

Investor's Presentation

February 2023





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International business ~75%

* Excludes Promoter loans

11 manufacturing facilities across world

2 R&D centres one each in India, UK, USA

~ 520 scientists with >80 PhD's, Drug Discovery team > 150 associates¹

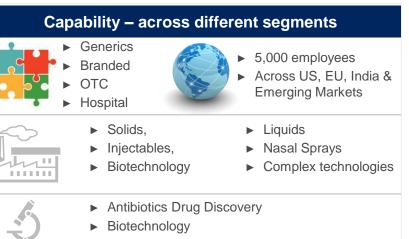
>20% of the 5,400 global associates based outside India¹



3,214 patents filed; 793 patents granted

Global Footprint : Significant Regulated market presence





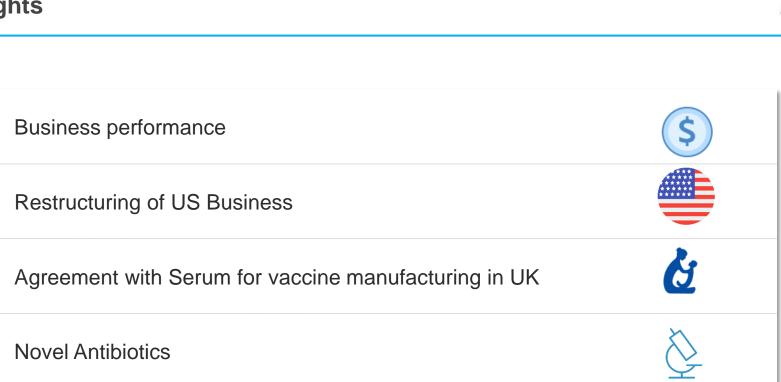
Pharma Generics



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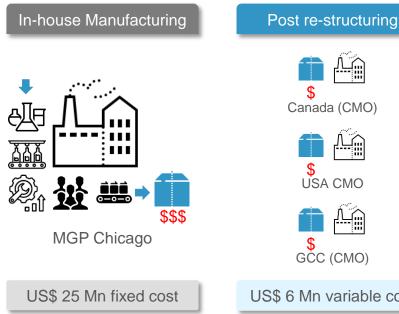
US Business Restructuring





US Cost Improvements

Site transfer from in-house Chicago site to CMOs











US\$ 6 Mn variable cost





Shut down of manufacturing facility at Morton Grove



Few products with high margin manufactured by 3rd party



Continue to maintain sales with ~ 40% gross margins

Annual savings of ~US\$ 12 Mn



Agreement with Serum for vaccine manufacturing at UK Facility 👌



- Joint venture of 51:49 in favor of Wockhardt
- Received £ 10 million as contribution for reserving capacity- 150 million doses across 15 years
- Serum has identified two vaccines

Wockhardt plans to manufacture post regulatory approvals and exhibit batches in next 8 - 12 months

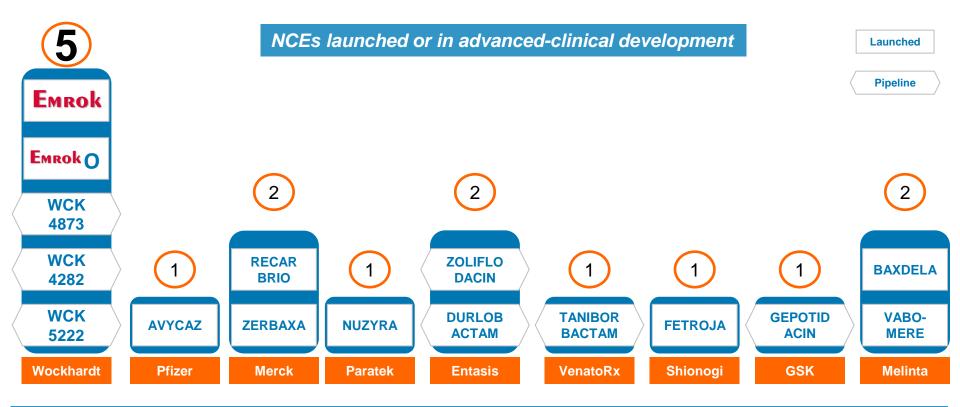




20 years	Committed to Novel Antibiotics research
#1	Probably only company to have comprehensive end to end drug discovery programs in Antibiotics
6	Programs granted QIDP status by US FDA denoting unmet needs; faster trials, quicker approvals by US FDA







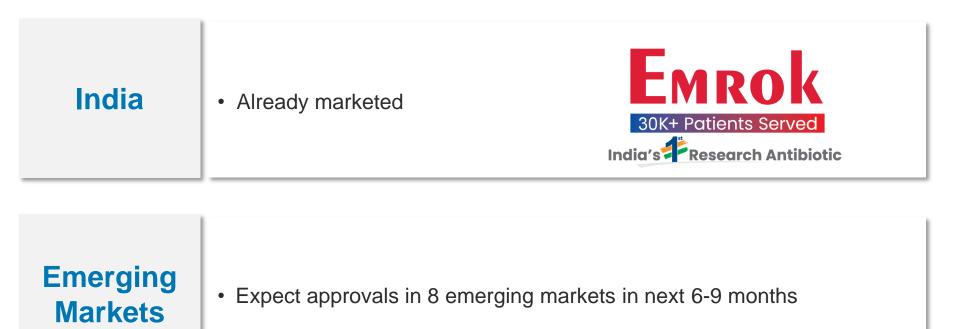
Shrinking pipeline - A result of scientific challenge and missing discovery talent base

trials

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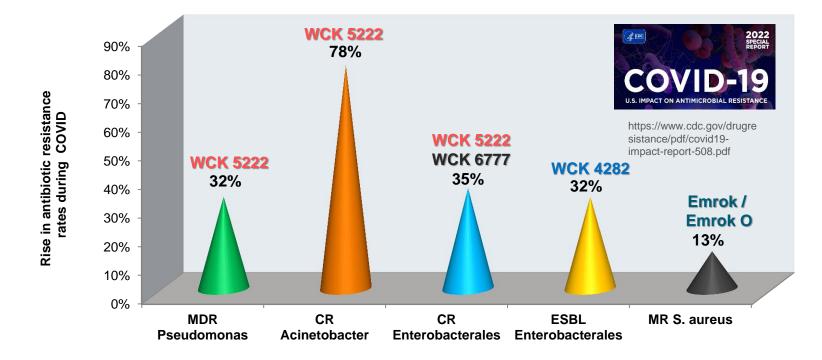
Licensing in China	 Signed licensing deal with Jemincare, China for development and commercialization in Greater China Received first milestone from Jemincare 		
India Phase III	Phase 3 clinical trial concluding by September 2023		

• Marketing in India in 2024 post DCGI approval



WCK 5222- Flagship novel antibiotic

Increase in	 2022 CDC report highlights rise in Anti-microbial Resistance during COVID-19
potential	Heightened vigilance reveals the rising level of threat



Global	 Global Phase III clinical trial initiated in August 2022 and expected to
Phase III	complete in next 15-18 months.
progress	 Expect to market product in USA, Europe, China & India in 2025

Saving	 3 critical patients; all other antibiotics options exhausted including recent global innovation product 		
lives	 Post DCGI approval for compassionate use, successfully cured and discharged on completion of treatment with WCK 5222 		



Recently under DCGI Permission 3 patients in India with life-threatening infection received WCK 5222 on compassionate ground since available antibiotics were not efficacious

Patient 1 (Sep. 2022)

- 50 y, female admitted in Medanta, Hospital Lucknow, treated by Dr. Dilip Dubey
- Post-abdominal surgery sepsis
- Infection of XDR Pseudomonas, resistant to ceftazidime/avibactam
- Though susceptible to colistin, even 4 weeks of therapy didn't improve the clinical condition
- 10 days of therapy with WCK 5222 rescued the patient paving for discharge in stable condition

Patient 2 (Nov-Dec. 2022)

- 11 y, boy admitted in Global Hospital, Chennai, treated by Dr. Ponni/Subramanian
- Post bone-marrow-transplant febrile neutropenia
- Infection of XDR *E. coli*, resistant to ceftazidime/avibactam
- Last-line therapies were not efficacious
- 2 weeks of WCK 5222 eradicated the pathogen; discharged in stable condition

Patient 3 (under treatment)

- 18 y, male admitted in AIG Hospital, Hyderabad, treated by Dr. Balasaheb
- Necrotic fasciitis after chemotherapy for acute leukemia
- Infection of XDR Pseudomonas, resistant to ceftazidime/avibactam
- Last-line therapies were not efficacious
- 10 days of WCK 5222 eradicated the pathogen; being treated for cytopenia

WCK 6777



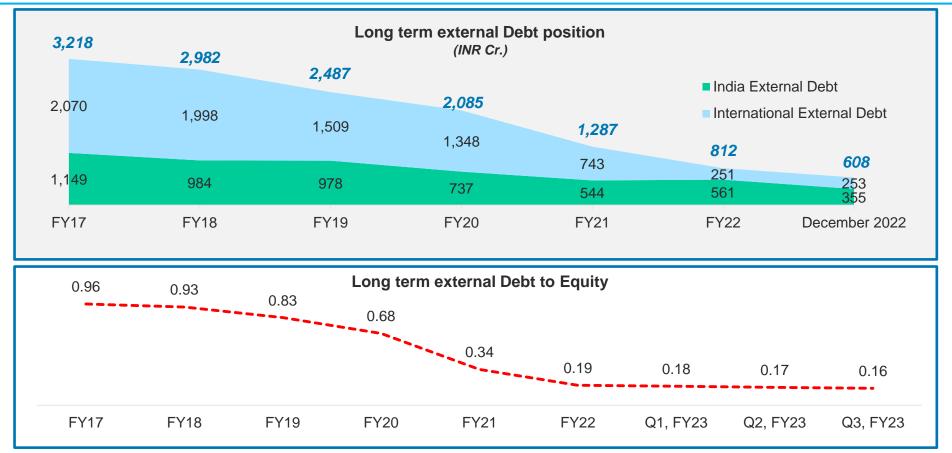
NIH National Institutes of Health

Funding • By National Institute of Health, USA

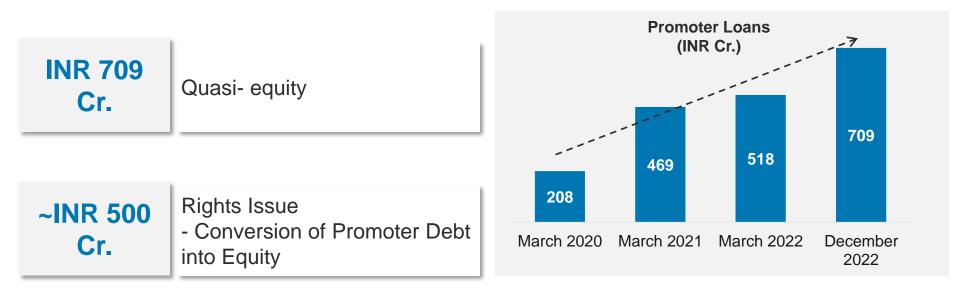
Stage	In Phase I US development	
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Financial / Business Highlights

Deleveraging: Reducing Long term external Debt



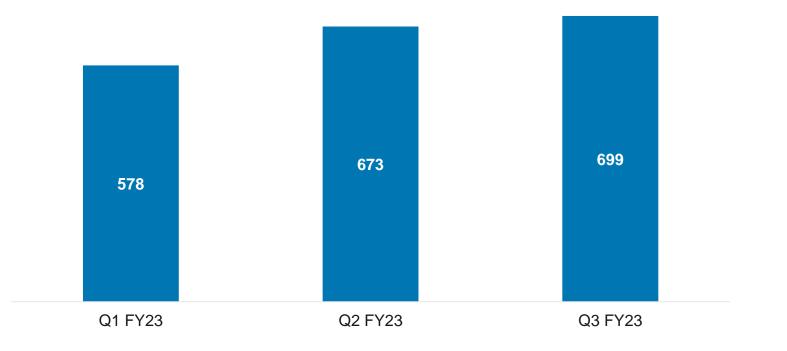


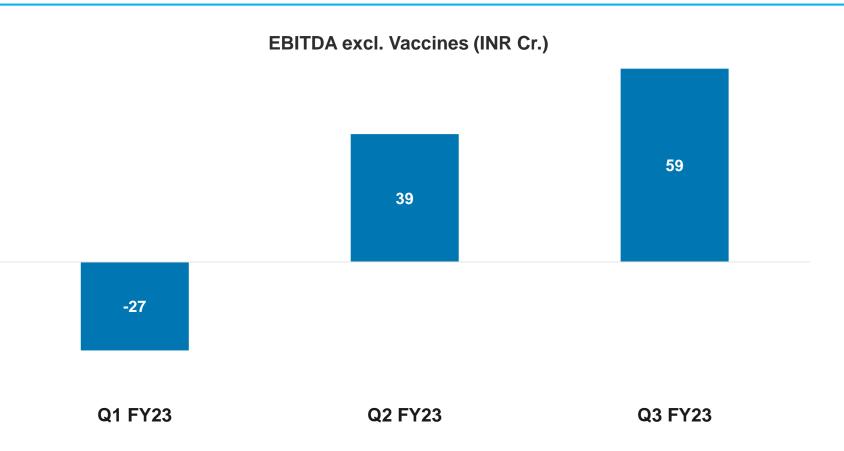


Promoter is fully committed to support the Company









Cash Flow Movement



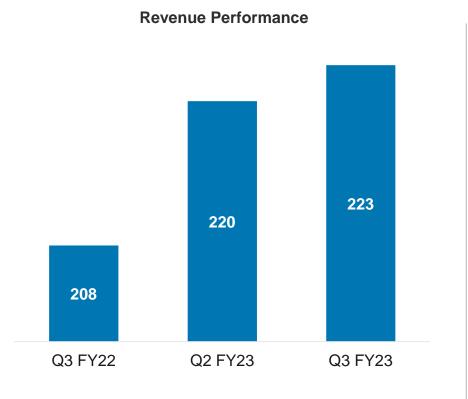
Q3 FY23 (INR Cr.)



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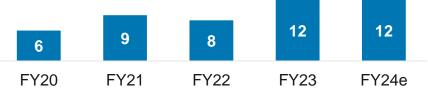




Key Highlights:

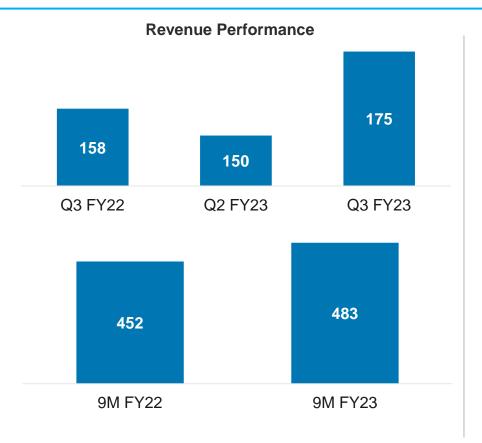
- Comprehensive, FDA-approved manufacturing facility for sterile injectables.
- Present in key therapies across Cardiac, Pain & Analgesics, Vitamins, Respiratory & Diabetes, Oncology, Anti-Infectives
- Collaboration with UK Government and Astrazeneca to manufacture COVID-19 Vaccines.

Accelerated new filings







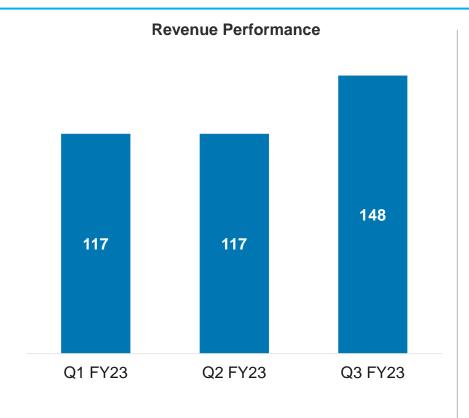


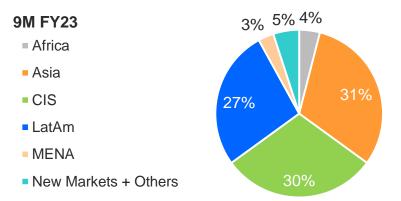
Key Highlights:

- Therapeutic focus across Diabetes, Nephro, Anti-Infectives, Pain / Analgesics, Vitamins & Nutrients, Neuro / CNS and others.
- One of the few Indian Companies who are into Manufacturing of Insulin, Glargine & OSD in Diabetes
- Robust Distribution Network CNF (20+)/ Stockiest / Distributors (~ 2000)
- Only Indian Pharma company to Launch novel NCE in India (Brand name – Emrok & Emrok O)
- On 9 months basis, Diabetes Biosimilar business grew >20%
- Launched Sitagliptin under brand name "Sitawok"

Emerging Markets Business







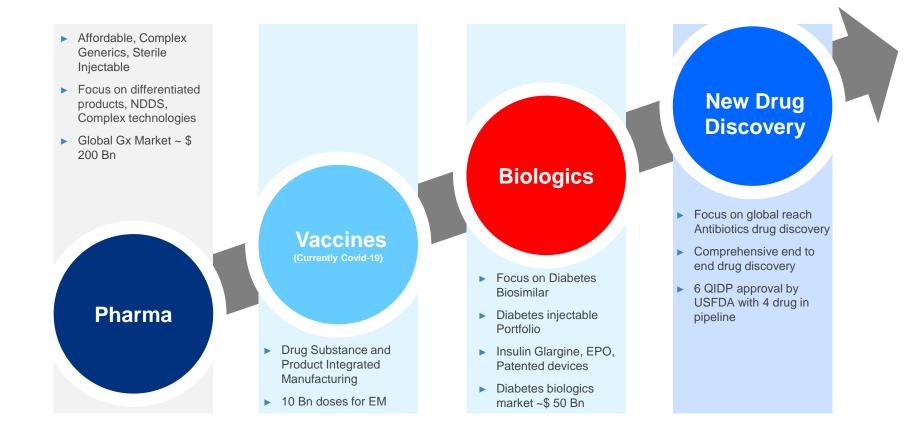
Key Highlights:

- De-risked model of Operations partnerships & relationship building.
- Differential & end to end Business model from Research to Manufacturing to Distribution with backward integration of API manufacturing.
- Strong Presence in Diabetes Segment Insulin, Glargine : both Vials & Cartridges
- Upcoming Focus Presence in New Markets / New Segments with high growth potential

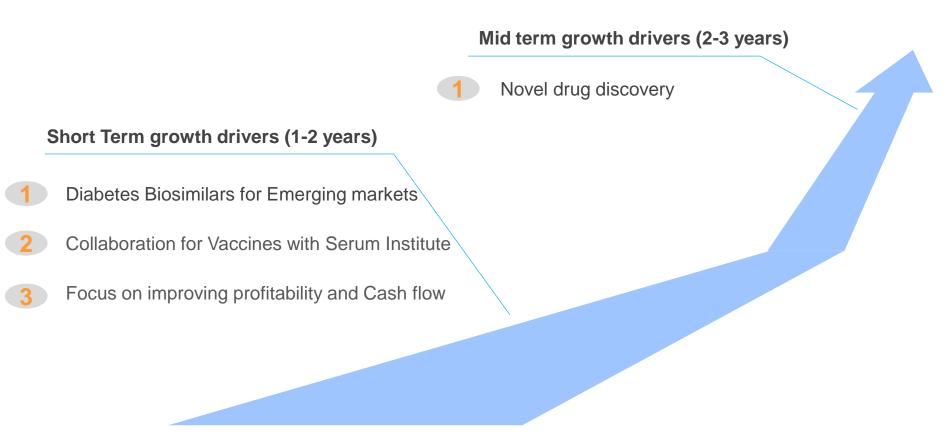


Growth Drivers







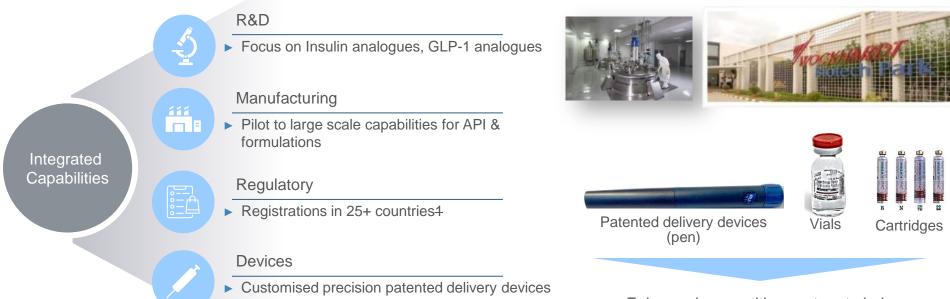






Competitive advantage by providing end to end solutions in Diabetes Biosimilars





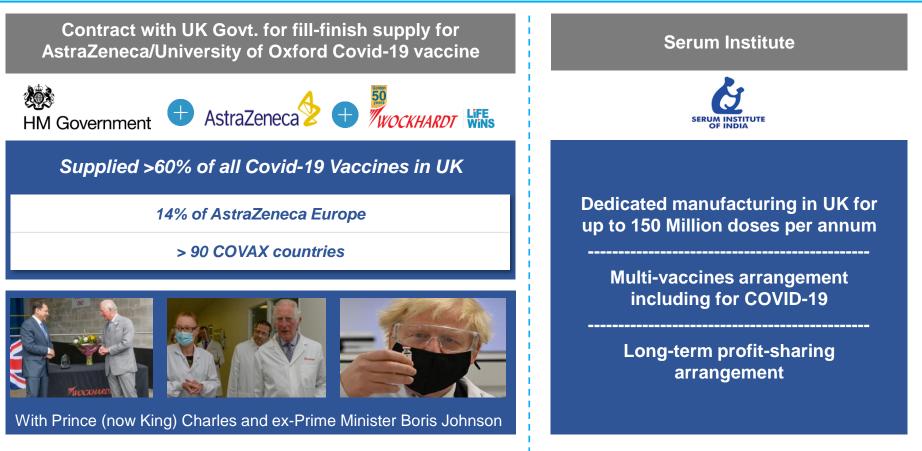
Enhanced competitive posture to help penetrate markets

Pipeline of Insulin Analogs including Insulin Aspart

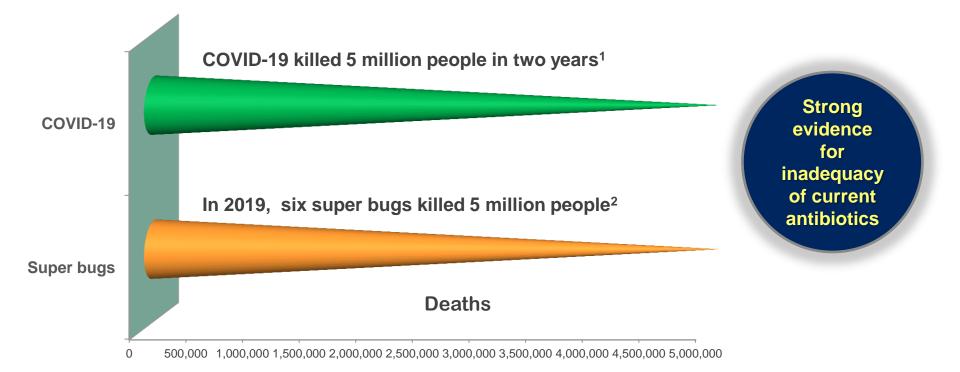
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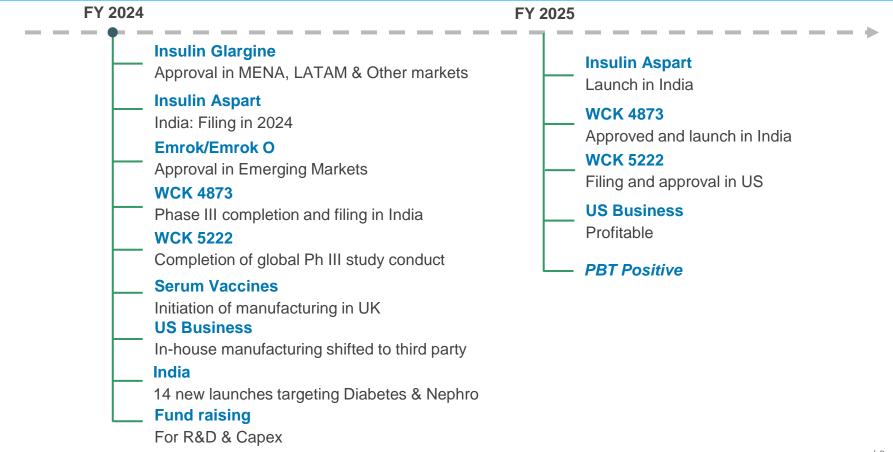
1.Nature, Vol 601, 20 January 2022; 2.Lancet. 2022 Feb 12;399(10325):629-655

3 Novel Antibiotics pipeline against Resistant Organisms

	WCK 5222	WCK 4282	WCK 4873	Emrok / Emrok O	WCK 6777
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Phase	Phase III (Global)	Phase III (Global)	Phase III (EM) Filing 2023	Launched in India EM Filing underway	Phase I
Potential Indication	cUTI HABP / VABP	cUTI HABP / VABP	CABP / RTI	ABSSSI	cUTI
			Partnered in China	Partnered in Russia	Funding by NIH
Positioning	Destination therapy for XDR Gram-ve Acinetobacter and Pseudomonas	Empiric-use; Carbapenem- sparing Gram-ve	Macrolide-resistant Respiratory Pathogens, Quinolone-Sparing	MDR Gram+ve Anti-MRSA	Out-patient therapy for MDR Gram -ve
Current Market Size for target n	At current a	generics	US\$ 5.6 Bn At current generics price	~US\$ 700 Mn At current generics price	~US\$ 3.4 Bn At current generics price

Our journey ahead – target key milestones





Life wins !





