	114%
COGS	(3,228)	(7,582)	
Gross Profits	1,672	2,913	74%
GM %	34%	28%	
Operating Expenses	(626)	(991)	
Operating Income	1,046	1,922	84%
OPM %	21%	19%	
Non-OPEX			
Financial costs	(54)	(109)	103%
Net of other gain/loss	32	27	-16%
Net Income Before Tax	1,024	1,840	80%
Net Income	750	1,402	87%
EPS (NTD)	10.04	18.52	84%

Consolidation of TWi starts from Sep 1st, 2022

- Licensed-in PAC GM <10%
- AG DLS – royalty + outsourced COGS
- Year-end plant shut-down in Q4

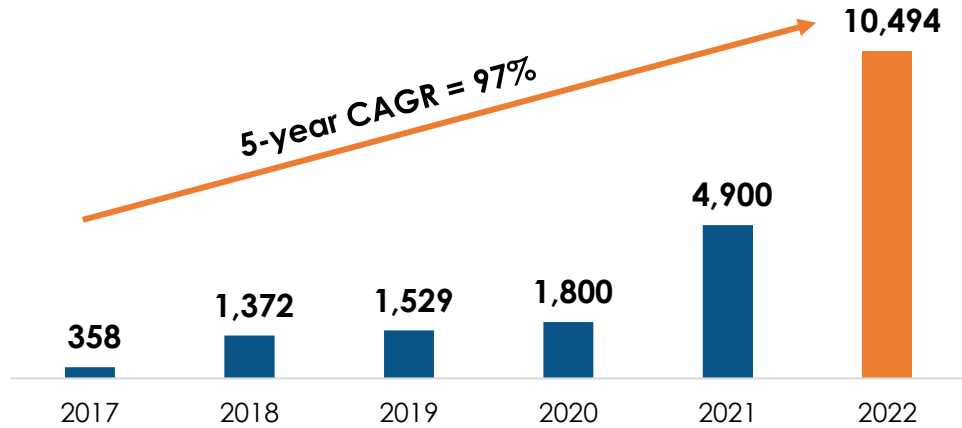
Mainly due to the inclusion of TWi starting from Sep 1st, 2022

Increased interest-bearing debt due to the drawn-down of the syndication loan

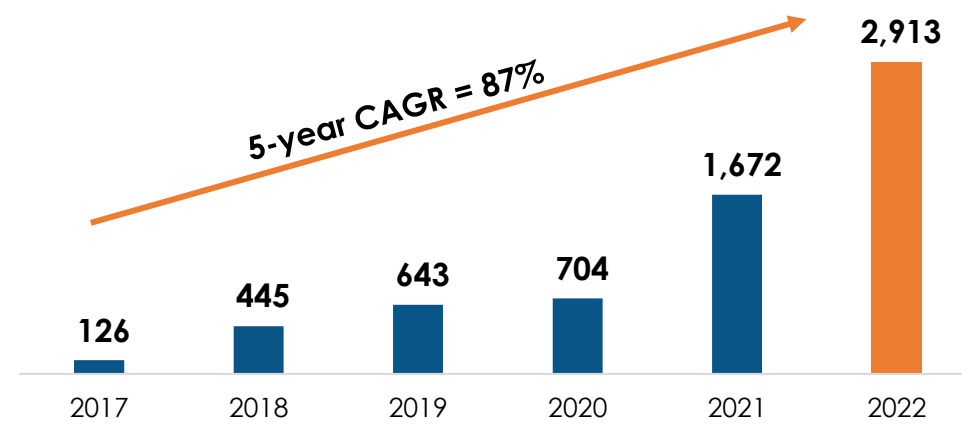
Bora Group's Rapid Growth Backed by Strong Financials



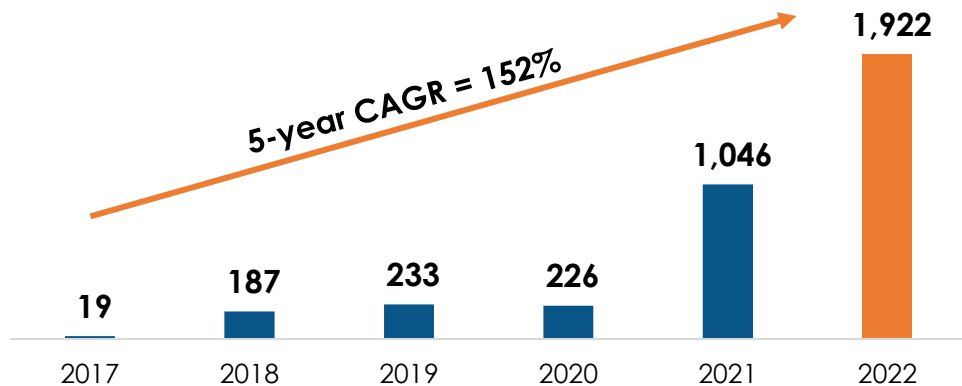
Revenue (NT\$'mn)



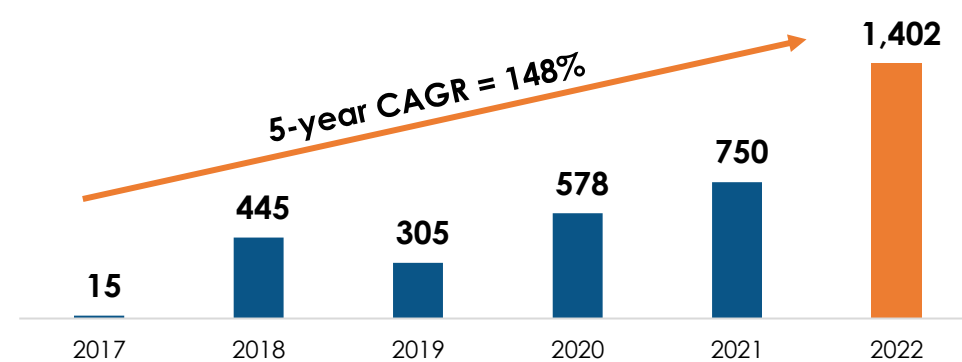
Gross Profits (NT\$'mn)



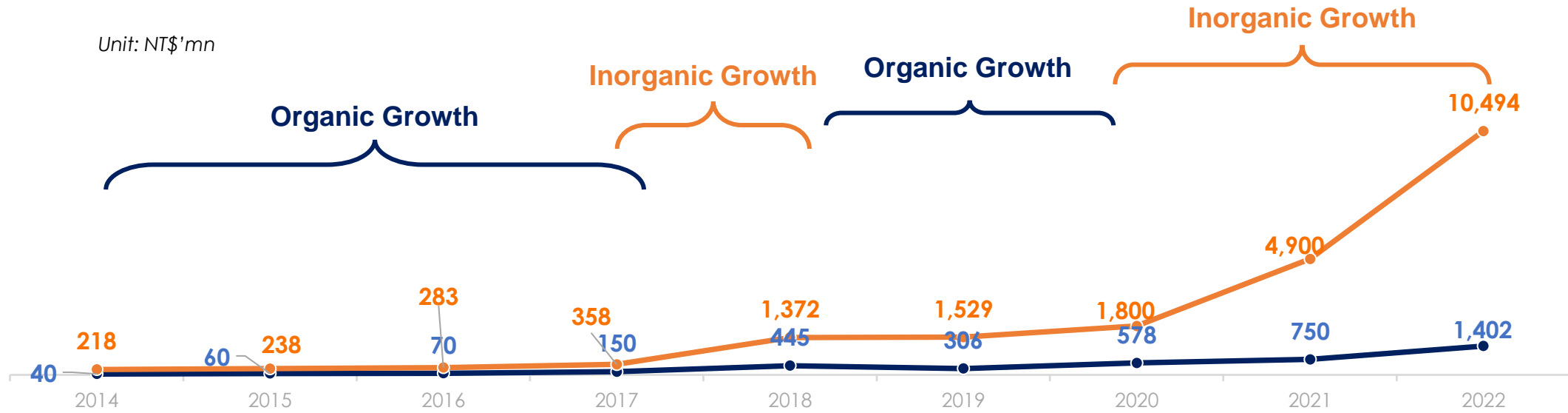
Operating Income (NT\$'mn)



Net Profits (NT\$'mn)

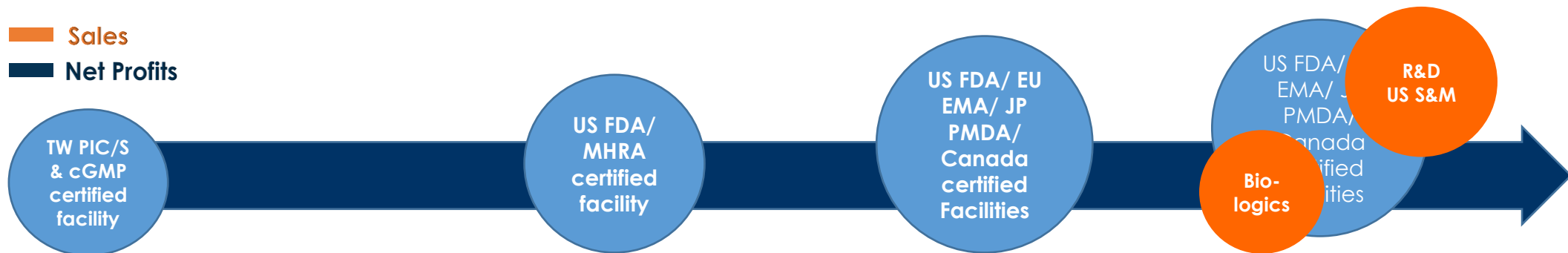


... Driven by both Organic & Inorganic Growth



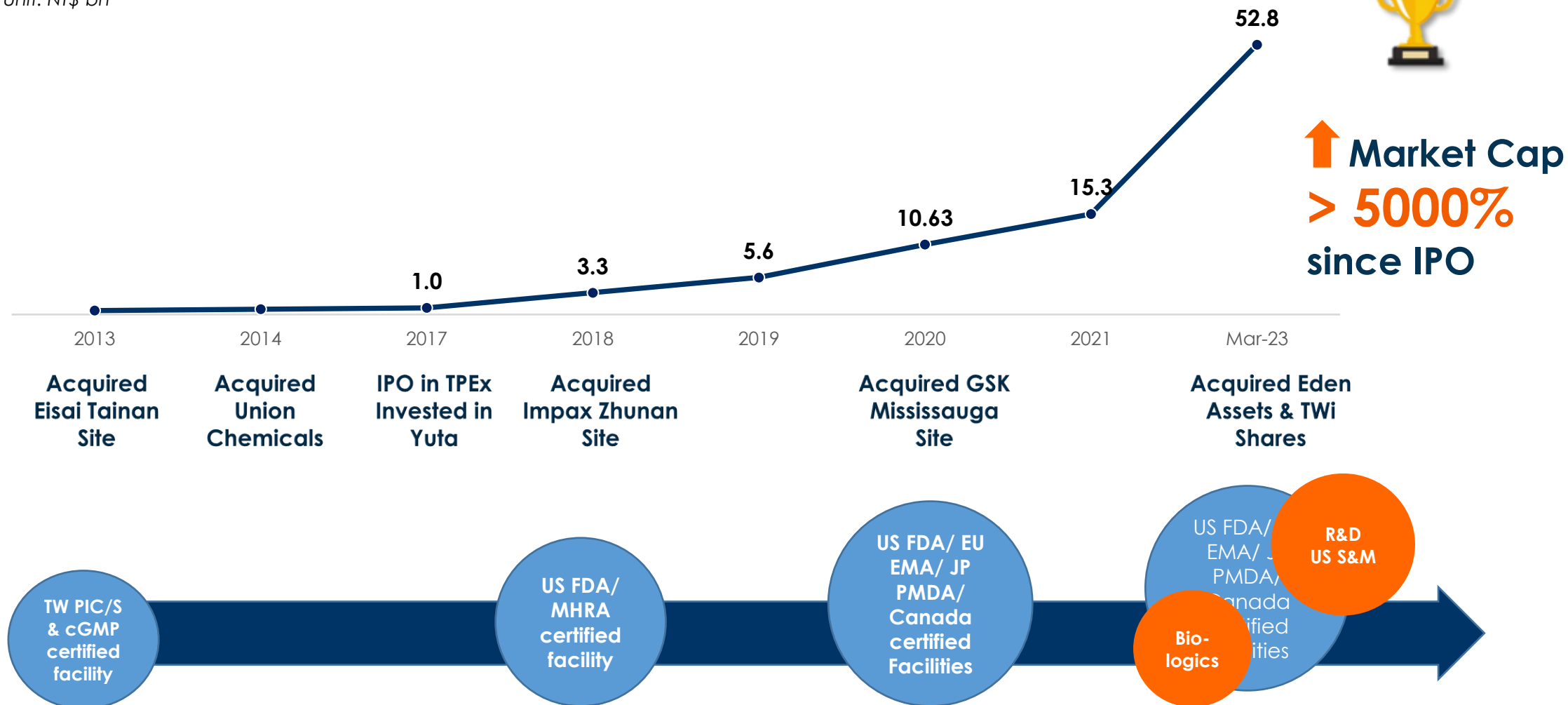
Acquired Eisai Tainan Site (2015)
 Acquired Union Chemicals (2016)
 Acquired Impax Zhunan Site (2017)
 Acquired GSK Mississauga Site (2020)
 Acquired Eden Assets & TWI Shares (2022)

■ Sales
■ Net Profits



Creating Substantial Value for All Shareholders

Unit: NT\$'bn





Strategic Deliverables & Future Outlook

Well-defined Strategic Deliverables to Fuel FY2023 Outlook



Global CDMO



Global Commercial Sale



Further expand business opportunities by capitalizing existing platforms

- 10+ new CDMO customers
- 10+ new molecules for global CDMO
- To support existing customers' expansion plan consistently

- 5+ new launches for the US market
- To expand out-licensing to ex-US countries



Continuously invest in portfolio & technology for sustainable growth

- Dedicated team to support clinical development & advanced manufacturing
- Investment in R&D CAPEX, equipment, and new technology

- 3+ US ANDA submissions
- 3+ US ADNA approvals
- 3+ licensing partnership deals



Efficiently execute the M&A integration to realize the synergies

- Production integration between different sites in Taiwan
- Expansion of Bora Biologics all-type biologics CDMO and SK ophthalmology CDMO business

- Integration between newly-acquired TWi and Bora Health to accelerate the growth of all product lines, including brand/generic, and CHC

- Current economic situation creates more opportunity to acquire high value assets for both global CDMO business as well as more catalysts for global commercial sale

Bora Group as the Leading Taiwan-based Pharma Company Fueled by Dual-engine Strategy with Global Reach



保瑞集團
Bora Group



Largest CDMO pharma company in Taiwan with well-established infrastructure



Dual-engine growth strategy covering the major fast-growing segments in global pharma industry with competitive edge



State-of-the-art manufacturing facilities, approved by most of the advance regulatory authorities, supplying 100+ countries globally



Superior execution of delivering the M&A synergies to maximize shareholders' value



Competent and experienced leadership team with proven track record of driving success globally



Contributing to
Better Health All
Over the World

Contributing to **Better** Health All Over the World

