

January 23, 2025

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

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 Code: 543322

Ref: Scrip Name: ALIVUS

Dear Sirs,

Sub: Investor Presentation

Pursuant to regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements), 2015, we are enclosing herewith the Investor Presentation – Q3 FY 24-25.

You are requested to take the same on record.

Thanking You.

Yours faithfully,
For Alivus Life Sciences Limited
(Formerly known as Glenmark Life Sciences Limited)

Rudalf Corriea
Company Secretary & Compliance Officer
Encl: As above

Alivus Life Sciences Limited (formerly Glenmark Life Sciences Limited)

Corporate Office:

4th Floor, OIA House, 470, Cardinal Gracious Road
Andheri (E), Mumbai 400 099, Maharashtra, India

Registered Office:

Plot No 170-172, Chandramouli Industrial Estate
Mohol Bazarpeth, Solapur 413 213, India

T: +91 22 6829 7979 | CIN: L74900PN2011PLC139963 | E: complianceofficer@glenmarklifesciences.com | W: www.alivus.com



Investor Presentation



Q3 & 9M FY25



Disclaimer

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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



Q3FY25 | Highlights



Dr. Yasir Rawjee
Managing Director &
Chief Executive Officer

“The start of the new year marks a notable shift for us as we transition to Alivus Life Sciences.

I am pleased to report that our Q3 performance reflects this renewed energy, with growth across both GPL and Non-GPL segments. Geographically, regions like India, Europe, ROW and Japan contributed to the growth.

Our quest for high-quality, innovative solutions and scalability continues, in order to build a sustainable business over the long term.”

REVENUE (IN ₹ MILLIONS)	6,418	26.6% QoQ	12.0% YoY
EBITDA (IN ₹ MILLIONS)	2,008	40.5% QoQ	15.2% YoY
PAT (IN ₹ MILLIONS)	1,370	43.7% QoQ	15.3% YoY

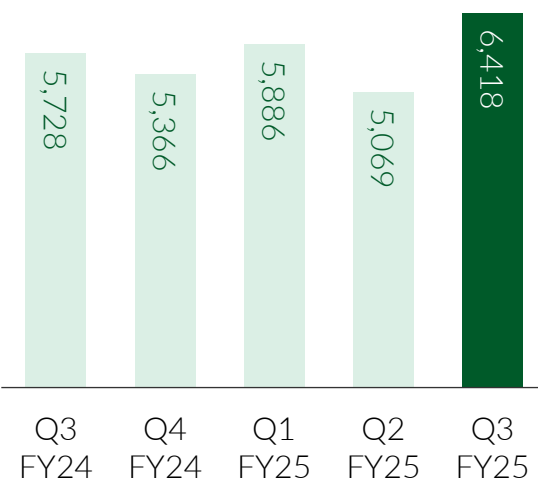
- Alivus registered a revenue from operations of ₹ 6,418 Mn for Q3FY25, recording a strong growth of 26.6% QoQ and 12.0% YoY.
- EBITDA margins for the quarter were at 31.3% up 310 bps QoQ and 90 bps YoY, driven by better product mix with stable expenses.
- Generic business grew by 16.9% YoY and 25.9% QoQ to ₹ 5,966 Mn, whereas CDMO business grew by 25.0% QoQ to ₹ 300 Mn.
- Non-GPL business grew by 10.9% YoY and 46.0% QoQ to ₹ 4,645 Mn; GPL business grew by 15.1% YoY at ₹ 1,773 Mn
- During 9MFY25, the company generated a strong free cash flow of Rs. 1,838 Mn leading to Cash and Cash Equivalents (including short term investments) of Rs. 4,993 Mn as of 31 December 2024.



Q3 & 9M FY25 Performance

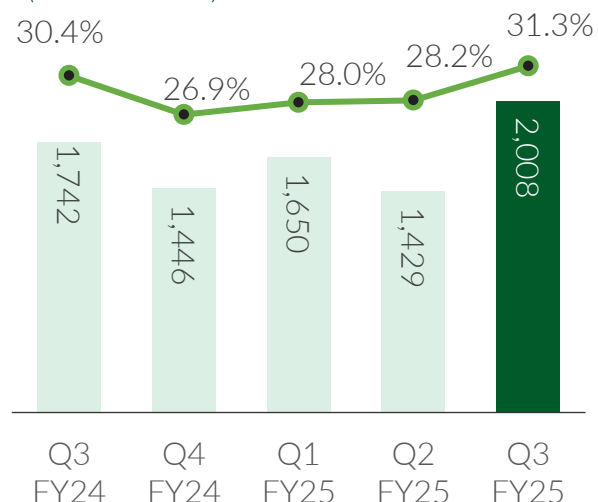
Revenue

(In ₹ Millions)



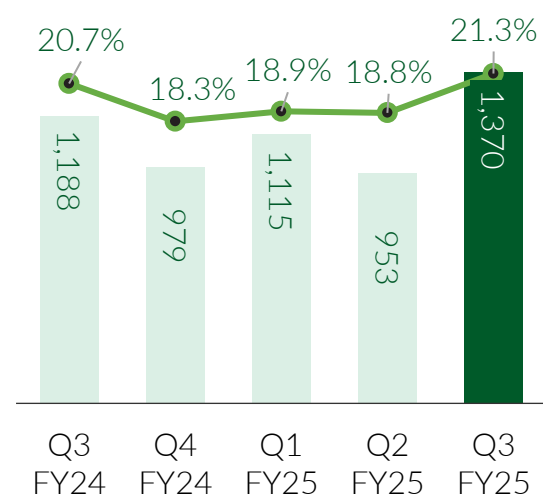
EBITDA

(In ₹ Millions)



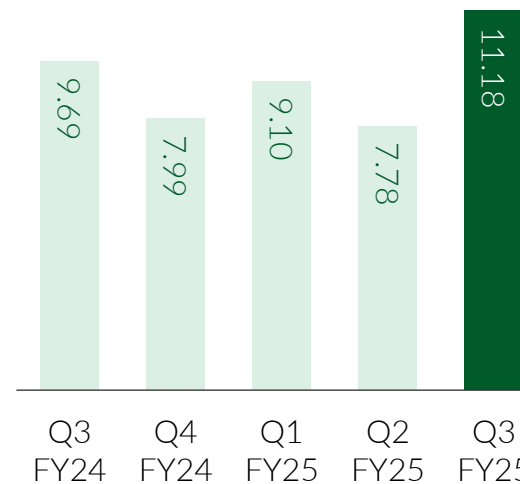
PAT

(In ₹ Millions)



EPS

(In ₹)



17,466

17,373

9M FY24

9M FY25

31.0%

5,417

29.3%

5,086

9M FY24

9M FY25

21.4%

3,730

19.8%

3,438

9M FY24

9M FY25

30.44

28.06

9M FY24

9M FY25

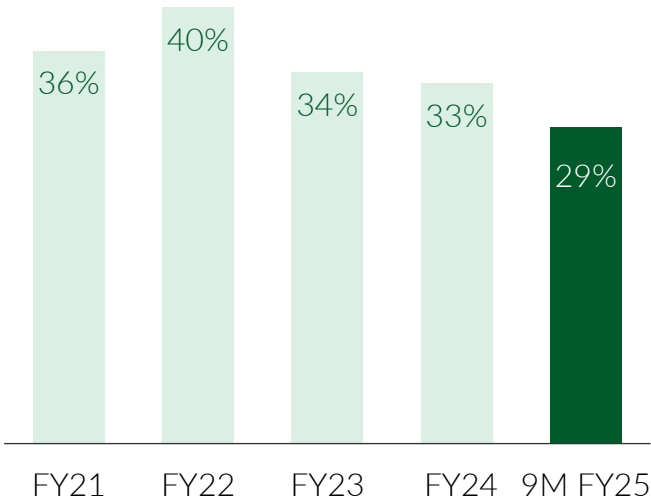
P&L Highlights | Q3 & 9M FY25

Particulars (In ₹ Millions)	Q3 FY25	Q2 FY25	QoQ	Q3 FY24	YoY	9M FY25	9M FY24	YoY	FY24
Revenue from Operations	6,418	5,069	26.6%	5,728	12.0%	17,373	17,466	-0.5%	22,832
Gross Profit	3,566	2,817	26.6%	3,306	7.9%	9,391	9,833	-4.5%	12,812
Gross Profit (%)	55.6%	55.6%		57.7%		54.1%	56.3%		56.1%
Other Income	105	85	23.0%	17	515.0%	245	89	175.1%	120
Employee Benefits Expense	657	624	5.3%	711	-7.6%	1,850	1,859	-0.5%	2,581
Other Expenses	1,006	849	18.5%	870	15.6%	2,700	2,646	2.0%	3,488
EBITDA	2,008	1,429	40.5%	1,742	15.2%	5,086	5,417	-6.1%	6,863
EBITDA Margin (%)	31.3%	28.2%		30.4%		29.3%	31.0%		30.1%
Depreciation and Amortisation Expense	152	151	0.4%	132	14.8%	446	389	14.8%	535
Finance Costs	5	3	52.7%	4	14.6%	12	12	-3.8%	15
PBT	1,851	1,275	45.2%	1,607	15.2%	4,628	5,016	-7.7%	6,313
PBT Margin (%)	28.8%	25.2%		28.1%		26.6%	28.7%		27.7%
PAT	1,370	953	43.7%	1,188	15.3%	3,438	3,730	-7.8%	4,709
Net Margin (%)	21.3%	18.8%		20.7%		19.8%	21.4%		20.6%

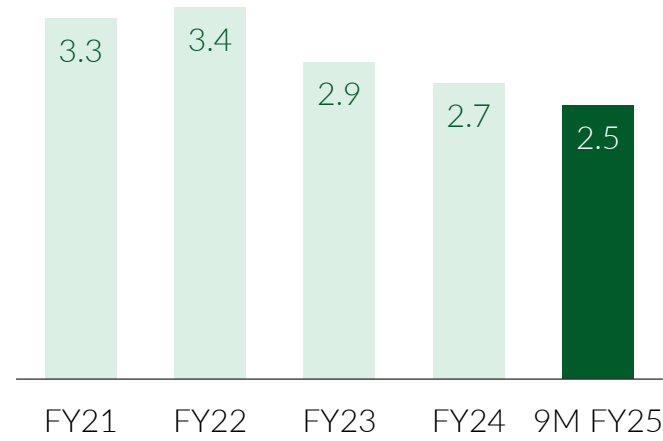


Healthy Return Indicators

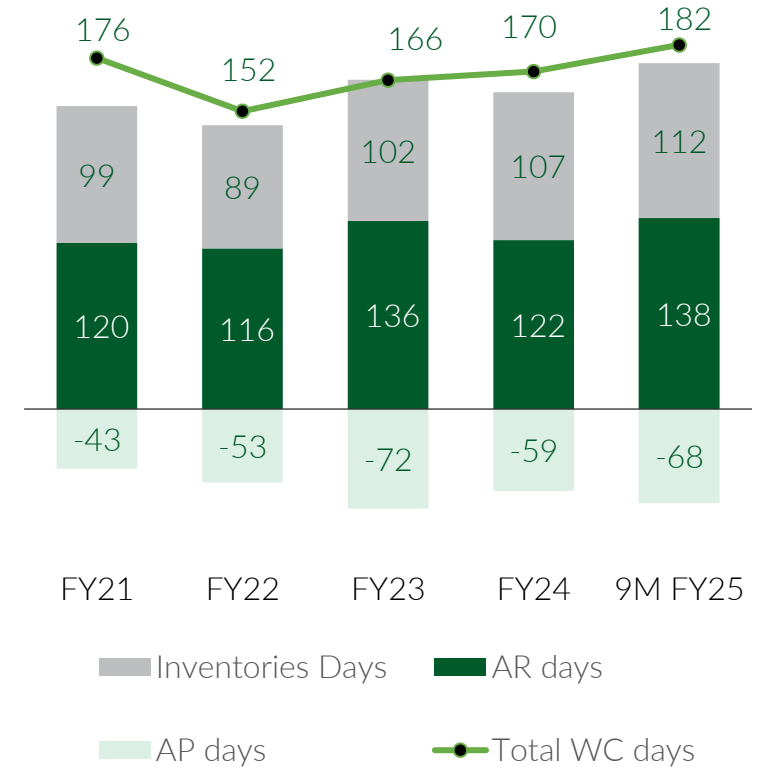
ROICE (%)



Fixed Assets Turnover (Times)



Working Capital Days



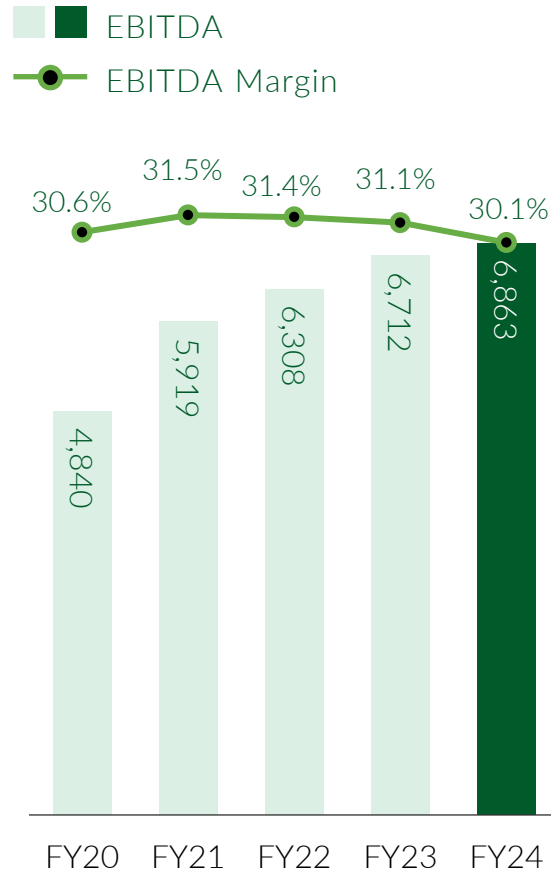
- ROICE is tracking at **~29%** – Capital employed driven by calibrated Capex strategy
- FATR is **2.5 times** – Asset turn trending slightly lower due to Capex cycle
- WC at **182 days** – Higher Day Sales Outstanding driven by business mix
- **Strong Balance Sheet** – Strong free cash generation of ₹ 1,838 Mn leading to Cash and Cash Equivalents (including short term investments) of ₹ 4,993 Mn as of 31 December 2024

Robust Growth & Steady Profitability

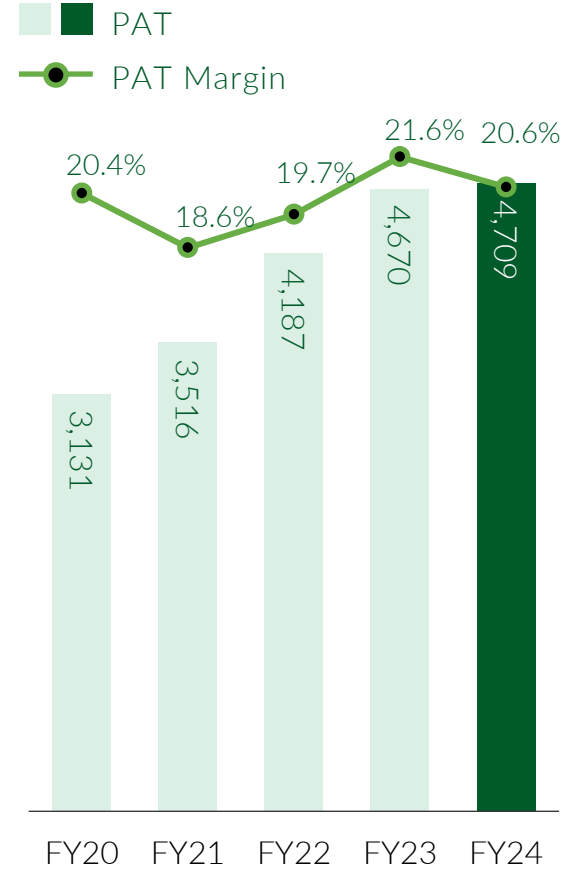
Revenue
(In ₹ Millions)



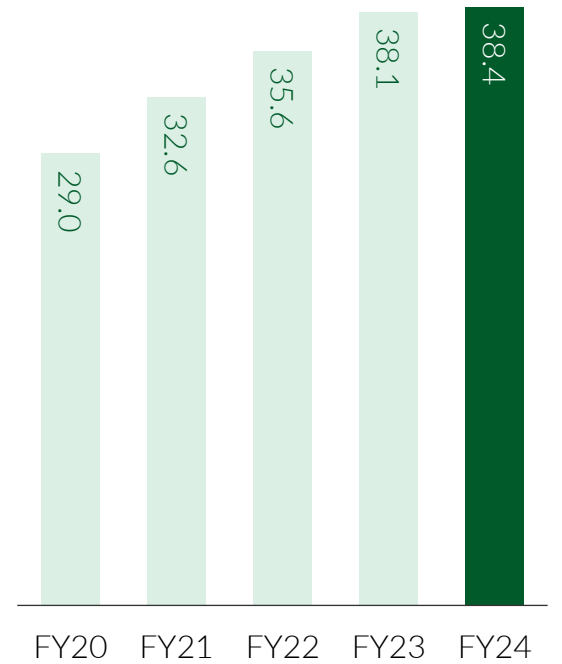
EBITDA
(In ₹ Millions)



PAT
(In ₹ Millions)



EPS
(In ₹)





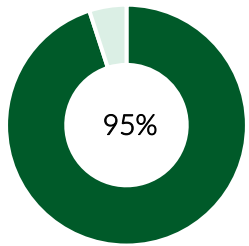
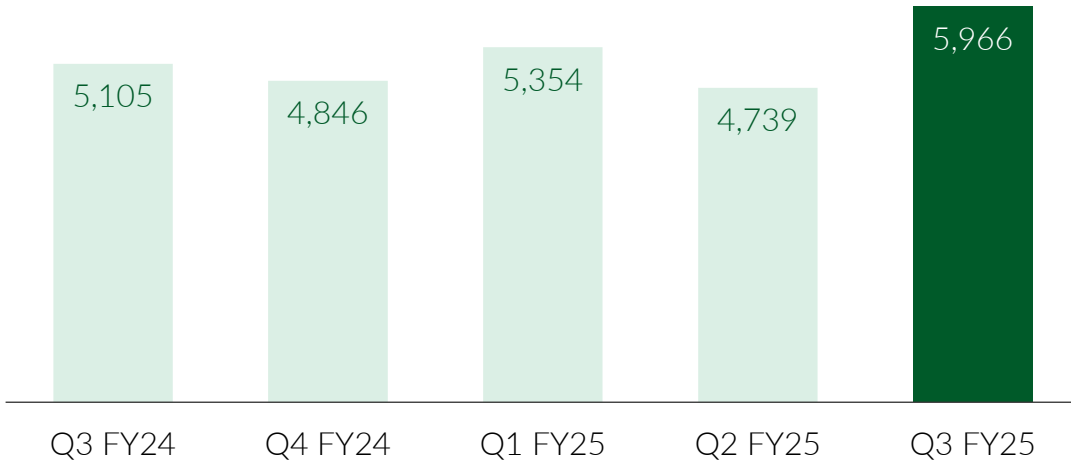
Business Performance Review



Segmental Performance | Generic API vs CDMO

Generic API

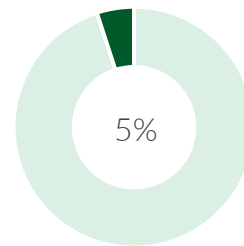
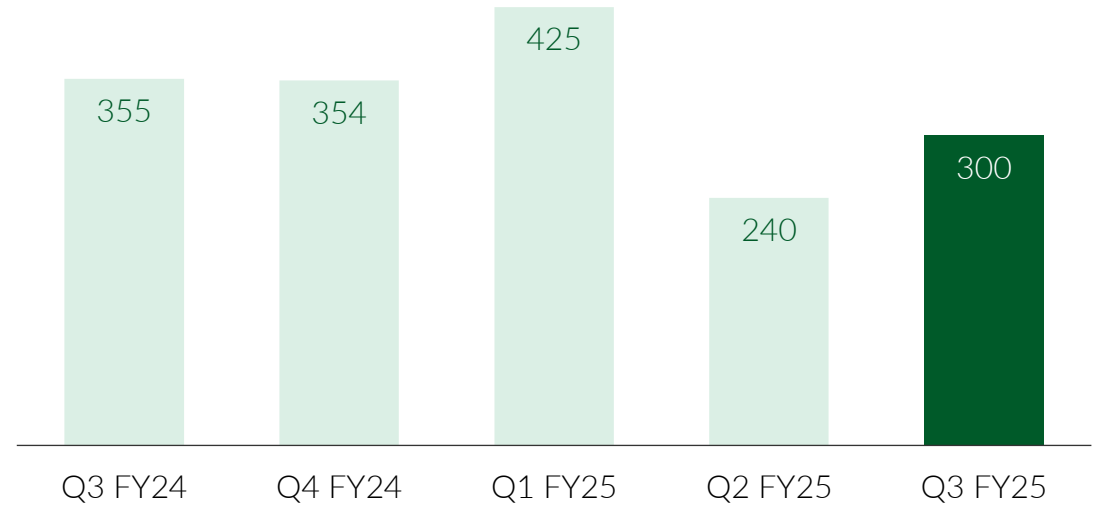
(Revenue In ₹ Millions)



- Generic API revenues in Q3FY25 grew by 25.9% QoQ and 16.9% YoY.
- Regions like Europe, ROW, Japan and India contributed to revenue growth.
- Generic API business was driven by strong growth in regulated markets.

CDMO

(Revenue In ₹ Millions)

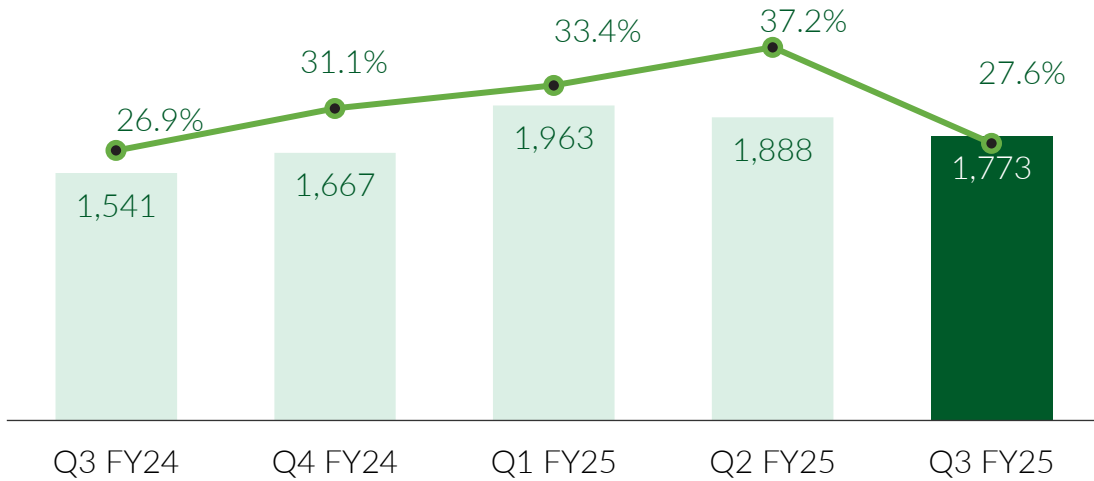


- CDMO business witnessed growth of 25% QoQ, driven by demand contribution from the 4th project.
- 5th project is expected to be commercialized by H1FY26.
- Multiple discussions are ongoing.

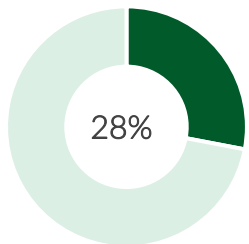
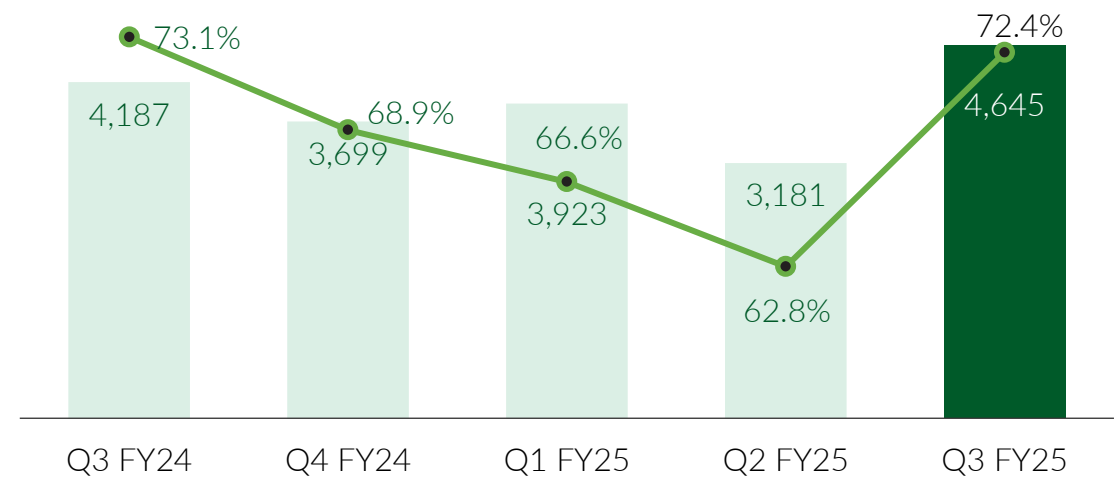


Segmental Performance | GPL vs Non-GPL

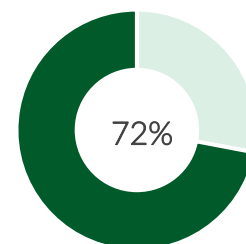
GPL
(In ₹ Millions) ● % of Total Revenue



Non-GPL
(In ₹ Millions) ● % of Total Revenue



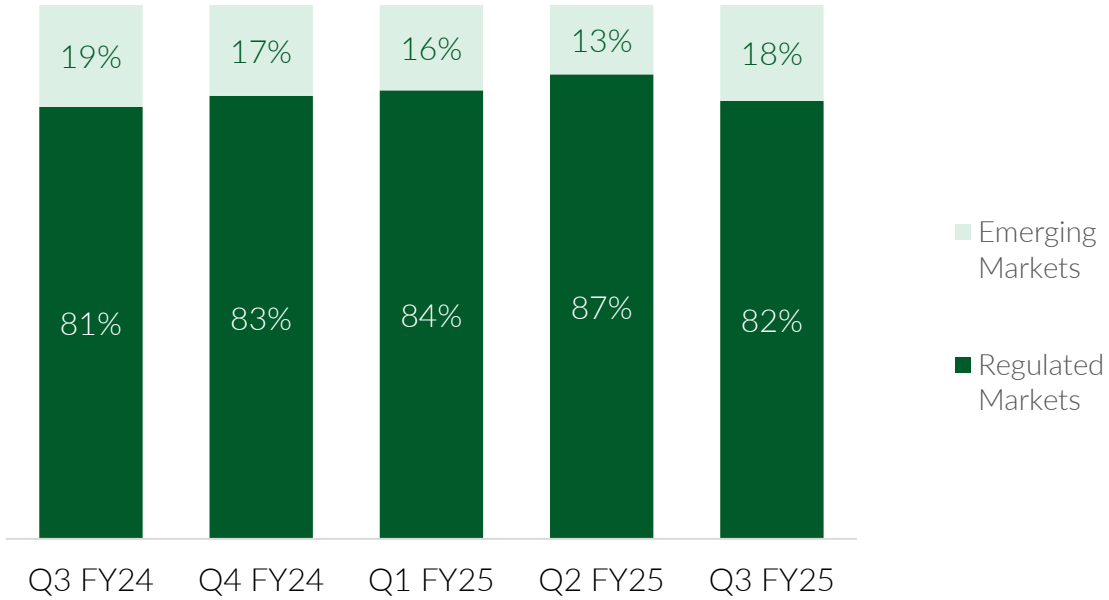
- In Q3FY25, GPL business witnessed a growth of 15.1% YoY and a de-growth of 6.1% QoQ.
- GPL business contributes ~28% of the total revenue from operations



- Non-GPL business saw a growth of 10.9% YoY and 46.0% QoQ.
- Non-GPL business was driven by strong growth in regions like Europe, ROW and Japan.

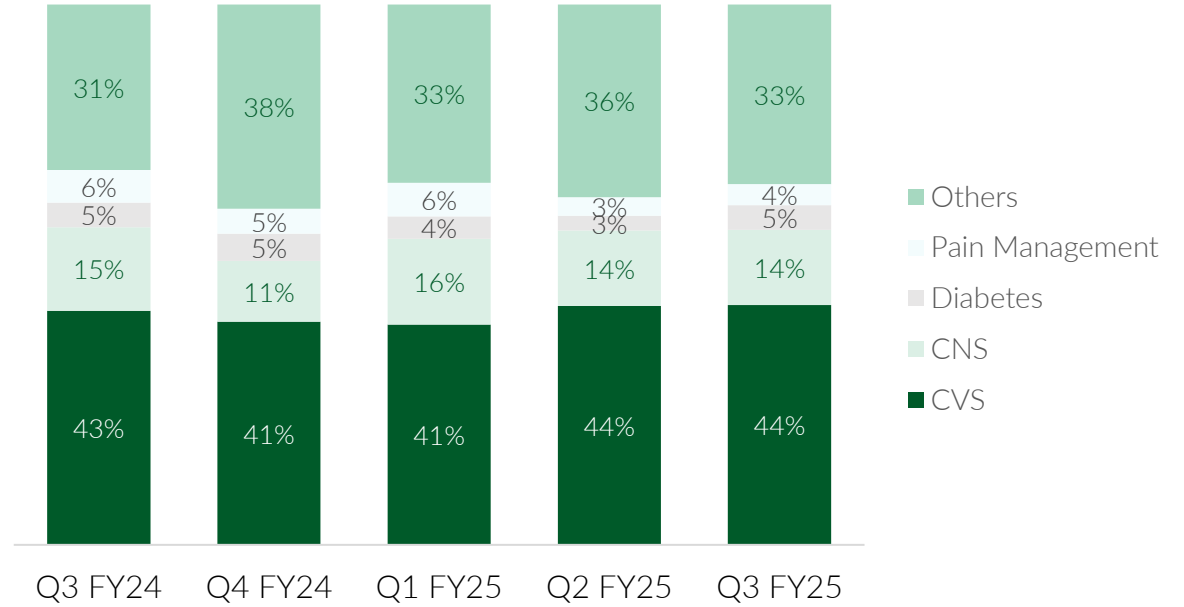
Market and Therapeutic Area Mix

Market Mix



- Regulated markets contributed 82% in Q3FY25
- The growth was primarily driven by Europe and Japan in the Regulated markets, whereas ROW and India led the growth in Emerging markets.

Therapeutic Area Mix



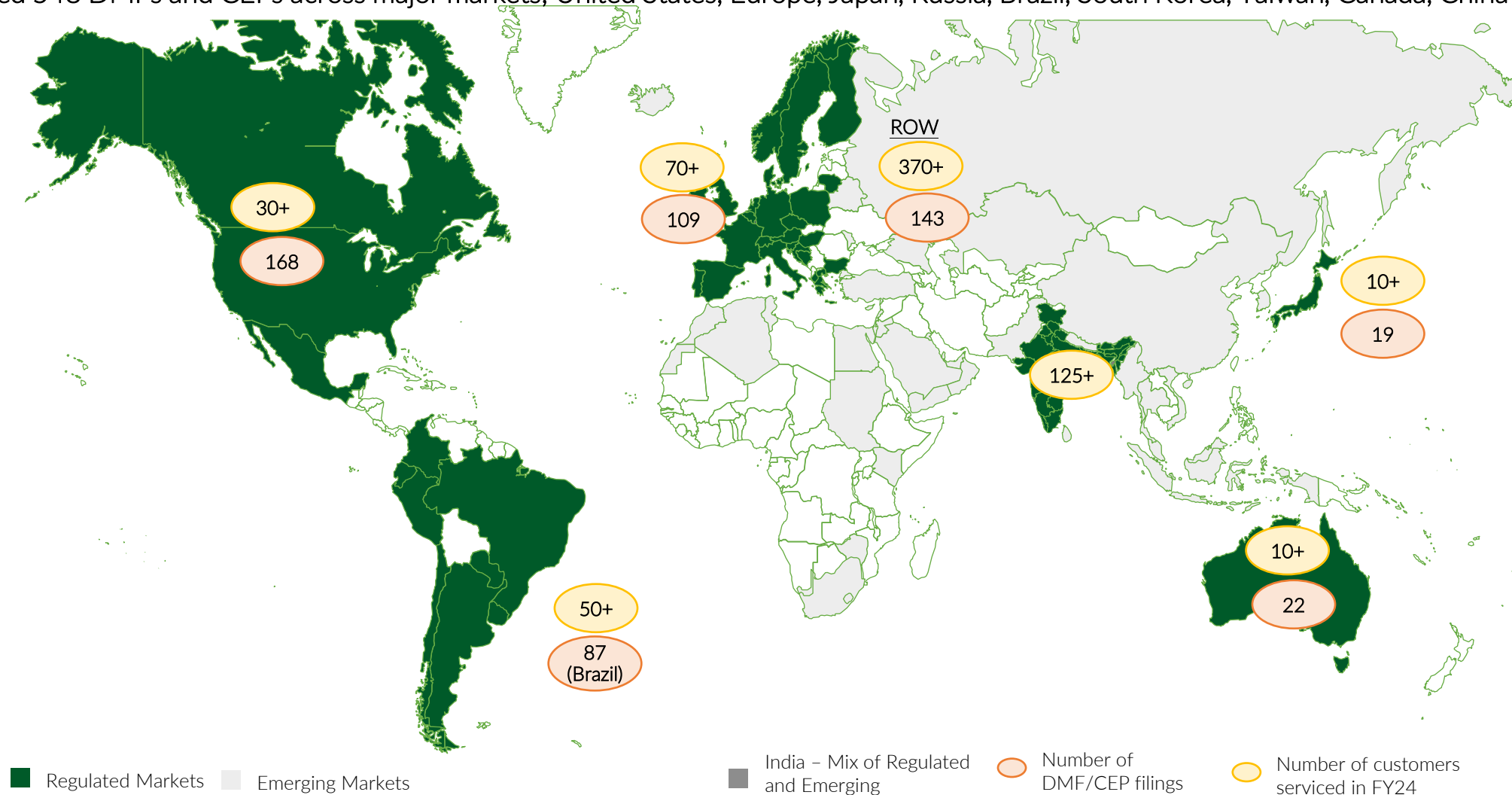
- CVS and CNS portfolio continued to deliver a strong growth
- Our key focused area of chronic therapies contributed 67% of the revenue in Q3FY25



Company Overview

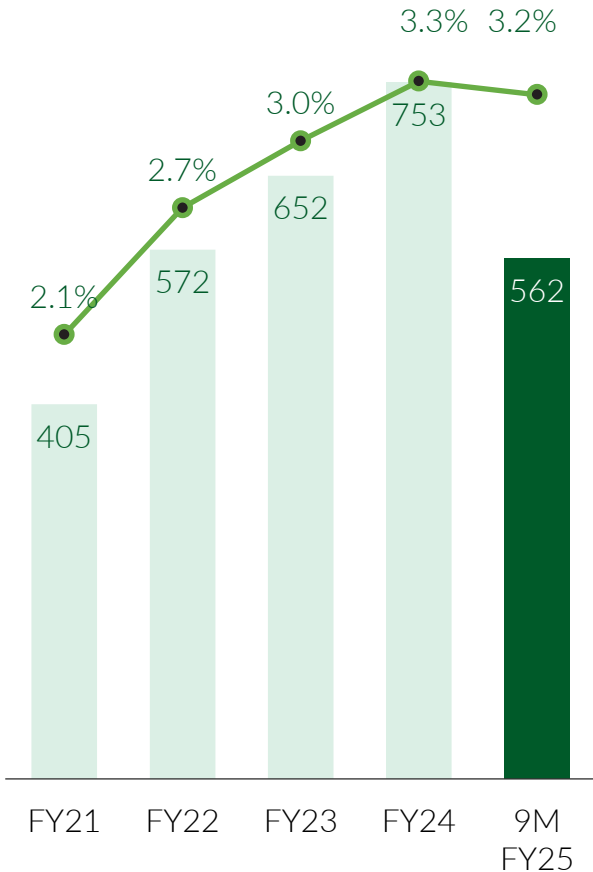
Global Footprint

- Filed 548 DMFs and CEPs across major markets; United States, Europe, Japan, Russia, Brazil, South Korea, Taiwan, Canada, China and Australia



As of December 31, 2024

R&D Spend (In ₹ Millions)



Cumulative Filing Status

Therapy	North America	Europe	Japan	Brazil	Australia	ROW	Total
CVS	38	37	4	21	10	38	148
CNS	39	25	8	16	2	20	110
Diabetes	10	5	-	9	-	14	38
Pain Management	1	2	-	4	1	9	17
Others	80	40	7	37	9	62	235
Total	168	109	19	87	22	143	548

- DMF/CEPs filing continue across major markets in Q3FY25, taking the total cumulative filings to 548 as on 31 December 2024.
- Two synthetic small molecules added to the development grid.
- The HP API portfolio remains on the development path with 21 products in the active grid representing market size of \$ 45 bn (Source: IQVIA, MAT Sep'24); six products are validated, six products are in advanced stages of development, remaining nine products progressing through earlier 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 \$2.2bn (Source: IQVIA, MAT Sep'24).



Quality Focused Manufacturing and R&D Infrastructure

Manufacturing Infrastructure

Location	Annual Installed Capacity	Last USFDA Inspection Date	Approvals
Ankleshwar, Gujarat	950.2 KL	July 2019	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

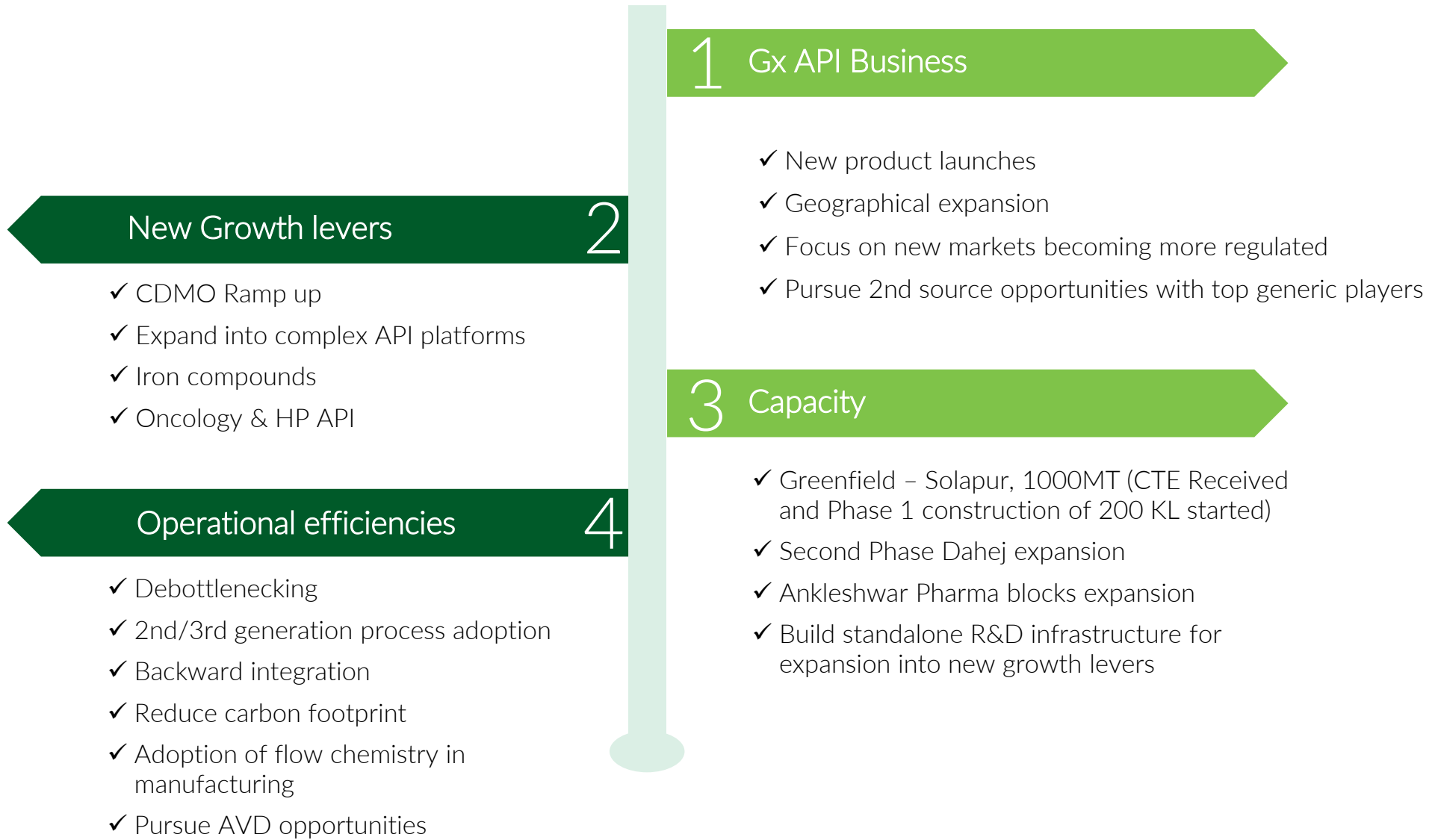
- Oncology R&D
- Cost improvement programs and process improvements



Strategy **Going Forward**



Strategy Growth Levers

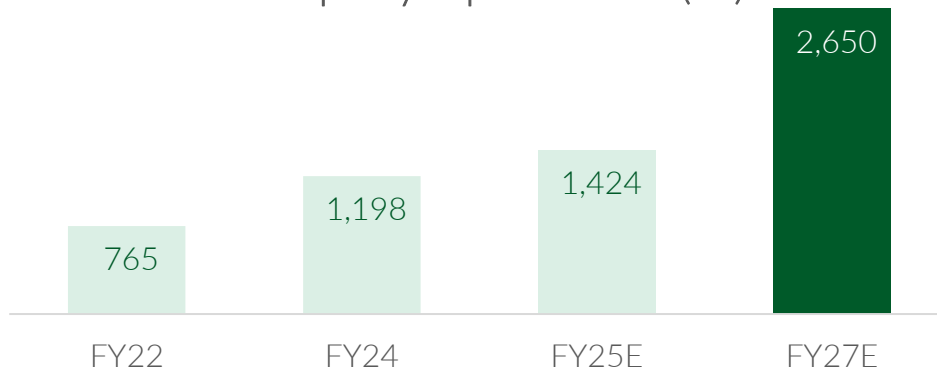


Future Capacity Expansion

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 started) Phase 1.1 – 400 KL (Backward Integration) Phase 2 - Planned addition of 400 KL	FY26 FY26 FY27/FY28

Total Reactor Capacity Expansion Plan (KL)

Capacity Progress by Year



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



Thank You

FOR FURTHER INFORMATION CONTACT

Email: complianceofficer@alivus.com

ERNST & YOUNG LLP – INVESTOR RELATIONS

DIWAKAR PINGLE

Email: Diwakar.Pingle@in.ey.com

RUNJHUN JAIN

Email: Runjhun.Jain1@in.ey.com

SNEHA SALIAN

Email: sneha2.salian@in.ey.com

CORPORATE OFFICE:

4th Floor, OIA House, 470,
Cardinal Gracious Road, Andheri
(E), Mumbai, 400 099, India.

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Plot No. 170-172, Chandramouli
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T: 91 22 68297979

CIN: L74900PN2011PLC139963

Website: www.alivus.com