

August 01, 2025

To,
Dy. General Manager
Department of Corporate Services,
BSE Ltd.,
P. J. Towers, Dalal Street,
Fort, Mumbai – 400 001

Ref: Scrip Code: 543322

Dear Sirs,

Sub: Investor Presentation

Pursuant to regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations 2015, we are enclosing herewith the Investor Presentation – Q1 FY 25-26.

You are requested to take the same on record.

Thanking You.

Yours faithfully,
For Alivus Life Sciences Limited
(formerly Glenmark Life Sciences Limited)

Rudalf Corriea
Company Secretary & Compliance Officer
Encl: As above

Alivus Life Sciences Limited (formerly Glenmark Life Sciences Limited)

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Investor Presentation



Q1 FY26





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Financial Performance **Review**



Q1 FY26 | Highlights



Dr. Yasir Rawjee
Managing Director &
Chief Executive
Officer

"Our performance this quarter was primarily driven by strong growth in the Non-GPL business, which expanded by 14.5% YoY. Key regions including India (Ex-GPL), Europe, ROW, LATAM, and Japan were major contributors to this growth. We remain confident of achieving high single-digit revenue growth in FY26, with stronger momentum expected in the second half. Margins are projected to remain healthy in the 28–30% range.

Our ongoing investments in capacity building and pipeline enrichment form the foundation of our future growth, and we believe these efforts, supported by an increasingly favourable market environment, will help unlock further value."

REVENUE

(IN ₹ MILLION)

6,018

-7.4%
QoQ

2.2%
YoY

EBITDA

(IN ₹ MILLION)

1,813

-13.0%
QoQ

9.9%
YoY

PAT

(IN ₹ MILLION)

1,215

-14.4%
QoQ

9.0%
YoY

- Alivus registered a revenue from operations of ₹ 6,018 Mn for Q1FY26, a growth of 2.2% YoY.
- Gross Margin in Q1FY26 were at 55.1% up 400 bps YoY, driven by rationalized input cost and leveraged operational efficiency.
- EBITDA margins for the quarter were at 30.1%, up by 210 bps YoY, resulting from better gross margins.
- Generic API revenues in Q1 FY26 grew by 3.0% YoY.
- GPL business was at ₹ 1,526 Mn, a de-growth 30.0% on QoQ and 22.3% on YoY, resulting from inventory rationalization at the customer's end; Non-GPL business was at ₹ 4,492 Mn, up by 4.1% QoQ and 14.5% YoY.
- During Q1FY26, the company generated a strong free cash flow of ₹ 1,000 Mn, leading to Cash and Cash Equivalents (including short term investments) of ₹ 6,604 Mn as of 30th June 2025.



Q1 FY26 Performance

Revenue

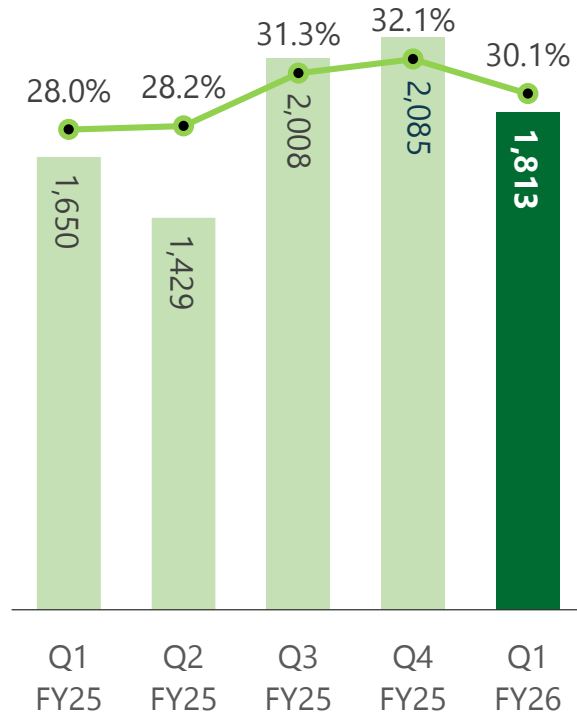
(In ₹ Million)



EBITDA

(In ₹ Million)

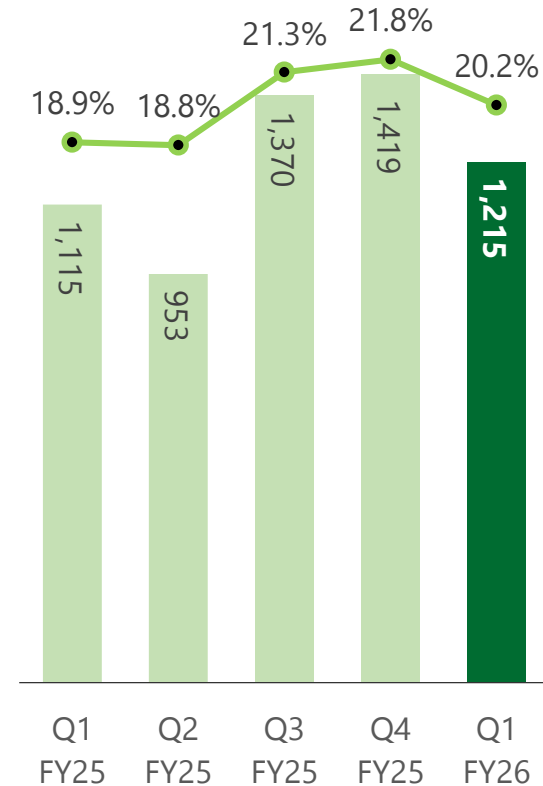
EBITDA Margin



PAT

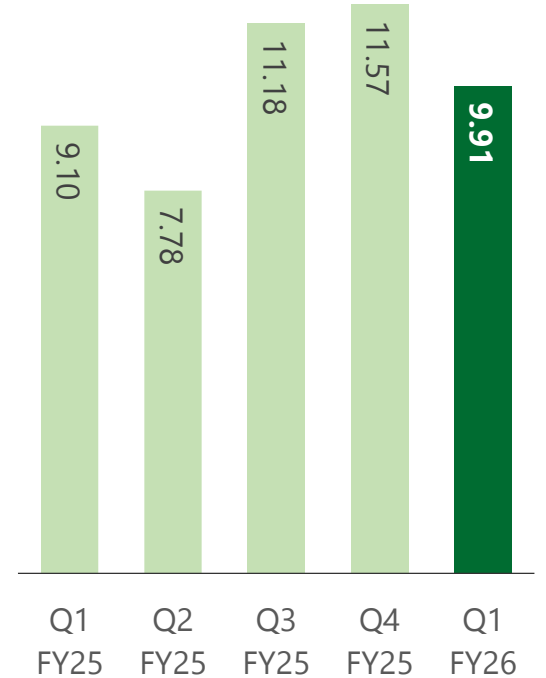
(In ₹ Million)

PAT Margin



EPS

(In ₹)



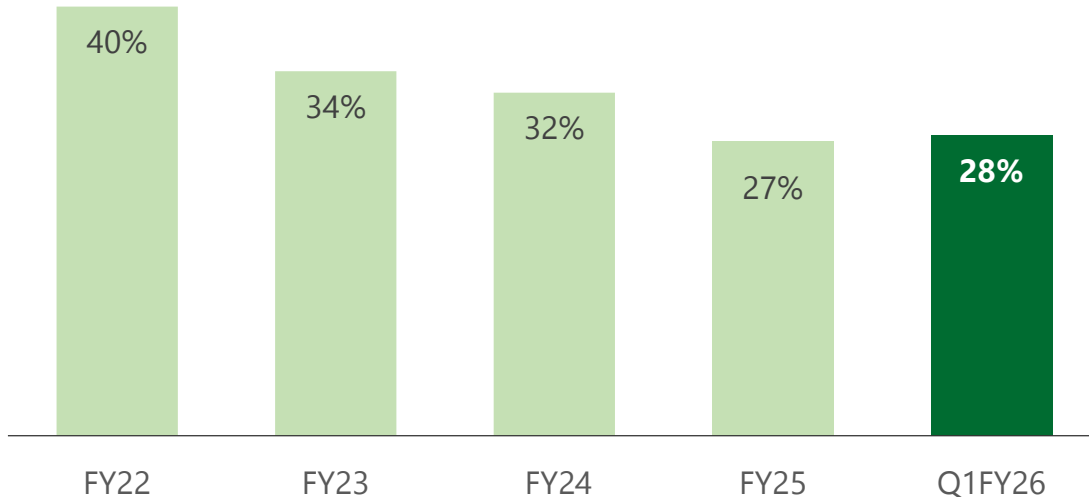
A P&L Highlights | Q1 FY26

Particulars (In ₹ Million)	Q1 FY26	Q4 FY25	QoQ	Q1 FY25	YoY	FY25
Revenue from Operations	6,018	6,496	-7.4%	5,886	2.2%	23,869
Gross Profit	3,315	3,670	-9.7%	3,008	10.2%	13,061
Gross Profit (%)	55.1%	56.5%	-140 bps	51.1%	400 bps	54.7%
Other Income	90	101	-10.9%	55	63.6%	346
Employee Benefits Expense	616	668	-7.8%	568	8.5%	2,517
Other Expenses	976	1,018	-4.1%	845	15.5%	3,718
EBITDA	1,813	2,085	-13.0%	1,650	9.9%	7,172
EBITDA Margin (%)	30.1%	32.1%	-200 bps	28.0%	210 bps	30.0%
Depreciation and Amortisation Expense	171	160	6.9%	144	18.8%	606
Finance Costs	12	12	-	4	200.0%	24
PBT	1630	1,913	-14.8%	1,502	8.5%	6,542
PBT Margin (%)	27.1%	29.4%	-230 bps	25.5%	160 bps	27.4%
PAT	1,215	1,419	-14.4%	1,115	9.0%	4,857
Net Margin (%)	20.2%	21.8%	-160 bps	18.9%	130 bps	20.3%

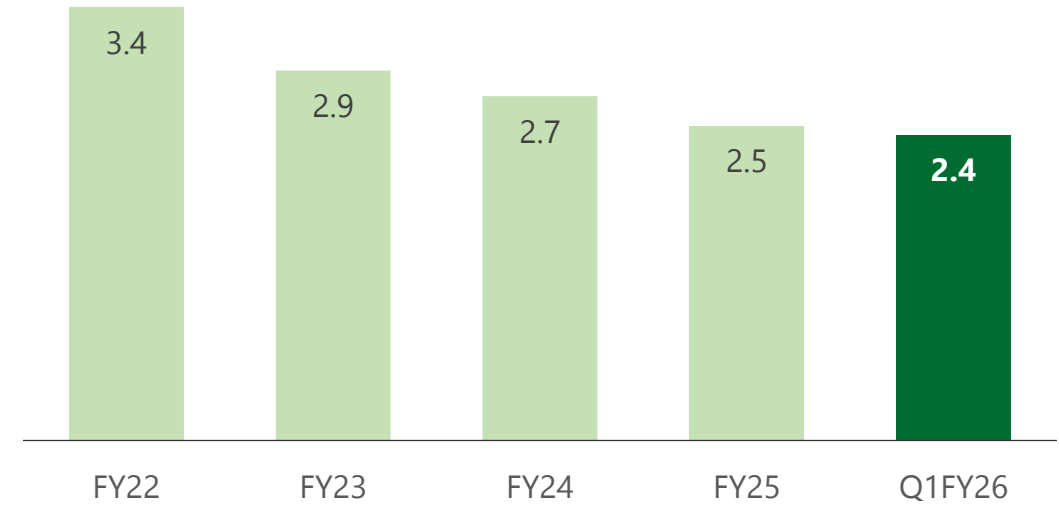


Healthy Return Indicators

ROICE (%)



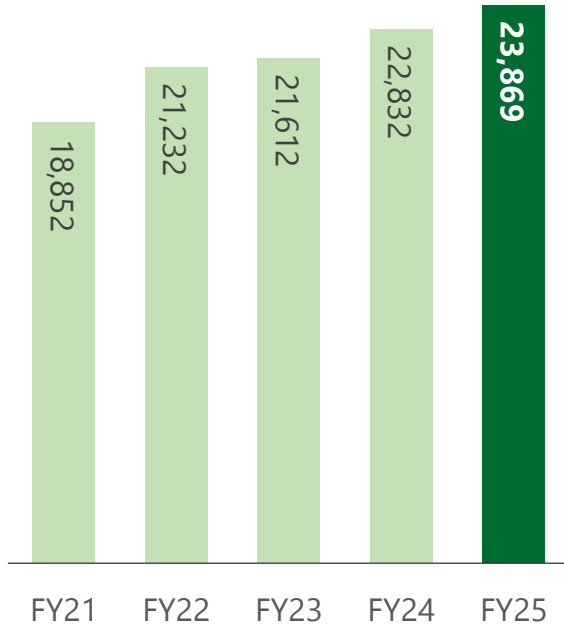
Fixed Assets Turnover (Times)



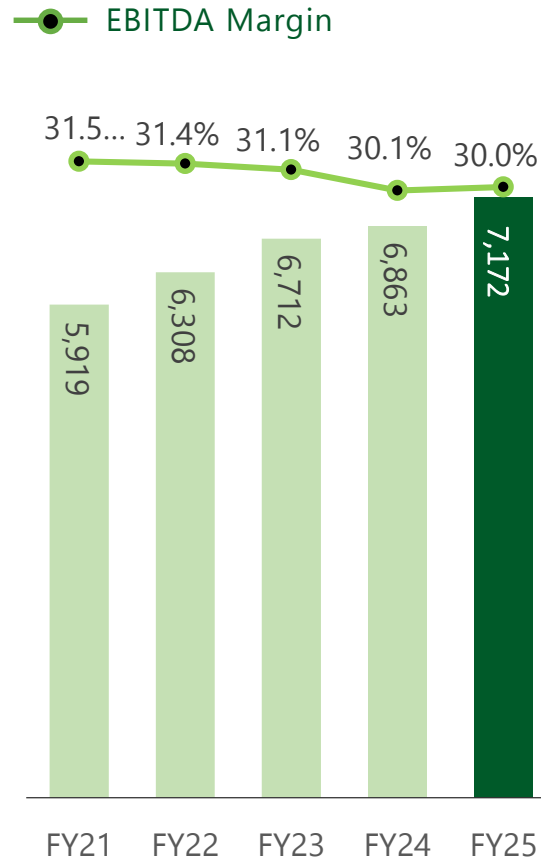
- **ROICE** is tracking at ~**28%** – Capital employed driven by calibrated Capex strategy
- **FATR** is **2.4 times** – Asset turn trending slightly lower due to Capex cycle
- **Strong Balance Sheet** – Strong free cash generation of ₹ 1,000 Mn leading to Cash and Cash Equivalents (including short term investments) of ₹ 6,604 Mn as of June 2025

Robust Growth & Steady Profitability

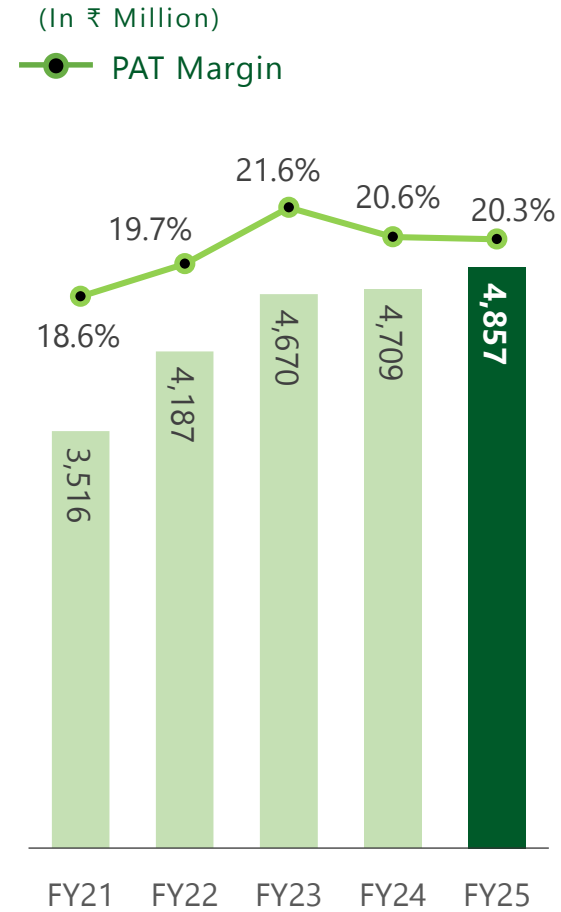
Revenue (In ₹ Million)



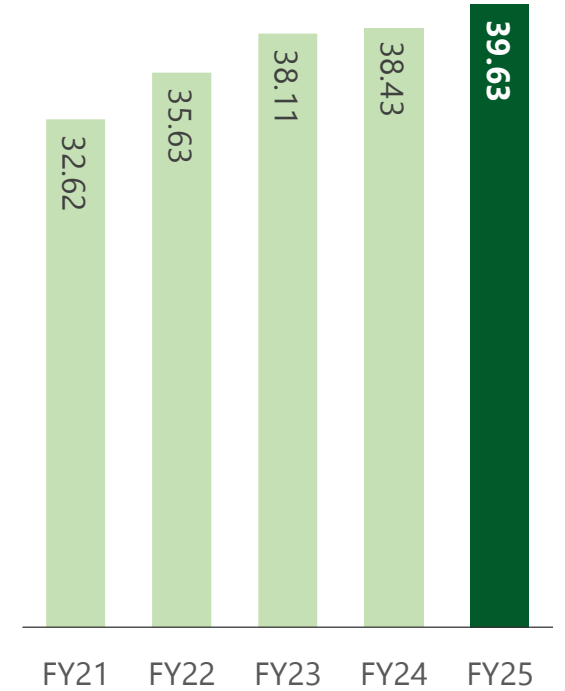
EBITDA (In ₹ Million)



PAT (In ₹ Million)



EPS (In ₹)



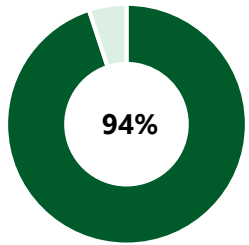
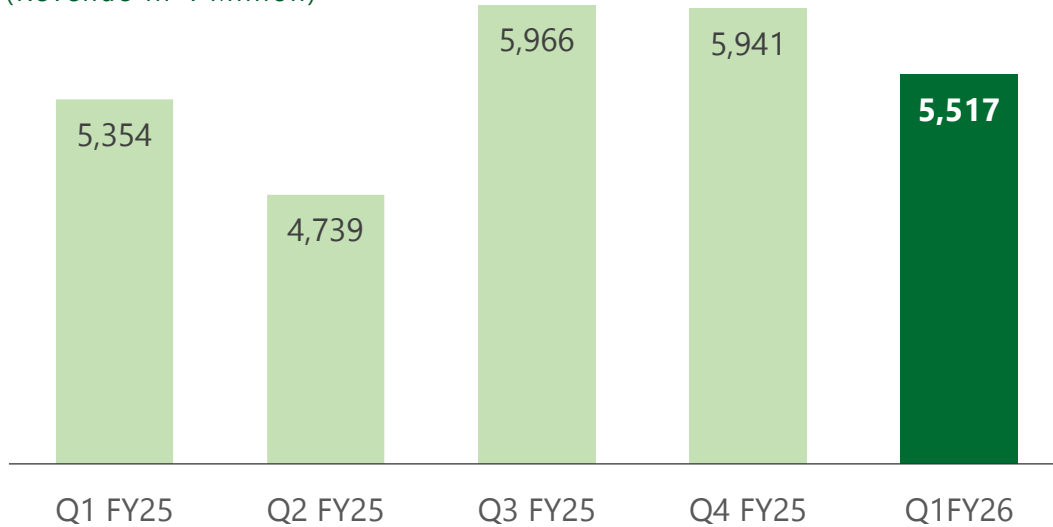
Business Performance Review



Segmental Performance | Generic API vs CDMO

Generic API

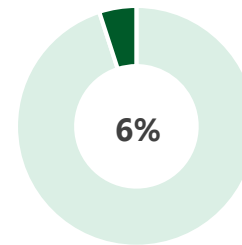
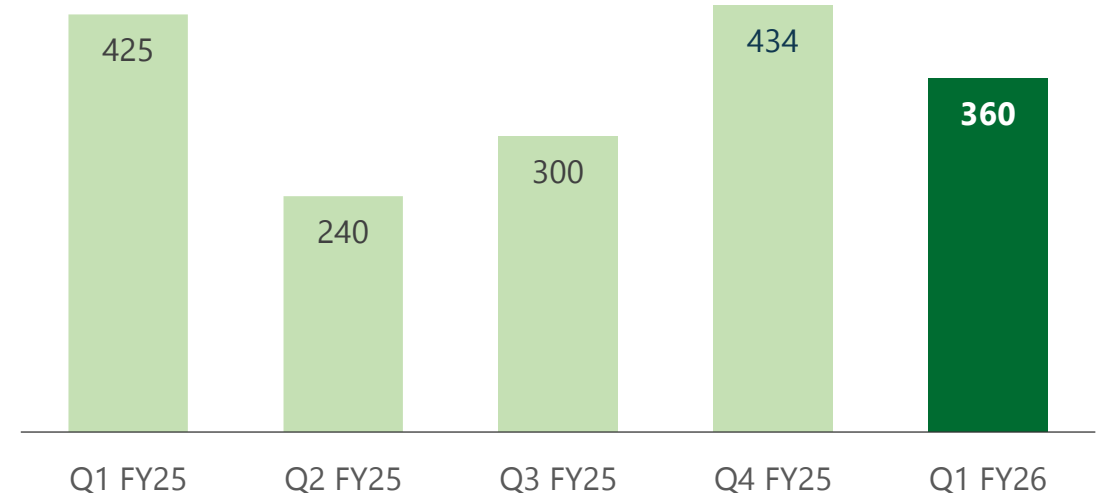
(Revenue In ₹ Million)



- Generic API revenues in Q1FY26 saw a subtle uptick of 3.0% YoY and de-grew by 7.1% QoQ.
- Regions like India (Ex-GPL), ROW, LATAM, and Japan contributed to YoY revenue growth.
- QoQ revenue decline was due to inventory rationalization by customers in North America, LATAM and in GPL business.

CDMO

(Revenue In ₹ Million)



- CDMO business witnessed a decline in revenue of 15.3% YoY and 17.1% QoQ due to temporary dip in demand across ongoing projects.
- 5th project is expected to be commercialized by H2FY26.
- Multiple discussions are ongoing.

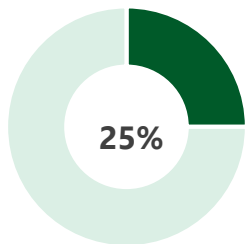
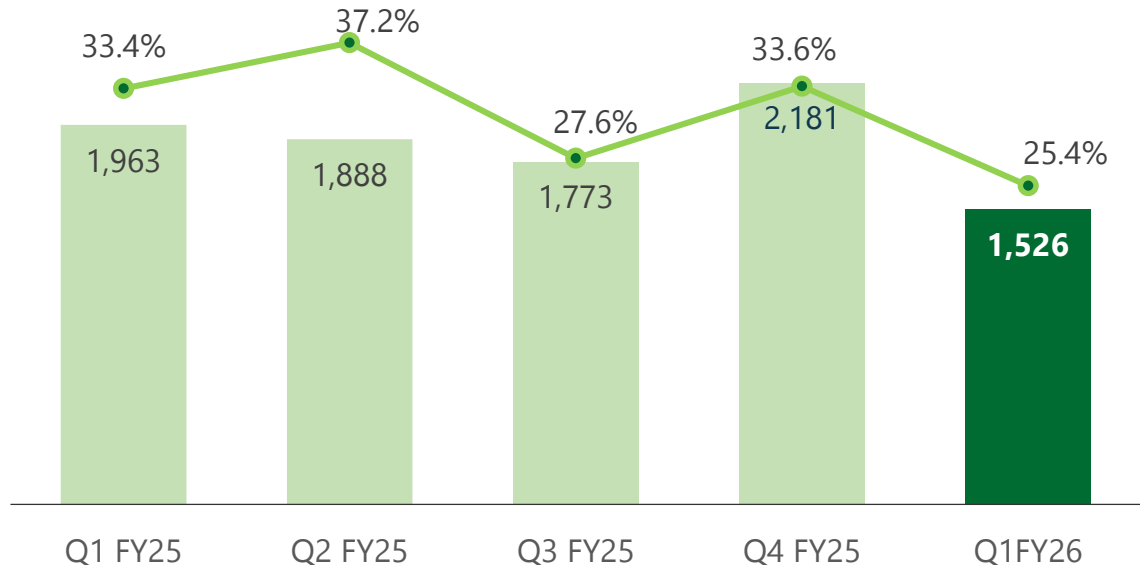


Segmental Performance | GPL vs Non-GPL

GPL

(In ₹ Million)

● % of Total Revenue

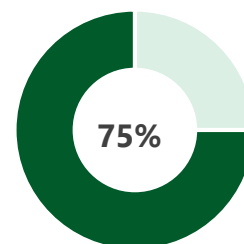
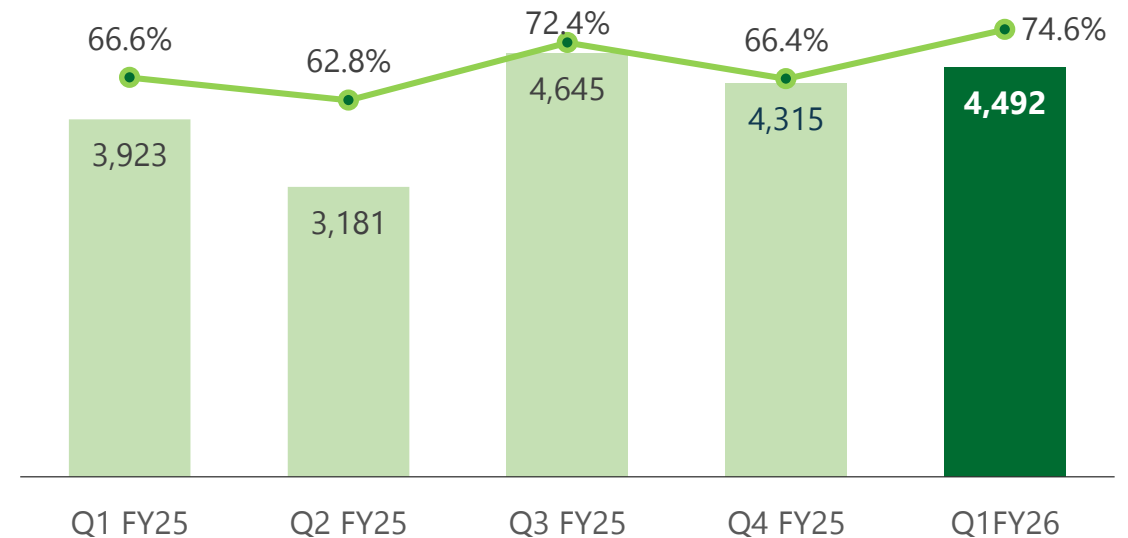


- GPL business witnessed a de-growth of 22.3% YoY and 30.0% QoQ mainly on account of inventory rationalization by the customer.
- In Q1FY26, GPL business contributed to ~25% of the total revenue from operations.

Non-GPL

(In ₹ Million)

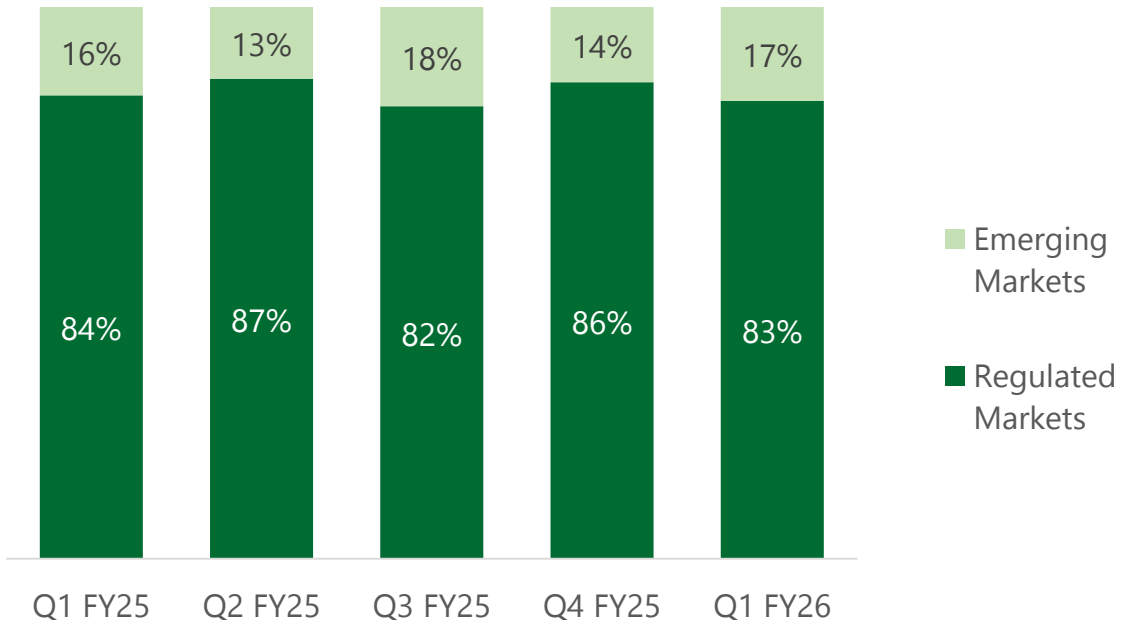
● % of Total Revenue



- Non-GPL business saw a strong growth of 14.5% YoY and 4.1% QoQ in Q1FY26.
- On YoY basis, Non-GPL business was driven by strong growth in regions like India, Europe, ROW, LATAM and Japan.

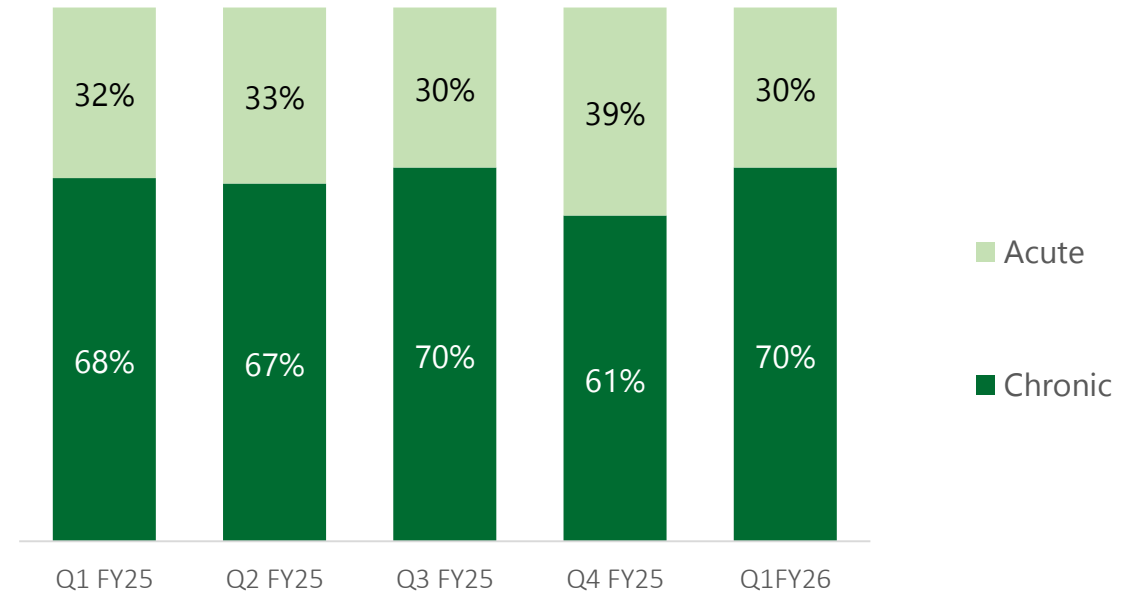
Market and Therapeutic Area Mix

Market Mix



- Regulated markets contributed 83% in Q1FY26 driven by robust performance in Non-GPL business.
- The growth was primarily driven by Europe and Japan in the Regulated markets, whereas India (Ex-GPL) and ROW led the growth in Emerging markets.

Therapeutic Mix



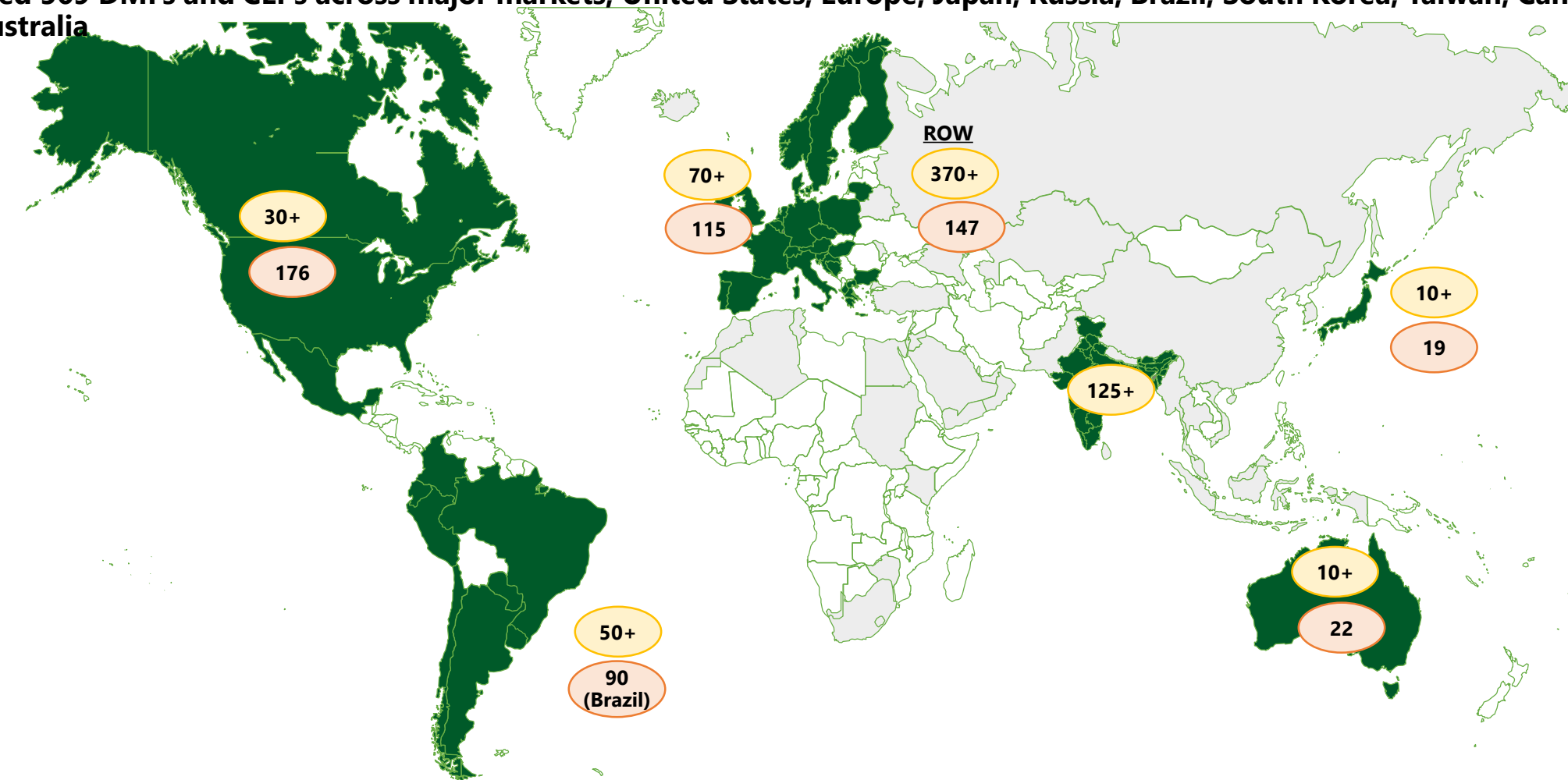
- Chronic portfolio continued to deliver a strong growth.
- The chronic therapies contributed 70% of the revenue in Q1FY26.

Company Overview



Global Footprint

- Filed 569 DMFs and CEPs across major markets; United States, Europe, Japan, Russia, Brazil, South Korea, Taiwan, Canada, China and Australia



■ Regulated Markets ■ Emerging Markets

■ India – Mix of Regulated and Emerging

○ Number of DMF/CEP filings

○ Number of customers serviced in FY25

As of June 30, 2025

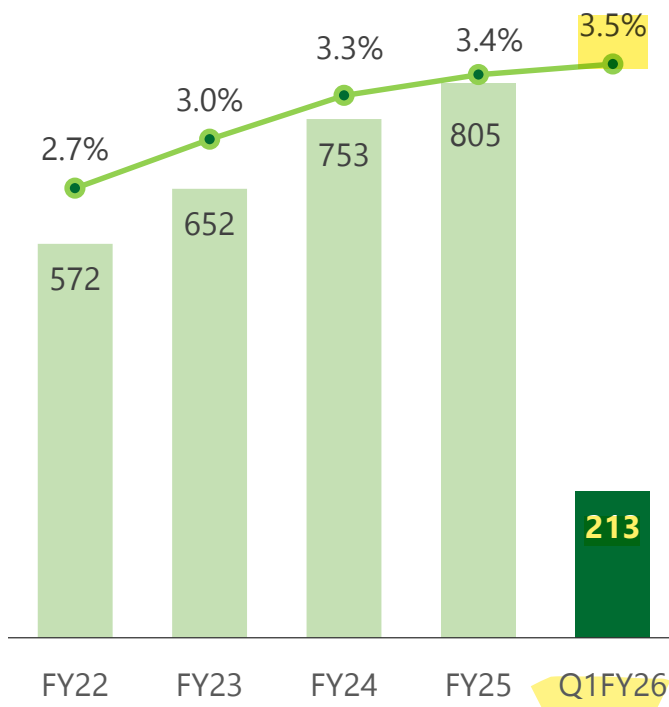


R&D Capabilities

R&D Spend

(In ₹ Million)

● % of Total Revenue



Cumulative Filing Status

Therapy	North America	Europe	Japan	Brazil	Australia	ROW	Total
CVS	39	40	4	23	10	41	157
CNS	42	26	8	17	3	21	117
Anti-Infective	20	11	2	3	3	13	52
Diabetes	10	5	-	9	-	15	39
Dermatology	8	6	1	12	1	9	37
Urology	10	7	2	5	1	9	34
Allergy	9	7	1	7	1	8	33
Others	38	13	1	14	3	31	100
Total	176	115	19	90	22	147	569

- DMF/CEP filings continue across major markets in Q1FY26, taking the total cumulative filings to 569 as on 30th June, 2025.
- Five synthetic small molecules were added to the development grid.
- The HP API portfolio remains on the development path with 26 products in the active grid representing market size of \$ 61 bn (Source: IQVIA, MAT March'25); nine products are validated, three products are in advanced stages of development, remaining 14 products progressing through lab development stages.
- Development progressing for iron complexes in the grid. Filing completed for 1 iron complex, 2 others are in advanced stages of development and preliminary development ongoing for 1 iron complex. Total addressable market of \$2.7bn (Source: IQVIA, MAT Mar'25).



Quality Focused Manufacturing and R&D Infrastructure

Manufacturing Infrastructure

Location	Annual Installed Capacity	Last USFDA Inspection Date	Approvals
Ankleshwar, Gujarat	950.2 KL	Jan 2025	USFDA, MHRA (UK), FIMEA (Finland), Romania (Europe) PMDA (Japan), COFEPRIS (Mexico), Health Canada, KFDA (South Korea), Gujarat FDCA, ANVISA (Brazil)
Dahej, Gujarat	399.9 KL	May 2025	USFDA, EDQM (Europe), PMDA (Japan), KFDA (South Korea), ANVISA (Brazil)
Mohol, Maharashtra	49.1 KL	March 2018	USFDA, Maharashtra FDA
Kurkumbh, Maharashtra	24.6 KL	-NA-	Maharashtra FDA

R&D Infrastructure

Mahape, Navi Mumbai

- R&D for new product development and complex molecules
- High-end analytical equipment for characterization

Ankleshwar, Gujarat

- Cost improvement programs and process improvements

Dahej, Gujarat

- Oncology R&D
- Cost improvement programs and process improvements

Strategy **Going Forward**



Strategy Growth Levers

New Growth levers

2

- ✓ CDMO Ramp up
- ✓ Expand into complex API platforms
- ✓ Iron compounds
- ✓ Oncology & HP API

Operational efficiencies

4

- ✓ Debottlenecking
- ✓ 2nd/3rd generation process adoption
- ✓ Backward integration
- ✓ Reduce carbon footprint
- ✓ Adoption of flow chemistry in manufacturing
- ✓ Pursue AVD opportunities

1 Gx API Business

- ✓ New product launches
- ✓ Geographical expansion
- ✓ Focus on new markets becoming more regulated
- ✓ Pursue 2nd source opportunities with top generic players

3 Capacity

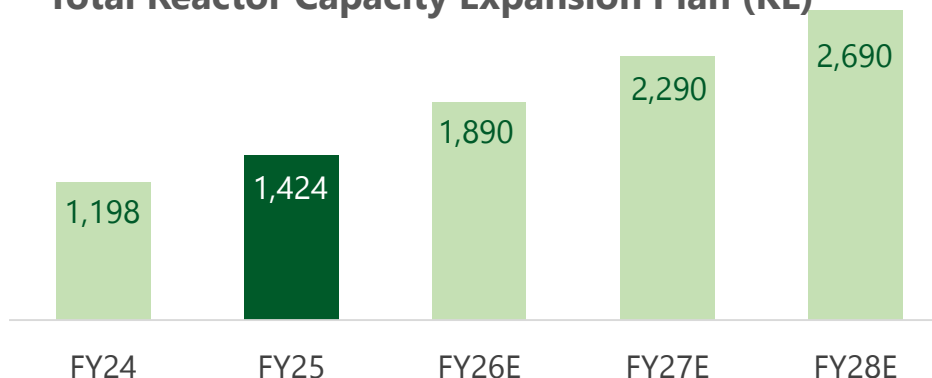
- ✓ Greenfield – Solapur, 1000MT (CTE Received and Phase 1 construction of 200 KL started)
- ✓ Second Phase Dahej expansion
- ✓ Ankleshwar Pharma blocks expansion
- ✓ Build standalone R&D infrastructure for expansion into new growth levers

Future Capacity Expansion

Expansion Type	Division	Location	Status & Planned Capacity	Operational Timelines
Brownfield	API / Intermediate	Ankleshwar	Planned addition of ~100KL Capacity	FY26
Brownfield	API	Dahej	Planned addition of ~160KL	FY26
Greenfield	API	Solapur	Phase 1 – ~200 KL (Construction is in process)	FY26
			Phase 1.1 – ~400 KL (Planned Backward Integration)	FY27
			Phase 2 - Planned addition of ~400 KL	FY28

Total Reactor Capacity Expansion Plan (KL)

**Capacity
Progress
by Year**



- ✓ **Construction work of 200 KL capacity (Phase 1) is in process at Solapur Plant**
- ✓ **Solapur's further capacity expansion will be calibrated as per the volume demand**

Thank You

FOR FURTHER INFORMATION CONTACT

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