

May 15, 2025

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

Ref: Scrip Code: 543322

To,
The Manager – Listing,
National Stock Exchange of India Ltd.,
Plot No. C/1, G Block,
Bandra Kurla Complex,
Bandra (E), Mumbai – 400 051

Ref: Scrip Name: ALIVUS

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 – Q4 & FY 24-25. 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



Q4 & FY25



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Important factors that could cause actual results to differ materially from our expectations include, amongst others, general economic and business conditions in India and abroad, ability to successfully implement our strategy, our research & development efforts, our growth & 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.

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Financial Performance Review

A Q4 FY25 | Highlights



Dr. Yasir Rawjee

Managing Director &
Chief Executive Officer

"FY25 was our first full year under the Nirma ownership. Our overall performance has remained consistent with past years, with sustained margins, despite facing external challenges, most notably the loss in PLI revenue. The road ahead has significant opportunities for which we have begun working in earnest via capacity investments and the integration of newer technologies into R&D and manufacturing operations.

One of the most encouraging outcomes this year was our ability to maintain EBITDA margins at 30%, even in the absence of PLI benefits and amid market headwinds. This reflects the strength of our diversified portfolio and our continued focus on operational efficiency.

Our full-year sales grew by 7.1%, normalizing for the PLI benefit last year and backed by healthy contributions from both GPL and Non-GPL segments. As we move into FY26, we remain committed to delivering sustainable growth with stable margins, while continuing to invest in our core growth drivers, expanding our product pipeline, and scaling our operations to create long-term value."

REVENUE
(IN ₹ MILLIONS)

6,496 1.2%
QoQ 21.1%
YoY

EBITDA
(IN ₹ MILLIONS)

2,085 3.8%
QoQ 44.2%
YoY

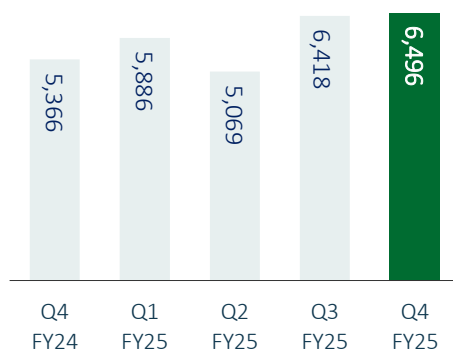
PAT
(IN ₹ MILLIONS)

1,419 3.6%
QoQ 44.9%
YoY

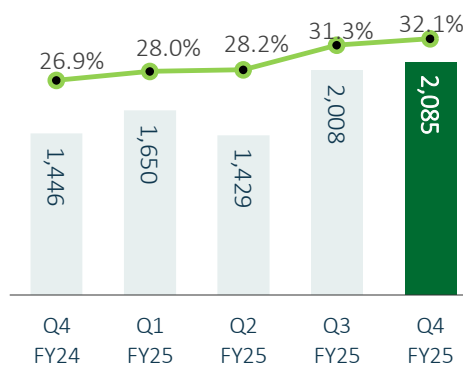
- Alivus registered a revenue from operations of ₹ 6,496 Mn for Q4FY25, recording a strong growth of 21.1% YoY.
- EBITDA margins for the quarter were at 32.1%, up 520 bps YoY and 80 bps QoQ, driven by better product mix and new product launches.
- Generic business grew by 22.6% YoY to ₹ 5,941 Mn, whereas CDMO business grew by 22.6% YoY and 44.4% QoQ to ₹ 434 Mn.
- Full year, GPL business grew by 8.8% YoY at ₹ 7,805 Mn, whereas, Non-GPL business grew by 6.3% YoY to ₹ 15,594 Mn. Overall, sales revenue growth (excluding other operating income) remains at 7.1%
- During FY25, the company generated a strong free cash flow of Rs. 2,328 Mn leading to Cash and Cash Equivalents (incl. short term investments) of Rs. 5,487 Mn as of 31 March 2025.

Q4 & FY25 Performance

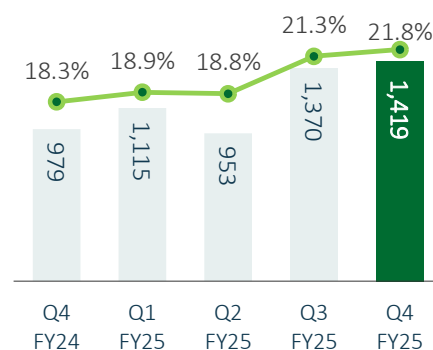
Revenue
(In ₹ Millions)



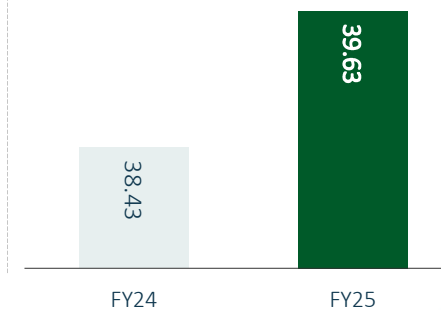
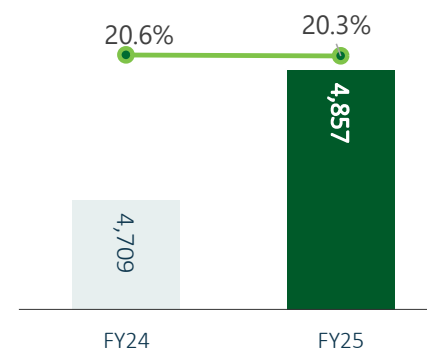
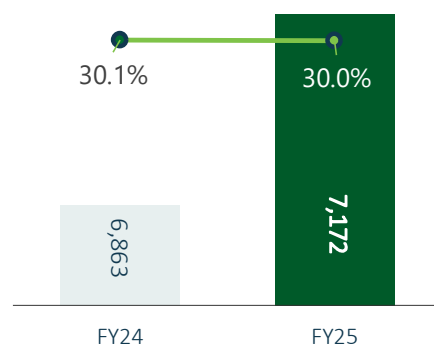
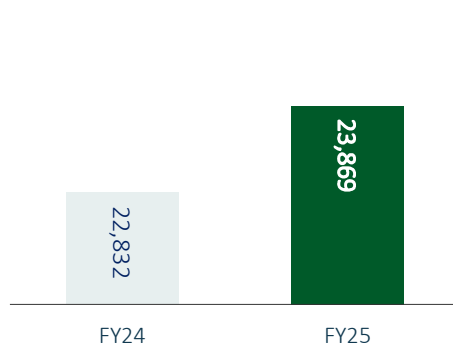
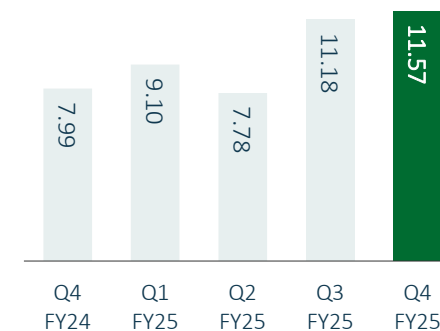
EBITDA
(In ₹ Millions)



PAT
(In ₹ Millions)



EPS
(In ₹)

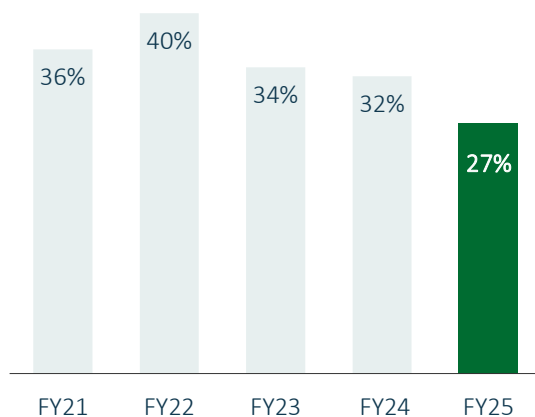


P&L Highlights | Q4 & FY25

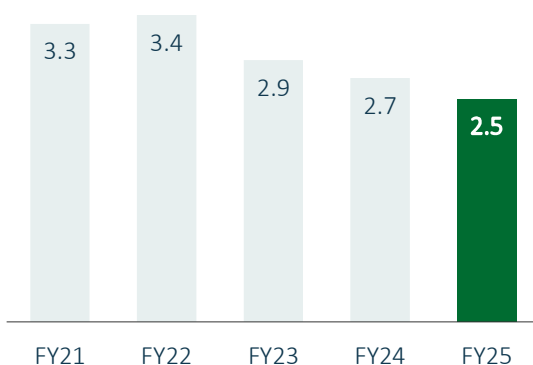
Particulars (In ₹ Millions)	Q4 FY25	Q3 FY25	QoQ	Q4 FY24	YoY	FY25	FY24	YoY
Revenue from Operations	6,496	6,418	1.2%	5,366	21.1%	23,869	22,832	4.5%
Gross Profit	3,670	3,566	2.9%	2,979	23.2%	13,061	12,812	1.9%
Gross Profit (%)	56.5%	55.6%		55.5%		54.7%	56.1%	
Other Income	101	105	-3.8%	31	225.8%	346	120	188.3%
Employee Benefits Expense	668	657	1.7%	723	-7.6%	2,517	2,581	-2.5%
Other Expenses	1018	1,006	1.2%	841	21.0%	3,718	3,488	6.6%
EBITDA	2,085	2,008	3.8%	1,446	44.2%	7,172	6,863	4.5%
EBITDA Margin (%)	32.1%	31.3%		26.9%		30.0%	30.1%	
Depreciation and Amortisation Expense	160	152	5.3%	145	10.3%	606	535	13.3%
Finance Costs	12	5	140.0%	4	200.0%	24	15	60.0%
PBT	1,913	1,851	3.3%	1,297	47.5%	6,542	6,313	3.6%
PBT Margin (%)	29.4%	28.8%		24.2%		27.4%	27.7%	
PAT	1,419	1,370	3.6%	979	44.9%	4,857	4,709	3.1%
Net Margin (%)	21.8%	21.3%		18.3%		20.3%	20.6%	

Healthy Return Indicators

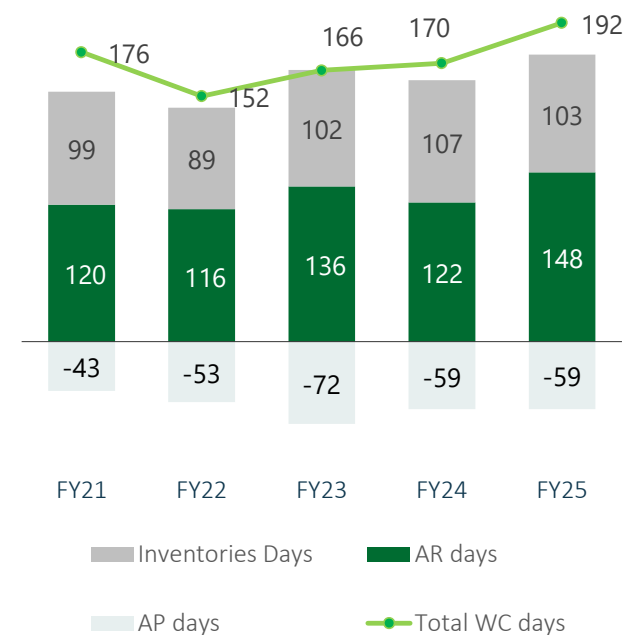
ROICE (%)



Fixed Assets Turnover (Times)



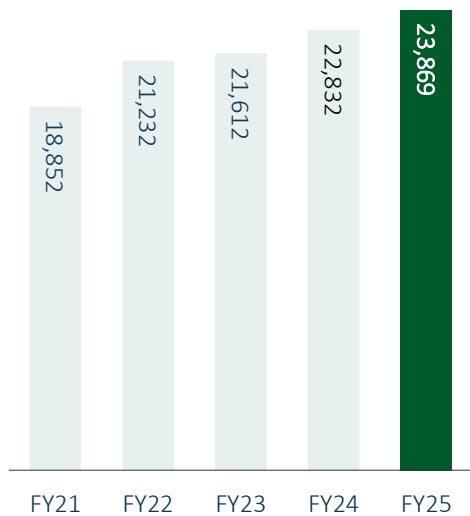
Working Capital Days



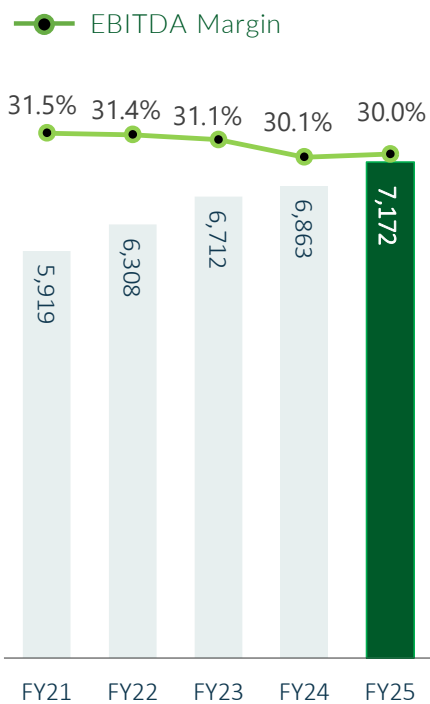
- ROICE is tracking at **~27%** – Capital employed driven by calibrated Capex strategy
- FATR is **2.5 times** – Asset turn trending slightly lower due to Capex cycle
- WC at **192 days** – Higher Day Sales Outstanding driven by business mix
- **Strong Balance Sheet** – Strong free cash generation of ₹ 2,328 Mn leading to Cash and Cash Equivalents (including short term investments) of ₹ 5,487 Mn as of 31 March 2025.

Robust Growth & Steady Profitability

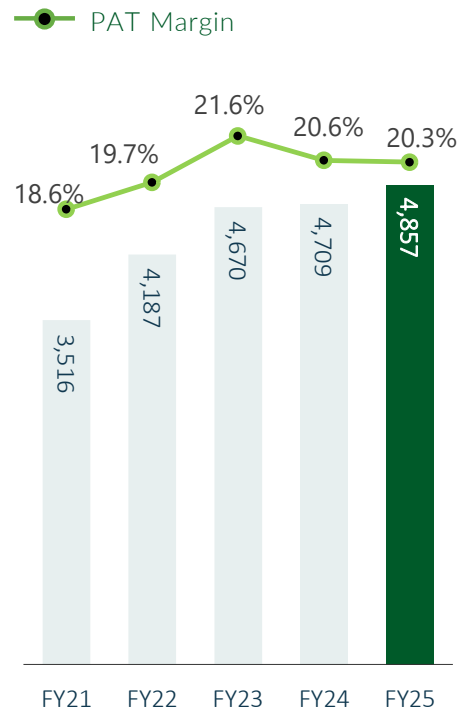
Revenue
(In ₹ Millions)



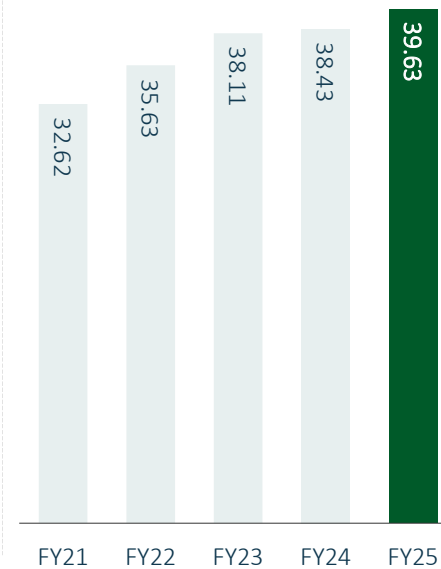
EBITDA
(In ₹ Millions)



PAT
(In ₹ Millions)



EPS
(In ₹)



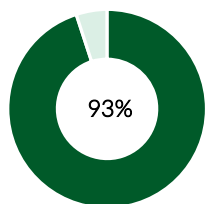
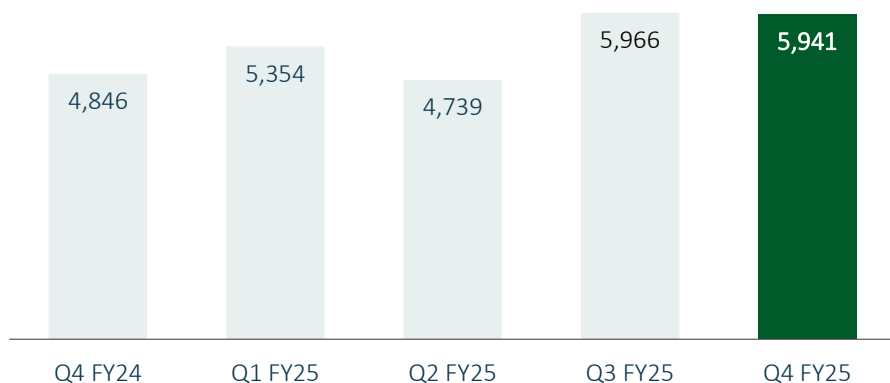


Business Performance Review

Segmental Performance | Generic API vs CDMO

Generic API

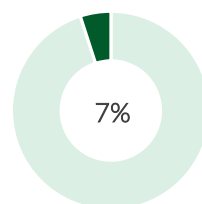
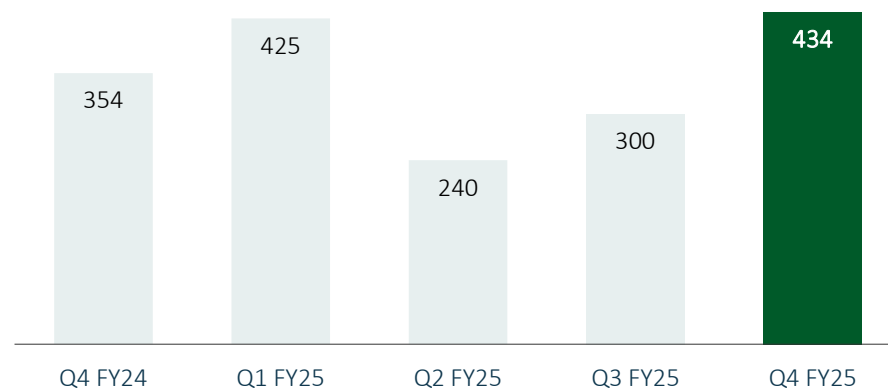
(Revenue In ₹ Millions)



- Generic API revenues in Q4FY25 grew by 22.6% YoY.
- Robust performance in GPL and Non-GPL business led the growth.
- Regions like India, Europe, ROW and Japan contributed to revenue growth.

CDMO

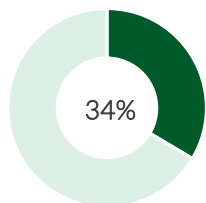
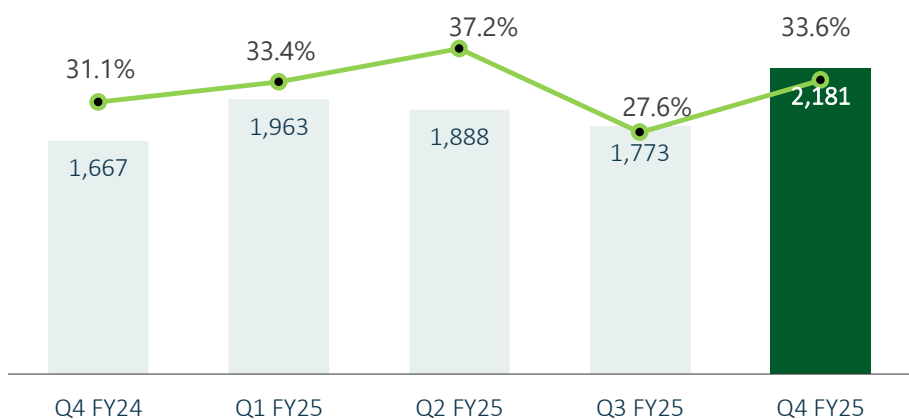
(Revenue In ₹ Millions)



- CDMO business witnessed growth of 22.6% YoY and 44.4% QoQ due to addition of 4th project during the year.
- 5th project is expected to be commercialized by H2FY26.
- Multiple discussions are ongoing.

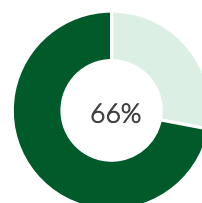
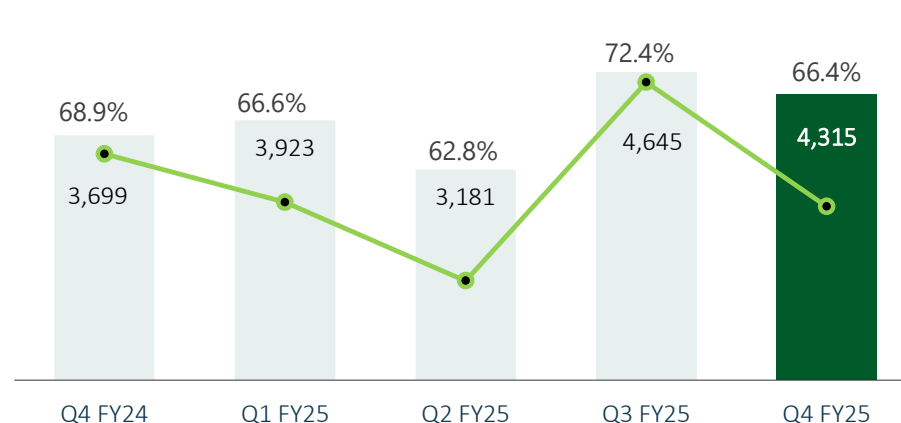
Segmental Performance | GPL vs Non-GPL

GPL
(In ₹ Millions)



- In Q4FY25, GPL business witnessed a strong growth of 23.0% QoQ and 30.8% YoY.
- GPL business contributes ~34% of the total revenue from operations

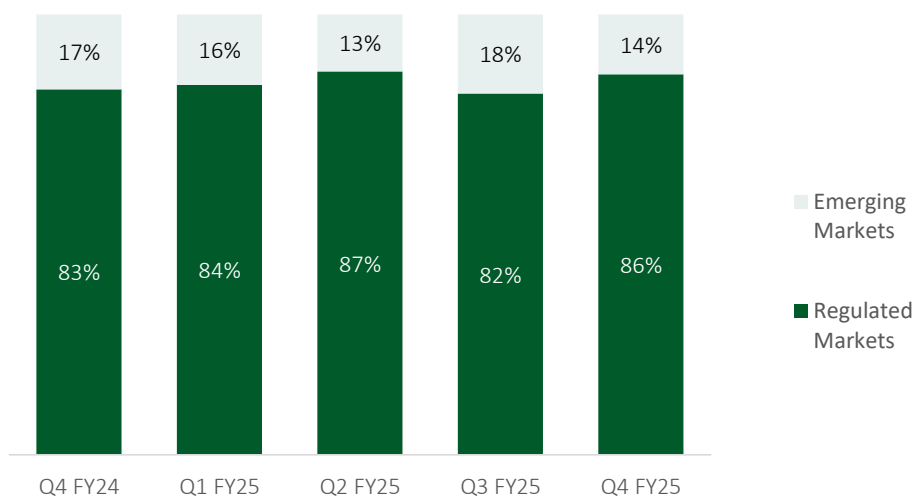
Non-GPL
(In ₹ Millions)



- Non-GPL business saw a growth of 16.6% YoY
- Non-GPL business was driven by strong growth in regions like India, Europe, ROW and Japan.

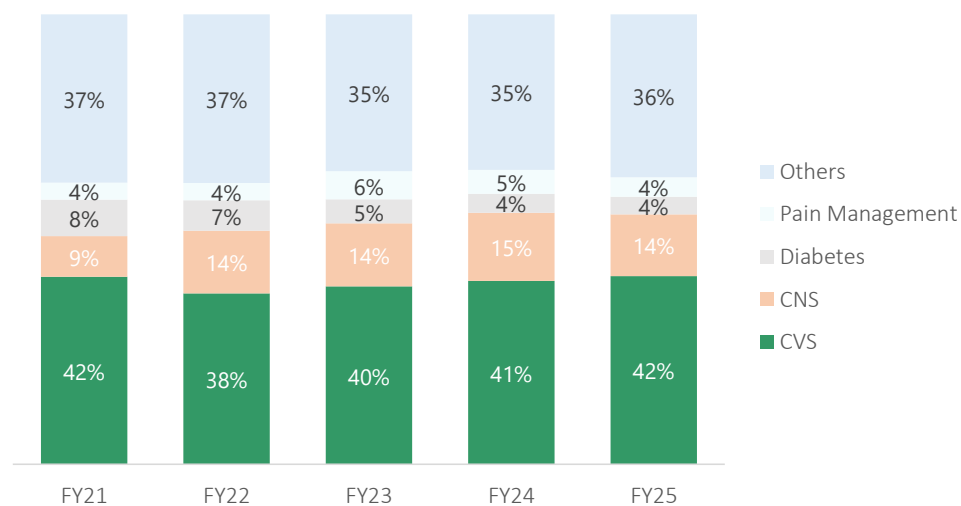
Market and Therapeutic Area Mix

Market Mix



- Regulated markets contributed 86% in Q4FY25 driven by robust performance in GPL and Non-GPL business.
- On a full year basis, regulated market contributed 85% of the total portfolio compared to 81% in FY24.

Therapeutic Area Mix



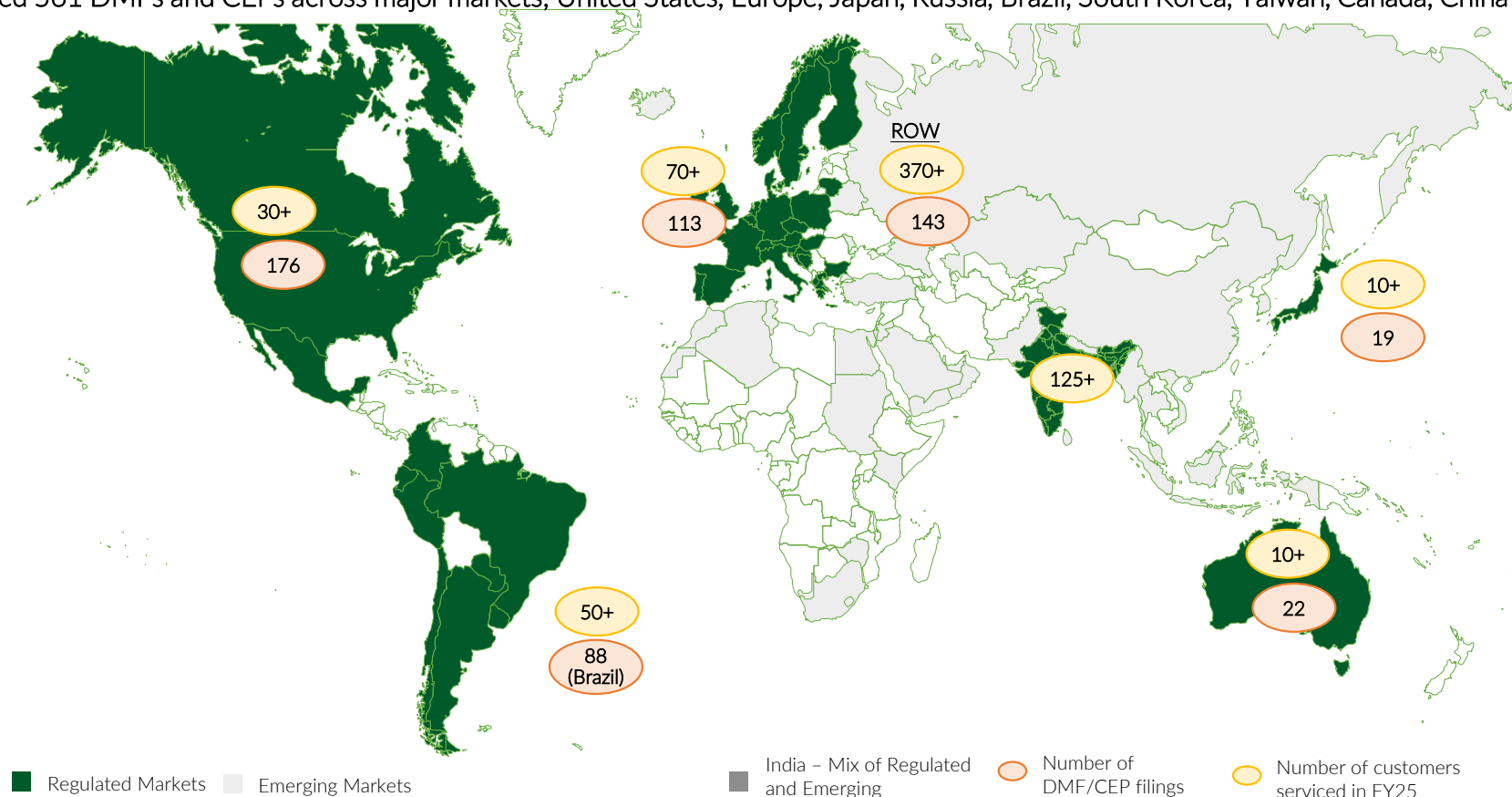
- CVS and CNS portfolio continued to deliver a strong growth.
- Our key focused area of chronic therapies contributed 64% of the revenue in FY25.



Company Overview

Global Footprint

- Filed 561 DMFs and CEPs across major markets; United States, Europe, Japan, Russia, Brazil, South Korea, Taiwan, Canada, China and Australia



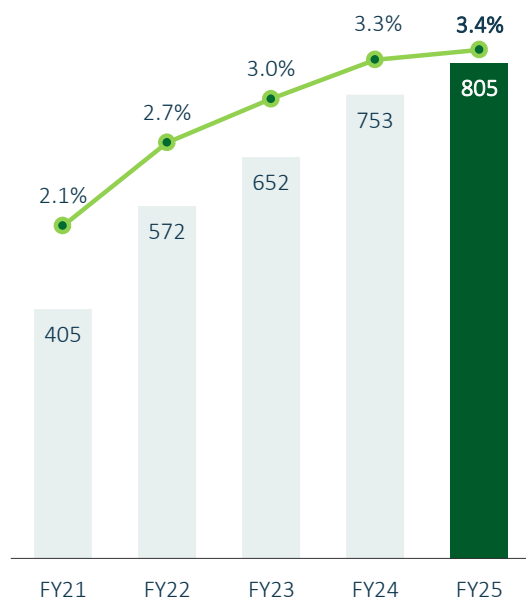
As of March 31, 2025

R&D Capabilities

R&D Spend

(In ₹ Millions)

● % of Total Revenue



Cumulative Filing Status

Therapy	North America	Europe	Japan	Brazil	Australia	ROW	Total
CVS	39	39	4	22	10	38	152
CNS	41	26	8	16	2	20	113
Diabetes	10	5	-	9	-	14	38
Pain Management	1	2	-	4	1	9	17
Others	85	41	7	37	9	62	241
Total	176	113	19	88	22	143	561

- DMF/CEP filings continue across major markets in Q4FY25, taking the total cumulative filings to 561 as on 31st March 2025.
- Four synthetic small molecules were added to the development grid.
- The HP API portfolio remains on the development path with 24 products in the active grid representing market size of \$ 49 bn (Source: IQVIA, MAT Dec'24); seven products are validated, five products are in advanced stages of development, remaining 12 products progressing through lab development stages.
- Development progressing for iron complexes in the grid. Filing completed for one iron complex with two others in advanced stages of development. Total addressable market of \$2.2 bn (Source: IQVIA, MAT Dec'24)

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	Oct 2018	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

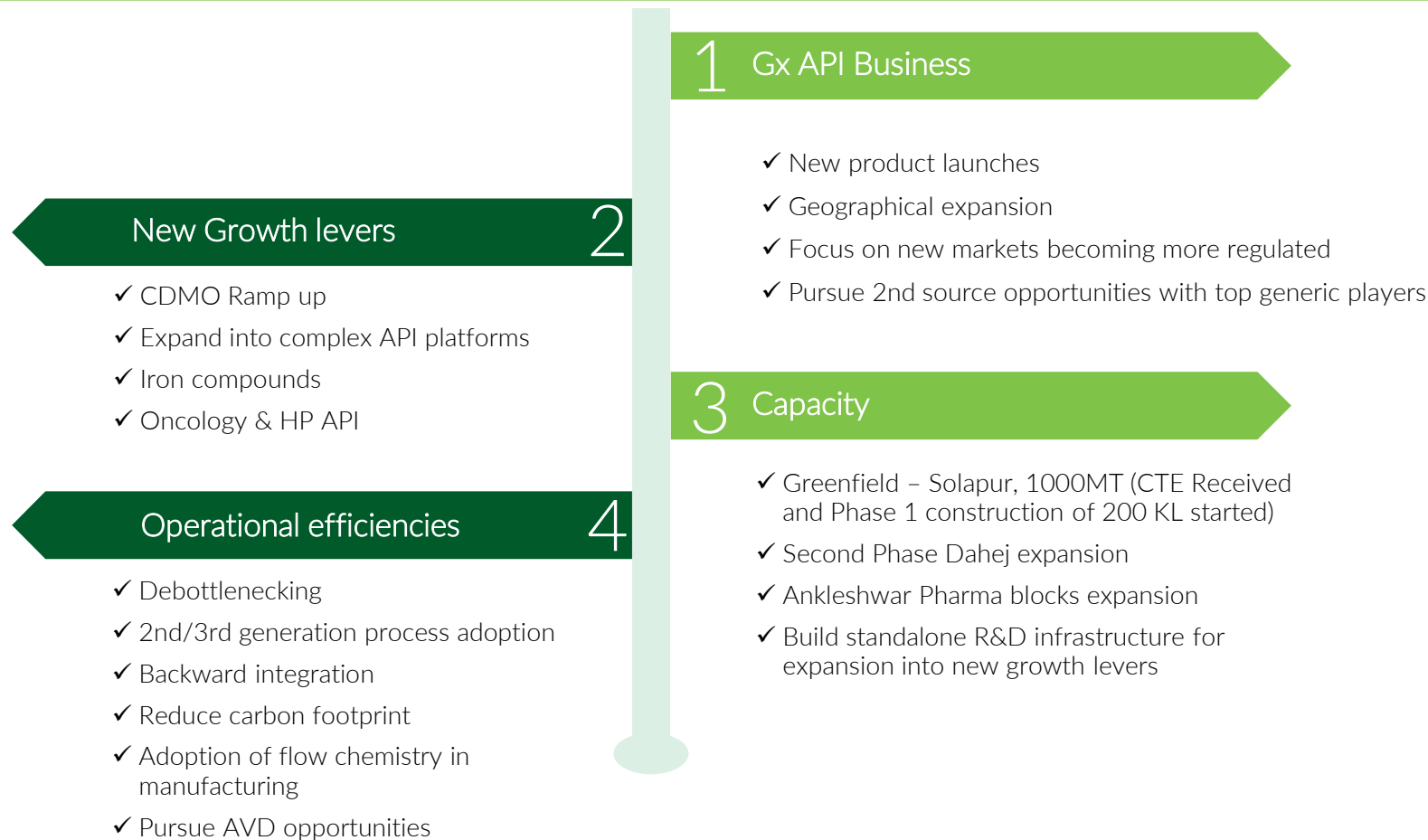
Dahej, Gujarat

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



Strategy **Going Forward**

Strategy Growth Levers

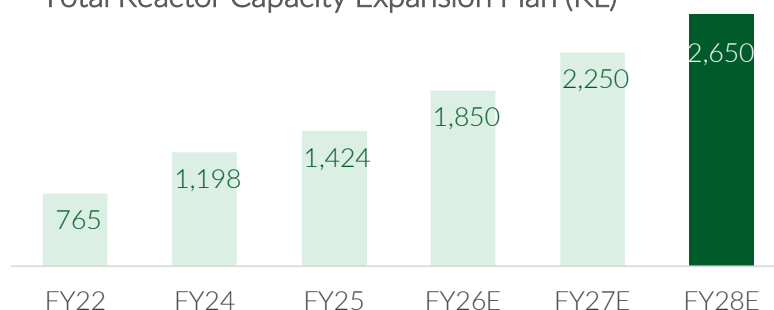


Future Capacity Expansion

Expansion Type	Division	Location	Status & Planned Capacity	Operational Timelines
Brownfield	API / Intermediate	Ankleshwar	Planned addition of ~60KL Pharma Capacity	FY26
Brownfield	API	Dahej	Planned addition of ~160KL	FY26
Greenfield	API	Solapur	Phase 1 – ~200 KL (Construction is in process) Phase 1.1 – ~400 KL (Planned Backward Integration) Phase 2 - Planned addition of ~400 KL	FY26 FY27 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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