

Lotus

LOTUS PHARMACEUTICAL

Asian-based Pharmaceutical Company with Global Reach

1795.TT

2025



Safe Harbor Statement

Except for historical information contained herein, the matters set forth in this presentation are forward looking statements that are subject to risks and uncertainties that could cause actual results to differ materially. These forward looking statements are not based on historical facts but rather on management's expectations regarding future growth, results of operations, performance, future capital and other expenditures, competitive advantages, business prospects and opportunities. Statements in this presentation about our future plans and intentions, results, level of activities, performance, goals or achievements or other future events constitute forward looking statements. Wherever possible, words such as "anticipate", "believe", "expect", "may", "could", "will", "potential", "intend", "estimate", "should", "plan", "predict", or the negative or other variations of statements reflect management's current beliefs and assumptions and are based on the information currently available to our management. **Investors are cautioned not to place undue reliance on these forward looking statements, which are made as of the date of this presentation** and we assume no obligation to update or revise any forward looking statements.

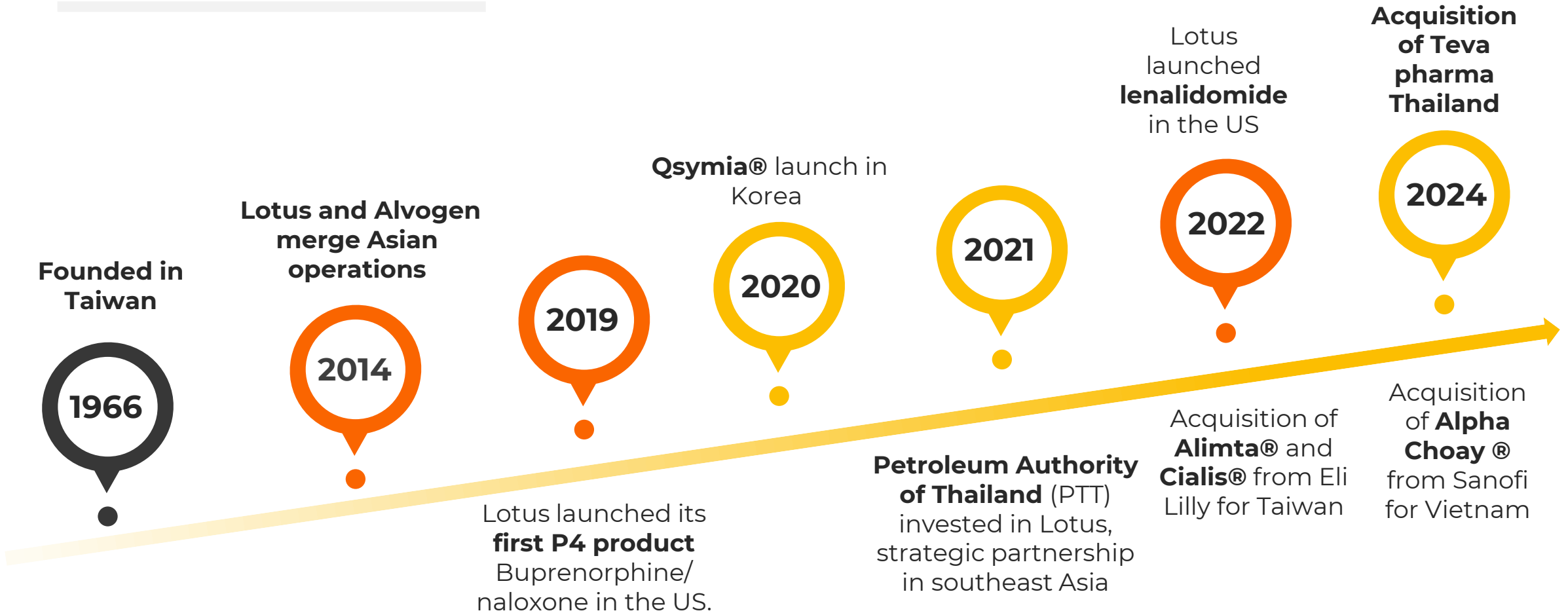
Lotus is well positioned on the growth path



From a single market company with \$20m sales 10 years ago to US\$500mn in 2024

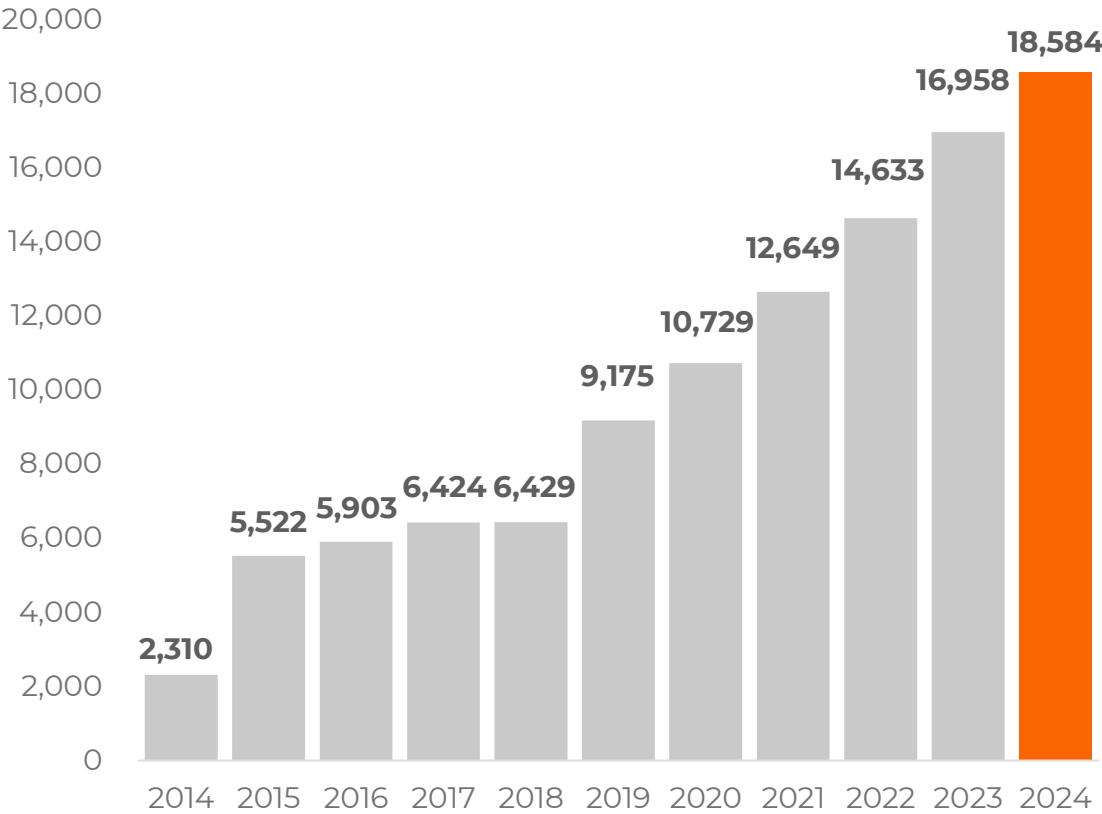
Actively seeking for Southeast Asia opportunities

Major milestones of Lotus

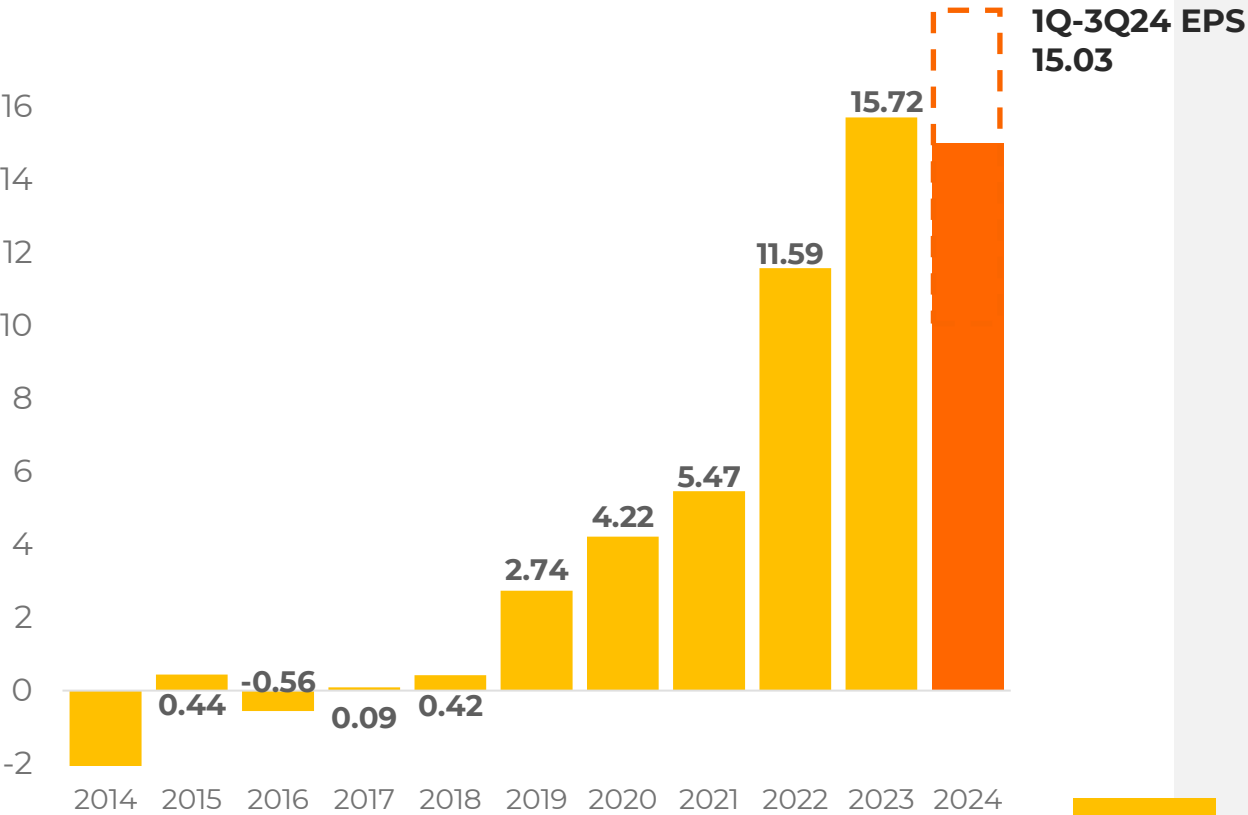


Lotus Transformation

Revenue in 2014-2023 (NT\$MN)



EPS in 2014-2023 (NT\$)



Experienced Management Team



20+

PETAR VAZHAROV
Chief Executive Officer





20+

MANISH CHAWLA
Vice President R&D





20+

EDIN BULJBASIC
General Counsel





20+

VALERIE LAU
Chief Commercial Officer





15+

JULIA LIN
Senior Director Regulatory Affairs

with Lotus since 2007 after graduated with BS in Biomedical Toxicology from University of Guelph



15+

VAMSI KIRAN KOSARAJU
Chief Operations Officer





15+

PRASHANT GODSE
Vice President Business Development





20+

JAVIER TORREJON NIETO
VP of Global Portfolio Management





8

SNAEVAR VIDISSON
Director Export B2B Business



20+

FIONA LI
Human Resource Senior director



 Years of Pharmaceutical industry experience

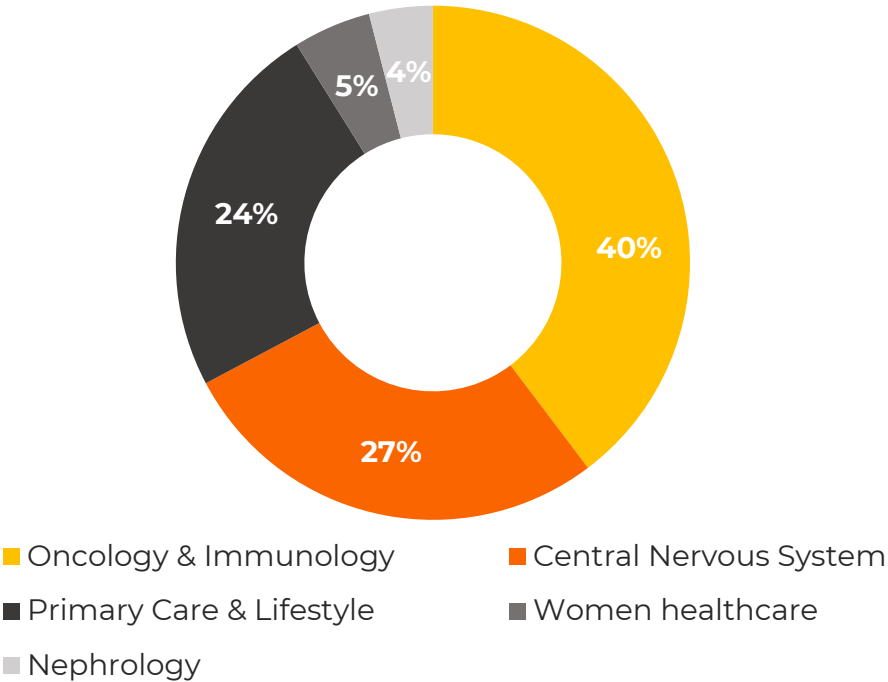
Strong Core Business supported by Two-Pronged Expansion Strategy



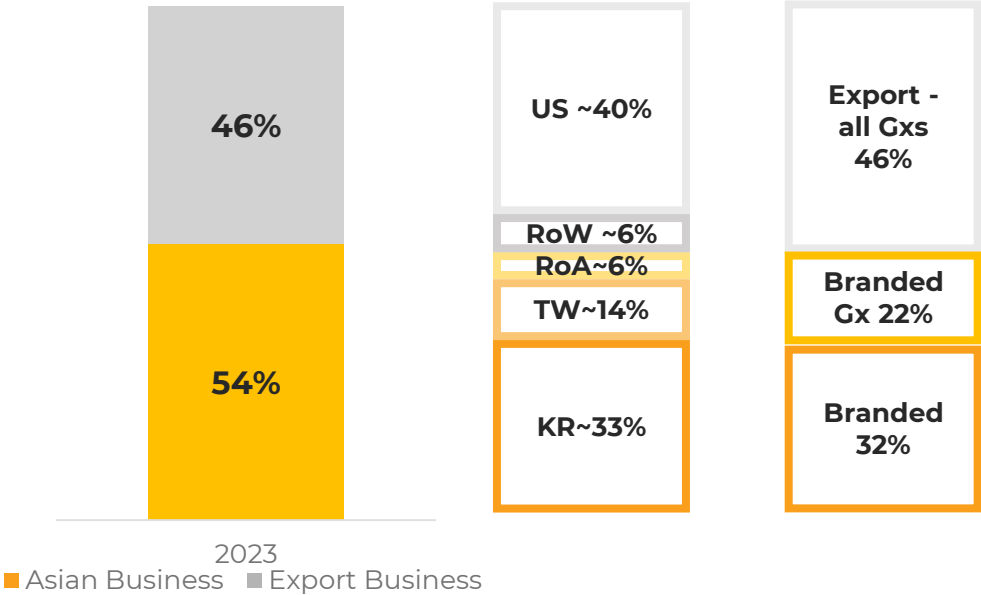
Diversification Across Therapies And Markets

Diversified Portfolio

2023 Revenue Breakdown by TA



Revenue Breakdown by Market



Commercial Infrastructure In APAC

Korea

- **12** sales offices with 216 S&M FTE
- Focused TAs - Oncology, anti-obesity, nephrology, CVS, CNS and WHC
- Exclusive distributor of AstraZeneca's Seroquel®, Zoladex®, Arimidex®, Casodex® and Orion's Stalevo® and Comtan®
- Successfully built a blockbuster weight-loss brand Qsymia®

Vietnam

- **Own Rep office** with GM, RA, SCM and other support functions for commercial operations
- Established team in place to handle both originator products like Stalevo® and generic products with oncology as priority

Malaysia

- **100% owned entity** and current operations via Hybrid partnership model
- Local team being enhanced for the upcoming new launches

Indonesia

- Acquired brands like Stalevo® and current operations via Hybrid model
- Pipeline includes biosimilars and generic products

Taiwan

- **3** sales offices with S&M FTE of 131
- **Global R&D center** focusing on oral oncology, complex Gx, and 505(b)2
- Focused TAs - CNS, osteoporosis, oncology and anti-addiction
- Successfully built block buster brands (e.g. Myseryl®, Arheuma®)
- Revitalized brands like Aclasta®, Cialis® and Alimta® post acquisition

Thailand

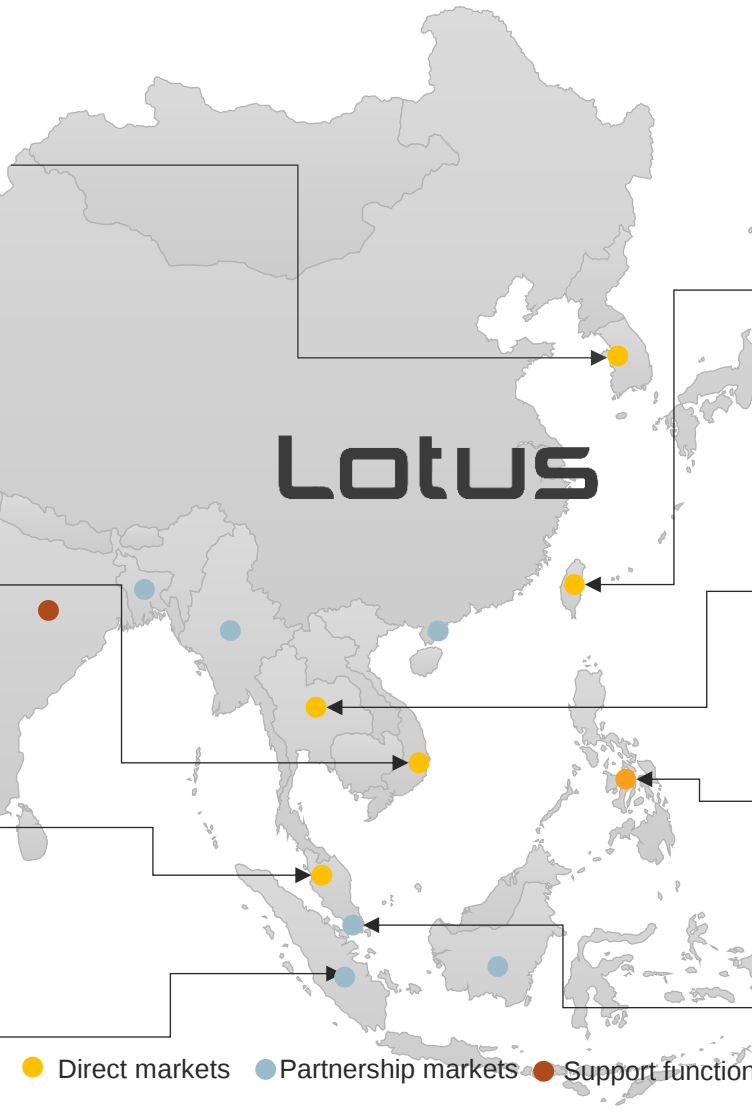
- **82 S&M FTE** covering all hospital and retail channels nationwide
- Focused TAs – Ophthalmology, respiratory and oncology

Philippines

- **Own entity** with 12 S&M FTE for Oncology and CNS portfolio
- Hybrid model for therapeutic area products

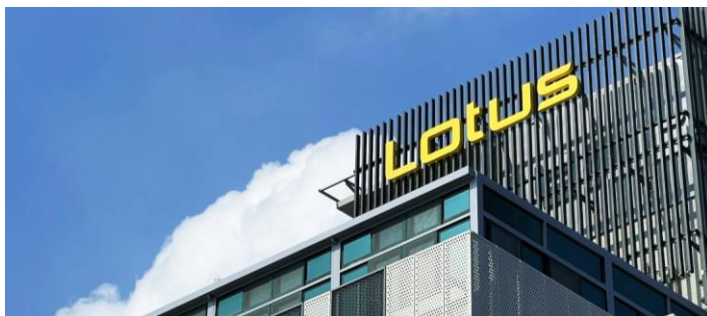
Singapore

- **Regional Hub** for majorly tender driven markets namely Singapore and Hong Kong **with own entity**, and partnerships in Pakistan, Bangladesh & Sri Lanka



High Quality Manufacturing Facilities

US FDA, EU EMA, PMDA,TFDA & PIC/S certified



Nantou Plant

Facility Size	84,000 sq. ft.
Production Capabilities	• Generic tablets, hard capsules and soft gel capsules • Generic high potency tablets, hard capsules and soft gel capsules
Current products	57 molecules / 480 SKU's
Capacity	• Maximum at c.500 million units (tablets/capsules) • Utilization rate ~70%



Hyangnam Plant

Facility Size	150,000 sq. ft.
Production Capabilities	• Tablets • Coated tablets • Hard capsules • Powders
Current products	54 molecules / 279 SKU's
Capacity	• Maximum at c.800 million units (tablets/capsules) • Utilization rate ~60%



Gongju Plant

Facility Size	75,700 sq. ft.
Production Capabilities	• Tablets • Coated tablets • Hard capsules • Powders, Granules
Current products	25 molecules / 83 SKU's
Capacity	• Maximum at c.500 million units (tablets/capsules/powders/granules) • Utilization rate ~50%

R&D capability

Two RD setups and talent 120 + scientists based in Taiwan/India/Korea

- › Fully functional site in Taiwan
- › New RD center (spread over 20000 sq feet) in India start functioning from Q3/Q4 2024.
- › Teams include Analytical, Product Development, Clinical, Intellectual property, Project Management, Development Quality
- › Fully-equipped laboratories with spray dryer, Hot melt extruder, Air jet mill, High shear granulator, HPLCs, GC, XRD, DSC, Lab scale soft-gel encapsulator, LCMS (under procurement) etc.

Well-defined product selection and robust pipeline w 10% revenue investment annually

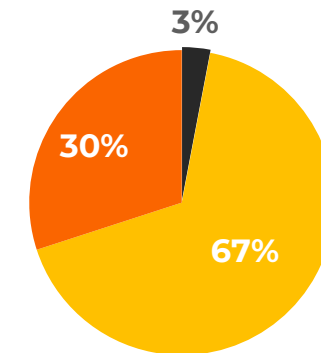
- › Close collaboration with Portfolio, business teams in identifying niche, difficult-to-develop oncological and non-oncological products
- › Product development for Global markets with a strong focus for US/EU/APAC
- › Portfolio encompasses a mix of Early development, FTF, generics and 505(b)(2)s

Integration of the R&D platform with solid oral (oncological and soft-gel) mfg facility certified by major regulatory authorities.

(US FDA/EMA/PMDA/TFDA/ANVISA/others)



Emp. Qualification



■ PhD ■ Post Grads ■ Grads and below

Strong B2B Network – ex-APAC

2019 To Date

265

Licensing
deals signed

160+

Markets

30+

Products

Cipla

europarma
sua vida move a nossa

glenmark
A new way for a new world

SANDOZ A Novartis
Division

JAMP
PHARMA GROUP

ZENTIVA

teva

STADA

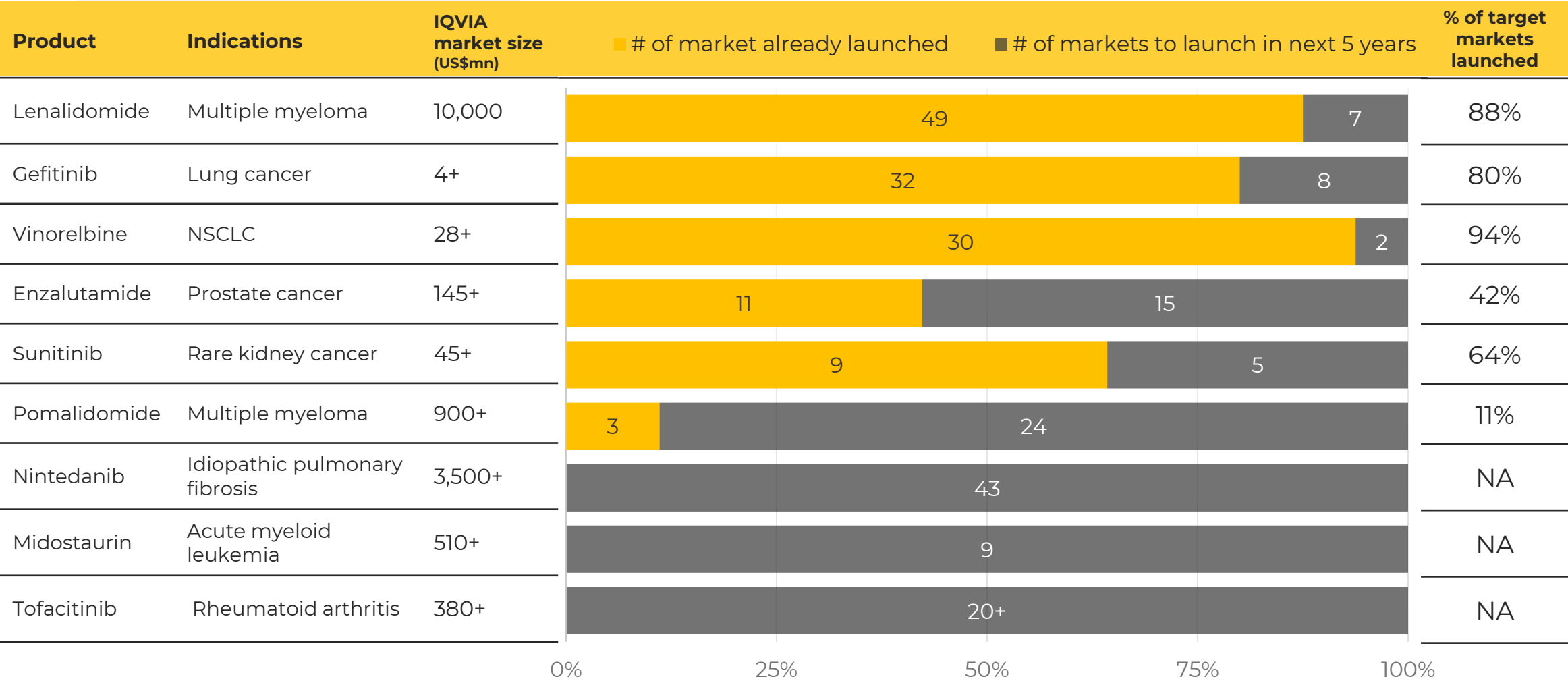
AUROBINDO
Committed to healthier life!

Dr.Reddy's

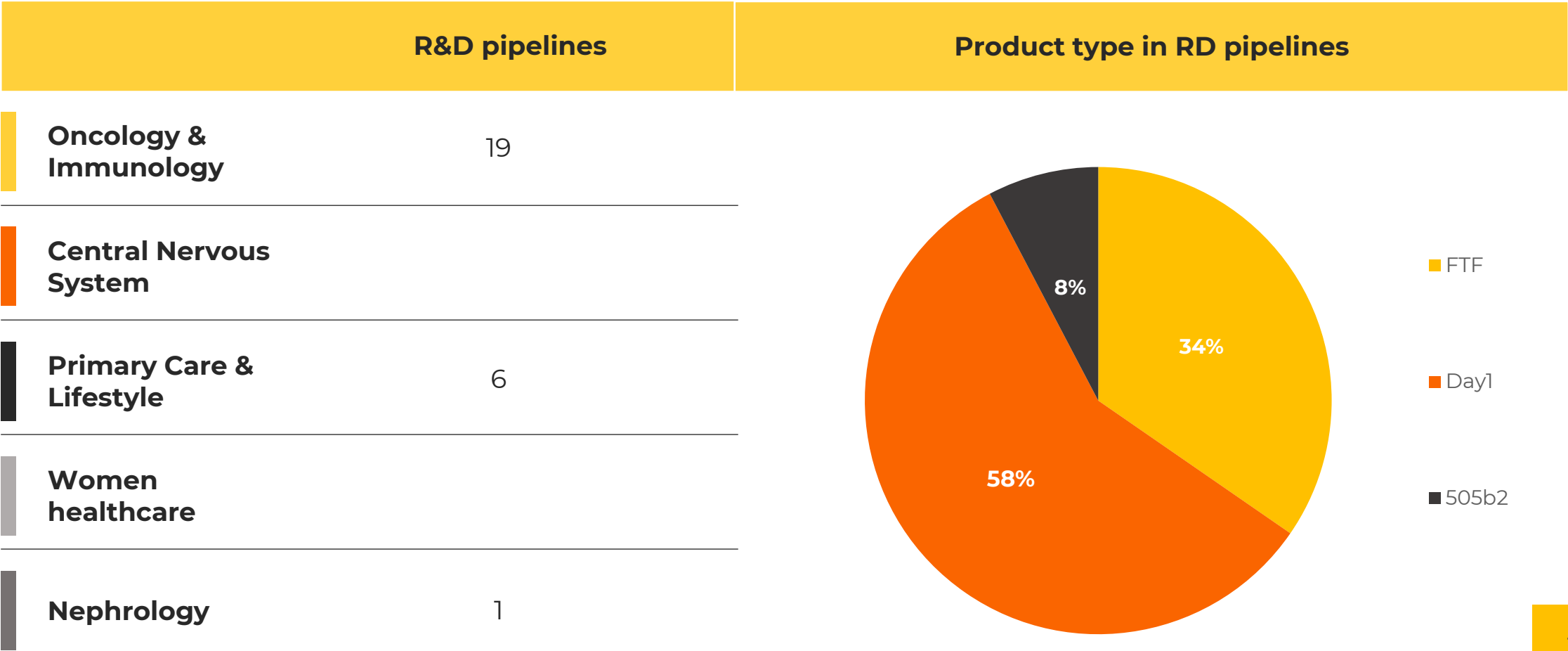
Alvogen

accord
We make it better

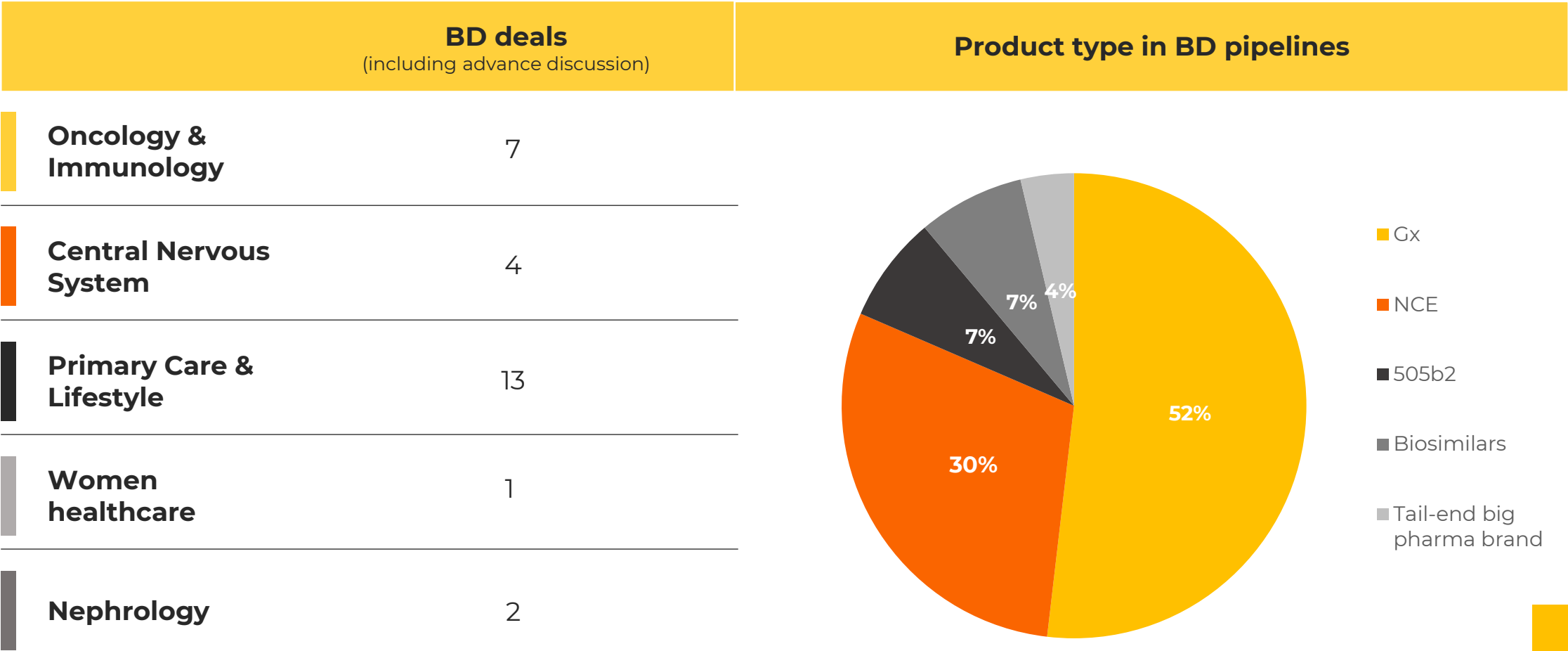
Global Rollout Of Commercial Products



High-barrier R&D pipelines focus on oncology products



Building Hybrid Pipeline Portfolio Through Strong Business Development Ability



Fueling Sustainable, Double-Digit Profit Growth

