



**Date: January 23, 2026**

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 the unaudited financial results for the third quarter and nine months ended on December 31, 2025, submitted to you today i.e., on January 23, 2026. We are now enclosing the presentation in this regard to the Analysts/Investors.

This is pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015.

Kindly take the above information on record.

**For GRANULES INDIA LIMITED**

CHAITANYA  
TUMMALA

Digitally signed by  
CHAITANYA TUMMALA  
Date: 2026.01.23 17:10:11  
+05'30'

**CHAITANYA TUMMALA  
(COMPANY SECRETARY &  
COMPLIANCE OFFICER)**



**REGISTERED OFFICE**

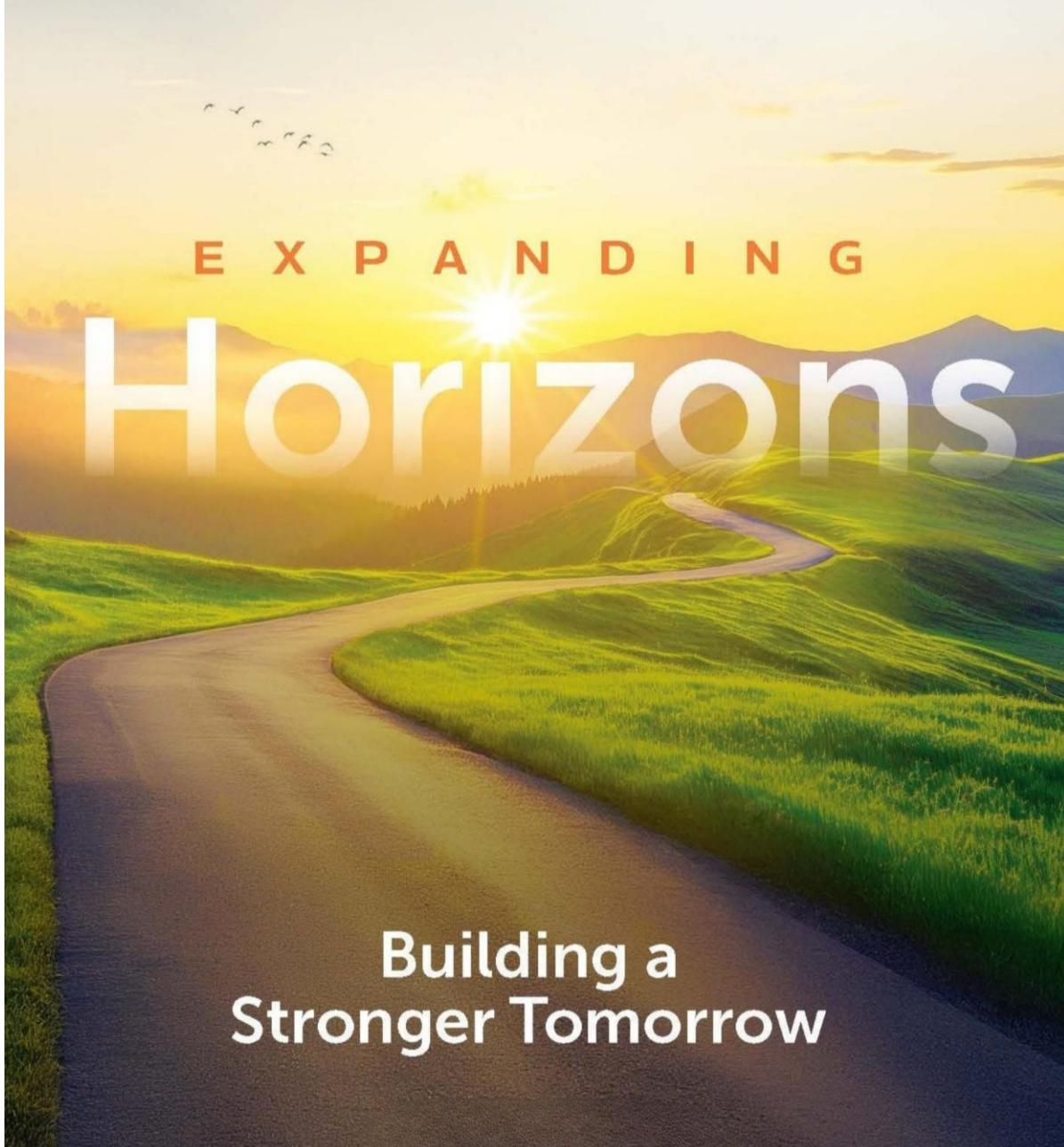
Granules India Limited

CIN: L24110TG1991PLC012471

15<sup>th</sup> Floor, Granules Tower, Botanical Garden Road, Kondapur, Hyderabad - 500084, Telangana, India

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# Granules India Limited

## Earnings Presentation

### Q3FY26

23<sup>rd</sup> January 2026

# Safe Harbour Statement

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.

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# Quarterly Performance

## Q3 FY 26

# Q3FY26: Executive Summary

## Financial Overview

- **Sales:** YoY revenue growth of 22% and QoQ growth of 7%
- **EBITDA:** YoY EBITDA increase by 196 bps (*22.2% from 20.2%*) and QoQ by 75 bps (*22.2% from 21.5%*) despite EBITDA loss of ₹248 Mn from Ascelis Peptides in Q3FY26

## Ratio's, Cashflow And CAPEX

- **ROCE:** 16.8% increased from 16.2% in Q2FY26 with the improvement of EBIT
- **Cash from operations :** Cash flow from operations of ₹2,187 Mn as compared to ₹1,937 Mn in Q2FY26
- **CAPEX:** ₹1,298 Mn as compared to ₹2,112 Mn in Q2FY26

## R&D

- **R&D expenses** stood at ₹689 Mn (5.0% of sales) in Q3 FY26 as compared to ₹705 Mn (5.4% of sales) in Q2 FY26. Inline with our long-term strategic growth.
- **Regulatory milestones:** 94\* approvals till date across the regions. Tentative approval received for Amphetamine ER (Adzenys) ANDA in Q3FY26

## Key Updates

- **Market Share % Rank #1 in 9/35 (25%) of US portfolio; Top-3 in 18/35 (50%) of US portfolio^**
- ANVISA Brazil GMP certificate received for Gagillapur site
- Granules has achieved highest “A” rating in climate change from “B” in 2024 from CDP
- ICRA upgraded credit rating from AA- to AA. Ind rating reaffirmed AA- positive outlook
- Granules India's Packaging Facility in US Completes FDA inspection with Zero Observations

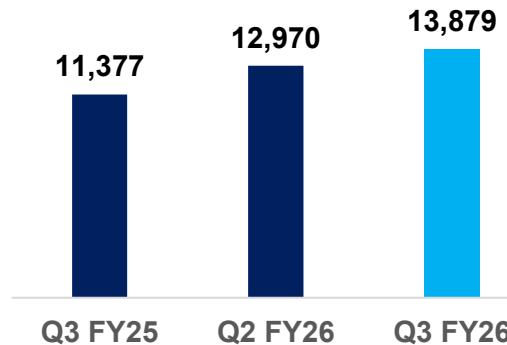
\*including 2 tentative approval

<sup>^</sup>Basis IQVIA MAT Nov'25

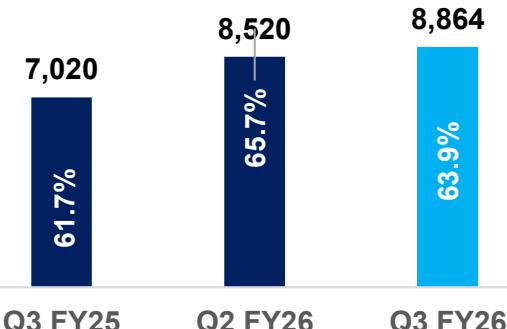
# Q3FY26: Financial Overview

Particular	Q3 FY26	Q2 FY26	Q-o-Q	Q3 FY25	Y-o-Y
Revenue	13,879	12,970	7%	11,377	22%
Gross Margin	8,864	8,520	4%	7,020	26%
<b>% of Gross Margin</b>	<b>63.9%</b>	<b>65.7%</b>	<b>183 bps</b> 	<b>61.7%</b>	<b>216 bps</b> 
Manpower Cost	2,260	2,192	3%	1,676	35%
R&D	689	705	-2%	568	21%
Other Expenses	2,834	2,841	0%	2,472	15%
EBITDA	3,081	2,782	11%	2,303	34%
<b>% of EBITDA</b>	<b>22.2%</b>	<b>21.5%</b>	<b>75 bps</b> 	<b>20.2%</b>	<b>196 bps</b> 
PBT	2,022	1,759	15% 	1,528	32% 
PAT	1,502	1,306	15%	1,176	28%
<b>% of PAT</b>	<b>10.8%</b>	<b>10.1%</b>	<b>75 bps</b> 	<b>10.3%</b>	<b>49 bps</b> 

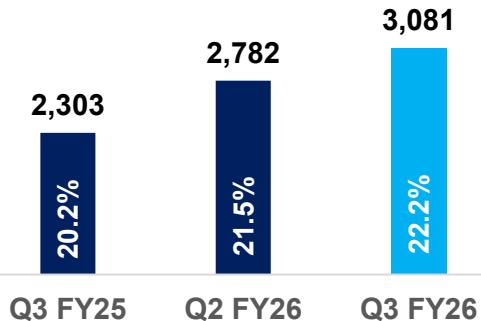
## Revenue



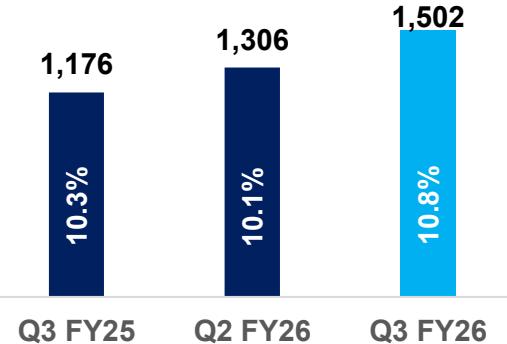
## Gross Margin



## EBITDA



## PAT



## Revenue

YoY revenue growth was driven by the Finished Dosage (FD) segment in North America and Europe.

QoQ revenue growth was led by higher sales in Europe.

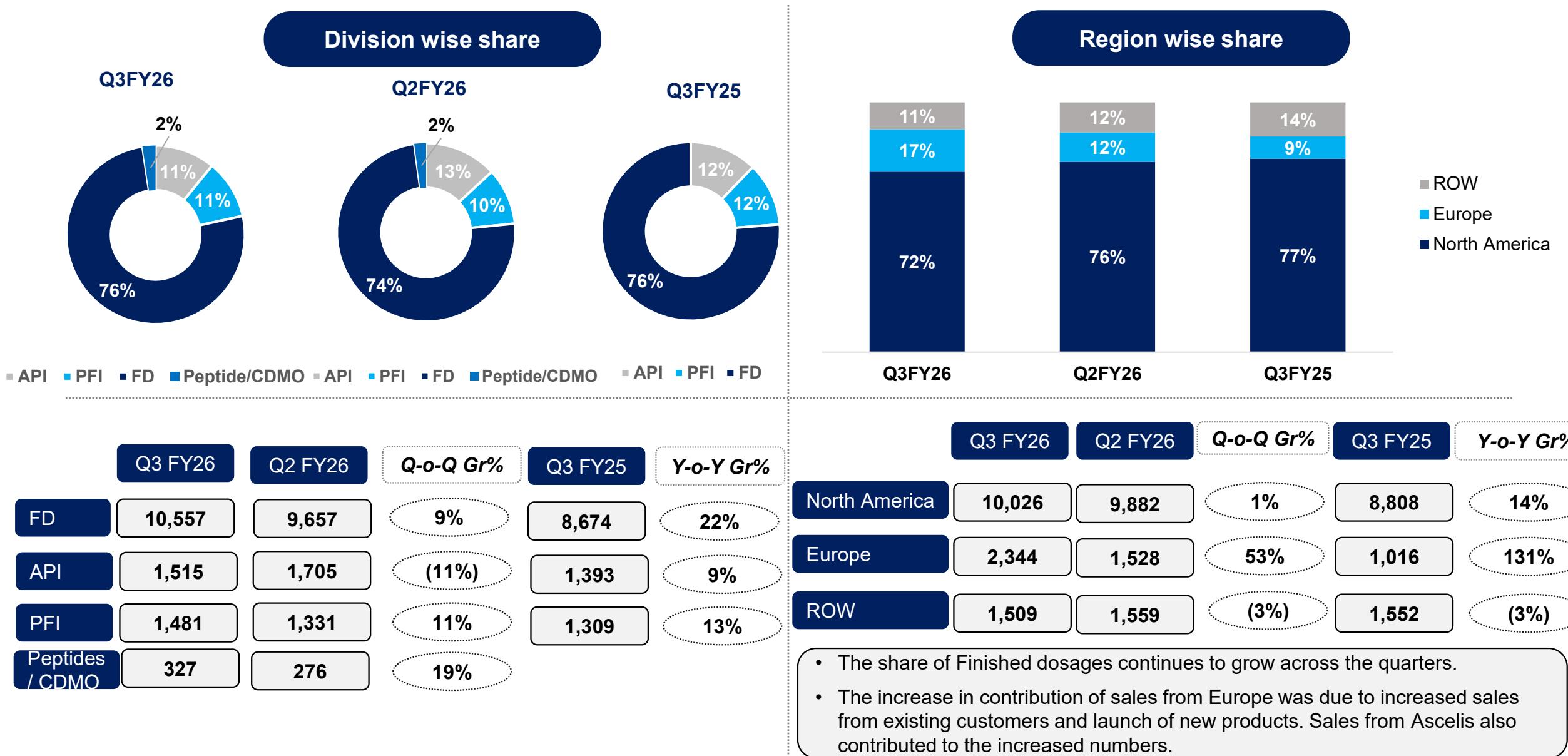
## Gross Margin

YoY Gross Margin expansion of 216 bps was due to a better product mix within the Finished Dosage segment

## EBITDA

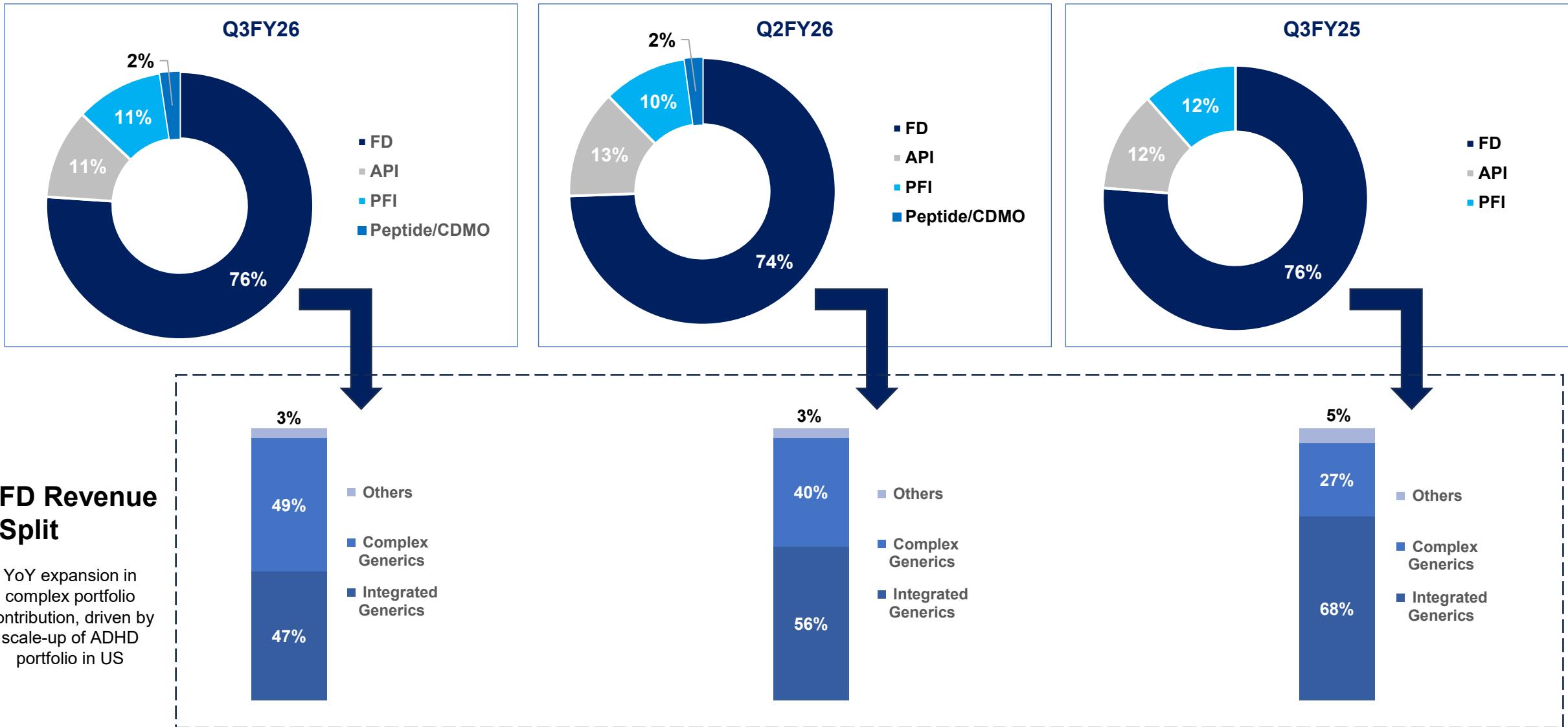
EBITDA growth was supported by improved sales and margins, partially offset by an EBITDA loss of ₹248 Mn from Ascelis Peptides.

# Q3FY26 : Revenue split by Segments and Market



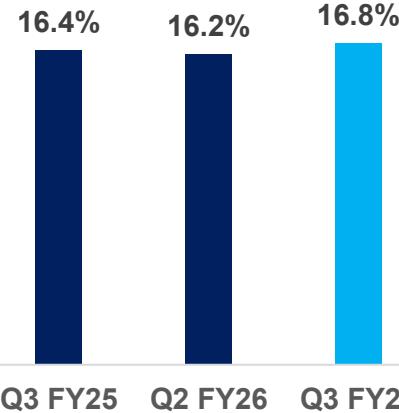
Segment	Definition
<b>Integrated Gx</b>	<p>Products where API is key; API forward integrated to value added PFI and FDs (mid to high volume)</p> <p><b>Value proposition:</b></p> <ul style="list-style-type: none"><li>Backward integration/ Strategic tie ups</li><li>Supply chain security and cost competitiveness</li></ul>
<b>Complex Gx</b>	<p>Products with high entry barriers due to various reasons;</p> <ul style="list-style-type: none"><li>Complexity in formulation development and manufacturing</li><li>Complexity in API development</li><li>Complexity in IP/ BE strategy</li><li>Complexity in Compliance</li><li>Complexity in route of administration or drug-device combination</li></ul>
<b>Others</b>	<p>Products that are time based/ opportunity products with an inherent capability of falling into the integrated basket.</p>

# Q3FY26 : FD Revenue Spilt by Portfolio Segments

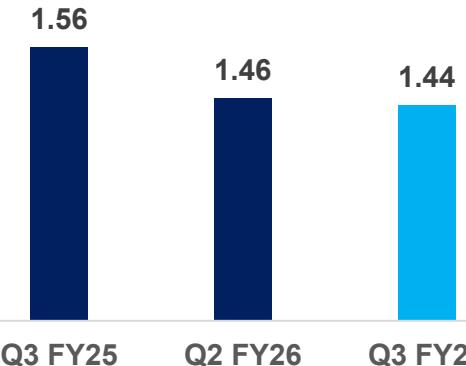


# Q3FY26: Cashflow and Ratios

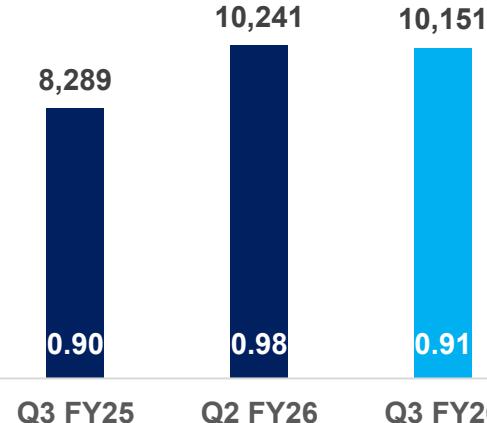
## ROCE %



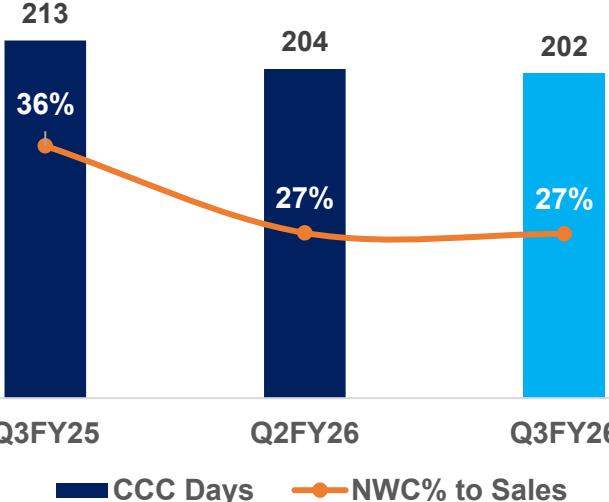
## Fixed Asset Turn



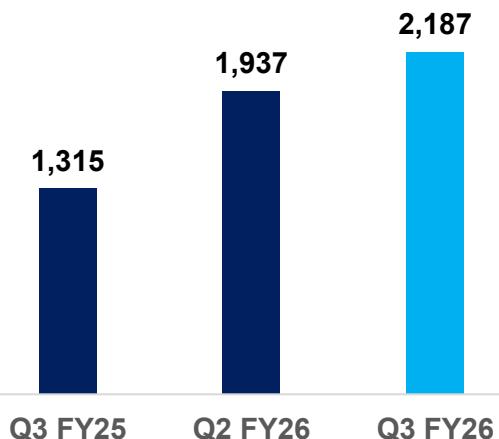
## Net Debt and Net Debt/EBITDA



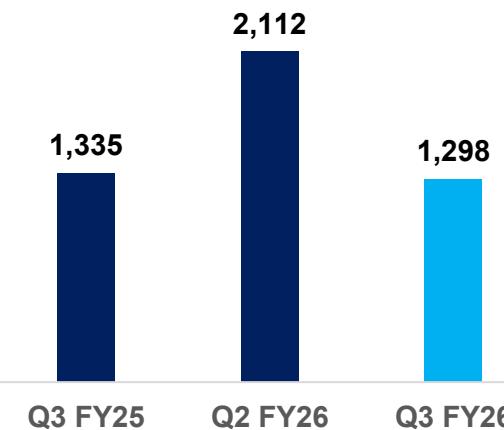
## CCC days and NWC% to sales



## Cash Flow from Operations



## CAPEX

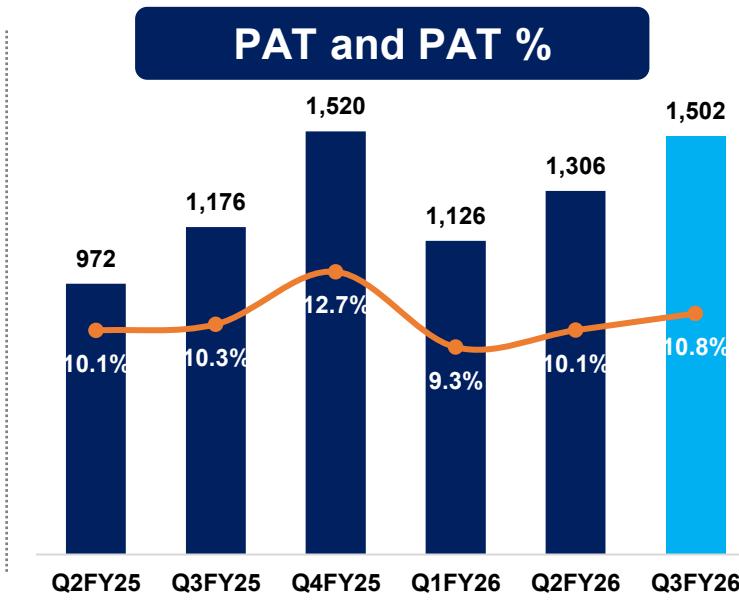
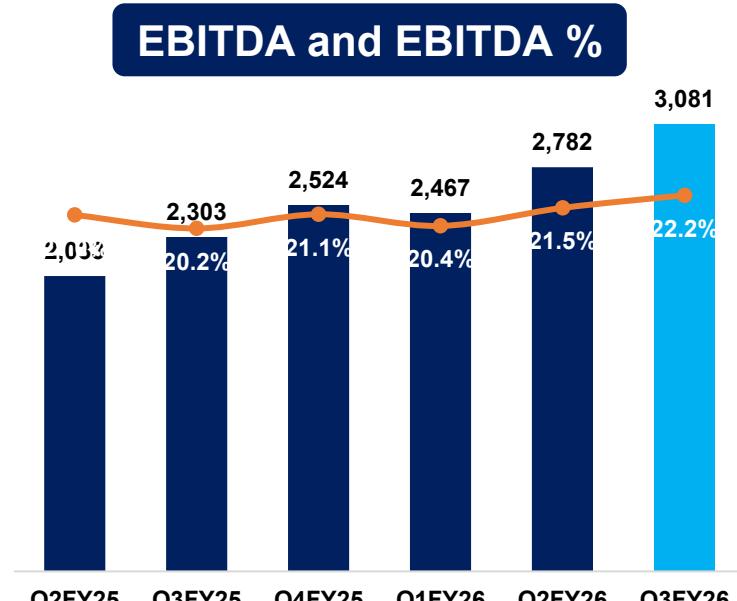
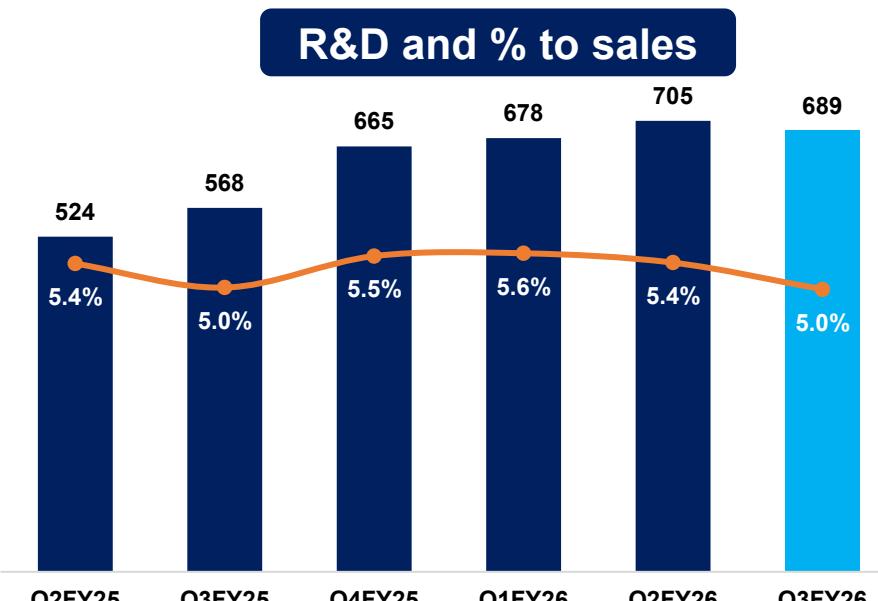
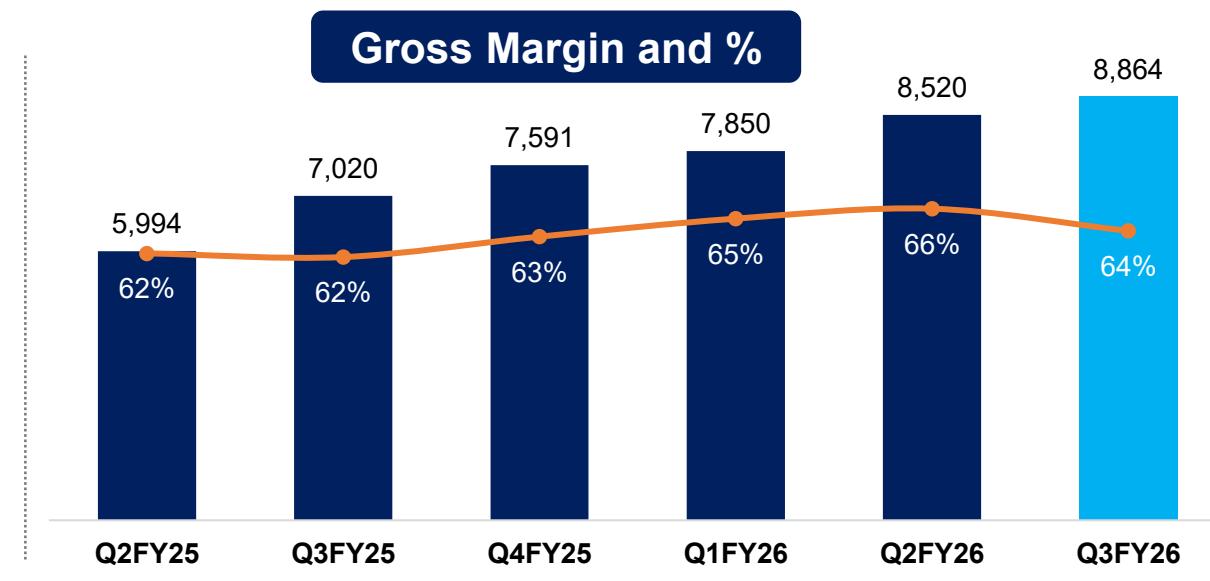
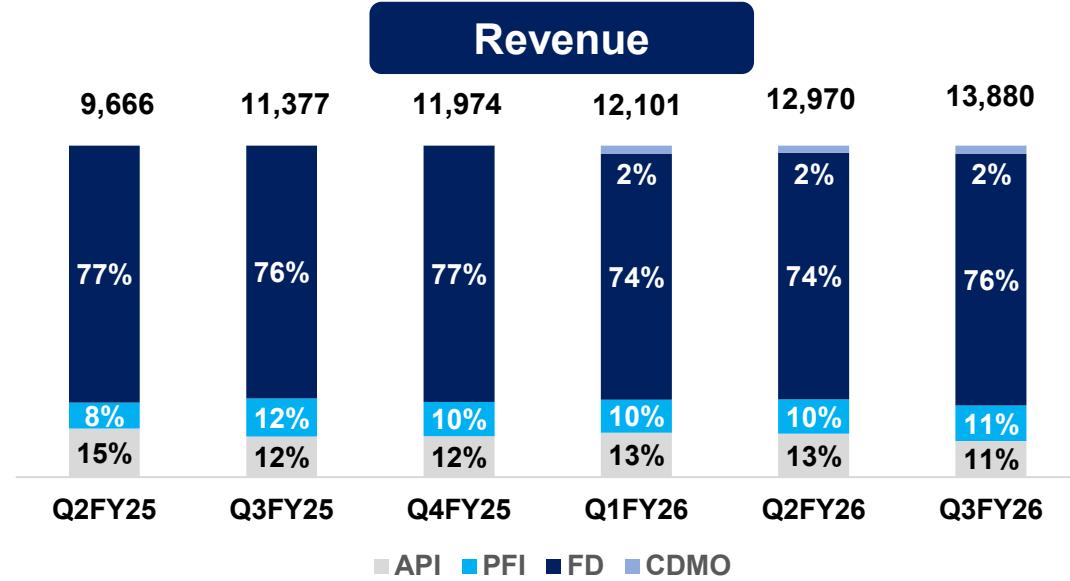


■ CCC Days ■ NWC% to Sales

## Quarterly Historical Performance

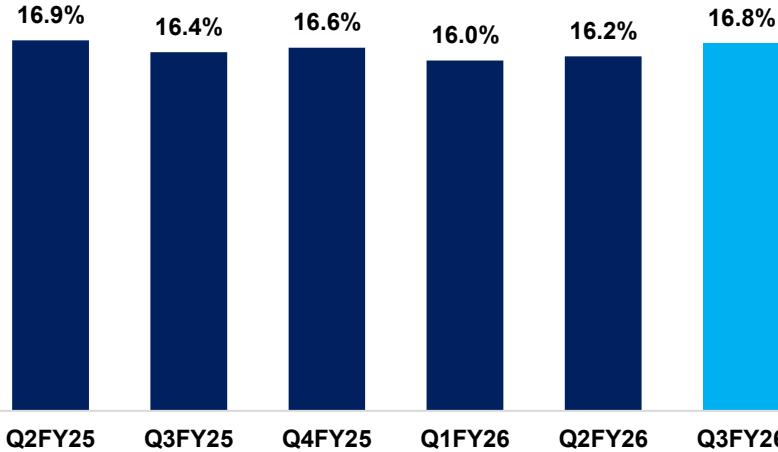


# Sales and Profitability: Last 6 quarters historical performance

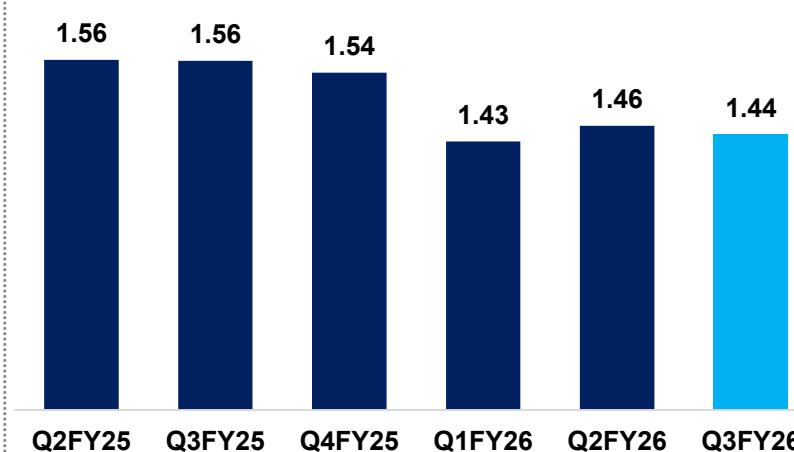


# Cashflow and Ratios: Last 6 quarters historical performance

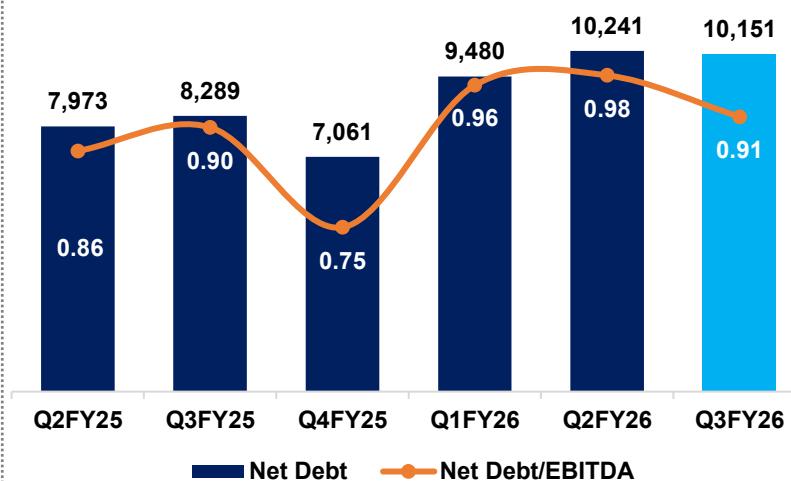
## ROCE%



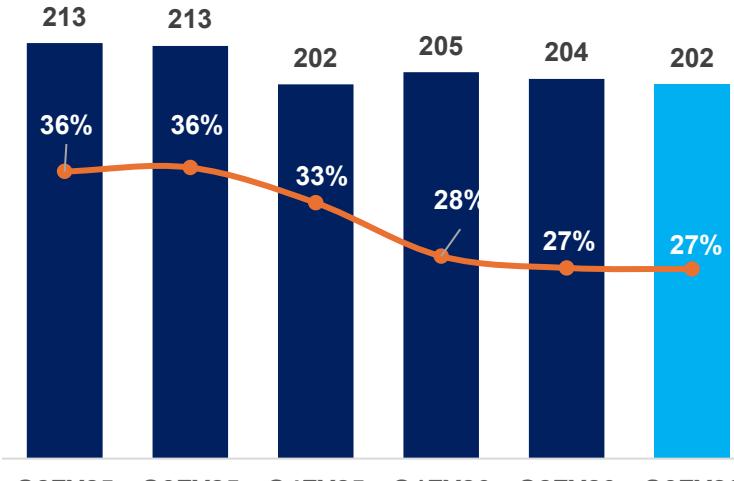
## Asset turns



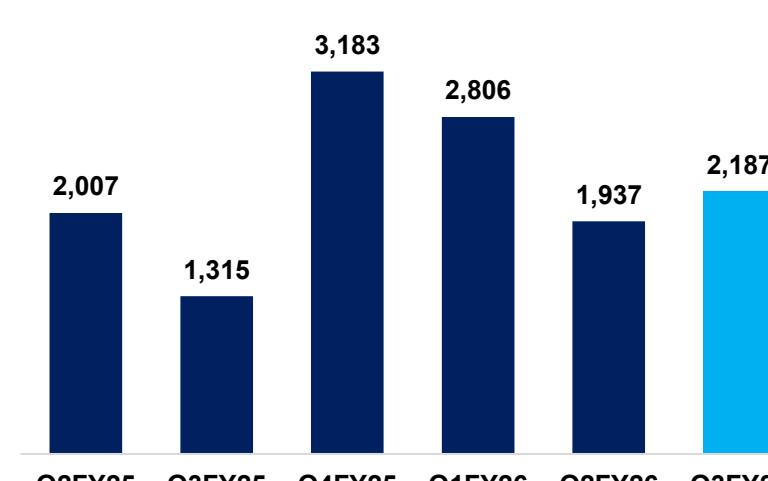
## Net debt and Net debt/EBITDA



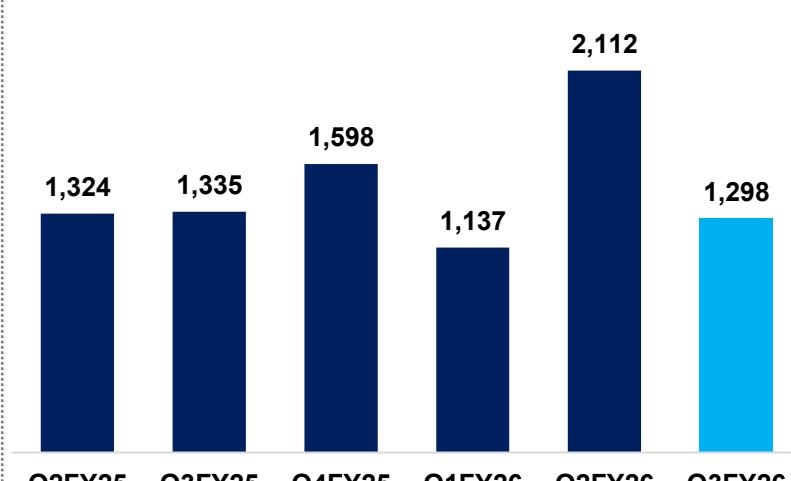
## CCC days and NWC% to sales



## Operating Cashflow



## Capex



■ CCC Days ■ NWC% to Sales

## R&D Capability



# R&D: Product pipeline and capability building



- ✓ Total of ~427 scientist across API, FD R&D and Peptides
- ✓ Received 94 dossier approvals till Q3FY26
- ✓ Filed 100+ DMF globally
- ✓ Robust integrated pipeline in the areas of focus



- ✓ R&D expenses for Q3FY26 Rs.689 Mn (5.0% of Sales) to support long term strategic growth

Dossier Filing Status	Approved	Tentatively Approved	Filed	Total Products
GPI IP	USA 31	1	4	36
GIL IP	USA 36	1	15	52
	Europe 8	0	10	18
	Canada 7	0	0	7
	ROW 8	0	7	15
	UK 2	0	0	2
	<b>Total 92</b>	<b>2</b>	<b>36</b>	<b>130</b>

## Integrated Product Development

### Genome Valley – MN Park

R&D facility for Integrated Product Development

- Fast tracking product development through seamless coordination & shared analytical resource

## Controlled Substances

### Pragathi Nagar R&D

- CoE for CII API Development
- Select KSMs

### GPI R&D

- CII Finished Dosage
- Complex technologies FD

## Peptides CDMO

Senn Chemical AG,  
Switzerland

Ascelis Peptides Lab in IIT  
Hyderabad

## New Technologies

### Bio Lab at Pragathi Nagar

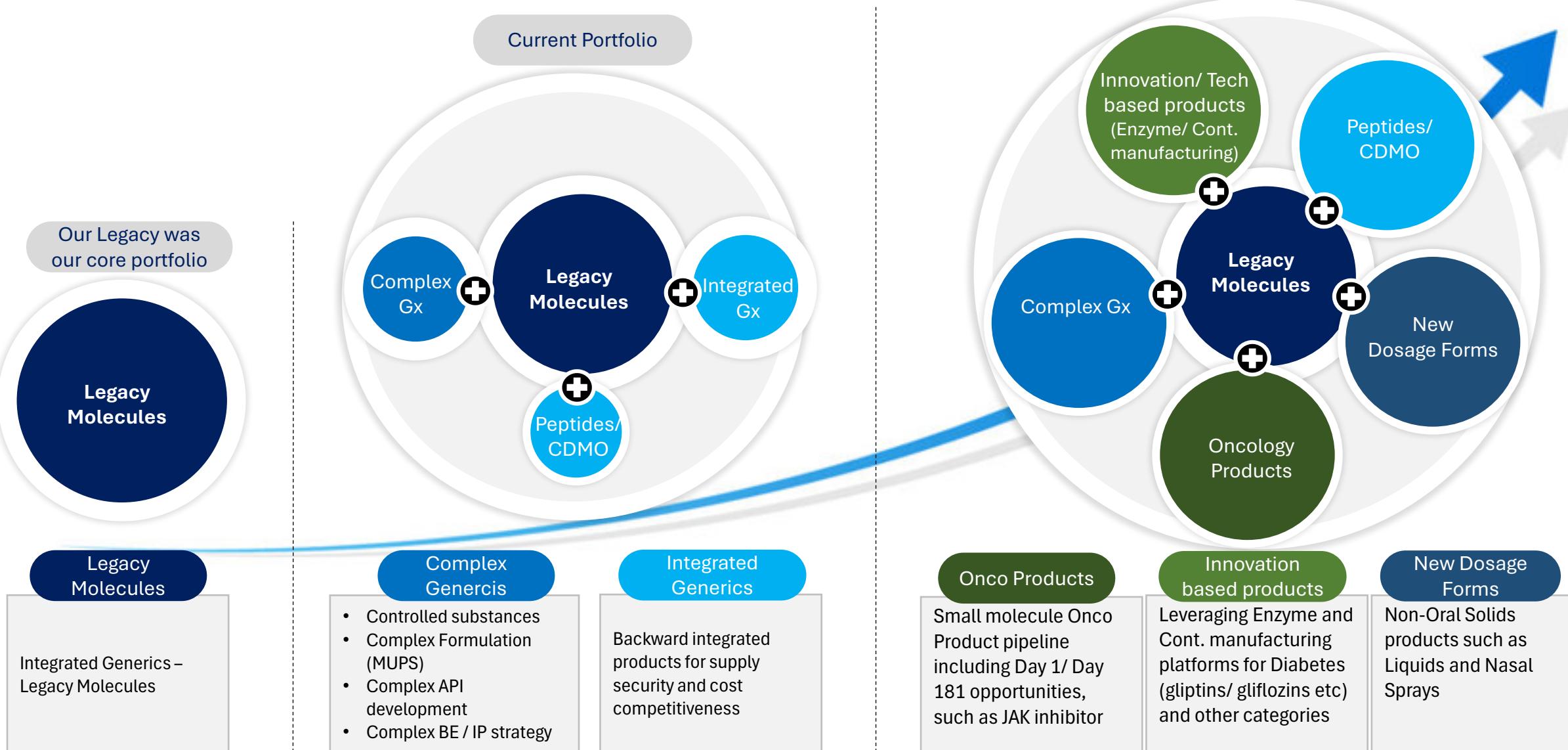
- Enzyme & Fermentation

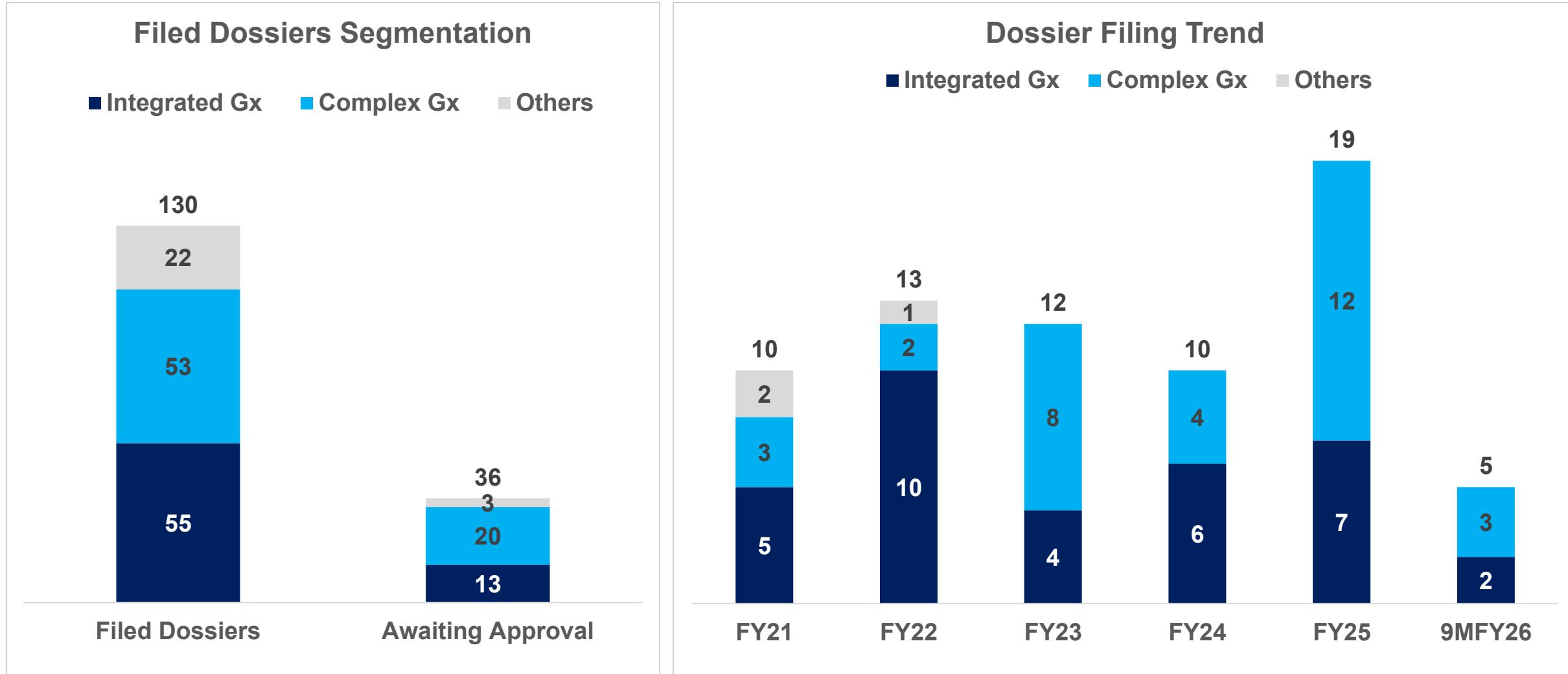
### Pune R&D

- New technologies
- KSM & Backward Integration

Market	Filed DMFs
US DMF	43
EU CEP	21
EU ASMF	9
Canada	5
Brazil	5
Korea	12
Japan	2
China	7
<b>Total</b>	<b>104</b>

## Portfolio Expansion: On Strength Of Focused R&D Investments & Leveraging Our Core Capabilities







**Granules at a glance**

# Granules Purpose, Vision & Values

## Purpose

Healing lives  
responsibly  
through pioneering  
green science

## Vision

To establish ourselves as a world  
leader in green chemical and  
pharmaceutical industry by  
harnessing cutting-edge technologies  
to enhance quality of life.

## Values

- 1 Challenging Limits
- 2 Futuristic Thinking
- 3 Empowering Employees
- 4 Customer Driven
- 5 Quality Everywhere
- 6 Environmental Stewardship

# An Illustrious Journey Spanning Four Decades

## Building A World Class, Vertically Integrated Pharmaceutical Manufacturing Platform



From being a paracetamol API manufacturing facility to a multi-faceted global pharmaceutical company, we have consistently achieved manufacturing success, set new standards of quality and expanded our presence across the world.



Triton Laboratories was formed to produce Paracetamol API at our Bonthapally facility in Hyderabad



A new large volume PFI facility in Gagillapur and a wholly owned subsidiary - Granules USA for marketing in the US



Acquired Auctus Pharma - An API Manufacturing Facility with regulatory approvals



Laid foundation for Oncology OSD Plant in Visakhapatnam



GPAK is a 79,000 sq. ft. packaging facility with 4 packaging suites & WH facility



CDMO specializing in Peptide development and manufacturing



Granules opened its second manufacturing facility at Jeedimetla to produce multiple APIs



Entry into the Finished Dosage segment



Sets up a wholly owned subsidiary in the US; Granules Pharmaceutical to focus on formulation R&D to forward integrate its APIs



Largest single manufacturing site for Multi-Unit Pellet system facility



Granules Life sciences (GLS) a Formulation facility



# Manufacturing Network & Capacity

**Bonthapally**



API: 34,560 TPA

**Jeedimetla**



API: 4,800 TPA  
PFI: 1,440 TPA

**Bonthapally II**



API (Intermediate):  
61.5 KL PA

**Vizag (Unit 4)**



API: 380 KL PA

**Vizag (Unit 5)**



API: 15 KL PA  
FD: 1.1 Bn

**Gagillapur**



FD: 26.8 Bn  
PFI: 23,200 TPA

**Virginia, USA**



FD: 2 Bn

**GPAK, USA**



2 OTC lines  
1 Rx line

**Genome Valley**



FD: 10Bn dosages

**Senn Chemicals AG, Switzerland**

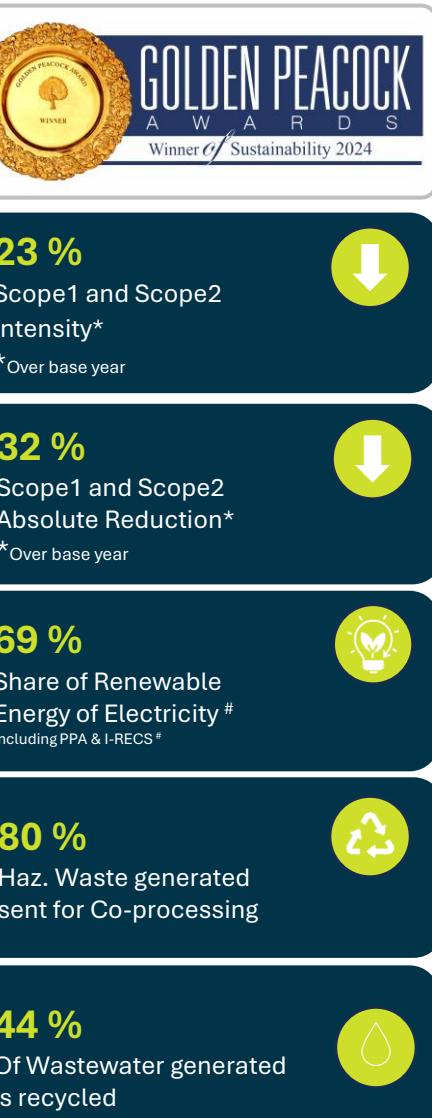


Peptides CDMO

# Sustainability at Granules : Pioneering Innovation for Complete Supply Chain Decarbonization of Pharmaceuticals



## Partner Collaboration Platforms



# Our CSR Journey : Empowering Communities, Enriching Lives

## OUR GOAL

Touch **1 Million**  
Lives by 2030

## OUR PROGRESS

**1,00,000 +**  
Lives Positively Impacted

## FOCUS AREAS



### Skill Development

**1600 +**  
trained through Pharma Pathashala  
since inception in 2017 with 250+  
in current year



### Health

**15000 +**  
Benefitted through Breast Cancer  
Screening Camps, Eye Screenings to  
school children



### Education

**1000+**  
Students are benefitting through  
Vidya Volunteers, Educational  
support through NGO partners



### Environment & Biodiversity

**18,000 +**  
Native trees have been planted.  
**1,00,000 +** Participants in Fourth Edition  
of Granules Green Heartfulness Run





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thank you!



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