AKUMS DRUGS & PHARMACEUTICALS LTD.

Plot No. 131 to 133, Block-C, Mangolpuri Ind. Area, Phase-I, (Adjoining CBSE Office) Delhi - 110083 (INDIA).

⊕ www.akums.in

akumsho@akums.net

+91-11-27023256



\(+91-11-69041000

CIN: L24239DL2004PLC125888

Ref: Akums/Exchange/2025-26/34

August 08, 2025

To, The Listing Department National Stock Exchange of India Ltd Exchange Plaza, C-1, Block G, Bandra Kurla Complex, Bandra (E),Mumbai – 400 051 To, The Listing Department BSE Limited Rotunda Building, Phiroze Jeejeebhoy Towers, Dalal Street, Fort, Mumbai – 400 001

Symbol: AKUMS

Scrip Code: 544222

Sub: Disclosure under Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 - Investor Presentation

Respected Sir/Madam,

In compliance of Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, a copy of investor presentation is enclosed herewith.

This is for your kind information and record.

Thanking You

For Akums Drugs and Pharmaceuticals Limited

Dharamvir Malik Company Secretary & Compliance Officer

Encl: as above



Earnings Presentation

August 2025



Disclaimer



Akums Drugs and Pharmaceuticals Limited may, from time to time, make written and oral forward-looking statements, in addition to statements contained in the company's filings with BSE Limited (BSE) and National Stock Exchange of India Limited (NSE), and its reports to shareholders. The company does not undertake to update any forward-looking statements that may be made from time to time by or on behalf of the Akums Drugs and Pharmaceuticals Limited.

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Managing Directors' Message

Akums

Dear Investors

We are pleased to declare the Q1 results for FY 26. This date marks just over one year since we began our journey as a listed company. We continue to work towards strengthening the organization with a focus on long term growth.

The long-term growth prospects of the company remain intact with growth in both revenues and profitability. R&D remains the back bone of the company. We achieved the milestone of 1,000 DCGI approvals. During this quarter we received 27 DCGI approvals as against 31 approvals for the entire last year.

As part of Akums' strategic vision to establish itself as a leading global CDMO, we received our first EU dossier approval for Rivaroxaban. We also filed our first dossier of Dapagliflozin combination in Switzerland. Both of these new products hold significant market potential. The commercialization of the EU contract continues to be on track. We will commence commercial supplies from April 2027.

As informed during the last call, Akums had received 100 mn Euros as part consideration for the EU contract in this quarter, consequently our cash surplus now stands at INR 1,518 cr. The strong liquidity position provides a robust foundation for Akums to strategically scale up its business operations through both organic growth initiatives and inorganic opportunities.

The first quarter of FY26 continued to see tepid volume growth, with market volume growth below 0.5% and IPM growth being driven largely by price growth. The weakness in API pricing continues.

Looking ahead, we remain focused on strengthening our CDMO leadership, scaling high-value capabilities, and driving operational excellence. Backed by a strong pipeline and prudent capital allocation, we are well-positioned to deliver sustainable and profitable growth in the years ahead.

- Sanjeev Jain & Sandeep Jain



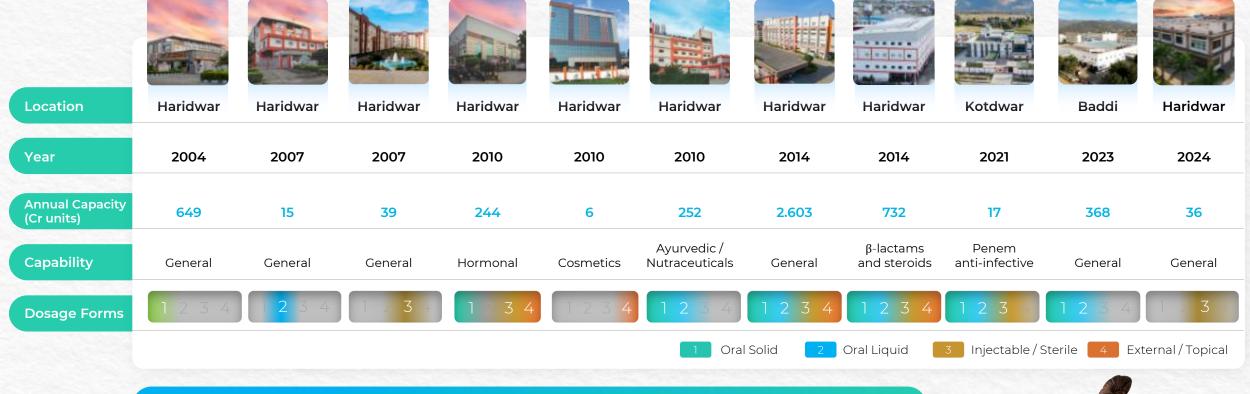
Two Decades' Legacy of Expertise, Experience & Trust





Technologically Advanced Manufacturing Capabilities





Key Accreditations*

























Illustrative

Differentiated Dosage Forms



Tablet In Tablet



Bi-layered, Sustained Release Tablet in Tablet



Inlay Tablet



Multiple Tablets in Capsule



Pre-filled syringes



Lyophilized Vials



Smart Tablets



Gummies



Mouth Melting Powders in Sachet



Tri Layered Tablet

Innovative In-house Technologies

















































Key Highlights



<u>1,518 Cr.</u> Cash Surplus (566 Cr in FY25)

1,000 DGCI Approvals (+27 in Q1FY26) € 100 Mn Received as advance against European CDMO contract

Zero Liquid Discharge (ZLD) implemented across 5 plants

Brazil ANVISA
approval for Plant 3
and Russia GMP
granted in Plant 4

20 New Approvals including 1st EU
Approval
(Rivaroxaban)

44 Dossiers filed – 1st submission in Switzerland, Iraq, and Jordan Strengthened senior leadership teams in IT, Finance and Strategy





Q1 FY 26

Financial Performance

Highlights

Consolidated Performance Highlights



Q1 FY26

1,051 cr

Total Income (+2.4% YoY)

156 cr 14.8% 65 cr 6.2% Adj. EBITDA (+208 bps YoY) Adj. PAT (+57 bps YoY)

Segment Break-up

	Business Vertical	Share in Q1 FY26 revenue
	CDMO	79.4%
	Domestic Branded Formulation	10.5%
(API)	API	4.4%
	International Branded Formulation	3.4%
	Trade Generics	2.3%

Consolidated Quarterly Performance Highlights





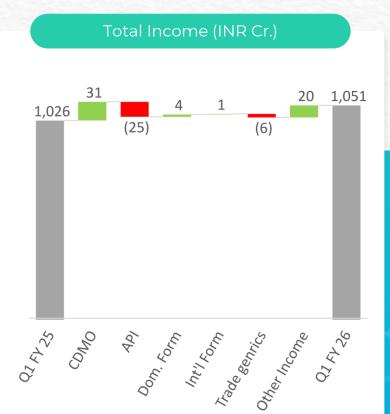




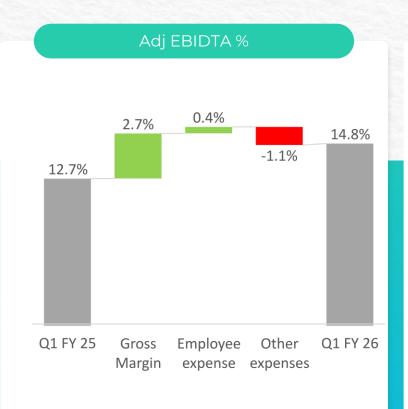
(1) Adjusted EBITDA = Profit before tax + fair value changes to financial instrument + finance cost + depreciation and amortization; (2) Adjusted PAT = PAT + Fair value changes to financial instrument - deferred tax created on brought forward losses (INR 106 cr) in Q4 FY 25)

Revenue and Margin Breakdown

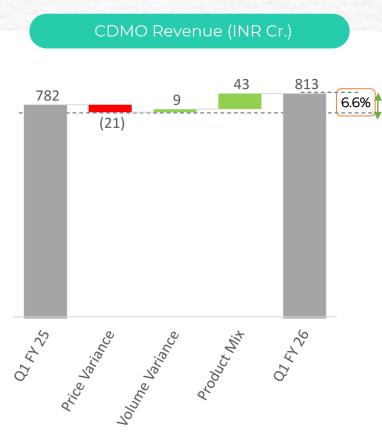




- CDMO growth slow on account of declining API prices and muted volume growth
- Trade Generics and API sales decline as focus was on to minimize losses



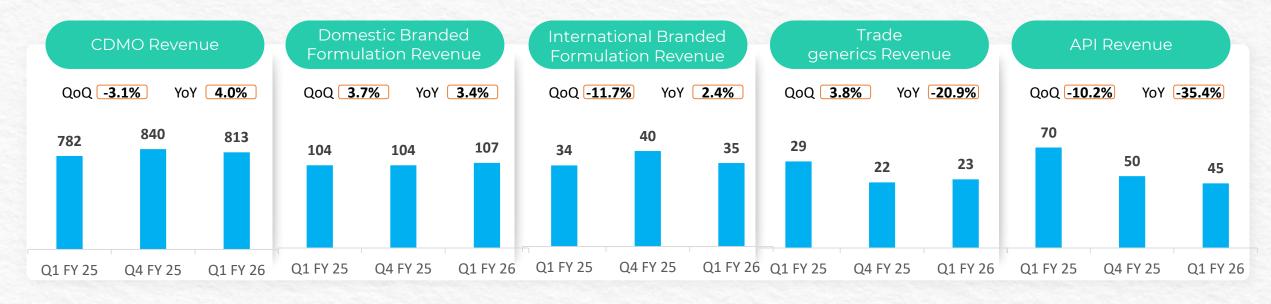
- Improved gross margins driven by continued focus on niche products.
- Other expenses driven by increased power and fuel costs and provisions.

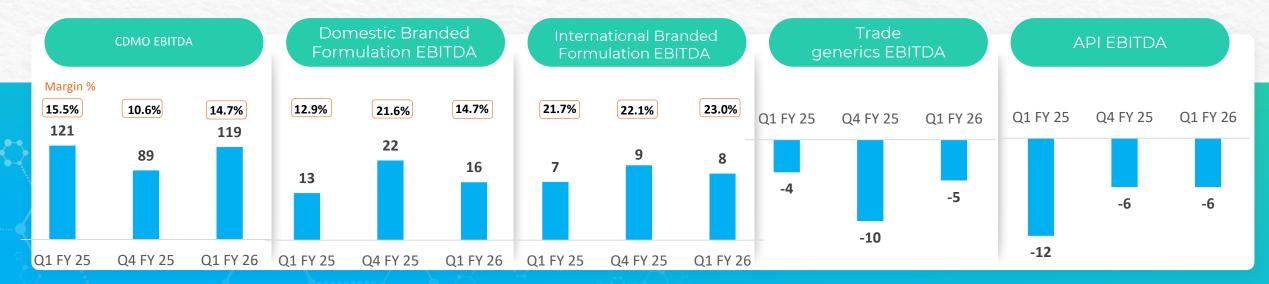


 Despite dip in API prices as well as muted volume growth, CDMO segment grew led by improved product mix

Segmental Quarterly Performance







Objects of the offer



Objects (INR Cr)	Amount as per offer document	Utilization as on June 30, 2025	Unutilized Amount
Repayment borrowings of Akums	159.91	159.91	-
Repayment of borrowings of our Subsidiaries	227.09	227.09	-
Funding incremental working capital requirements of our Company	55.00	55.00	-
Pursuing inorganic growth initiatives through acquisitions	27.87	27.87	-
General corporate purposes	167.50	167.50	-
Total	637.37	637.37	-



Thank You



For further information, contact:

IR Desk

Akums Drugs and Pharmaceuticals Ltd

Email: investors@akums.net/ankit.jain@akums.net

Siddharth Rangnekar / Shruti Joshi
CDR, India

Tel: +91 97699 19966 / 75065 67349

Email: siddharth@cdr-india.com / shruti@cdr-india.com