

**** LUPIN

LUPIN LIMITED

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Materials and information provided during this presentation may contain 'forward-looking statements'. These statements are based on current expectations, forecasts and assumptions that are subject to risks and uncertainties which could cause actual outcomes and results to differ materially from these statements.

Risks and uncertainties include general industry and market conditions, and general domestic and international economic conditions such as interest rate and currency exchange fluctuations. Risks and uncertainties particularly apply with respect to product-related forward-looking statements. Product risks and uncertainties include, but are not limited to, technological advances and patents obtained by competitors. Challenges inherent in new product development, including completion of clinical trials; claims and concerns about product safety and efficacy; obtaining regulatory approvals; domestic and foreign healthcare reforms; trends toward managed care and healthcare cost containment; and governmental laws and regulations affecting domestic and foreign operations.

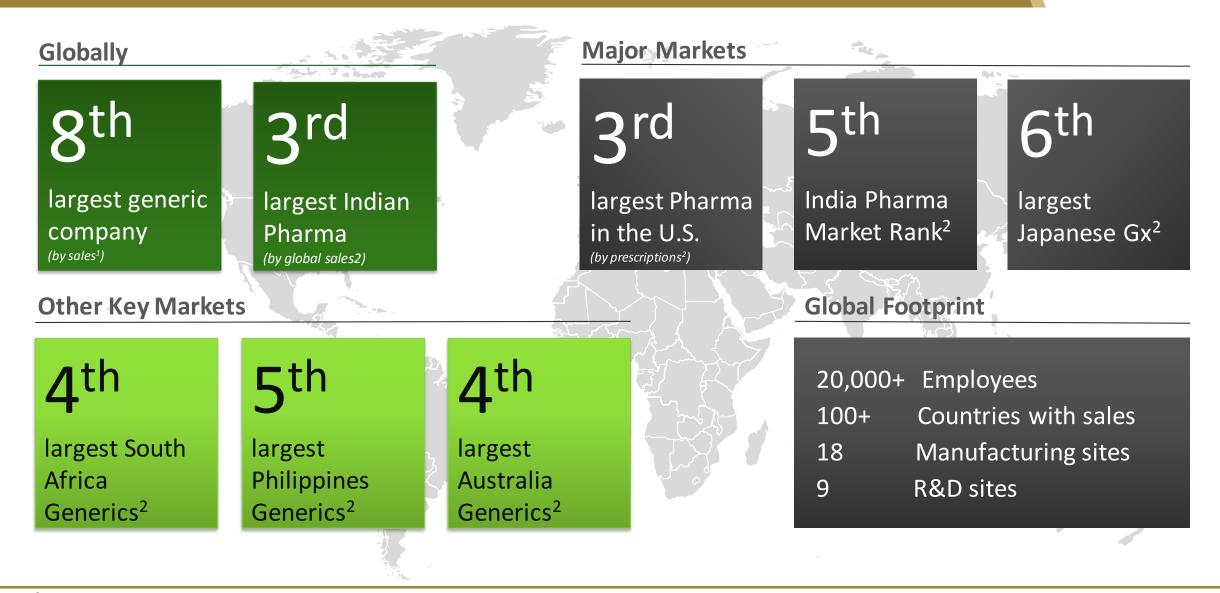
Also, for products that are approved, there are manufacturing and marketing risks and uncertainties, which include, but are not limited to, inability to build production capacity to meet demand, unavailability of raw materials, and failure to gain market acceptance.

You are cautioned not to place undue reliance on these forward-looking statements, which reflect our opinions only as of the date of this presentation.

The Company disclaims any intention or obligation to update or revise any forward-looking statements whether as a result of new information, future events, or otherwise.

Lupin today - Leading global pharmaceutical player





Strategic Vision





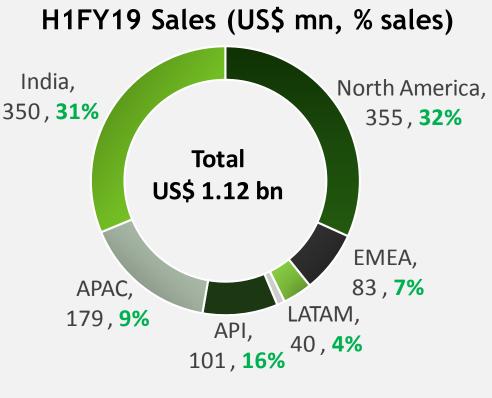
- Major revenue contributor currently
- Maximize on capability to maintain leadership in US generics
- Continue growth momentum in India and other emerging markets

- Deliver on key complex generics, esp. Inhalation and Injectables
- Continue filing of P4 and semiexclusive generics
- Successfully file and partner biosimilars

- Create a meaningful women's health franchise in US
- Neurology / CNS focus in other developed markets

Strong Foundation







2018 Highlights

US

- $\circ~$ US business starting to stabilize, and getting back on growth mode
- Average market share for our products 32.8%¹ (for Sep qtr.)
- Filed 35+ ANDAs and received 25+ approvals in 2018
- Cumulatively 165+ products marketed, 150+ ANDAs awaiting approval
- 41 FTFs incl. 14 exclusive FTFs awaiting approval
- Successfully launched Solosec on the Specialty front

India

- Continue to outpace industry growth and gain market share
- Leadership across cardiac (#3), diabetes (#3), and respiratory (#2)
- >15 alliances, supports faster chronic segment (58% of revenue)growth;
- Expanded diabetes partnership with Boehringer Ingelheim and Eli Lilly

Other markets

- Focus to drive organic growth, attain scale and self-sustain
- Japan (market leader in CNS¹), South Africa, Mexico, Brazil (launched derm/aesthetics business), Australia, Germany are key markets

Complex Generics Pipeline



17Inhalation13• First MDI & DPI filed with more (6 MDI/2 DPI) under development • Albuterol MDI (gProAir) filing under FDA review • Tiotropium DPI (gSpiriva) filed, FTF confirmed33Biosimilars6• bEtanercept filed in EU and Japan • Partnered with Mylan (EU & other markets) and Nichi-Iko (Japan) • Pegfilgrastim- US clinical studies underway; advancing other programs12Injectables (incl. Complex Inj.)>30 (10 depot inj.)• 4 injectable products approved in 2018 • Advancing multiple complex Inj. (depot, peptides, iron products)	Target Market Size ¹ (US\$ bn)	Product segment	No. of Products	Pipeline Progress
 Biosimilars Biosimilars Partnered with Mylan (EU & other markets) and Nichi-Iko (Japan) Pegfilgrastim-US clinical studies underway; advancing other programs Injectables (incl. Advancing multiple complex lpi (depet perioducts) 	17	Inhalation	13	 Albuterol MDI (gProAir) filing under FDA review
12 (incl. 30 • 4 injectable products approved in 2018 (incl. (10 denot ini.) • Advancing multiple complex ini. (denot, poptides, iron products)	33	Biosimilars	6	 Partnered with Mylan (EU & other markets) and Nichi-Iko (Japan)
	12 (incl. (10			

Products in development target US\$ 104 bn sales, of which complex categories account for ~70% (incl. complex orals, other dosage forms, biosimilars)

Specialty - U.S. Women's Health





Specialized salesforce

- 133 dedicated WH Sales Reps
- 70% of Reps with an average 7 years WH experience

Unique positioning

- Solosec[™] First and only single oral-dose treatment for Bacterial Vaginosis
- Designated as a Qualified Infectious Disease Product (QIDP) with 10 years of marketing exclusivity

Managed Care traction

- 93% Payer coverage 68% unrestricted
- Continued focus on generating pull through at Physician offices and Pharmacy

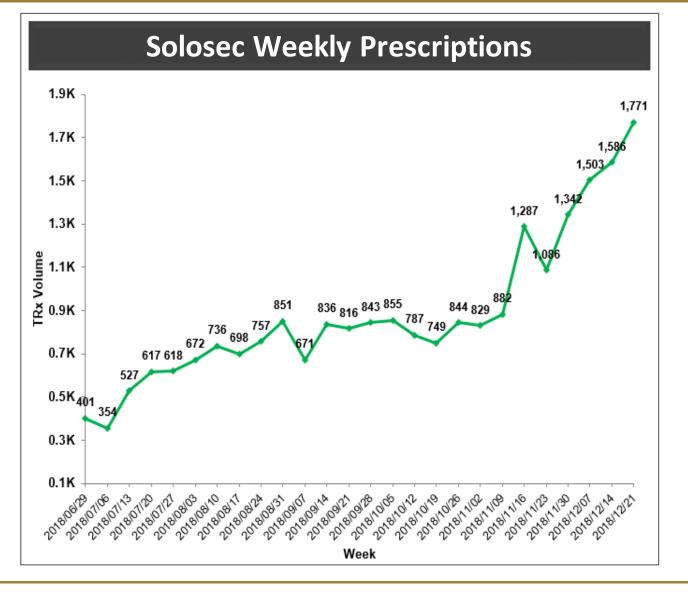
Business Development / Medical

- Experienced BD and Medical team actively focused on acquiring WH assets
- LCM/ label expansion efforts for Solosec underway

Established top notch Women's Health leadership team, Solosec launched successfully Foundation in place to build a leading Women's Health Business

Solosec Rx Trends





21,678 Solosec prescriptions dispensed since launch

1,700+ Solosec prescriptions dispensed per week (Dec 21) with a strong upward trajectory

6,258 Solosec unique prescribers since launch

3,282 Solosec repeat prescribers

ONE PACKET

ONE DOSE.

ONE TIME.

Specialty ex US





NaMuscla[®]'s Orphan drug designation ratified as first EU treatment for myotonia
 Obtained EMA Marketing Authorization in December 2018 for symptomatic treatment of myotonia in adults with non-dystrophic myotonic (NDM) disorders
 Planned launch in UK and Germany in Q1 2019. Partnering discussions ongoing for commercialization in other European territories



- Bipresso[®] was launched as the first specialty new drug from Lupin Japan in October 2017, indicated for Bipolar Depression
- Bipresso[®] listed in the formularies of the top university hospitals
- Q2FY19 sales grew 36% QoQ



AbbVie / MALT1 Partnership

abbvie

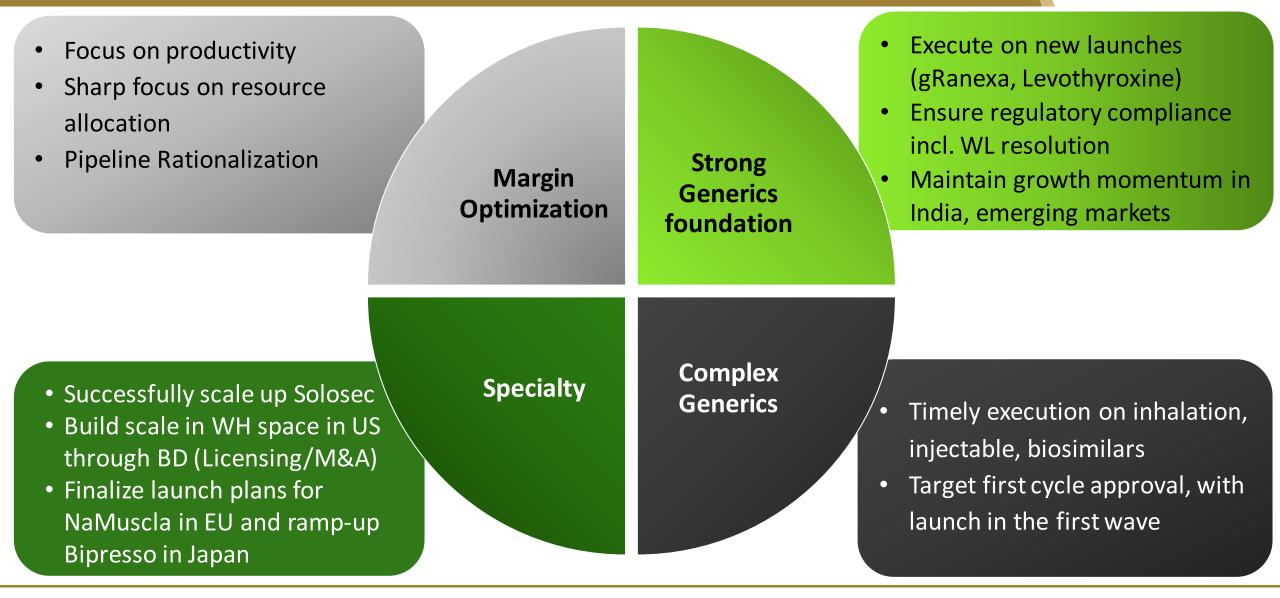
- In December 2018, AbbVie licensed Lupin's MALT1 (Mucosa-Associated Lymphoid Tissue Lymphoma Translocation Protein 1) Inhibitor Program
- AbbVie intends to pursue development across a range of hematological cancers
- AbbVie will pay Lupin US\$ 30 mn Upfront for an exclusive license
- Lupin is eligible to receive milestone payments of up to US\$ 947 mn and double digit royalty on sales

NCE Portfolio

Therapeutic Area	Product	Development Stage	Target Indication
Endocrine	Calcium Sensing Receptor PAM	Phase IIa PoC	1 st and 2 nd Hyperpara- thyroidism
Oncology	MEK Inhibitor NAM	Phase IIa PoC	Solid Tumors
Oncology	STING Agonist	Lead Identification	Solid Tumors & Lymphomas
Oncology	PRMT5 Inhibitor	Lead Identification	Mantle Cell Lymphoma Pancreatic Cancer

2019 Priorities









THANK YOU!