

November 06, 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) Regulations 2015, we are enclosing herewith the Investor Presentation – Q2 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)

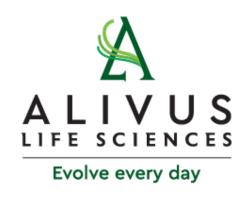
RUDALF Digitally signed by RUDALF JOSEPH CORRIEA Date: 2025.11.06 19:19:55 +05'30'

Rudalf Corriea

Company Secretary & Compliance Officer

Encl: As above

Alivus Life Sciences Limited (formerly Glenmark Life Sciences Limited)



Investor Presentation





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



Q2FY26 | Highlights





Dr. Yasir RawjeeManaging Director &
Chief Executive
Officer

"Our performance, this quarter, was fuelled by strong momentum in the non-GPL business, which delivered an impressive 39.7% YoY growth. Non-GPL business has shown a robust performance over last four quarters, driven by healthy demand across key geographies including ROW, LATAM, Japan, Europe, and India, all of which contributed meaningfully to the overall performance.

As we look forward, we remain confident of delivering a stronger performance in second half. This outlook is supported by continued robustness in non-GPL business and a better visibility on GPL business. Moreover, a continued ramp-up of our CDMO projects gives us confidence in the overall growth of our business.

We remain confident of maintaining margins at around 30%, despite the absence of PLI benefits. The margins are reinforced by a robust pipeline of new launches and operational efficiencies."

REVENUE (IN ₹ MILLION)	5,880	-2.3% QoQ	16.0% YoY
EBITDA (IN ₹ MILLION)	1,939	6.9% QoQ	35.7% Yo Y
PAT (IN ₹ MILLION)	1,301	7.1% QoQ	36.5% YoY

- Alivus registered a revenue from operations of ₹ 5,880 Mn for Q2FY26, a growth of 16% YoY.
- Gross Margin in Q2FY26 were at 57.7%, up by 260 bps QoQ & 210 bps YoY, driven by strong launches and product mix.
- EBITDA margins for the quarter were at 33%, up by 290 bps QoQ & 480 bps YoY.
- Generic API revenues in Q2FY26 grew by 15.3% YoY and CDMO revenues grew by 15.8% YoY.
- GPL business was at ₹ 1,437 Mn, a de-growth 5.8% on QoQ and 23.9% on YoY, resulting from inventory rationalization at the customer's end; Non-GPL business was at ₹ 4,443 Mn, up by 39.7% YoY.
- During Q2FY26, the company generated a strong free cash flow of ₹ 1,477 Mn, leading to Cash and Cash Equivalents (including short term investments) of ₹ 6,526 Mn as of 30th Sep 2025.

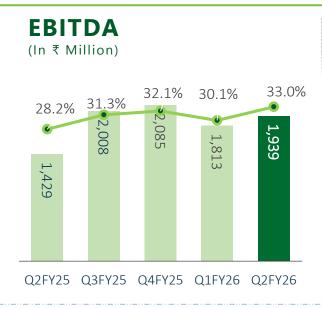
GPL: Glenmark Pharmaceuticals Ltd.

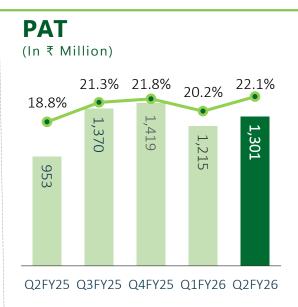


Q2 & H1 FY26 Performance



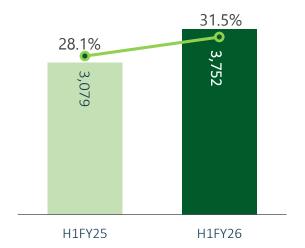


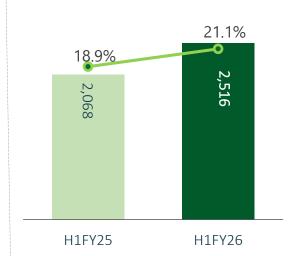
















P&L Highlights | Q2 & H1 FY26



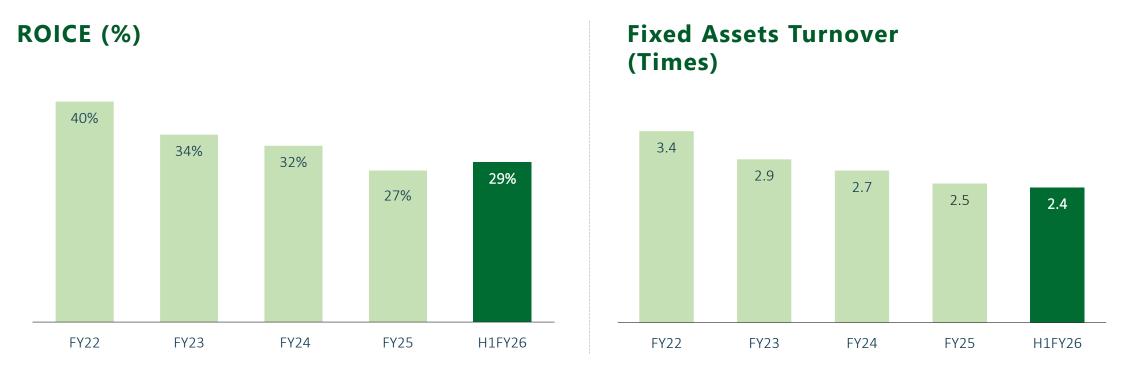
Particulars (In ₹ Million)	Q2 FY26	Q1 FY26	QoQ	Q2 FY25	YoY	H1 FY26	H1 FY25	YoY	FY25
Revenue from Operations	5,880	6,018	-2.3%	5,069	16.0%	11,898	10,955	8.6%	23,869
Gross Profit	3,391	3,315	2.3%	2,817	20.4%	6,706	5,825	15.1%	13,061
Gross Profit (%)	57.7%	55.1%	+260 bps	55.6%	+210 bps	56.4%	53.2%	+320 bps	54.7%
Other Income	147	90	63.3%	85	72.9%	237	140	69.3%	346
Employee Benefits Expense	664	616	7.8%	624	6.4%	1,280	1,192	7.4%	2,517
Other Expenses	935	976	-4.2%	849	10.1%	1,911	1,694	12.8%	3,718
EBITDA	1,939	1,813	6.9%	1,429	35.7%	3,752	3,079	21.9%	7,172
EBITDA Margin (%)	33.0%	30.1%	+290 bps	28.2%	+480 bps	31.5%	28.1%	+340 bps	30.0%
Depreciation and Amortisation Expense	184	171	7.6%	151	21.9%	355	295	20.3%	606
Finance Costs	13	12	8.3%	3	333.3%	25	7	257.1%	24
PBT	1,742	1,630	6.9%	1,275	36.6%	3,372	2,777	21.4%	6,542
PBT Margin (%)	29.6%	27.1%	+250 bps	25.2%	+440 bps	28.3%	25.3%	+300 bps	27.4%
PAT	1,301	1,215	7.1%	953	36.5%	2,516	2,068	21.7%	4,857
Net Margin (%)	22.1%	20.2%	+190 bps	18.8%	+330 bps	21.1%	18.9%	+220 bps	20.3%





Healthy Return Indicators





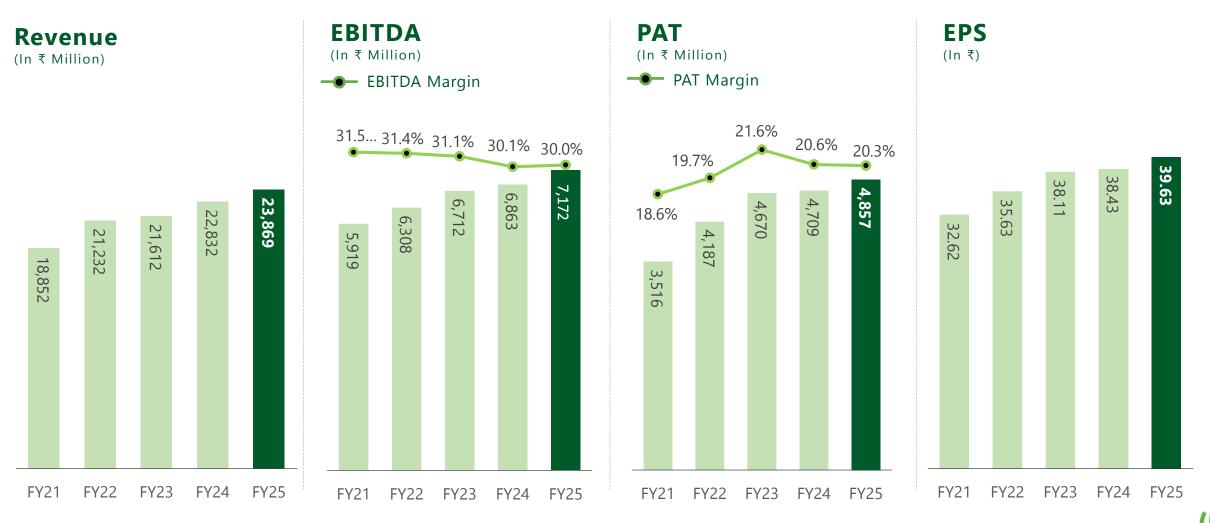
- **ROICE** is tracking at ~29% Capital employed driven by calibrated capex strategy and better margins
- **FATR** is **2.4 times** Asset turn trending slightly lower due to Capex cycle
- **Strong Balance Sheet** Strong free cash generation of ₹ 1,477 Mn leading to Cash and Cash Equivalents (including short term investments) of ₹ 6,526 Mn as of September 2025





Robust Growth & Steady Profitability







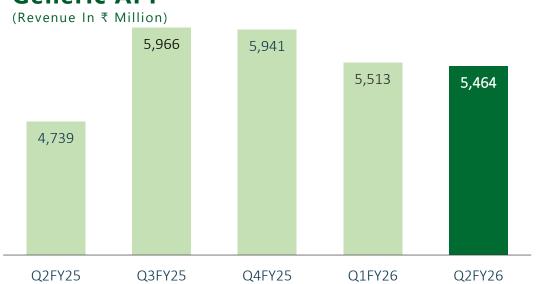
Business Performance Review

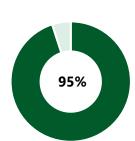


Segmental Performance | Generic API vs CDMO







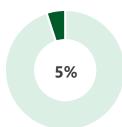


- Generic API revenues in Q2FY26 saw a growth of 15.3% YoY.
- Regions like ROW, LATAM, Japan, Europe and India (Ex-GPL) contributed to YoY revenue growth.

CDMO

(Revenue In ₹ Million)





- CDMO business saw a growth of 15.8% YoY.
 CDMO business is likely to witness a strong growth in H2FY26 led by addition of new project.
- Multiple discussions are ongoing for the CDMO pipelines.



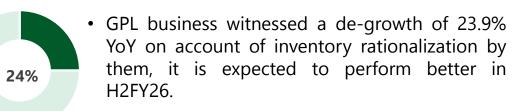


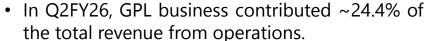
Segmental Performance | GPL vs Non-GPL

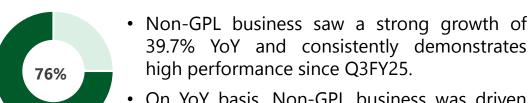


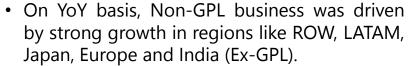










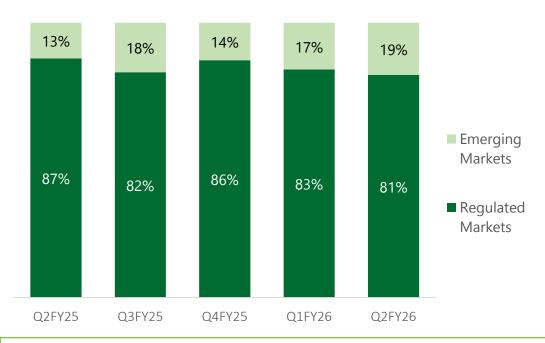




Market and Therapeutic Area Mix

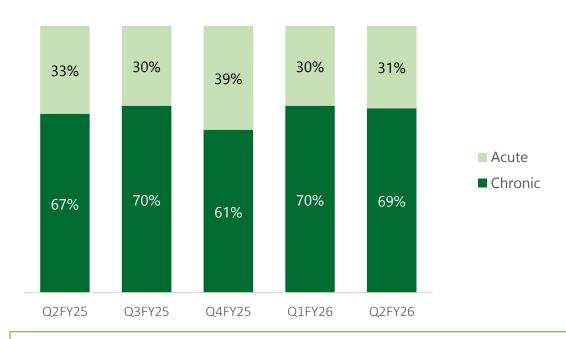


Market Mix



- Regulated markets contributed 81% in Q2FY26 driven by robust performance in Non-GPL business.
- The growth was primarily driven by LATAM, Japan and Europe in the Regulated markets, whereas ROW and India (Ex-GPL) led the growth in Emerging markets.

Therapeutic Mix



- Chronic portfolio continued to deliver a strong growth.
- The chronic therapies contributed 69% of the revenue in Q2FY26.



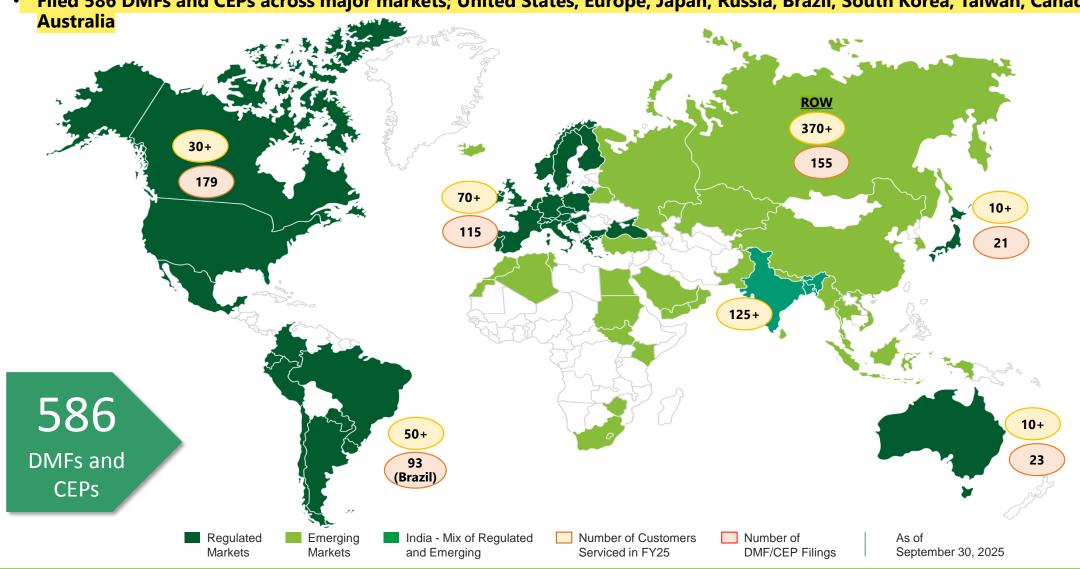
Company Overview



Global Footprint



Filed 586 DMFs and CEPs across major markets; United States, Europe, Japan, Russia, Brazil, South Korea, Taiwan, Canada, China and





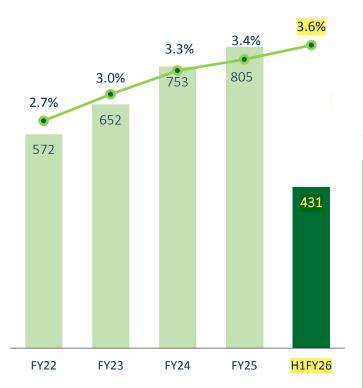
R&D Capabilities



R&D Spend

(In ₹ Million)





Cumulative Filing Status

Therapy	North America	Europe	Japan	Brazil	Australia	ROW	Total
CVS	39	40	4	23	11	41	158
CNS	43	26	8	18	3	21	119
Anti-Infective	21	11	3	3	3	14	55
Diabetes	10	5	-	9	-	16	40
Dermatology	9	6	1	13	1	11	41
Urology	10	7	2	6	1	11	37
Allergy	9	7	1	7	1	8	33
Others	38	13	2	14	3	33	103
Total	179	115	21	93	23	155	586

- DMF/CEP filings continue across major markets in Q2FY26, taking the total cumulative filings to 586 as on 30th September, 2025.
- Development grid remains steady and future ready with mix of near term launches, NCE-1 and patent cliff opportunities for target markets.
- The HP API portfolio remains on the development path with 26 products in the active grid representing market size of \$66 bn (Source: IQVIA, MAT Jun'25); 10 products are validated, Seven products are in advanced stages of development, remaining Nine products progressing through lab development stages.
- Development progressing for iron complexes in the grid. Filing completed for One iron complex, Two others are in advanced stages of development and preliminary development ongoing for One iron complex. Total addressable market of \$2.7 bn (Source: IQVIA, MAT June'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

4

- √ 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

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

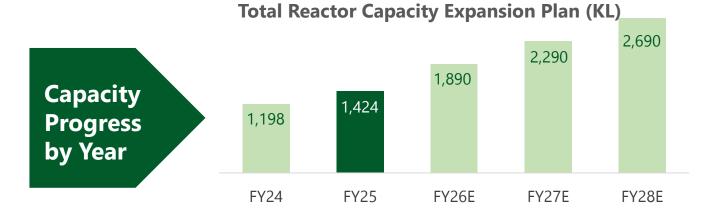
- ✓ Greenfield Solapur, 1000MT (CTE Received and Phase 1 construction of 200 KL capacity ongoing)
- √ Second Phase of Dahej expansion is on track
- ✓ Ankleshwar Pharma blocks expansion is in process
- ✓ Build standalone R&D infrastructure for expansion into new growth levers



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 Capacity	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



- ✓ 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



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