

CSD/BSE&NSE/BM/2024-25 February 12, 2025

To
The Manager
Department of Corporate Services
BSE Limited
25th Floor, P. J. Towers,
Dalal Street, Mumbai - 400 001

To
The Manager
Listing Department
National Stock Exchange of India Limited
Exchange Plaza, Bandra Kurla Complex
Bandra (E), Mumbai – 400 051

Scrip Code: 543064 Scrip Symbol: SUVENPHAR

Dear Sir/Madam,

**Sub: Outcome of the Board Meeting** 

Pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure requirements) Regulations, 2015 ("SEBI Listing Regulations"), we wish to inform you that the Board of Directors of the Company ("Board") at its meeting held today, i.e., on February 12, 2025, has, *inter alia*, approved the Unaudited Standalone and Consolidated Financial Results prepared under Ind AS for the quarter and nine months ended December 31, 2024, pursuant to Regulation 33 of the SEBI Listing Regulations. In this connection, we annexed herewith the following documents:

- a) Unaudited Standalone and Consolidated Financial Results under Ind AS for quarter and nine months ended December 31, 2024;
- b) Limited Review Reports on the above financial results;
- c) Press Release on the financial results; and
- d) Investor Presentation

We request you to take these documents in your records. The Board Meeting commenced at 03.25 pm IST and concluded at 04.15 pm IST.

This is for your information and record.

Yours faithfully, For **Suven Pharmaceuticals Limited** 

#### Kundan Kumar Jha

Company Secretary, Compliance Officer and Head-Legal

Encl: as above

#### Suven Pharmaceuticals Limited

**Registered Office:** # 215 Atrium, C Wing, 8th Floor, 819-821, Andheri Kurla Road, Chakala, Andheri East, Chakala Midc, Mumbai- 400093, Maharashtra, India Tel: 91 22 61539999

**Corporate Office:** # 202, A-Wing, Galaxy Towers, Plot No.1, Hyderabad Knowledge City, TSIIC, Raidurg, Hyderabad - 500081 Telangana, India Tel: 91 40 2354 9414 / 3311

Email: info@suvenpharm.com | Website: www.suvenpharm.com | CIN: L24299MH2018PLC422236



#### SUVEN PHARMACEUTICALS LIMITED

Regd. Off: 215 Atrium, C Wing, 8th Floor, 819-821, Andheri Kurla Road, Chakala, Andheri East, Chakala Midc, Mumbai, Mumbai, Maharashtra, India, 400093

STATEMENT OF UNAUDITED STANDALONE AND CONSOLIDATED FINANCIAL RESULTS FOR THE QUARTER AND NINE MONTHS ENDED 31
DECEMBER 2024

PART -				STAND	ALONE		
C1 . 11	DARTICIU ADC		Quarter ende	d	Nine mor	nths ended	For the yea
Sl. No.	PARTICULARS	31-Dec-24	30-Sep-24	31-Dec-23	31-Dec-24	31-Dec-23	31-Mar-24
		Unaudited	Unaudited	Unaudited	Unaudited	Unaudited	Audited
1	Income						
	Revenue from operations	275.39	236.06	212.82	733.54	781.47	1,024.99
	Other income	15.18	13.43	13.97	46.56	38.23	55.09
	Total income	290.57	249.49	226.79	780.10	819.70	1,080.08
2	Expenses						
	a) Cost of materials consumed	55.11	39.17	60.65	129.01	185.22	248.38
	b) Changes in inventories of finished goods and work-	(7.97)	6.84	6.39	20.84	36.86	53.88
	in-progress						
	c) Employee benefits expense	50.34	44.25	32.57	137.46	89.39	126.88
	d) Finance costs	2.60	1.47	2.15	5.67	5.14	7.43
	e) Depreciation and amortisation expense	13.44	12.29	11.38	37.73	33.13	48.79
	f) Other expenses	62.45	44.79	44.15	146.72	132.24	185.9
	Total expenses	175.97	148.81	157.29	477.43	481.98	671.3
3	Profit before tax (1-2)	114.60	100.68	69.50	302.67	337.72	408.77
4	Tax expenses						
	a) Current tax	35.22	27.91	13.85	83.15	79.45	98.14
	b) Current tax - earlier years	6.57		(0.78)	6.57	(0.78)	(0.78
	c) Deferred tax	(13.54)	(3.56)	4.07	(14.80)	7.66	6.59
5	Net Profit for the period/year (3-4)	86,35	76,33	52.36	227.75	251.39	304.82
6	Other comprehensive income/ (loss)						
6.a	(i) Items that will not be reclassified to profit or loss	(0.11)	(0.10)	(1.13)	(0.21)	(1.12)	(0.39
	(ii) Income tax relating to items that will not be	0.02	0.03	0.28	0.05	0.28	0.10
	reclassified to profit or loss						
6.b	(i) Items that will be reclassified to profit or loss	- 1		-			
	(ii) Income tax relating to items that will be	-	-			-	
	reclassified to profit or loss						
	Total other comprehensive loss	(0.09)	(0.07)	(0.85)	(0.16)	(0.84)	(0.29
7	Total comprehensive income for the period/year	86.26	76.26	51.51	227.59	250.55	304.53
8	Paid-up equity share capital	25.46	25.46	25,46	25.46	25,46	25.46
	Face Value of the Share	Re.1.00	Re.1.00	Re.1.00	Re.1.00	Re.1.00	Re.1.00
9	Other equity						2,030.44
	Earning Per Share (EPS)-Face value of Rs. 1/- each)						
1000000	a) Basic	3.39	3.00	2.06	8.95	9.88	11.97
	b) Diluted	3.37	2.98	2.06	8.91	9.88	11.97
		(not	(not	(not	(not	(not	(annualised
	·	annualised)	annualised)	annualised)	annualised)	annualised)	***************************************



SIGNED FOR IDENTIFICATION PURPOSES







#### SUVEN PHARMACEUTICALS LIMITED

Regd. Off: 215 Atrium, C Wing, 8th Floor, 819-821, Andheri Kurla Road, Chakala, Andheri East, Chakala Midc, Mumbai, Mumbai, Maharashtra, India, 400093

PART -		CONSOLIDATED					
SI. No.	PARTICULARS		Quarter ende	d	Nine mor	nths ended	For the yea ended
31. NO.	PARTICULARS	31-Dec-24	30-Sep-24	31-Dec-23	31-Dec-24	31-Dec-23	31-Mar-24
		Unaudited	Unaudited	Unaudited	Unaudited	Unaudited	Audited
1	Income						
	Revenue from operations	307.15	257.72	219.82	795.56	798.42	1,051.3
	Other income	15.71	14.12	14.34	47.99	44.85	61.9
	Total income	322.86	271.84	234.16	843.55	843.27	1,113.2
2	Expenses						
	a) Cost of materials consumed	67.04	47.08	63.52	152.42	194.45	265.8
	b) Changes in inventories of finished goods and work- in-progress	(11.41)	4.98	8.91	18.87	36.86	49.1
	c) Employee benefits expense	63.29	51.87	34.79	160.48	96.14	135.9
	d) Finance costs	3.33	1.67	2.16	6.60	5.15	7.4
	e) Depreciation and amortisation expense	20.35	16.96	12.79	50.71	37.35	54.6
	f) Other expenses	70.53	49.78	46.52	162.15	138.50	194.5
	Total expenses	213.13	172.34	168.69	551.23	508.45	707.5
3	Profit before tax and share of profit/(loss) of Associate (1-2)	109.73	99.50	65.47	292.32	334.82	405.6
4	Add: Share of profit/(Loss) of Associate	-	-	-		-	
5	Profit before tax (3+4)	109.73	99.50	65.47	292.32	334.82	405.6
6	Tax expenses						
	a) Current tax	35.03	27.91	15.43	82.96	81.03	99.5
	b) Current tax - earlier years	6.57	-	(0.78)	6.57	(0.78)	(0.7
	c) Deferred tax	(15.16)	(10.39)	4.07	(23.25)	7.66	6.5
7	Net Profit for the period/year (5-6)	83.29	81.98	46.75	226.04	246.91	300.2
	Net Profit for the period/year attributable to						
- 1	a) Shareholders of the company	82.88	82.21	46.75	225.86	246.91	300.2
	b) Non-controlling interest	0.41	(0.23)		0.18		
	Other comprehensive income		10.207				
		(0.00)	(0.10)	(1.12)	(0.19)	(1 12)	0.1
9.a	(i) Items that will not be reclassified to profit or loss	(0.09)	(0.10)	(1.12)	(0.19)	(1.12)	0.1
	(ii) Income tax relating to items that will not be reclassified to profit or loss	0.02	0.03	0.28	0.05	0.28	0.1
	(i) Items that will be reclassified to profit or loss	2.81	0.01		3.41		13.0
		2.01	0.01		3.77		13.0
	(ii) Income tax relating to items that will be reclassified to profit or loss	-	-	•	-	-	
	Total other comprehensive income/(loss)	2.74	(0.06)	(0.84)	3,27	(0.84)	13.2
	Total comprehensive income for the period/year (7+9)	86.03	81.92	45.91	229.31	246.07	313.4
11	Total comprehensive Income for the period/year attributable to						
	a) Shareholders of the company	85.62	82.15	45.91	229.13	246.07	313.4
	b) Non-controlling interest	0.41	(0.23)		0.18		
	Paid-up equity share capital	25.46	25.46	25.46	25.46	25.46	25,4
	Face Value of the Share	Re.1.00	Re.1.00	Re.1.00	Re.1.00	Re.1.00	Re.1.0
	Other equity Earning Per Share (EPS)- (Face value of Rs.1/-						2,025.2
	a) Basic -	3.26	3.23	1,84	8,87	9.70	11.8
	b) Diluted	3.23	3,21	1.84	8.83	9.70	11.8
		(not	(not	(not	(not	(not	(annualis
		annualised)	annualised)	annualised)	annualised)	annualised)	





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#### SUVEN PHARMACEUTICALS LIMITED

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#### Notes

- 1) The above results have been reviewed and recommended to the Board of Directors by the Audit Committee and subsequently approved by the Board of Directors at its meeting held on 12 February 2025. These results have been subjected to limited review by statutory auditors who have expressed an unmodified review conclusion. The financial results for the quarter and nine months ended 31 December 2023 and year ended 31 March 2024 were reviewed/audited by Karvy & Co., Chartered Accountants ('predecessor auditors').
- 2) The above financial results are prepared in accordance with the Indian Accounting Standard prescribed under section 133 of the Companies Act, 2013 and are in compliance with the presentation and disclosure requirements of Regulation 33 of the SEBI (listing Obligations and Disclosure Requirements) Regulations, 2015 (as amended).
- 3)The Company reportable activity falls under single operating segment i.e. Manufacturing (CRAMS) Bulk Drugs & Intermediates & Services, hence segment reporting as per Ind AS 108 (Operating Segment) is not presented.
- 4) Pursuant to definitive agreements entered by the company with Sapala Organics Private Limited ("Sapala"), the Company has acquired 51% of the share capital on a fully diluted basis (i.e., 67.5% of the present equity share capital) of Sapala on 12 July 2024 for a consideration of ₹258.00 crore and gained control of Sapala Organics Private Limited ("Sapala") as a subsidiary. As at 12 July 2024, the fair value of assets and liabilities acquired had been determined by the Group and accounted for in accordance with IND AS 103 - "Business Combination" based on provisional purchase price allocation.

Consolidated financial results for the quarter ended 30 September 2024, 31 December 2024 and nine months ended 31 December 2024, include the impact of the above transaction with effect from 12 July 2024 and are not comparable with previous corresponding periods.

5) Pursuant to definitive agreements entered by the Company with NJ Bio Inc ("NJ Bio"), the Company has acquired 56% of the share capital of NJ Bio Inc on 20 December 2024 for a consideration of ₹547.96 crore and gained control of NJ Bio Inc ("NJ Bio") as a subsidiary. As at 20 December 2024, the fair value of assets and liabilities acquired have been determined by the Group and accounted for in accordance with IND AS 103 "Business Combination".

As per the Share Purchase Agreement, NJ Bio has issued a put option to acquire the shares held by minority shareholders. The Put option obligation has been accounted for as a liability on the acquisition date at its fair value of ₹426.31 crore with a corresponding debit to other equity.

The fair value of assets and liabilities acquired have been provisionally determined by the Company and accounted for in accordance with IND AS 103 - "Business Combination". Consolidated financial results for the quarter and nine months ended 31 December 2024, include the impact of the above transaction with effect from 20 December 2024 and are not comparable with previous corresponding periods.

Total consideration has been allocated based on provisional purchase price allocation as under:

Particulars	Amount
Par ticulars	(₹ In crores)
Fair value of assets acquired including intangible assets	539.58
Fair value of liability assumed	(235.13)
Deferred tax liabilities on fair value of net assets acquired	(28.38)
Fair value of net assets acquired (A)	276.07
Non-controlled interest in the acquired entity, based on their proportionate interest in the recognised amounts of identifiable net assets of NJ Bio (B)	121.46
Total consideration paid (C)	547.96
Goodwill ((B+C)-A)	393.35

6) The Board of Directors had approved on 29 February 2024, the Scheme of Amalgamation of Cohance Life Sciences Limited (Transferor Company) into and with Suven Pharmaceuticals Limited ('The Company').

Based on the NCLT order dated 22 October 2024, meetings of the equity shareholders of both the Transferor Company and the Transferee Company were held on 28 November 2024 to consider and approve the Scheme. The Scheme has been approved by the Members of the Company with

The Scheme of Amalgamation remains subject to applicable approvals, including approvals from the Hon'ble NCLT, and such other approvals, permissions, and sanctions of regulatory and other authorities, as may be applicable.

7) The Board of directors of Suven Pharmaceuticals Limited ("Company" /"Transferee Company") has approved on 29 February 2024, the scheme of amalgamation of Casper Pharma Private Limited ("Transferor Company") (a wholly owned subsidiary of the Company) into and with the Company under the provisions of Sections 230 to 232 of the Companies Act, 2013 subject to receipt of applicable approval including approval from Hon'ble NCLT (" Scheme of Amalgamation").

The Hon'ble NCLT, Mumbai vide its Order dated 24 October 2024 has sanctioned the Scheme of Amalgamation. The Company has filed the certified copy of the Order with Registrar of Companies on 4 December 2024. As per the Scheme, the Appointed date which is also the effective date of the Scheme has been determined as 1 January 2025. Accordingly, the Scheme shall be accounted from the Appointed/ Effective date i.e. 1 January 2025 and in the manner prescribed under the scheme.

- 8) The figures of the previous year/periods have been regrouped/recast to render the classification comparable with that of current period.
- 9) The financial results for the quarter and nine months ended 31 December 2023 and for the year ended 31 March 2024 were presented in INR Lakhs. With effect from quarter ended 30 June 2024, the Company has presented the financial results in INR crores. Consequently, the results for Juen Pharmaceu the comparative periods have also been presented in INR Crores.



SIGNED FOR IDENTIFICATION **PURPOSES** 



For and on behalf of the Board Suven Pharmaceuticals Limited

Vivek Sharma **Executive Chairman** 

Place: Hyderabad Date: 12 February 2025

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Walker Chandiok & Co LLP 21st Floor, DLF Square Jacaranda Marg, DLF Phase II Gurugram – 122 002 India

T +91 124 4628099 F +91 124 4628001

Independent Auditor's Review Report on Standalone Unaudited Quarterly Financial Results and Year to Date Results of the Company pursuant to the Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 (as amended)

To the Board of Directors of Suven Pharmaceuticals Limited

- 1. We have reviewed the accompanying statement of standalone unaudited financial results ('the Statement') of Suven Pharmaceuticals Limited ('the Company') for the quarter ended 31 December 2024 and the year to date results for the period 01 April 2024 to 31 December 2024, being submitted by the Company pursuant to the requirements of Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 (as amended) ('Listing Regulations').
- 2. The Statement, which is the responsibility of the Company's management and approved by the Company's Board of Directors, has been prepared in accordance with the recognition and measurement principles laid down in Indian Accounting Standard 34, Interim Financial Reporting ('Ind AS 34'), prescribed under section 133 of the Companies Act, 2013 ('the Act'), and other accounting principles generally accepted in India and is in compliance with the presentation and disclosure requirements of Regulation 33 of the Listing Regulations. Our responsibility is to express a conclusion on the Statement based on our review.
- 3. We conducted our review of the Statement in accordance with the Standard on Review Engagements (SRE) 2410, Review of Interim Financial Information Performed by the Independent Auditor of the Entity, issued by the Institute of Chartered Accountants of India. A review of interim financial information consists of making inquiries, primarily of persons responsible for financial and accounting matters, and applying analytical and other review procedures. A review is substantially less in scope than an audit conducted in accordance with the Standards on Auditing specified under section 143(10) of the Act, and consequently, does not enable us to obtain assurance that we would become aware of all significant matters that might be identified in an audit. Accordingly, we do not express an audit opinion.
- 4. Based on our review conducted as above and the consideration of the review reports of the branch auditor referred to in paragraph 5 below, nothing has come to our attention that causes us to believe that the accompanying Statement, prepared in accordance with the recognition and measurement principles laid down in Ind AS 34, prescribed under section 133 of the Act, and other accounting principles generally accepted in India, has not disclosed the information required to be disclosed in accordance with the requirements of Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 (as amended), including the manner in which it is the disclosed, or that it contains any material misstatement.

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5. We did not review the interim financial information of one branch (Suven Pharmaceuticals Limited – Branch Office, USA) included in the Statement, where such interim financial information reflects total revenues of ₹Nil, total net loss after tax of ₹(12.44) crores and ₹(20.36) crores, total comprehensive loss of ₹(12.44) crores and ₹(20.36) crores, for the quarter and year-to-date period ended on 31 December 2024, respectively, as considered in the Statement. Such interim financial information has been reviewed by the branch auditor, whose reports have been furnished to us by the management, and our conclusion, in so far as it relates to the amounts and disclosures included in respect of this branch, is based solely on the review report of branch auditor.

Further, the above branch is located outside India whose interim financial information have been prepared in accordance with accounting principles generally accepted in their respective countries and which have been reviewed by branch auditor under Statement on Standards for Accounting and Review Services promulgated by the Accounting and Review Services Committee of the AICPA. The Company's management has converted the financial information of the branch from accounting principles generally accepted in its country to accounting principles generally accepted in India. We have reviewed these conversion adjustments made by the Company's management. Our conclusion, in so far as it relates to the balances and affairs of this branch is based on the review report of branch auditors and the conversion adjustments prepared by the management of the Company and reviewed by us.

Our conclusion on the Statement is not modified in respect of the above matter with respect to our reliance on the work done by and the reports of the branch auditors.

6. The review of standalone unaudited quarterly and year-to-date financial results for the period ended 31 December 2023 and audit of standalone financial results for the year ended 31 March 2024 included in the Statement was carried out and reported by M/s. Karvy & Co., Chartered Accountants who has expressed unmodified conclusion vide their review report dated 5 February 2024 and unmodified opinion vide their audit report dated 30 May 2024, respectively, whose reports have been furnished to us and which have been relied upon by us for the purpose of our review of the Statement. Our conclusion is not modified in respect of this matter.

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For Walker Chandiok & Co LLP

**Chartered Accountants** 

Firm Registration No: 001076N/N500013

Ashish Gupta

Partner

Membership No. 504662

UDIN: 25504662BMOODT1621

Place: New Delhi Date: 12 February 2025

Walker Chandiok & Co LLP 21st Floor, DLF Square Jacaranda Marg, DLF Phase II Gurugram – 122 002 India

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Independent Auditor's Review Report on Consolidated Unaudited Quarterly Financial Results and Year to Date Results of the Company pursuant to the Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 (as amended)

To the Board of Directors of Suven Pharmaceuticals Limited

- 1. We have reviewed the accompanying statement of unaudited consolidated financial results ('the Statement') of Suven Pharmaceuticals Limited ('the Holding Company') and its subsidiaries (the Holding Company and its subsidiaries together referred to as 'the Group') and its associate (refer Annexure 1 for the list of subsidiaries and associate included in the Statement) for the quarter ended 31 December 2024 and the consolidated year to date results for the period 01 April 2024 to 31 December 2024 being submitted by the Holding Company pursuant to the requirements of Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 (as amended) ('Listing Regulations').
- 2. This Statement, which is the responsibility of the Holding Company's management and approved by the Holding Company's Board of Directors, has been prepared in accordance with the recognition and measurement principles laid down in Indian Accounting Standard 34, Interim Financial Reporting ('Ind AS 34'), prescribed under section 133 of the Companies Act, 2013 ('the Act'), and other accounting principles generally accepted in India and is in compliance with the presentation and disclosure requirements of Regulation 33 of the Listing Regulations. Our responsibility is to express a conclusion on the Statement based on our review.
- 3. We conducted our review of the Statement in accordance with the Standard on Review Engagements (SRE) 2410, Review of Interim Financial Information Performed by the Independent Auditor of the Entity, issued by the Institute of Chartered Accountants of India. A review of interim financial information consists of making inquiries, primarily of persons responsible for financial and accounting matters, and applying analytical and other review procedures. A review is substantially less in scope than an audit conducted in accordance with the Standards on Auditing specified under section 143(10) of the Act, and consequently, does not enable us to obtain assurance that we would become aware of all significant matters that might be identified in an audit. Accordingly, we do not express an audit opinion.

We also performed procedures in accordance with the Circular issued by the SEBI under Regulation 33 (8) of the Listing Regulation, to the extent applicable.

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- 4. Based on our review conducted and procedures performed as stated in paragraph 3 above and upon consideration of the review reports of the branch auditors and other auditors referred to in paragraph 5 below, nothing has come to our attention that causes us to believe that the accompanying Statement, prepared in accordance with the recognition and measurement principles laid down in Ind AS 34, prescribed under section 133 of the Act, and other accounting principles generally accepted in India, has not disclosed the information required to be disclosed in accordance with the requirements of Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 (as amended), including the manner in which it is to be disclosed, or that it contains any material misstatement.
- 5. We did not review the interim financial information of one subsidiary included in the Statement and one branch (Suven USA branch) included in the unaudited interim standalone financial statements of the entities included in the Group, whose financial information reflects total revenues of ₹Nil crores and ₹Nil crores, total net loss after tax of ₹(13.12) crores and ₹(21.29) crores, total comprehensive loss of ₹(13.12) crores and ₹(21.29) crores, for the quarter and year-to-date period ended on 31 December 2024, respectively, as considered in the respective unaudited interim standalone financial information of the entities included in the Group. These interim financial information have been reviewed by other auditors and branch auditors whose review reports have been furnished to us by the management, and our conclusion in so far as it relates to the amounts and disclosures included in respect of the subsidiary and a branch is based solely on the review reports of such other auditors and branch auditors and the procedures performed by us as stated in paragraph 3 above.

Further, the above subsidiary and a branch are located outside India, whose interim financial information have been prepared in accordance with accounting principles generally accepted in their respective countries and which have been reviewed by other auditors and branch auditors under Statement on Standards for Accounting and Review Services promulgated by the Accounting and Review Services Committee of the AICPA. The Holding Company's management has converted the financial statements of such subsidiary/branch from accounting principles generally accepted in their respective countries to accounting principles generally accepted in India. We have reviewed these conversion adjustments made by the Holding Company's management. Our conclusion, in so far as it relates to the balances and affairs of the subsidiary/branch is based on the review report of other auditors and branch auditors and the conversion adjustments prepared by the management of the Holding Company and reviewed by us.

Our conclusion is not modified in respect of this matter with respect to our reliance on the work done by and the reports of the other auditors.

6. The review of unaudited consolidated quarterly and year-to-date financial results for the period ended 31 December 2023 and audit of standalone financial results for the year ended 31 March 2024 included in the Statement was carried out and reported by M/s. Karvy & Co., Chartered Accountants who have expressed unmodified conclusion vide their review report dated 5 February 2024 and unmodified opinion vide their audit report dated 30 May 2024, respectively, whose reports have been furnished to us and which have been relied upon by us for the purpose of our review of the Statement. Our conclusion is not modified in respect of this matter.

CHANDIOK

For Walker Chandiok & Co LLP

Chartered Accountants

Firm Registration No. 001076N/N500013

Ashish Gupta

Partner

Membership No. 504662

UDIN: 25504662BMOODU6892

Place: New Delhi

Date: 12 February 2025

#### Walker Chandiok & Co LLP

#### Annexure 1

#### List of entities included in the Statement

#### **Subsidiaries**

- 1. Casper Pharma Private Limited, India

- Caspel Fhama Frivate Limited, India
   Sapala Organics Private Limited, India (with effect from 11 July 2024)
   Suven Pharma Inc, USA
   NJ Bio Inc, USA (with effect from 20 December 2024)
   NJ Bio India Pharmaceuticals Private Limited, India (with effect from 20 December 2024)
   NJ Bio India Pharmaceuticals Private Limited, India (with effect from 20 December 2024)
- 6. NJ Biotherapeutics LLC, USA (with effect from 20 December 2024)

#### **Associates**

1. Aruka Bio Inc, USA (with effect from 20 December 2024)

#### **Branch Office**

1. Suven Pharmaceuticals Limited - Branch office, USA





#### **Suven Pharmaceuticals Announces Q3 and 9MFY25 Results**

Revenue Growth of 40% YoY in Q3 for proforma merged entity; Maintaining Growth Momentum for FY25

**Hyderabad, February 12, 2025** Suven Pharmaceuticals Ltd. (BSE: 530239, NSE: SUVENPHARM), a technology-led global Contract Development and Manufacturing Organization (CDMO), today announced its financial results for the third quarter and nine months ended December 31, 2024.

On a proforma merged basis, along with Cohance, Suven reported revenue growth of 40% YoY in-line with previously communicated expectations of growth in H2FY25. The key growth driver, the Pharma CDMO segment reported 101% YoY growth driven by R&D and BD efforts. Gross margins for the quarter stood at 71.5% and adjusted EBITDA margins stood at 38.7%.

#### Q3 and 9MFY25 Financial Performance Highlights

#### • Q3 Performance

#### Suven+Cohance (proforma basis)

 Revenue growth of 40% YoY to 6.76bn and Adjusted EBITDA growth of 85% YoY with margins of 38.7%

#### Suven

 Revenue Growth of 40% YoY to 3.07bn and Adjusted EBITDA growth of 70% YoY with margins of 44.9%

#### • 9mFY25 Performance

- Given the lumpy nature of the industry, the business performance reviewed on an annual basis provides better assessment. We continue to build a diversified business with multiple modalities to deliver steady growth across the year
- Suven+Cohance (proforma basis)

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 Revenue growth of 5% YoY to 17.7bn and Adjusted EBITDA growth of 3% YoY with margins of 34.8%

#### Suven

- Revenue of Rs 7.96bn and Adjusted EBITDA margins of 42.4%
- Capex for 9MFY25 was Rs 938mn. Generated free cash flow of Rs 1.33bn. and maintained a cash & bank balance of Rs 2.82bn

#### **Strategic and Operational Highlights:**

#### Pharma CDMO:

- Pipeline expansion: 2x YoY increase in RFQs for 9MFY25, reflecting continued strong demand across stages on back of secular macro tailwinds.
- Two molecules added this quarter: One molecule advanced to Phase 3 and one directly added to Phase III (laterals addition), increasing total active Phase 3 projects to 15 with 9 molecules.
- A previously highlighted positive readout on Phase III molecule has now met the primary endpoint for a second indication.
- Onboarded a new strategic customer one of the top five global pharma leaders for early-to-mid-phase projects.

#### • SpecChem CDMO:

 Witnessing recovery as expected, following the bottoming out in Q2.

#### **Executive Commentary:**

Mr. Vivek Sharma, Executive Chairman, commented: "Suven Pharma has reaffirmed its growth trajectory in Q3 FY25 as we had expected. We continue to make BD efforts and remain focused on high-value CDMO offerings and expansion in niche technology platforms in line with our strategic vision. Our investments in ADCs and oligonucleotides with fast underlying market growth position us well for sustained mid and long-term growth. As we gear

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to deliver our \$1bn revenue guidance with higher CDMO and niche technology share"

**Dr. V. Prasada Raju, Managing Director,** added: "We continue to witness a strong momentum in our Pharma CDMO business, with increased RFQ inflows and a robust late-stage pipeline. As our Phase 3 pipeline has expanded now to 15 projects with 9 molecules, gives us comfort on our strategic endeavors moving in the right direction. Our ability to drive innovation with addition of Oligos, deepening our ADC presence, and serve as a trusted partner to global innovators continues to enhance our competitive edge."

#### **Outlook:**

- Expect to deliver YoY growth for the combined platform in FY25 on a full year basis, with acceleration expected in FY26.
- Suven, being a technology-driven CDMO with a global footprint remains uniquely positioned to be one of the global leaders in the CDMO space from India which continues to witness secular tailwinds continue driven by supply chain de-risking priorities for innovators.
- Suven targets to reach \$1bn revenue, led by an expanding CDMO share, niche tech investments and continued strategic M&A.

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#### **Earnings call details**

Suven Pharmaceuticals Ltd will conduct a conference call to discuss its Q3 and 9MFY25 results performance. The management team will be represented by Mr. Vivek Sharma (Executive Chairman), Dr. V Prasada Raju (Managing Director), Dr. Sudhir Kumar Singh (Chief Executive Officer), Mr. Himanshu Agarwal (Chief Financial Officer) and Cyndrella Carvalho (Head IR).

The conference call will be initiated with a brief discussion after which the floor will be opened for Q&As. The financial results will be announced earlier on February 12, 2025.

In order to pre-register: Copy this URL in your browser:

https://services.choruscall.in/DiamondPassRegistration/register?confirmationNumber=2820130&linkSecurityString=134300f00e

#### Details of the conference call are as follows:

Timing	:	7.30 pm IST on Wednesday, February 12, 2025
Conference dial-in Primary number	:	+ 91 22 6280 1141/+91 22 7115 8042
Hong Kong Local Access Number	:	800 964 448
Singapore Local Access Number	:	800 101 2045
UK Local Access Number	:	0 808 101 1573
USA Local Access Number	:	1 866 746 2133

#### -ENDS-

#### For more information, please visit www. suvenpharm.com OR contact:

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Email: cyndrella.carvalho@suvenpharm.com Email: gavin@cdr-india.com

rishab@cdr-india.com

**Disclaimer.** Except for historical information, all of the statements, expectations and assumptions, including expectations and assumptions, contained in this presentation may be forward-looking statements that involve a number of risks and uncertainties. Although Suven attempts to be accurate in making these forward-looking statements, it is possible that future circumstances might differ from the assumptions on which such statements are based. Other important factors which could cause these statements to differ materially including outsourcing trends, economic conditions, dependence on collaborative partnership programs, retention of key personnel, technological advances and continued success in growth of sales that may make our products/services offerings less competitive; Suven may not undertake to update any forward-looking statements that may be made from time to time.

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## Suven Pharmaceuticals Ltd.

...Going Above and Beyond

**Investor Presentation** Q3 and 9MFY25



## Safe Harbour

Except for historical information, all of the statements, expectations and assumptions, including expectations and assumptions, contained in this presentation may be forward-looking statements that involve a number of risks and uncertainties. Although Suven attempts to be accurate in making these forward-looking statements, it is possible that future circumstances might differ from the assumptions on which such statements are based. Other important factors which could cause these statements to differ materially including outsourcing trends, economic conditions, dependence on collaborative partnership programs, retention of key personnel, technological advances and continued success in growth of sales that may make our products/services offerings less competitive; Suven may not undertake to update any forward-looking statements that may be made from time to time.



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## Q3\* and 9MFY25\* Proforma Operating and Financial Performance

\*Proforma basis subject to the proposed amalgamation of Cohance Lifesciences

### **Executive Summary**



#### Strategic Vision and Key Updates

- Suven, a technology-driven CDMO with a global footprint, is well-positioned to emerge as a global leader from India, leveraging structural tailwinds: advancing technological capabilities, China+1, EU+1, and the BioSecure Act.
- Targeting \$1bn in revenue, driven by: a) A diversified growth strategy built on three key pillars —Pharma CDMO, Specialty Chemicals CDMO, and APIs+, ensuring steady and predictable growth, b) Increasing mix of differentiated modalities, including ADCs, oligonucleotides, and other emerging technologies, to accelerate growth and enhance business defensibility c) A programmatic M&A approach to acquire differentiated assets d) A professionally managed organization with a strong leadership
- Over the past few quarters, we have prioritized team building, strengthening our R&D capabilities, built global commercial presence across the US, EU, & Japan, and streamlining backend operations. We are confident in the foundation we have built to drive long-term growth

#### Performance Highlights on proforma merged basis\*

- Growth trajectory reaffirmed with Q3 and 9M FY25, in line with our communicated expectations
  - o Q3: 40% YoY revenue growth, with adjusted EBITDA margins at 38.7%.
  - o 9M: 5% YoY revenue growth, with adjusted EBITDA margins at 34.8%.
  - o Given the lumpy nature of the Industry, business performance reviewed on an annual basis provides better assessment.
- Business Segment Performance (individual business details on following pages)
  - Pharma CDMO
    - Q3: 101% YoY revenue growth; 9M: 11% YoY growth.
    - Robust pipeline expansion with 2x YoY increase in RFQs (9M), including new customers, laterals and new product categories; expanded commercial team across US & Japan
    - 16 commercial molecules with large pharma; Phase III pipeline strengthened added two molecule: One molecule successfully advanced to Phase III, One new
      addition to Phase III brings the total to 15 Phase III projects across 9 molecules.
    - New customer onboarded: A top-five global pharma leader for early-to-mid-phase projects
    - We remain focused on expansion in ADC & Oligonucleotides with fast underlying market growth; leveraging front-end synergies.
      - Investments in cGMP facilities initiated to scale up both businesses
  - SpeChem CDMO
    - In line with our communicated expectation, segment bottomed out in Q2; sequential recovery seen in Q3. Strategic efforts yielding early results.
  - API+ & Others
    - Q3: 29% YoY growth; 9M: 17% YoY growth

Outlook: FY25E expected to grow YoY on a combined basis, with growth acceleration expected in FY26.

Merger Update: Shareholder approval received (99.99% in favor); Final NCLT hearing on February 18; Merger expected to be effective in Q1FY26, subject to regulatory approvals (incl DoP)

### Proforma Merged Earnings - Pharma CDMO drives growth



#### Proforma Suven + Cohance - Q3FY25 Performance:

- Combined platform reported revenue growth of 40% YoY driven by Pharma CDMO.
- The Pharma CDMO business grew by 101% YoY to Rs 2.9bn
- API+ segment revenue grew 29%YoY.
- Spec Chem segment lower by 22%YoY. Significant recovery on a sequential basis.
- Gross margins on combined basis at 71.5%.
- Adjusted EBITDA margins expanded by 960bps.

#### Other Key highlights

- 9M Free cash flow of Rs 3.2 bn
- 9M Cash on books at Rs 2.99 bn
- In 9M, we have spent Rs 2.31 bn on capex.
- We have recently been honored with the title "WORLD's BEST companies sustainable growth 2025" award by Times and Statista.
- Our facilities are audited for SA 8000:2014 certification, expect to receive the status soon.
- On ESG front, we have submitted SBTi commitment for all three types of emissions.

#### Q3FY25 Consolidated Financial Highlights

40%#

Revenue from operations (YoY)

**INR 6.8 bn** 

**Total Revenue** 

38.7%

EBITDA% excl. one time

56%#

Revenue Excl. Spec Chem(YoY)

INR 2.6bn\*

Adjusted EBITDA

24.8%

Adjusted PAT %

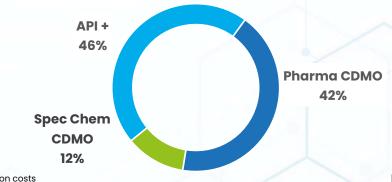
101%#

Pharma CDMO (YoY)

INR 1.7 bn\*

Adjusted Profit after Tax

#### Segmental Revenue Q3FY25 - CDMO share at 54%



<sup>\*</sup> Adjusted EBITDA and PAT includes one-time adjustments of INR 169Mn which comprises largely ESOP charges, merger and acquisition costs # Q3 includes consolidation of Sapala INR 113 mn and 12 days consolidation of NJ Bio INR 116Mn

Proforma basis subject to the proposed amalgamation of Cohance Lifesciences

## Q3FY25 Proforma Consolidated Financial results – Adjusted EBITDA margins at 39%



#### Suven + Cohance\* Consolidated Financials

#### **INR Million**

Particulars	Q3FY24	Q3FY25	YoY
Revenue from Operations	4,846	6,764	39.6%
Material costs / COGS	(1,718)	(1,925)	
Material Margin	3,128	4,839	54.7%
Material Margin %	64.6%	71.5%	
Manufacturing Expenses	(675)	(675)	
Employee Cost	(891)	(1,193)	
Other Expenses	(440)	(575)	
Total Expenses	2,007	2,443	
EBIDTA (Reported)	1,122	2,396	113.6%
EBIDTA (Reported) %	23.1%	35.4%	
FX MTM gain	32	55	
Onetime expenses	258	169	
EBIDTA (Adjusted)	1,412	2,620	85.5%
EBIDTA (Adjusted) %	29.1%	38.7%	
Depreciation & Amortization	(319)	(409)	
Finance costs	(104)	(108)	
Other income	148	139	
PBT (Adjusted)	1,137	2,242	97.1%
Tax(Adjusted)	(299)	(561)	
PAT (Adjusted)	838	1,681	100.5%
PAT Margin	17.3%	24.8%	
PAT (Reported)	811	1,660	104.8%
PAT Margin	16.7%	24.5%	

- Q3FY25 reported 40% YoY growth supported by Pharma CDMO and API+ revenue growth.
- Gross margins expanded 699 bps YoY to 71.5%, supported by a favourable business mix and robust growth in Pharma CDMO and API+.
- Adjusted EBITDA margins at 38.7%, reflecting the strength of R&D and BD collaboration in accelerating growth and focus on operational efficiencies along with business mix.
- Strategic talent investments continue to drive sustainable growth and long-term value creation.
   These costs have a lead effect and are yet to fully contribute to earnings.

### 9MFY25 Proforma Consolidated Financial results



#### **Suven + Cohance Consolidated Financials**

#### **INR Million**

Particulars	9MFY24	9MFY25	YoY
Revenue from Operations	16,903	17,691	4.7%
Material costs / COGS	(5,667)	(5,279)	
Material Margin	11,236	12,412	10.5%
Material Margin %	66.5%	70.2%	
Manufacturing Expenses	(1,980)	(1,847)	
Employee Cost	(2,779)	(3,310)	
Other Expenses	(1,189)	(1,552)	
Total Expenses	5,948	6,709	
EBIDTA (Reported)	5,288	5,702	7.8%
EBIDTA (Reported) %	31.3%	32.2%	
FX MTM gain	98	118	
Onetime expenses#	594	329	
EBIDTA (Adjusted)	5,979	6,149	2.8%
EBIDTA (Adjusted) %	35.4%	34.8%	
Depreciation & Amortization	(844)	(1,044)	
Finance costs	(276)	(306)	
Other income	561	430	
PBT (Adjusted)	5,420	5,229	(3.5)%
Tax(Adjusted)	(1,401)	(1,251)	
PAT (Adjusted)	4,019	3,978	(1.0)%
PAT Margin	23.8%	22.5%	
PAT (Reported)	3,953	3,916	(0.9)%
PAT Margin	23.4%	22.1%	

Note: # Adjusted EBITDA includes One-time adjustment for ESOP, Merger and acquisition costs Rs.594 mn & 329 mn for 9MFY24 & 9MFY25 respectively.

- Gross margins expanded by 365 bps to 70.2%.
- Adjusted EBITDA margins stood at 34.8% with Adjusted EBITDA of Rs 6.15bn.

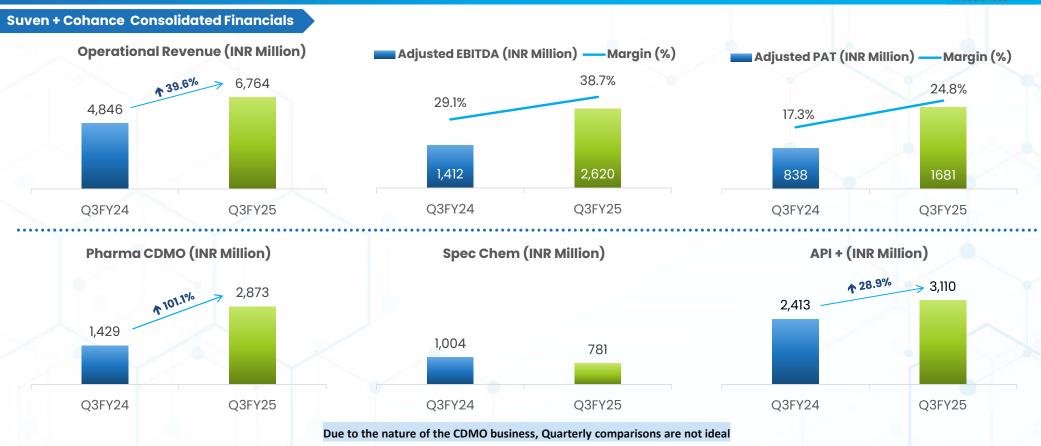
**INR Million** 

Balance Sheet Highl	lights
As on 31st December 2024	
Shareholders' funds	28,519
NCI- Sapala + NJ Bio	1,720
Net Fixed assets	28,498
Other net assets 1	(1,486)
Net cash/(debt) <sup>2</sup>	137
Total Use of Funds	3,090

1) Other assets calculated as Inventories + Trade receivables + Non-current investments + Current tax assets + Other assets less Trade payables + deferred tax liabilities + Other liabilities at the end of the year. 2) Net cash/(debt) calculated as the Cash & cash equivalents (Cash and bank balances + current Investments) less Total debt (Short-term and Long-term borrowings) at the end of the period.

## Q3FY25 Proforma Business performance overview combined platform





Note: 1) Adjusted EBITDA includes One-time adjustment for ESOP, Merger and acquisition costs of Rs. 258 mn and Rs.169 Mn respectively for Q3FY24 and Q3FY25.

2) Segment revenue 's are Restated.

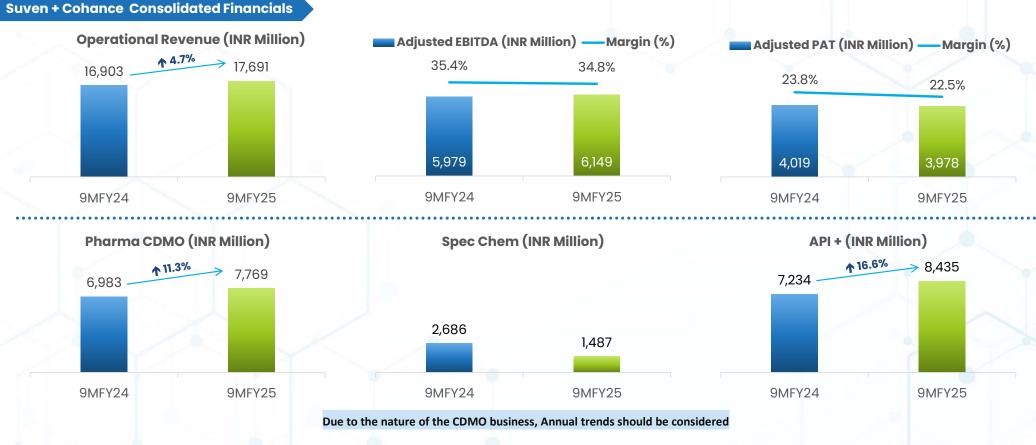
3) Q3 includes consolidation of Sapala INR 113 mn and 12 days consolidation of NJ Bio INR 116Mn

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## 9MFY25 Proforma Business performance overview







Note: 1) Adjusted EBITDA includes One-time adjustment for ESOP, Merger and acquisition costs Rs.594 mn & 329 mn for 9MFY24 & 9MFY25 respectively.

2) Segment revenues are 'restated. 3) 9M FY25 includes consolidation of Sapala INR 113 mn and 12 days consolidation of NJ Bio INR 116Mn

### Q3FY25 Suven Pharma - Business performance overview





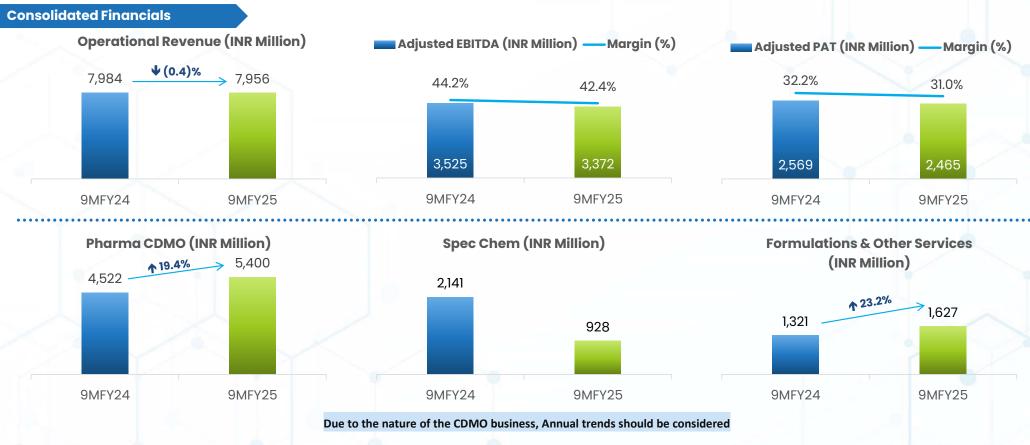
Note: 1) Adjusted EBITDA includes one-time adjustments of INR 163Mn in Q3FY25, comprising largely an ESOP charge

2) Segment revenue 's are Restated.

3) Q3 includes consolidation of sapala INR 113 mn and 12 days consolidation of NJ Bio INR 116Mn

### 9MFY25 Suven Pharma - Business performance overview





Note: 1) Adjusted EBITDA includes one-time adjustments of INR 274Mn in 9MFY25, comprising largely an ESOP charge

2) Segment revenue 's are Restated.

3) 9m iFY25 ncludes consolidation of Sapala INR 113 mn and 12 days consolidation of NJ Bio INR 116Mn







# Combined Business: Proforma Metrics



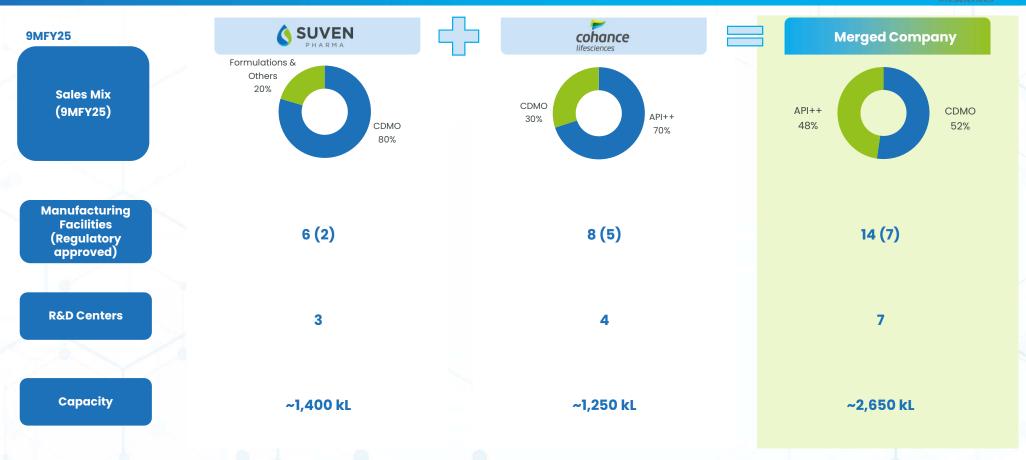
## **Proforma Merged Entity**

9MFY25 INR Mn	SUVEN	cohance	Merged Company
Revenue	7,956	9,735	17,691
Adjusted EBITDA	3,372	2,777	6,141
Adjusted EBITDA margin %	42.4%	28.5%	34.8%
Adjusted PAT	2,465	1,513	3,978
Adjusted PAT margin %	31.0%	15.5%	22.5%
RoCE	24.9%	26.8%	25.8%
RoE	13.4%	23.0%	13.4%
(Net Debt) / Net Cash to Adj. EBITDAx	0.5X	(0.4)X	0.0X

Note: 1) Adjusted EBITDA includes One-time adjustment for ESOP, Merger and acquisition costs Rs.594 mn and Rs.329 mn for 9MFY24 & 9MFY25 respectively. Source: Cohance LifeSciences Website published Investor Presentation

### SUVEN PHARMA Cohance

## Proforma Merged Entity - Combined business mix



Source: Cohance LifeSciences Website published Investor Presentation





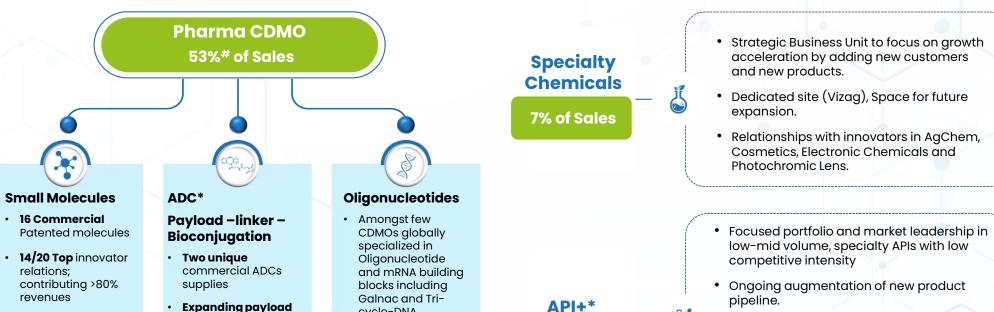


## Combined Business: Key segment wise strategy

### Our Growth Engines – Pharma CDMO key driver



Small Molecules, ADCs, Oligonucleotides and Peptides constitute ~52% of New Drug Additions to the Global Preclinical and Clinical Pipeline in 2024<sup>1</sup>



40% of Sales

7 molecules in Phase-3 translating into 12 intermediates; RFQs growing 2.2x

- **Expanding payload** and products Portfolio and Clinical Collaborations added 3 customers and new products
- **Drug Discovery to** commercial full chain exposure

# % sales - 9M Revenue Dec'24 Suven and Cohance combined + Sapala proforma FY24 + CY24 NJ Bio proforma

Galnac and Tricyclo-DNA

pipeline.

- Built deep cost position through backward integration.
- Top 3 player in 8 out of 10 top molecules in the API portfolio.
- · Offering end to end vertically integrated solutions including pellets and formulations.

Source: 1 Citeline Pharmprojects, \*Proforma basis subject to the proposed amalgamation of Cohance Lifesciences

### Pharma CDMO - Small molecules



#### • Phase III pipeline moving with higher conversions

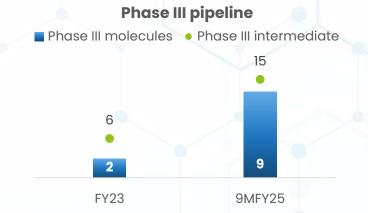
- o Active pipeline of 100+ projects spanning Phase I to Phase III.
- o We have 16 commercial Pharma molecules across combined platform

#### • Phase III pipeline

- Our Phase III pipeline has strengthened further now comprising 9 molecules with 15 intermediates.
- The two recent additions: one molecule has successfully advanced to phase III, while another has been directly added to Phase III.
- As previously highlighted, positive readout on a Phase III molecule has now met the primary endpoint for a second indication.

## • Highest streak of RFQs inflow persists; Higher mix of laterals, RFQs from new customers and category expansion.

- 9M RFQs 2x or doubled year-on-year.
- Product mix: Contribution from Late-Stage and Mid-Stage RFQs continues to grow, strengthening our position as a strategic partner for developments of laterals.
- Product type mix: incremental contribution continue to increase from niche technology projects like ADCs, Peptides, Oligonucleotide Fragments.
- Customer mix: new RFQs received from select Biotech companies; Increasing share of new customers, aligning with our strategic focus on R&D efforts and expanding our customer base, progressing up the value chain(from intermediates to APIs).

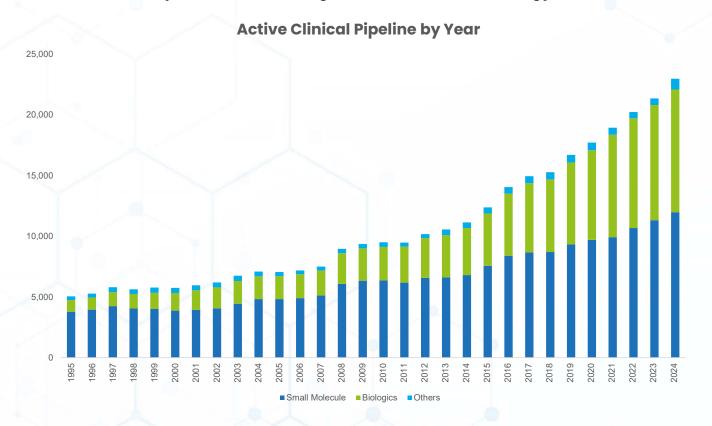




## Favorable Industry Macros leading to growth in Small Molecule Pipeline



#### Small Molecule Pipeline continues to grow on the back of Oncology contributes more than >50%



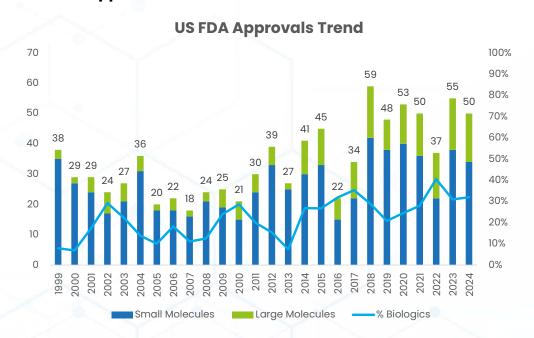
- The active clinical trial pipeline keeps growing and has reached 22,936 active drugs.
- Small Molecules is 52% of the current pipeline, Biologics 43% and Others (including Natural Substances) is 5%
- By far the largest chunk of drugs fall into the oncology bucket.

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## Small molecules: Increasing proportion of US FDA approvals



#### US FDA Approvals lean towards small molecules



**Record Number of Approvals in 2024:** 50 novel drugs were approved in 2024.

Other notable statistics include:

- Cancer remains the dominant focus of drug developers, with 30% novel approvals in 2024.
- Biologic approvals stay constant at 30-35% per year
- 52% received Orphan Drug Designation for treating rare diseases.
- 36% were designated Breakthrough designation.
- 56% received priority review, a regulatory designation for therapies that the FDA expects to offer 'significant improvements over the standard of care.

The positive trend continues in 2024: In 2024, 50 novel drugs were approved by FDA, of which 34 were small molecules (68%), which includes two Oligos, one Peptide and one Radiopharma

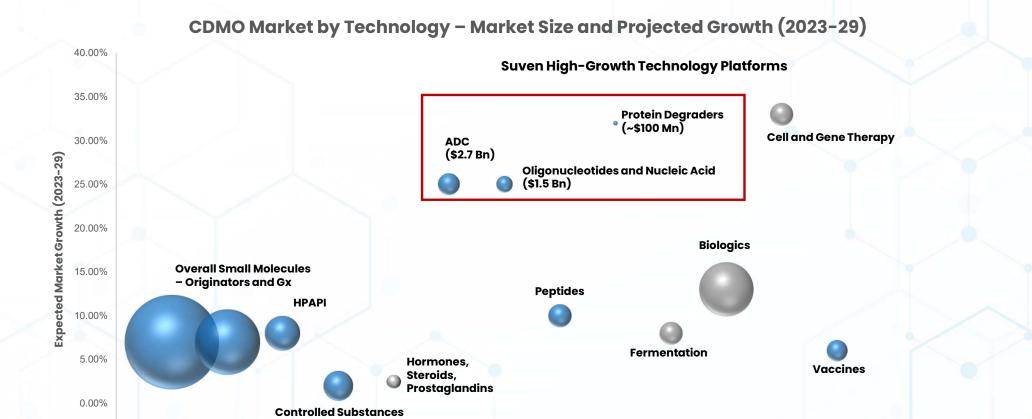
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# Suven is present in the fast growing tech platforms of ADCs & Oligos

-5.00%

Source: Industry data



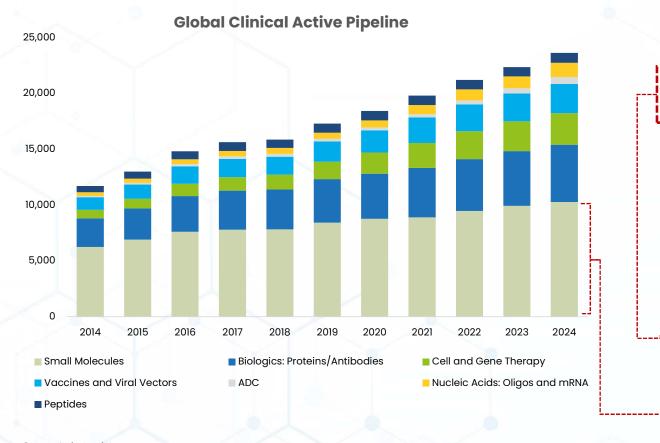


Suven + Cohance Platform presence

## Significant R&D investments in ADC and Oligos/mRNA in the clinical pipeline







R&D pipeline growth (CAGR)	2019-24	
Peptides	2%	
 Nucleic Acids (Oligos/mRNA)	19%	
Antibody-Drug Conjugates (ADC)	20%	
Vaccines and Viral Vectors	8%	
Cell and Gene Therapy	12%	
Biologics- Proteins/Antibodies	6%	
Small Molecules - General	4%	
Overall Clinical Pipeline	6%	

#### Pharmaceutical Drug R&D Trends

Surging Interest in Targeted Therapies and Genetic Treatment leading to uptake in ADCs, Nucleic Acids and Cell/Gene Therapies

Presence in small molecules, contributing >50% of total addressable R&D pipeline (incl. Oligos, ADCs)

Source: Industry data

# Suven uniquely placed to achieve leadership position in the fast-growing ADC/XDC segment



Expanded ADC offerings to become an integrated End to End CRDMO post acquisition of NJ Bio

#### Our unique capabilities in ADCs and XDCs

End-to-end CRDMO Partner from Drug Discovery to Commercialization

**Extensive Library of Payload-Linkers for Discovery based on biology** of the ADC target from a library of **500+** Payload-Linkers

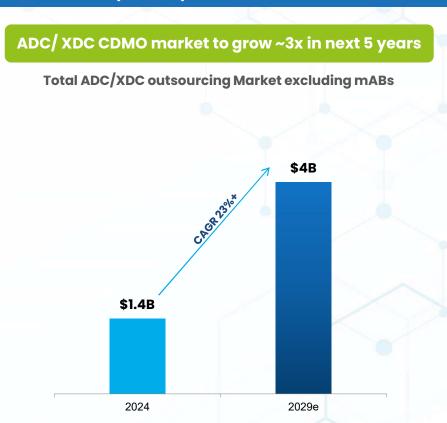
**Integrated Service Offerings:** across variety of standard and custom Payloads, Linkers, Analytical and Bioconjugation

**Global leadership in Camptothecin payloads**; supplying to **2 commercial ADCs**; leadership in **S-Trione** - a key intermediate in camptothecin derivatives

**Uniquely positioned as a Pureplay Payload Supplier:** covering **75%** of Payload market

Unique breadth of **XDC** and different payload capabilities – **Oligonucleotides**, **Radionuclides**, **Protein Degraders** 

Capacity augmentation in US & India; Portfolio expansion in new payloads and linker



Source: Industry data

# Oligonucleotides is the emerging modality





# Amongst the few CDMOs globally, supplying complex building blocks for Oligonucleotides

## Our Niche in Oligonucleotide segment

Capable of synthesizing a **spectrum of modified amidites and nucleosides** with excellent purity with high level of backward integration (15+ steps)

**Diversified innovator customer** (CDMO and Diagnostic) **base** with a strong Japan presence

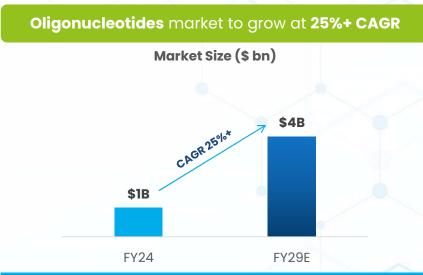
Only supplier of Tricyclo-DNA Amidites in the world

**Multi-kilo scale synthesis of wide variety of GalNAc compounds** supplied to Innovators with highest purity profile.

**Mastered the chemistry of conformationally constrained nucleic acids** and supply to innovators

**Capacity augmentation**: Investing in a **cGMP facility** to enhance capacity and drive R&D growth

Forward integrating to oligonucleotide drug substance manufacturing



Amidite and Galnac segments to grow significantly faster than oligonucleotides market itself

Nucleic acids & oligos vital for R&D in therapeutics, diagnostics, and synthetic biology.

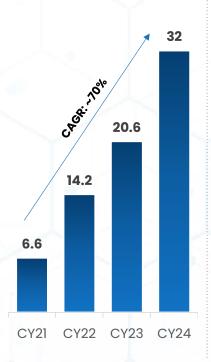
- Market Growth: Moving from rare diseases to high prevalence chronic indications. Rising use in molecular diagnostics and clinical applications.
- Increased Investments: Pharma and Biotech driving expansion

Source: : Industry data

# Acquisition of NJ Bio gives Suven End to end ADC/XDC capabilities; makes Suven Platform a major ADC player



# **Revenue Growth** Trajectory2 (US\$mn)



### **Business and service offerings**

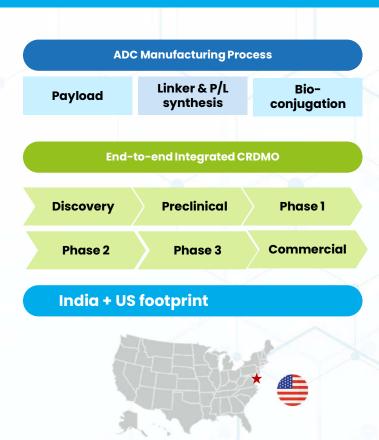
- End-to-end ADC chemistry capabilities (P-L1 synthesis, bioconjugation, bioanalytical services). Also has capabilities in the broader XDC segment (radio conjugates, oligo conjugates, peptide conjugates etc) and mRNA
- · Initially started with focus on preclinical services including proof-ofconcept, process development, scale up, IND batches. Delivered 500+ projects so far
- · Developed an extensive library of payload-linkers and offers 'Express Conjugation' service that allows to establish proof of concept for a novel ADC. Additional investments being made towards R&D for novel payload linkers and new technologies (e.g. protein degraders)
- Recently forayed into GMP Ph1/Ph2 P-L manufacturing. Plans to further expand of GMP Ph1/Ph2 capacities (including bioconjugation) in the near

### **Manufacturing and Operations**

- Headquartered in Princeton, NJ, USA (a key ADC innovation hub). 80,000 sq ft of lab space and GMP suites in Princeton with ~100 employees (including 80+ scientists).
- Also has India operations: 6,500 sq ft space in Mumbai; ~40 employees involved in creating payload-linker library and R&D innovation work

### **Financials**

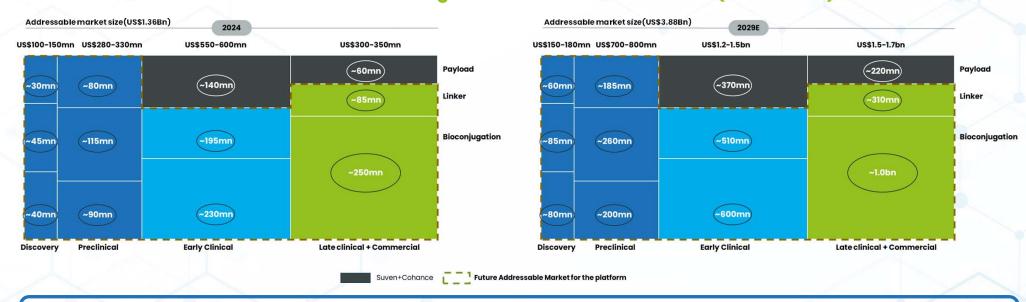
- Has grown sales robustly: US\$7mn in CY21 to US\$32mn in CY24; significant margin expansion potential as business expands
- Company is in a high growth phase foray into cGMP compliant manufacturing in CY24 has led to uptick in opex. Operating leverage will drive EBITDA margin expansion in medium term



# Expansion of ADC - CDMO Market share for Suven+Cohance platform via NJ Bio



Suven's Addressable Market expands 7x (US\$200mn to US\$1.4bn), post-acquisition. Suven Platform and NJ Bio's relevant addressable market is slated to grow from US\$1.4bn to US\$4bn (23%+ CAGR)



## **Expansion of Addressable Market**

- Acquisition expands the serviceable market for Cohance which is presently in late phase/Commercial Payloads
- Supply to Early-Stage Payload along with addition of novel offerings like Linkers and Bioconjugation

## **Market Share within Existing Segments**

Enables Suven+Cohance to tap the customer early and maintain continuity of supply

# New Opportunities: Bioconjugation and Linkers at Commercial Phase

- NJ Bio's Linker and Bioconjugation capabilities are confined to early stage due to lack of GMP experience
- Combination with Cohance will enable entry to the late phase Linkers and Bioconjugation

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# API+ delivers double digit growth in 9M

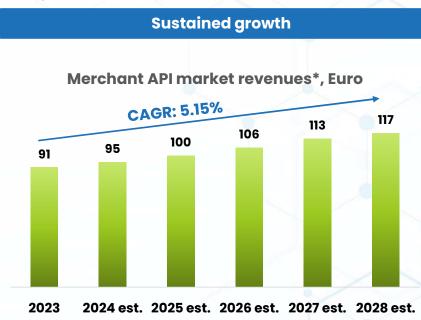


# API+ reported 17% YoY growth in 9M with healthy order book

# What will accelerate the base growth:

- Portfolio is unique and can drive sustained growth
  - Business model focus is on small-mid volume APIs. These products segments have less concentration risk and limited pricing pressure.
  - Focus on expanding market share on the back of deep cost position backed by backward integration
    - Continue to be amongst the Top 3 players for most top molecules (8/10)
  - Capabilities to handle a drug end-to-end throughout its lifecycle
- Higher product validations over 18-24 months; well supported by our BD efforts; target to add 7+ new products in FY25
- Outlook: In-line with earlier expectation, expected to deliver full year growth backed by new product launches and market demand recovery.

# \$101+ Bn Total Addressable Market



Small molecules continue to be a significantly large proportion of Merchant API market revenues

<sup>\*</sup>Source: Industry/Market data

# Specialty Chemical recovering as envisaged

## Ag-chem:

- As indicated, we have seen strong sequential recovery in this business segment. Our concerted BD efforts and early benefits of SBU strategic focus yielding results.
- We're seeing new product discussions and fresh RFQs including from potential new customers and existing strategic partnerships.
- Development and Commercial manufacturing with focus on intermediates and Als
- Flexible capacity Dedicated site for AgChem (Vizag), Space for future expansion,
   Kilo / Pilot scale facility available
- Improved processes, introducing EHS Best Practices

# **Spec Chem**

- Relationships with Originators in Cosmetics, Electronic Chemicals, Photochromic Lens and Energy Industries
- Successfully delivered innovator projects from gram to multi kilo scale
- Amongst India's leading manufacturers of high purity electronic chemicals;
- Highly backward integrated



Source: Cohance investor presentation









Update on Amalgamation of Suven Pharmaceuticals with Cohance Lifesciences

# Suven Pharma – Cohance Merger – Approval update

- Received shareholders' approval, with 99.99% votes in favour
- Final hearing at NCLT scheduled for February 18
- Merger expected to be effective in Q1FY26 subject to regulatory approvals including DoP.









# **ESG Aspirations**



# We have set multi-dimensional ESG goals for the next 5 years

#### **Our achievement** To be achieved **EHS** Social Governance ISO 45001:2018 Reduce absolute Occupational Health Gold in Eco Vadis Employees undergo Scope 1+2 Ethical non-35% 100% **ZERO** and Safety Health and Safety Sustainability ecovadis CO2 emissions from compliance training Management System assessment - 2025 current level (2030) **Business continuity** Transition to management System Representation of Regulatory nonrenewable energy 20+% **ZERO** ISO 22301: 2019 Women workforce by compliance or Signing third party sources of total 2030 fines purchase agreement energy use by 2027 SOLAR for renewable power for all the facilities British safety council five-star certification Reduce, reuse and Employees and Board Members to recycle specific Reduce Attrition by 25% 100% <10 water consumption FY28 acknowledge the Pharmaceutical Code Of Conduct by 2028 SILVER Silver in EcoVadis Supply Chain PSCI SUPPLY CHAIR Sustainability Initiative (PSCI) ecovadis assessment membership - 2025 Promote public Reportable Loss health education Zero **CSR** time injury and disease prevention Committed to Science **Based Target initiative** (SBTi) 97% score in ESG report for FY 2023-24 **ESG Profile** TFS audit







# Financial Performance Q3 & 9MFY25



# Proforma P&L Suven + Cohance Combined - Snapshot

INR million										CAGR	<u>Y</u> c	<u>Y</u>
Combined Proforma P&L Snapshot	FY20	FY21	FY22	FY23	FY24	Q3FY24	Q3FY25	<u>9MFY24</u>	<u>9MFY25</u>	FY20-FY24	<u>Q3</u>	<u>9M</u>
Revenue	16,969	20,140	26,004	26,779	23,922	4,846	6,764	16,903	17,691	9.0%	39.6%	4.7%
COGS	(5,997)	(7,024)	(9,291)	(9,283)	(8,006)	(1,718)	(1,925)	(5,667)	(5,279)			
Material Margin	10,972	13,116	16,713	17,496	15,916	3,128	4,839	11,236	12,412	9.7%	54.7%	10.5%
Material Margin%	64.7%	65.1%	64.3%	65.3%	66.5%	64.6%	71.5%	66.5%	70.2%			
Manufacturing Expenses	(1,994)	(2,461)	(3,009)	(3,242)	(2,506)	(675)	(675)	(1,980)	(1,847)			
Employee cost	(1,924)	(2,195)	(2,719)	(3,038)	(3,771)	(891)	(1,193)	(2,779)	(3,310)			
Other expenses	(1,197)	(1,266)	(1,559)	(1,541)	(1,959)	(440)	(575)	(1,189)	(1,552)			
Adjusted EBITDA (pre Fx)	5,857	7,194	9,426	9,675	7,680	1,122	2,396	5,288	5,702	7.0%	113.6%	7.8%
Operating Forex gain / (loss)	224	261	208	415	102	32	55	98	118			
One time Expenses					752	258	169	594	329			
Adjusted EBITDA (post Fx)	6,080	7,455	9,635	10,089	8,534	1,412	2,620	5,979	6,149	8.8%	85.5%	2.8%
EBITDA%	35.8%	37.0%	37.1%	37.7%	35.7%	29.1%	38.7%	35.4%	34.8%			
Depreciation & Amortization	(679)	(786)	(900)	(1,002)	(1,139)	(319)	(409)	(844)	(1,044)			
Finance costs	(396)	(137)	(173)	(283)	(406)	(104)	(108)	(276)	(306)			
Other income	335	216	309	349	731	148	139	561	430			
Adjusted PBT	5,340	6,748	8,871	9,153	7,720	1,137	2,242	5,420	5,229	9.7%	97.1%	-3.5%
Tax	(1,322)	(1,710)	(2,961)	(2,380)	(1,981)	(299)	(561)	(1,401)	(1,251)			
Adjusted PAT	4,018	5,038	5,910	6,773	5,739	838	1,681	4,019	3,978	9.3%	100.5%	-1.0%
PAT%	23.7%	25.0%	22.7%	25.3%	24.0%	17.3%	24.8%	23.8%	22.5%			

- Q3 Revenue grew by 40% YoY, as guided earlier on higher growth in 2HFY25.
   Strong growth by Pharma CDMO and API+ with healthy sequential recovery in Spec Chem business.
- The gross and EBITDA margins were at 71.5% and 38.7%, respectively, driven by business mix and our BD and R&D teams efforts.
- At a combined platform level, we anticipate growth in the second half of FY25, and growth acceleration from FY26 onwards.

#### Note:

Depreciation and amortization

PAT (post consol adjustments)

Tax impact of above

1) Till FY23, proforma and adjusted financials of Cohance entities (RAC, ZCL and Avra) have been extracted from report issued by Deloitte Touche Tohmatsu India LLP. Adjusted P&L numbers are reported numbers adjusted out for one-time expenses and income; FY24 numbers as per audited financials of the merged entity (Cohance).

(28)

7

1,660

(83)

21

3,916

(88)

22

3,953

2) RoU and Intangible assets Includes RoU under development and intangibles under development respectively

5,039

(185)

47

5,772

(75)

19

6,718

3) Adjusted EBITDA includes One-time adjustment for ESOP, Merger and acquisition costs of Rs. 258 mn and Rs. 169 Mn respectively for Q3FY24 and Q3FY25 and Rs. 594 mn & 329 mn for 9MFY24 & 9MFY25 respectively.

(37)

9

(102)

26

5,662

4) Suven Q3FY25 and 9MFY25 includes consolidation of Sapala and NJ BIO.

Accounting entries relating to merger of AI Pharma and RA Chem

5) PAT attributable to NCI is of Rs. 1.7 mn and Rs. 3.8 mn in 9MFY25 and Q3FY25 respectively in Suven.

4,018



# Proforma BS Suven + Cohance Combined - Snapshot

### INR million

INR MIIIION							
Combined Balance Sheet Snapshot <sup>1</sup>	<u>FY20</u>	FY21	FY22	FY23	<u>FY24</u>	<u>9MFY24</u>	<u>9MFY25</u>
Property, plant and equipment (PPE)	7,354	8,499	9,396	10,059	10,273	10,169	15,527
Right of use asset (RoU) <sup>2</sup>	22	105	193	372	762	732	2,401
Capital work-in-progress	1,114	1,116	758	2,818	4,082	3,541	3,004
Intangible Assets <sup>2</sup>	76	77	146	740	728	728	7,566
Fixed Assets	8,566	9,797	10,492	13,988	15,845	15,170	28,498
Inventories	3,643	4,562	6,100	6,769	5,986	6,243	6,006
Trade receivables	4,326	4,241	6,018	5,356	6,469	5,237	7,104
Trade payables	(2,016)	(2,546)	(2,729)	(2,940)	(2,418)	(2,087)	(3,510)
Core Net Working Capital (Core NWC)	5,953	6,257	9,389	9,185	10,038	9,393	9,600
Other net assets	2,947	3,549	965	1,626	1,002	1,355	(1,486)
	0	0	0	0	0	0	(6,510)
Borrowings	(3,531)	(2,742)	(2,693)	(3,359)	(5,274)	(5,253)	(2,844)
Cash and Cash equivalents (including liquid investments)	3,918	5,820	9,396	5,843	9,440	8,969	2,981
Net (debt) / cash	387	3,078	6,703	2,484	4,167	3,717	137
Net assets	17,853	22,682	27,549	27,283	31,052	29,635	30,239
Shareholder's funds	17,853	22,682	27,549	27,282	31,052	29,635	28,519
Non Controlling Interests							1,720

- The combined balance sheet remained net cash despite strategic acquisition two through funded internal accruals in 9MFY25.
- Working capital as guided has seen improvement and we are positive on sustaining the improving trend as growth traction sustains for the combined business.
- The indicative ROCE business is in 9MFY25 at 25.8%.

RoU and Intangible assets Includes RoU under development and intangibles under development respectively.

Suven 9MFY25 includes consolidation of Sapala and NJ BIO.

Till FY23, proforma and adjusted financials of Cohance entities (RAC, ZCL and Avra) have been extracted from report issued by Deloitte Touche Tohmatsu India LLP. Adjusted P&L numbers are reported numbers adjusted out for one-time expenses and income; FY24 numbers as per audited financials of the merged entity (Cohance). Figures are after adjusting accouting entries relating to merger of AI Pharma and RA Chem.

# **Suven + Cohance Combined Ratios**

Key Ratios <sup>#</sup>	<u>FY20</u>	<u>FY21</u>	<u>FY22</u>	<u>FY23</u>	FY24	<u>9MFY24</u>	<u>9MFY25</u>	<u>Basis</u>
Net Working Capital (as days of sales)	128	113	132	125	153	139	142	NWC / Revenue * 365 days
PPE (as % of sales)	43.3%	42.2%	36.1%	37.6%	42.9%	41.3%	62.8%	PPE / Revenue
Capex spend during the year (INR M)	1,527	1,918	1,663	4,203	2,607	1,688	2,311	
Capex spend (as % of sales)	9.0%	9.5%	6.4%	15.7%	10.9%	6.9%	9.4%	Capex spend / Revenue
(Net Debt)/ Net Cash to adjusted EBITDA (x times)	0.1x	0.4x	0.7x	0.2x	0.5x	0.4x	0.0x	Net Debt / Adjusted EBITDA
Adjusted EBIT (INR M)	5,402	6,670	8,735	9,087	7,394	5,135	5,105	Adjusted EBITDA - Depreciation and Amortization
Adjusted EBIT (INR M) - LTM basis						8,419	7,364	Avg of opening and closing Capital employed (Net fixed assets + NWC + other net assets)
Avg Capital employed (INR M)	13,949	15,192	17,833	21,350	24,001	25,358	28,493	
ROCE (%)	38.7%	43.9%	49.0%	42.6%	30.8%	33.2%	25.8%	Adjusted EBIT / Avg. Capital employed
Avg Shareholder's funds (INR M)	14,460	16,924	22,724	25,944	27,326	28,459	29,785	Avg of Opening and closing shareholder's funds
ROE (%)	27.8%	29.8%	26.0%	26.1%	21.0%	14.1%	13.4%	Adjusted PAT / Avg Shareholder's funds

# calculated based on Proforma P&L and Balance Sheet of Suven + Cohance combined

Note:

New ratios computed on LTM basis for 9MFY25

The above ratios for 9MFY25 are after considering Sapala and NJBIO consolidation

# Suven P&L - 9M Adjusted EBITDA margins at 42%



INR million										CAGR	<u>Y</u>	οY
Consolidated P&L Snapshot	FY20	FY21	FY22	FY23	FY24	Q3FY24	Q3FY25	<u>9MFY24</u>	<u>9MFY25</u>	FY20-FY24	<u>Q3</u>	<u>9M</u>
Revenue	8,338	10,097	13,202	13,403	10,514	2,198	3,072	7,984	7,956	6.0%	39.7%	-0.4%
COGS	(2,292)	(3,019)	(3,991)	(4,091)	(3,150)	(724)	(556)	(2,313)	(1,713)			
Material Margin	6,046	7,078	9,211	9,312	7,364	1,474	2,515	5,671	6,243	5.1%	70.7%	10.1%
Material Margin%	72.5%	70.1%	69.8%	69.5%	70.0%	67.0%	81.9%	71.0%	78.5%			
Manufacturing Expenses	(1,038)	(1,338)	(1,732)	(1,763)	(1,224)	(330)	(347)	(969)	(930)			
Employee cost	(651)	(762)	(1,005)	(1,105)	(1,359)	(348)	(633)	(961)	(1,605)			
Other expenses	(540)	(573)	(680)	(702)	(722)	(135)	(358)	(416)	(691)			
EBITDA (pre Fx)	3,817	4,405	5,794	5,742	4,059	661	1,177	3,325	3,016	1.5%	78.1%	-9.3%
EBITDA%	45.8%	43.6%	43.9%	42.8%	38.6%	30.1%	38.3%	41.6%	37.9%			
Operating Forex gain / (loss)	50	115	138	268	81	15	40	67	82			
Onetime expenses	0	0	0	(134)	211	134	163	134	274			
Adjusted EBITDA (post Fx)	3,867	4,520	5,932	5,876	4,351	810	1,380	3,525	3,372	3.0%	70.5%	-4.3%
EBITDA%	46.4%	44.8%	44.9%	43.8%	41.4%	36.8%	44.9%	44.2%	42.4%			
Depreciation & Amortization	(235)	(316)	(391)	(480)	(502)	(128)	(204)	(373)	(507)			
Finance costs	(199)	(91)	(62)	(128)	(75)	(22)	(33)	(52)	(66)			
Other income	131	27	123	195	538	129	117	382	398			
Adjusted PBT	3,564	4,140	5,602	5,463	4,312	789	1,261	3,482	3,197	4.9%	59.8%	-8.2%
Tax	(875)	(1,053)	(2,138)	(1,451)	(1,118)	(221)	(306)	(913)	(732)			
Adjusted PAT	2,689	3,087	3,464	4,012	3,194	568	955	2,569	2,465	4.4%	68.2%	-4.1%
PAT%	32.2%	30.6%	26.2%	29.9%	30.4%	25.8%	31.1%	32.2%	31.0%			

- Pharma CDMO reported robust growth in Q3 as guided earlier on a heavy 2HFY25, we remain certain on reporting growth in FY25 in Suven consolidated numbers as guided.
- Gross margins improved by 14.52 ppt YoY, purely driven by the business mix.
- Adjusted EBITDA margins were 44.9% an expansion of 8.05 ppt YoY, reflecting our current investments aimed at steering Suven towards the next growth orbit and supported by business mix.
- PAT margins stood at 31.1%.

Note: 1) Adjusted EBITDA includes One-time adjustment for ESOP, Merger and acquisition costs of Rs. 134 mn and Rs.163 Mn respectively for Q3FY24 and Q3FY25 and Rs.134mn & 273 mn for 9MFY24 & 9MFY25 respectively.

<sup>2)</sup> Q3FY25 and 9MFY25 includes consolidation of Sapala and NJ BIO.

<sup>3)</sup> PAT attributable to NCI is of Rs. 1.7 mn and Rs. 3.8 mn in 9MFY25 and Q3FY25 respectively.

# Suven Balance Sheet – Healthy cash rich B/S



#### **INR** million

INR million							
Consolidated Balance Sheet Snapshot	<u>FY20</u>	<u>FY21</u>	FY22	FY23	FY24	<u>9MFY24</u>	<u>9MFY25</u>
Property, plant and equipment (PPE)	3,531	4,371	5,306	5,842	5,672	5,611	8,390
Right of use asset (RoU)	9	17	14	169	406	403	2,111
Capital work-in-progress	1,016	961	300	1,651	1,790	1,813	2,386
Intangible Assets (Including Goodwill)	29	26	22	622	619	619	7,448
Fixed Assets	4,584	5,375	5,642	8,284	8,487	8,447	20,335
Inventories	1,749	2,011	2,834	3,128	2,312	2,532	2,466
Trade receivables	1,172	1,024	2,364	1,109	1,337	1,264	2,865
Trade payables	(711)	(829)	(1,059)	(701)	(424)	(420)	(1,362)
Core Net Working Capital (Core NWC)	2,210	2,205	4,139	3,537	3,225	3,376	3,968
Other net current assets	196	399	424	763	480	0	(1,558)
Other net non current assets	2,863	3,339	738	591	457	801	(252)
Forward Liability	0	0	0	0	0	0	(6,510)
Borrowings	(1,853)	(1,412)	(956)	(692)	(386)	(345)	(677)
Cash and Cash equivalents (including liquid investments)	447	1,902	5,285	4,869	8,244	7,535	2,819
Net (debt) / cash	(1,405)	490	4,330	4,178	7,858	7,190	2,142
Net assets	8,448	11,808	15,272	17,352	20,507	19,814	18,125
Shareholder's funds	8,448	11,808	15,272	17,352	20,507	19,814	16,406
Non Controlling interests					)		1,720

- Working capital under control.
- Free Cash generation in 9MFY25 was Rs 1.33bn.
- Cash and bank balance of Rs 2.82bn, post the payment on the account of NJ Bio acquisition.

Note: 1) PPE includes carved out land of Rs. 375mn in sapala acquistiion & classified as held for sale as per sharepurchase agreement 2) 9MFY25 includes consolidation of Sapala and NJ BIO.

# Suven – Key Ratios



Key Ratios	<u>FY20</u>	<u>FY21</u>	FY22	FY23	FY24	<u>9MFY24</u>	<u>9MFY25</u>	<u>Basis</u>
Net Working Capital (as days of sales)	97	80	114	96	112	106	138	Core NWC / Revenue * 365
PPE (as % of sales)	42.3%	43.3%	40.2%	43.6%	54.0%	48.1%	80.0%	Closing PPE / Revenue
Capex spend during the year (INR M)	1,029	1,108	752	2,857	518	360	938	
Capex spend (as % of sales)	12.3%	11.0%	5.7%	21.3%	4.9%	3.1%	8.9%	Capex spend / Revenue
(Net Debt)/ Net Cash to adjusted EBITDA (x times)	-0.4x	0.1x	0.7x	0.7x	1.8x	1.3x	0.5x	(Net Debt) or Net Cash/ Adjusted EBITDA
Adjusted EBIT (INR M)	3,631	4,203	5,541	5,396	3,848	4,941	3,562	Adjusted EBITDA - Depreciation and Amortization
Avg Capital employed (INR M)	6,655	7,242	8,739	10,586	11,070	12,899	14,316	Avg of Opening and Closing Capital employed (excluding Goodwill, Non- current investments and Cash & CE)
ROCE (%)	54.6%	58.0%	63.4%	51.0%	34.8%	38.3%	24.9%	Adjusted EBIT / Avg. Capital employed
Avg Shareholder's funds (INR M)	5,638	6,785	11,148	14,840	17,088	18,583	18,456	Avg of Opening and closing shareholder's funds (excluding Goodwill and Non-current investments)
ROE (%)	47.7%	45.5%	31.1%	27.0%	18.7%	13.8%	13.4%	Adjusted PAT / Avg Shareholder's funds

Note: 1) Key ratios computed on LTM basis for 9MFY24 and 9MFY25
2) The Ratios for 9MFY25 are after considering Sapala and NJBIO consolidation

# Cohance Proforma P&L - Snapshot



IR million												Yo	Υ
roforma P&L Snapshot	<u>FY19</u>	<u>FY20</u>	<u>FY21</u>	<u>FY22</u>	<u>FY23</u>	FY24	Q3FY24	Q3FY25	<u>9MFY24</u>	<u>9MFY25</u>	CAGR FY19-FY24	<u>Q3</u>	<u>9M</u>
evenue	7,272	8,631	10,043	12,802	13,375	13,408	2,648	3,692	8,919	9,735	13.0%	39.4%	9.1%
OGS	(2,900)	(3,705)	(4,004)	(5,300)	(5,058)	(4,990)	(993)	(1,369)	(3,354)	(3,566)			
aterial Margin	4,372	4,926	6,039	7,502	8,317	8,418	1,655	2,324	5,565	6,169	14.0%	40.4%	10.9%
aterial Margin%	60.1%	57.1%	60.1%	58.6%	62.2%	62.8%	62.5%	62.9%	62.4%	63.4%			
anufacturing Expenses	(1,058)	(955)	(1,123)	(1,277)	(1,480)	(1,282)	(345)	(328)	(1,011)	(917)			
mployee cost	(1,137)	(1,273)	(1,433)	(1,714)	(1,933)	(2,447)	(544)	(560)	(1,818)	(1,705)			
ther expenses	(584)	(657)	(693)	(879)	(839)	(1,279)	(305)	(217)	(773)	(861)			
BITDA (pre Fx)	1,593	2,041	2,790	3,633	4,066	3,410	461	1,219	1,963	2,686	16.4%	164.5%	36.8%
BITDA%							17.4%	33.0%	22.0%	27.6%			
perating Forex gain / (loss)	19	174	146	69	147	21	18	15	31	36			
ne-time Expenses(ESOP&Merger)						752	124	5	460	55			
djusted EBITDA (post Fx)	1,612	2,214	2,936	3,702	4,213	4,183	603	1,240	2,454	2,777	21.0%	105.6%	13.1%
BITDA%	22.2%	25.7%	29.2%	28.9%	31.5%	31.2%	22.8%	33.6%	27.5%	28.5%			
epreciation & Amortization	(479)	(444)	(469)	(509)	(522)	(637)	(191)	(205)	(471)	(537)			
nance costs	(169)	(197)	(45)	(110)	(154)	(332)	(83)	(75)	(224)	(240)			
ther income	157	204	189	186	154	193	19	21	179	32			
djusted PBT	1,121	1,777	2,610	3,269	3,691	3,408	349	981	1,938	2,032	24.9%	181.3%	4.8%
xx	(282)	(447)	(657)	(823)	(929)	(863)	(78)	(255)	(488)	(519)			
djusted PAT	839	1,330	1,953	2,446	2,762	2,544	270	726	1,450	1,513	24.8%	168.5%	4.4%
AT%	11.5%	15.4%	19.4%	19.1%	20.6%	19.0%	10.2%	19.7%	16.3%	15.5%			

- As guided earlier, H2
   FY25 was expected to be
   strong, and this is
   reflected in Q3FY25
   performance, with the
   CDMO segment
   reporting 58% YoY
   growth and the API+
   segment growing 31%
   YoY.
- With a strong order book visibility, Cohance remains well-positioned to drive growth in FY25.
- In Q3, EBITDA margins expanded by 10.8 ppt YoY to 33.6%, driven by a higher share of CDMO and improved utilization.
- Proforma for acquisitions, organic growth for the platform

Note: 1) Till FY23, proforma and adjusted financials of Cohance entities (RAC, ZCL and Avra) have been extracted from report issued by Deloitte Touche Tohmatsu India LLP. Adjusted P&L numbers are reported numbers adjusted out for one-time expenses and income; FY24 numbers as per audited financials of the merged entity (Cohance) 2) Manufacturing expenses include power and fuel, consumption of stores & spares, repairs & maintenance, EHS expenditure, etc. 3) Employee costs include on-payroll employee benefit expenses and contract employee expenses 4) Other expenses include Freight outward, Commission and brokerage, Legal and professional fees, Rates and taxes, Insurance, etc. 5) Adjusted EBITDA includes One-time adjustment for ESOP,merger and other costs of Rs. 124 mn and Rs.5 Mn respectively for Q3FY24 and Q3FY25 and Rs.460mn & 55mn for 9MFY25 respectively.

# Cohance Proforma Balance Sheet - Snapshot



#### **INR** million

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Proforma Balance Sheet Snapshot <sup>1</sup>	Mar19	Mar20	Mar21	Mar22	Mar23	<u>Mar-24</u>	<u>9MFY24</u>	<u>9MFY25</u>
Property, plant and equipment (PPE)	3,699	3,824	4,128	4,090	4,217	4,601	4,557	7,137
Right of use asset (RoU) <sup>2</sup>	0	13	89	179	202	356	328	290
Capital work-in-progress	45	99	155	458	1,167	2,292	1,728	618
Intangible Assets <sup>2</sup>	47	47	51	123	118	109	109	118
Fixed Assets	3,790	3,982	4,422	4,850	5,704	7,358	6,722	8,163
Inventories	1,674	1,894	2,551	3,266	3,641	3,674	3,711	3,540
Trade receivables	2,434	3,154	3,218	3,654	4,202	5,133	3,973	4,240
Trade payables	(852)	(1,305)	(1,716)	(1,670)	(2,141)	(1,994)	(1,666)	(2,148)
Core Net Working Capital (Core NWC)	3,256	3,743	4,052	5,250	5,703	6,813	6,017	5,632
Other net assets	(70)	(111)	(189)	(196)	218	65	555	323
Borrowings	(2,059)	(1,678)	(1,330)	(1,738)	(2,668)	(4,888)	(4,907)	(2,167)
Cash and Cash equivalents (including liquid investments)	3,323	3,470	3,918	4,111	974	1,197	1,434	163
Net (debt) / cash	1,264	1,793	2,588	2,373	(1,694)	(3,692)	(3,473)	(2,005)
Net assets	8,239	9,406	10,874	12,277	9,931	10,545	9,821	12,113
Shareholder's funds	8,239	9,406	10,874	12,277	9,931	10,545	9,821	12,113
Accounting entries relating to merger of AI Phar	med and R	1 Chem						
Goodwill	mod dila ki	· Onom	5,800	5,800	5,800	5,800	5,800	5,800
Tangible assets			397	389	382	376	377	370
Intangible assets			803	624	556	454	468	377
Tax impact			(297)	(137)	(99)			
Other reconciling items			(21)	(20)	`o´			
Net assets (post consol adjustments)	8,239	9,406	17,556	18,932	16,569	17,174	16,466	18,659
Shareholder's funds	8,239	9,406	17,556	18,932	16,569	17,174	16,466	18,659

- Capex stood at ₹1.37bn in 9M FY25, with capacity expansions across multiple plants and a strategic focus flexibility enhancing through backward integration. During the period, we acquired a new facility from Avra Synthesis for capitalized and ₹415mn Ankleshwar Block V with ₹1.36bn.
- Debt reduction remains a priority, with cash flows directed towards repayment.
   Free cash flow generation stood at ₹1.91bn in 9M FY25
- New capacity utilization is set to improve further, supported by synergies from the merger and cross-pollination opportunities gaining momentum.

#### Note:

2) RoU and Intangible assets Includes RoU under development and intangibles under development respectively

Till FY23, proforma and adjusted financials of Cohance entities (RAC, ZCL and Avra) have been extracted from report issued by Deloitte Touche Tohmatsu India LLP. Adjusted P&L numbers are reported numbers adjusted out for one-time expenses and income; FY24 numbers as per audited financials of the merged entity (Cohance). Figures are after adjusting accouting entries relating to merger of AI Pharma and RA Chem.

# Cohance Proforma – Key Ratios



Delivered significant improvement in the net working capital days, to sustain the ongoing effort.

Key Ratios	<u>FY19</u>	FY20	FY21	FY22	<u>FY23</u>	FY24	<u>9MFY24</u>	<u>9MFY25</u>	<u>Basis</u>
Net Working Capital (as days of sales)	163	158	147	150	156	185	170	145	NWC / Revenue * 365
PPE (as % of sales)	51%	44%	41%	32%	32%	34%	35%	50%	PPE / Revenue
Capex spend during the year (INR M)	313	498	810	911	1,346	2,089	1,328	1,373	As per proforma cashflows
Capex spend (as % of sales)	4.3%	5.8%	8.1%	7.1%	10.1%	15.6%	10.3%	9.7%	Capex spend / Revenue
Net Debt)/ Net Cash to adjusted EBITDA (x times)	0.8x	0.8x	0.9x	0.6x	-0.4x	-0.9x	-0.8x	-0.4x	Net Debt / Adjusted EBITDA
Adjusted EBIT (INR M)	1,133	1,771	2,466	3,193	3,691	3,546	3,478	3,802	Adjusted EBITDA - Depreciation and Amortization
avg Capital employed (INR M)		7,294	7,949	9,095	10,764	12,931	12,460	14,177	Avg of opening and closing Capital employed (Net fixed assets + NWC + other net assets)
POCE (%)		24.3%	31.0%	35.1%	34.3%	27.4%	27.9%	26.8%	Adjusted EBIT / Avg. Capital employed
avg Shareholder's funds (INR M)		8,822	10,140	11,576	11,104	10,238	9,876	11,329	Avg of Opening and closing shareholder's funds
ROE (%)		15.1%	19.3%	21.1%	24.9%	24.9%	25.7%	23.0%	Adjusted PAT / Avg Shareholder's funds

ROCE in FY24 and 9MFY25 reflects higher investments in growth capex

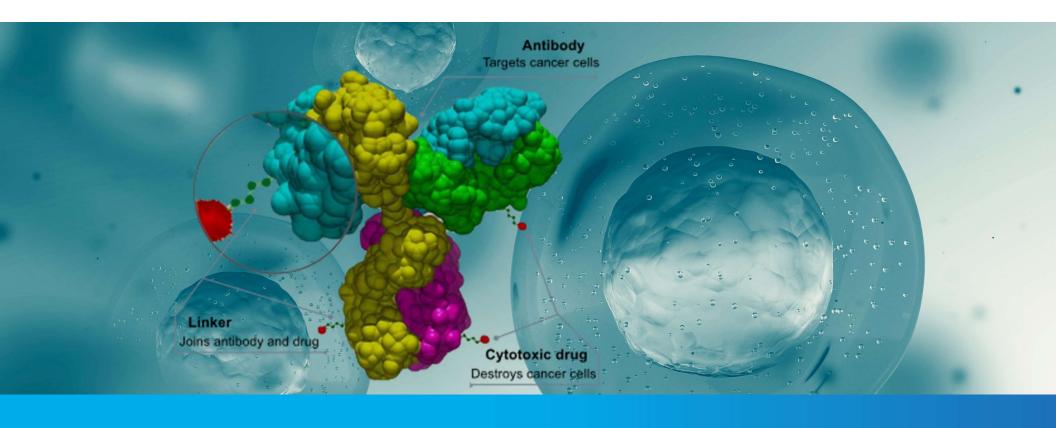
#### Note:

<sup>1)</sup> Key ratios computed on LTM basis for 9MFY24 and 9MFY25.





**Annexure** 



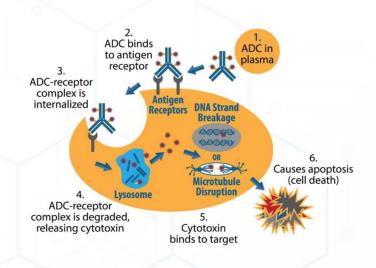


The ADC Segment

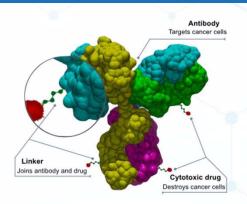
# Building blocks of Antibody Drug Conjugates (ADC) and Mechanism of Action



Rapidly growing class of drugs intended at targeted delivery of highly potent and cytotoxic agents selectively to tumor tissue



Complex Products with sophisticated interplay of variables: The monoclonal antibodies, payloads, and linkers, form a trimolecular prodrug achieving precise and efficient elimination/suppression of target cells and minimize the off-target effects on normal tissues



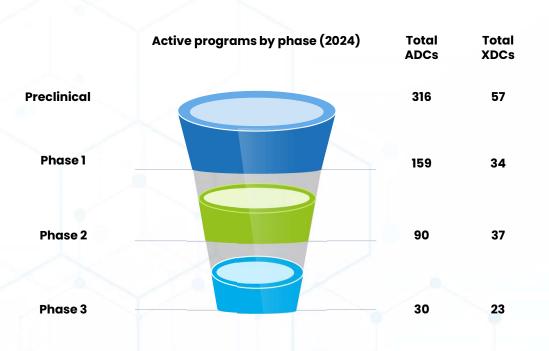
# **Exploring ADC composition**

- Antibody (mAB) is the targeting component of an ADC. It must be highly specific to an antigen that is abundantly expressed on cancer cells but minimally present on healthy cells
- **Linker** is a critical component that connects the antibody to the drug payload. It must be stable in circulation to prevent premature drug release but able to release the drug once inside cancer cells
- **Cytotoxic Payload** is typically a highly potent cytotoxic agent that would be too toxic for systemic administration on its own

# Transforming Therapies: Rapid expansion of ADC R&D Pipelines



ADCs / XDCs as a class continue to see strong growth in R&D investments, providing strong foundation for CRDMO market



- ~750 active ADC / XDC programs in clinics or in preclinical phases.
   Increasing ADC approvals over the recent years
- Volume of clinical trials have experienced a steep growth between '18 and '24: ~120 trials/year up to ~280 trials/year
- Approvals of two blockbuster products (Kadcyle and Enhertu) in 2019 created a pathway for future ADCs
- Growing proportion of Novel conjugates (outside ADC) forms part of XDCs. These include Radioconjugates, Oligo Antibody Conjugates, Protein Degraders and others
- ~85% of the ADC pipeline development originates from Biotechs
- ADC & XDC have shown stronger deal flow (VC investments and Big Pharma in-licensings/M&A) compared to broader Pharma and Biotech industry

# Appendix Adequate Capacity to serve current and future demand



### Vizag, Andhra Pradesh, India

### API's/Advanced Intermediate's/CMO

- o 706 KL reactor volume
- o 3KL to 12KL Reactors
- GL/SS (45No's)



### Pashamylaram, Telangana, India

### API & Formulation Facility

- o 406 KL reaction volume
- o 50L 6000 L GL/SS (45)
- o R&D



### Suryapet, Telangana, India

### Intermediate Facility

- o 300 CM reactors (93)
- o 6651 KL GL/SS
- GMP Intermediates



### Hyderabad Knowledge City, Hyderabad, India Corporate Office





Jeedimetla, Telangana, India

### R&D-Pilot Plant

- Process Research
- o Discovery R&D, Analytical R&D
- o Killo lab, 30L CM Reactors (32)
- o 27 KL GL/SS



Genome Valley, Hyderabad, India

#### R&F

- Synergy Square I, Genome Valley,
- Shamirpet, Hyderabad,
- o Telangana 500078

1) 410KL new capacity in Suryapet included **Source:** Internal



**USA**, New Jersey

### **Business Office**

- Business Development
- Project Management
- o Intellectual Property Management

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# COHANCE'S Specialized manufacturing capabilities



### API Unit-1, Andhra Pradesh, India

- o 120 reactors, > 520Kl capacity
- o USFDA (latest in 2019)
- o EDQM (latest in 2023)
- Others: Korea-FDA, PMDA-Japan, COFEPRIS-Mexico, ANVISA-Brazil, MOH-Russia, CDSCO, WHO GMP



### API Unit-2, Andhra Pradesh, India

- 46 reactors, >140Kl capacity
- o EDQM (latest in 2023)



### API Unit-3, Gujrat, India

- o 68 reactors, >420Kl capacity
- o USFDA (latest in 2023)
- EDQM (latest in 2017)
- Others: PMDA-Japan, COFEPRIS-Mexico, Korea-FDA, ANVISA-Brazil



### FDF Unit-1, Telangana, India

- o 1.8Bn OSDs and 350MT Pellets per annum
- o USFDA (latest in 2019)
- o EU GMP (latest in 2023)
- Others: MHRA, Health Canada, EU GMP, PMDA-Japan, MOH-Russia, WHO GMP, DCGI, Saudi-FDA, Taiwan-FDA





API Unit-4, , Telangana, India

- 60 reactors, >40Kl capacity, Unit with Oncology facility
- USFDA (latest in 2019)
- o EDQM (latest in 2024)
- o Others: WHO GMP

### API Unit-5, Andhra Pradesh, India

- o 49 reactors, >130Kl capacity
- o GMP



FDF Unit-2, Telangana, India

- o 480MT Pellets per annum
- o WHO GMP



# **Contact Information**



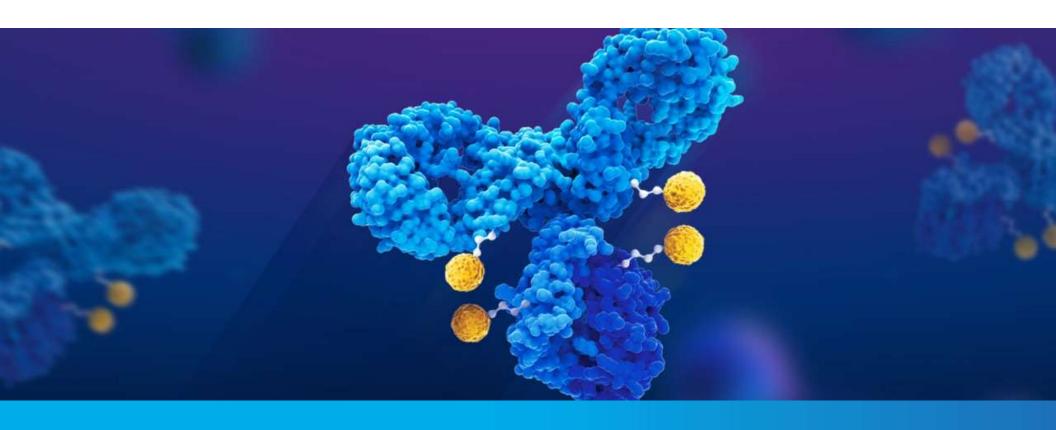
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**Thank You**