

REGISTERED OFFICE

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CIN: L24110TG1991PLC012471

Dated October 23, 2019

To,

National Stock Exchange of India Limited

BSE Limited

Symbol: NSE: GRANULES; BSE: 532482

Dear Sir,

Sub: Presentation to the Analysts/Investors

We refer to Un-audited financial results for the second quarter and half-year ended September 30, 2019 submitted to you on 22nd October 2019.

We are now enclosing the presentation in this regard to the Analysts/Investors which is also being uploaded on our website.

This is pursuant to Regulation 46 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015.

Kindly take the above information on record.

Thanking you.

Yours faithfully,

Y. (hait our flower

FOR, GRANULES INDIA LIMITED

(CHAITANYA TUMMALA) COMPANY SECRETARY &

COMPLIANCE OFFICER



Earnings Presentation

Q2FY20



Agenda



- 1 Q2FY20 update
- **2** About Granules
- **3** Performance metrics
- 4 Way forward

Safe harbour



The Presentation is to provide the general background information about the Company's activities as at the date of the Presentation. The information contained herein is for general information purposes only and based on estimates and should not be considered as a recommendation that any investor should subscribe / purchase the company shares. The Company makes no representation or warranty, express or implied, as to, and does not accept any responsibility or liability with respect to, the fairness, accuracy, completeness or correctness of any information contained herein.

This presentation may include certain "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. Important factors that could cause actual results to differ materially from our expectations include, amongst others general economic and business conditions in India, ability to successfully implement our strategy, our research and development efforts, our growth and expansion plans and technological changes, changes in the value of the Rupee and other currencies, changes in the Indian and international interest rates, change in laws and regulations that apply to the Indian and global pharmaceuticals industries, increasing competition, changes in political conditions in India or any other country and changes in the foreign exchange control regulations in India. Neither the company, nor its directors and any of the affiliates or employee have any obligation to update or otherwise revise any forward-looking statements. The readers may use their own judgment and are advised to make their own calculations before deciding on any matter based on the information given herein.

No part of this presentation may be reproduced, quoted or circulated without prior written approval from Granules India Limited.

Agenda

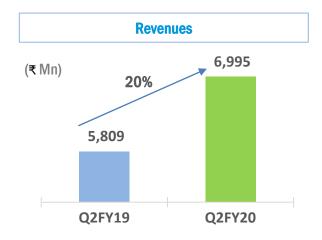


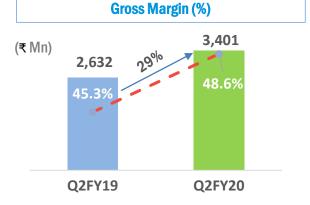
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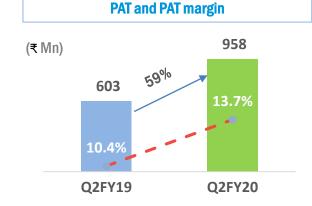
Q2FY20 Financial highlights







EBITDA and EBITDA margin (₹ Mn) 1,436 17.3% Q2FY19 Q2FY20



Key Highlights

· Gross Revenues:

- Revenues were up 20% YoY on account of PFI sales growth of 37% and FD sales growth of 34%
- Increase in regulated markets' sales by 34%

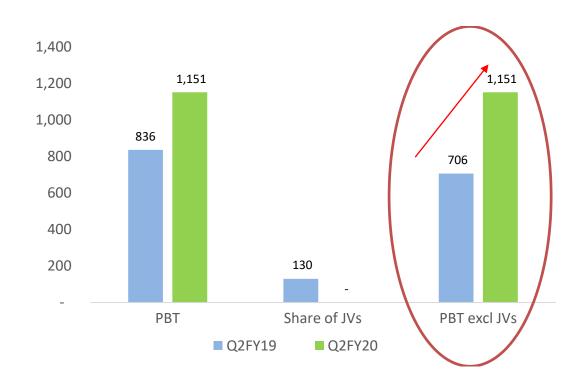
• PAT:

 PAT has increased due to an increase in overall sales volumes and a reduction in tax rate

Profit Reconciliation



(₹Mn)



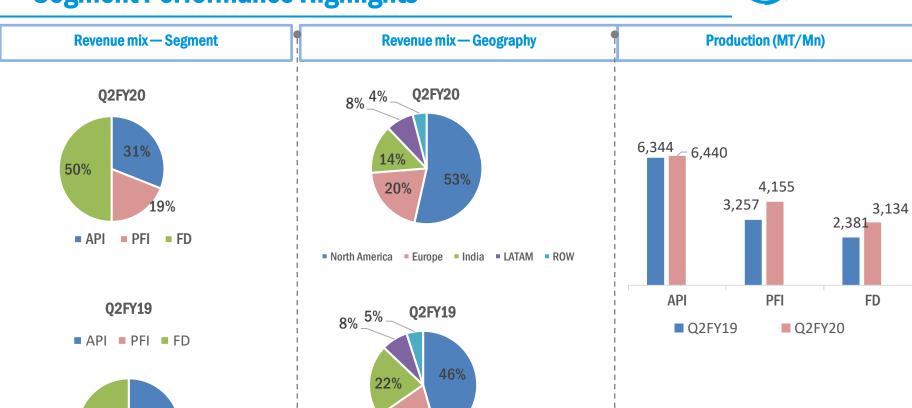
63% YoY growth in PBT despite divestment of JVs

Segment Performance Highlights

45%

16%





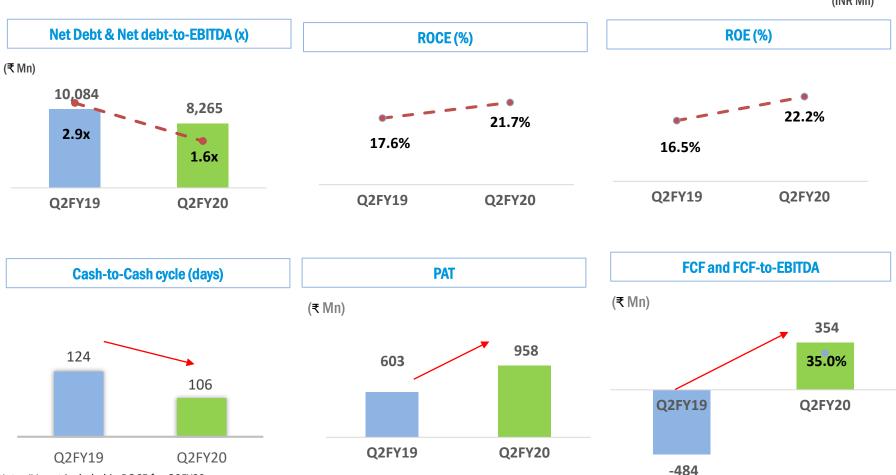
North America
 Europe
 India
 LATAM
 ROW

20%

Continued focus on improvement



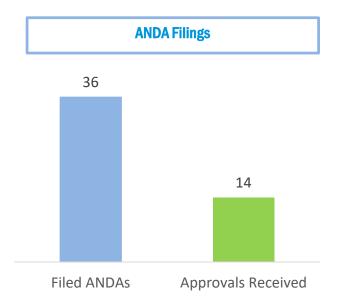
(INR Mn)



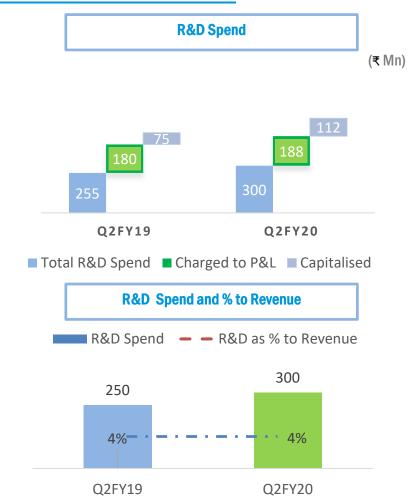
Note: JVs not included in ROCE for Q2FY20

Other Business Highlights





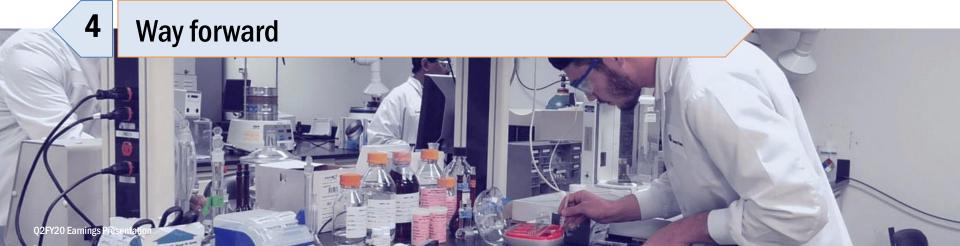
- Two ANDAs filed during the quarter One each from GPI and GIL
- One ANDA approved during the quarter



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Granules at a glance





Building Blocks

Regulatory compliance
Operational efficiency
Customer centricity

Scales of operation



Installed Capacity

39,360 TPA of API 290 KL 24,640 TPA of PFI 21.3 Bn dosages of FD



Regulatory Approvals

USFDA, EDQM, EU GMP, COFEPRIS, WHO GMP, TGA, KFDA, DEA, MCC, HALAL



Business Divisions

Core business

US generics

Multi API and Onco



Global Reach

60+ countries

250+ customers



Revenue Mix

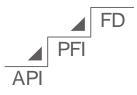
80% of revenues are exports

66% of revenues from regulated markets of US, Europe and Canada

LATAM contributes nearly 10% of revenues



Business Verticals



Vertically integrated across value chain



People Strength

2,789 people as on 31 March 2019 in GIL



Intangible Assets

36 ANDA filings; 14 approved

17 US & 6 European DMFs

14 CEPs with EDQM

10 other country's DMFs



Manufacturing Base

6 Operational facilities; 5 in India, 1 in US

One of the world's largest paracetamol API facilities

World's largest PFI facility

One of the largest singlesite FD facilities

Business Structure — Granules India









Focus on volume based products such as Paracetamol, Metformin HCl, Ibuprofen, Guaifenesin, Methocarbamol, Naproxen and newly added Cetrizine and Fexofenadine

Large scale manufacturing enables sustained competitive advantage & secured supply source for the customers

Pioneered the concept of commercializing PFIs — suits large volume drugs

Auxiliary growth engine to secure the future

Multiproduct and multi-stage API-to-FD manufacturing facilities in Vizag with special focus on Oncology

Fully-integrated facility to offer APIs and FDs in onco therapeutics, a top growth segment

Fully integrated infrastructure including R&D, manufacturing and marketing to enable "Make in America"

Business-to-Consumer model (B2C)

Focus on developing niche/differentiated modified and immediate-release products in varied dosage forms

Excellence in quality & being best-in-class



Granules India

Incorporated in 1991, this is the only listed entity in the group, with 5 operational plants located at Hyderabad (Jeedimetla, Bonthapally and Gagillapur) and one at Vizag Pharma City Granules USA Inc

Granules Pharmaceuticals Inc

Granules Europe Ltd

100% subsidiary with front-end marketing for OTC products in US

100% subsidiary focused on advanced formulation development, with manufacturing set-up and front-end marketing in Rx products at Chantilly, US

100% subsidiary for front-end marketing in Europe

Value chain	Facility location	Installed capacity	Approvals	
API	Bonthapally	34,560 TPA	US FDA, EDQM, WHO, COFEPRIS, INFARMED	
	Jeedimetla	4,800 TPA	US FDA, EDQM, COFEPRIS, WHO, CDCSO	
	Vizag	Vizag 290 KL US FDA, KFDA, EU GMP,		
PFI	Gagillapur	Gagillapur 23,200 TPA US FDA, COFEPRIS, TGA,		
	Jeedimetla	1,440 TPA	WHO GMP, COFEPRIS, INFARMED	
FD	Gagillapur	19.8 Bn	US FDA, MCC, COFEPRIS, TGA, INFARMED	
	Virginia, USA	1.5 Bn	US FDA, DEA	
API Intermediates	Bonthapally	61.5 KL		

Core Business



Overview

Presence across the entire pharmaceutical manufacturing value chain from API to finished dosages

Large-scale manufacturing enables not only sustain competitive advantage but also build operational efficiencies into the system

Focus on first line of defence portfolio products such as Paracetamol, Metformin HCl, Ibuprofen, Guaifenesin,

Methocarbamol, Cetirizine and Fexofenadine

Pioneered the concept of commercializing PFIs which is for large volume molecules

3 US FDA-approved plants located in Hyderabad (Jeedimetla, Bonthapally and Gagillapur)

Drivers		Value enablers		
		Among the world's largest PFI and single-site FD facilities API capacity 39,360 MT; PFI 24,640 TPA; finished dosages 19.8 billion		
Scale		DMFs, Dossiers, ANDAs for APIs and finished dosages, which can be repacked and marketed by customers under their own brand		
Presence across value Chain		Fully integrated in key molecules from APIs to finished Dosages		
	>	Selective addition of molecules to the portfolio		
Expanding portfolio		Enhance existing products by introducing controlled released products		
portiono		Access to value-added products such as rapid-release caplets and bi-layer tablets		
		All plants are approved by FDA and other regulatory bodies		
Regulatory complianc		Regular customer audits ensures continuous compliance		
		Dedicated resource to monitor and improve compliances		

Multi APIs/Oncology



Overview

Multiproduct and multi-stage API-to-FD manufacturing facilities in Vizag with special focus on Oncology

Fully integrated facility to offer APIs and FDs in onco therapeutics, a top growth segment

Value enablers **Drivers** Multiproduct API and oncology manufacturing facility on over 18 acres of land Infrastructure Integrated API-to-FD facility for oncology IT-enabled infrastructure to ensure efficiency and compliance Supplying to domestic and international markets with own and **Customer centricity** customer products Regulatory Products are under stability and expect faster site approval once filed record Robust product pipeline for other limited competition APIs. **Portfolio** 30 techpacks ready for onco block

US Generics



Overview

Fully integrated infrastructure from R&D, manufacturing to marketing enabling "Make in America"

Balanced product portfolio, combination of internal capabilities and market dynamics

Strategic portfolio selection focusing on limited competition in Para 3

US FDA approved facility in Virginia with a team strength of 90+

Access to talent pool in the US

Business to Consumer — front end sales and marketing for Rx & OTC products

Drivers

Value enablers

"Niche products," hidden opportunities with different layers of complexities

Portfolio



Differentiated dosage forms; limited competition

Complex generics; products within the modified release solid oral dosage forms

Controlled substances; Schedule 2 to Schedule 5

Differentiated technology platforms across the solid oral category

Capabilities



Competent teams across all departments; combination of experiences from global to more small to mid tier pharmaceuticals companies

100,000 sft facility set up with established R&D and manufacturing capabilities with all necessary approvals required for the development and manufacturing of controlled substances

Balanced portfolio with a combination of high volume and "niche," products

Integration across the value chain for development; from API, PFI to FD

Integration across the value chain; from R&D to front end commercialization

Focus on quality and compliance

Granules' advantage

Joint Venture divestments



Granules Omnichem

Investment sold to JV partner for INR 1,098.5 mn

Entered in 2012

INR 500 mn: Book value of investment as on 30 June 2019

Reason for exit

Does not fit in our evolving business model

JV resulted in lower capacity utilization and lower return on capital

Granules Biocause

Decision to sell investment to JV partner

Entered in 2007

INR 1436.5 Mn: Book value of investment as on 30 June 2019

Reason for exit

Large investments required for Chinese environmental compliances

Improved availability of Ibuprofen API in domestic and international markets makes JV unnecessary

Core management team





C Krishna Prasad Chairman and Managing Director

Mr Prasad's journey as pharma entrepreneur began in 1984, when he set up a paracetamol API manufacturing facility that focused on capital and process efficiency. Pharmaceutical formulations intermediates (PFIs) as a cost efficient product for global formulations manufacturer is a concept pioneered and popularized by Mr Prasad.



Uma Devi Chigurupati
Executive Director

Experienced over 30 years in various fields, Ms Uma cofounded with Mr Prasad Triton Laboratories in 1984, which was later amalgamated with Granules. Currently, she heads Granule's CSR activities and HR initiatives.



Harsha Chigurupati
Executive Director

Responsible for Sales, Marketing and Manufacturing Operations of Granules India Limited. He will drive business performance through operationalising Company strategy into business plan, conducting periodic operations reviews and driving corrective - preventive measures to bridge gap or enhance performance.



Priyanka Chirugupati
Executive Director
Granules Pharmaceuticals. Inc.

Responsible for the US Generics business including commercial and strategic initiatives. Ms Chigurupati's numerous roles in her 5 years at Granules' divisions in the US and India include the Core Business, Emerging Business and Consumer Health.



GSR PrasadChief Operating Officer

Responsible for all the manufacturing operations of GIL. He will continue to lead Projects related responsibilities too. GSR Prasad is Masters in Pharmacy and possess more than 30 years of rich experience in Manufacturing, Engineering, SCM and Projects. He joined us as Head of Projects in January 2017. Earlier, he worked with various pharmaceutical companies like Astra Zeneca, Themis, Sangfroid Granules, Pharmeng, Biovail, and Novast Laboratories in India, US, Canada and China.



Mr. Stefan Lohle
Chief Marketing Officer

Mr. Lohle has over two decades of experience in the Pharmaceutical industry. Mr Lohle has been associated with Granules since 2001 and previously was Head of Latin American Operations, where he primarily focused on the PFI Business. Mr Lohle previously served at Kimberly Clark Corporation for New Project Development



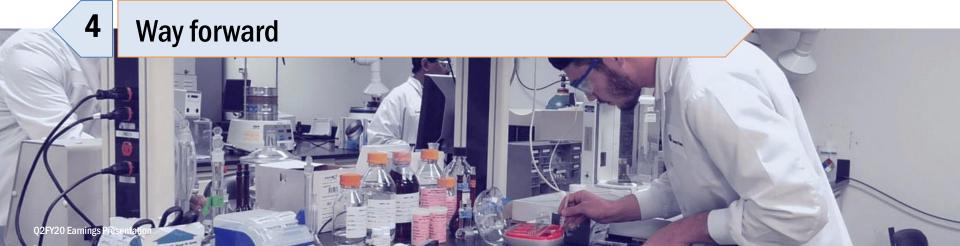
Mr. Atul Dhavale
Chief Human Resources Officer

Mr. Atul Dhavle is a Graduate in Production Engineering from Nagpur University and holds a Postgraduate certificate in Business Management from XLRI. He has over 26 years of experience with reputed organizations like Mahindra & Mahindra, DuPont, Welspun, Dr. Reddy's and handled various assignments in Human Resources and Operational Excellence functions.

Agenda

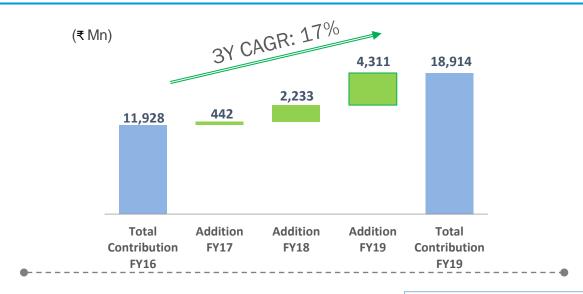


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High volume molecules showing stable growth

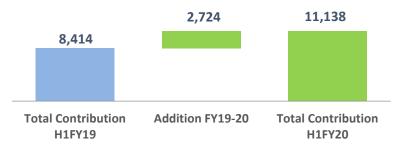




High volume based molecule business contribution

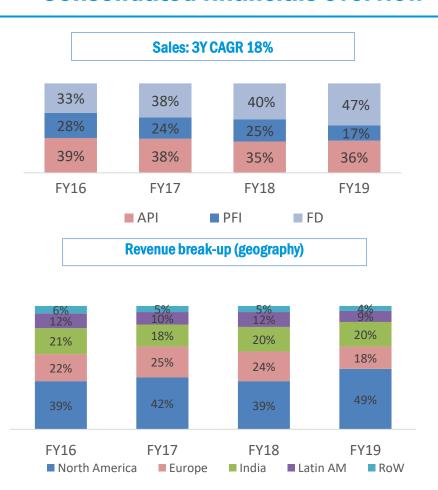
(₹ Mn)

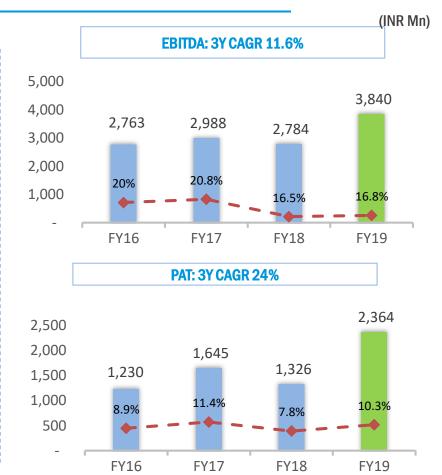
Revenue contribution from core molecules



Consolidated financials overview



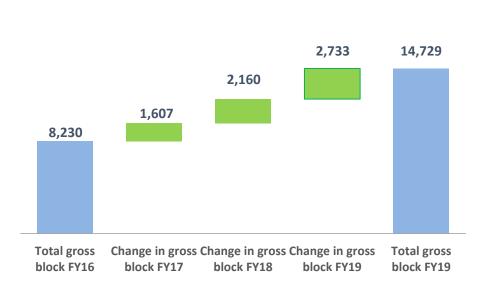




Balance sheet trends

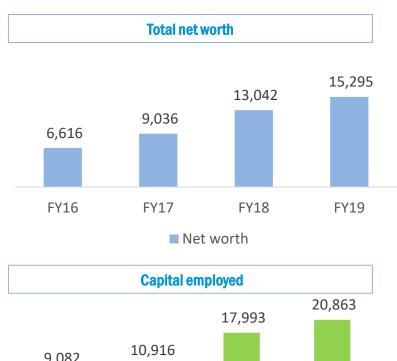






	FY16	FY17	FY18	FY19	YTD
Cash outflow for Capex	1,622	3,166	4,464	2,788	1,156

Moderation in Capex and reduction in debt going forward

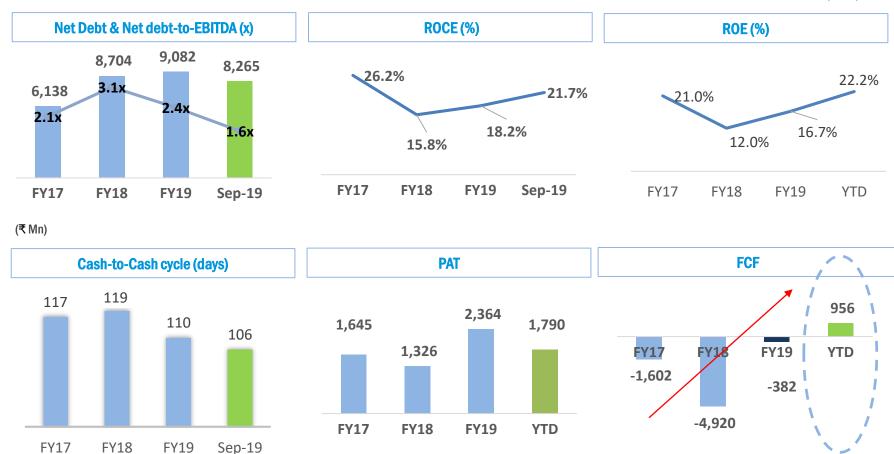




Continued focus on improvement



(₹ Mn)



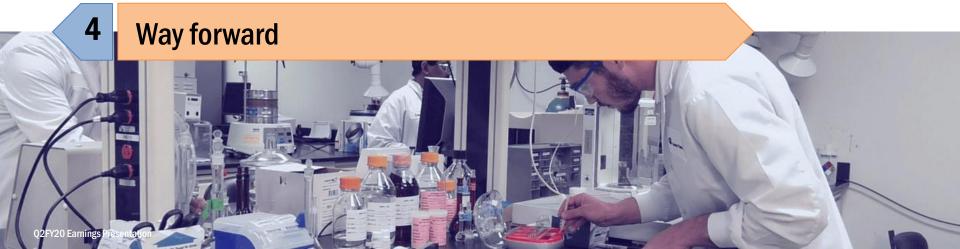
Note: JVs not included in ROCE for Sep-2019

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Way forward



Huge Economies of Scale

Foundation of Quality & Compliance

Experienced and Proven Management team



Building blocks in place

Integrated platform in place to address the entire value chain from intermediates to finished dosages

Investments in FD including pulsatile drug release, orally disintegrating modified release and abuse deterrent technology platform for controlled substances



Profitable growth through execution

Expanding portfolio range with addition of differentiated and limited competition products in API/FPI

Building portfolio of strong brands in regulated markets



Steady returns to stakeholders

Financial discipline is a key focus

Focus on generating operating leverage

Identifying newer growth avenues to deliver sustainable growth



Thank you

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