



October 27, 2023

BSE Limited,
Floor 25, P. J. Towers
Dalal Street,
Mumbai - 400 001

National Stock Exchange of India Limited,
Exchange Plaza, 5th Floor,
Bandra-Kurla Complex,
Bandra (E),
Mumbai – 400051

Scrip Code: 530019

Symbol: JUBLPHARMA

Dear Sirs,

Sub: Press Release alongwith Earnings Presentation
Ref: Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 ("Listing Regulations")

Pursuant to Provisions of Regulation 30 of Securities and Exchange Board of India (Listing Obligations and Disclosure Requirements) Regulations, 2015, please find herewith the Press Release along with presentation on the financials and performance of the Company for the quarter ended September 30, 2023.

The above mentioned documents will be simultaneously posted on the Company's website at www.jubilantpharmova.com.

We request you to take the same on record.

Thanking you,

Yours faithfully,
For Jubilant Pharmova Limited

Naresh Kapoor
Company Secretary

Encl.: as above

A Jubilant Bhartia Company

OUR VALUES



Jubilant Pharmova Limited

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PRESS RELEASE

Noida, Friday, Oct 27, 2023

JUBILANT PHARMOVA – Q2'FY24 RESULTS

The Board of Jubilant Pharmova Limited met today to approve financial result for the quarter ended Sep 30, 2023.

Financial Results Overview Q2'FY24 - Consolidated (Rs Crs)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24
Total Revenue	1,600	1,587	1,680
Adjusted Revenue¹	1,582	1,587	1,704
Reported EBITDA	232	178	252
	14.5%	11.2%	15.0%
Adjusted EBITDA¹	224	178	289
	14.2%	11.2%	16.9%
Exceptional Items on Bonds Refinancing	(57)	0	0
Profit Before Tax	36	25	98
Adjusted Profit Before Tax²	86	25	135
Reported PAT	5	6	62
Adjusted PAT²	54	6	90

1. Adjustments include non-recurring or one-off revenues related to Remdesivir, Covid related revenues in CDMO Sterile Injectables business and one-time discount to customers of Generics business in US market due to the pricing pressure and shelf stock adjustment in certain products
2. Adjusted Profit Before Tax and Adjusted PAT is after adjusting for exceptional items and adjustments factored in EBITDA

Financial Results Overview H2'FY24 - Consolidated (Rs Crs)

Particulars	H1'FY23	H1'FY24
Total Revenue	3,051	3,267
Adjusted Revenue¹	2,963	3,291
Reported EBITDA	436	430
	14.3%	13.2%
Adjusted EBITDA¹	361	467
	12.2%	14.2%
Exceptional Items on Bonds Refinancing	(57)	0
Profit Before Tax	105	123
Adjusted Profit Before Tax²	88	160
Reported PAT	52	68
Adjusted PAT²	48	96

- Adjustments include non-recurring or one-off revenues related to Remdesivir, Covid related revenues in CDMO Sterile Injectables business and one-time discount to customers of Generics business in US market due to the pricing pressure and shelf stock adjustment in certain products
- Adjusted Profit Before Tax and Adjusted PAT is after adjusting for exceptional items and adjustments factored in EBITDA

Key Ratios – Consolidated

Particulars	Mar 31, 2023	Sep 30, 2023
Net Debt (Constant Currency)	2,426	2,475
Net Debt / EBITDA	2.98	3.06
Net Debt / Equity	0.45	0.46

- EBITDA is on TTM basis
- Exchange rate: 1USD = INR 83.05 as on Sep 30, 2023 and 1USD = INR 82.17 as on Mar 31, 2023



Financial Highlights – Radiopharmaceuticals (Rs Crs)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	248	204	251	444	455
% of Company Revenue	16%	13%	15%	15%	14%
EBITDA	163	93	132	256	226
EBITDA Margin (%)	66%	46%	53%	58%	50%

Key Priorities – Radiopharmaceuticals

- Maintain leadership position in stable high margin core portfolio in North America, e.g., lung functional imaging and thyroid targeted radiotherapeutics
 - Innovation leader in PET cardiac imaging through proprietary RUBY-FILL (best in class cardiac imaging product). Further accelerate Ruby-Fill installs in US and other global markets.
 - Timely execution of roadmap to enable CY 25 launch of MIBG
 - Targeting pediatric patients with high-risk Neuroblastoma. Incidence in the US is 800 (orphan drug) cases per year
 - Peak potential market size for MIBG is around USD 240 Mn
 - Continue launch of high-growth innovation products. Launched Technetium Mertiatide Injection in Q1'FY24. One more launch planned in FY24 subject to regulatory approvals
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Financial Highlights – Radiopharmacies (Rs Crs)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	410	487	490	806	977
% of Company Revenue	26%	31%	29%	26%	62%
EBITDA	-17	2	6	-38	8
EBITDA Margin (%)	-4%	0%	1%	-5%	1%

Key Priorities – Radiopharmacies

- Accelerate sales of high growth new products, e.g., Ga-PSMA, and to further gain market share in existing SPECT products
 - Maintain current momentum of strong growth in 3rd party sales
 - Leverage existing cyclotrons to capture share of PET product growth
 - Additionally, explore opportunity to further expand presence into PET radiopharmacies, due to strong demand of PET products, such as PET-PSMA
 - Continue to enhance operational and procurement efficiencies leading to improvement in financial performance in FY24
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-



Financial Highlights – Allergy Immunotherapy (Rs Crs)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	156	151	179	286	330
% of Company Revenue	10%	10%	11%	9%	21%
EBITDA	53	50	86	97	136
EBITDA Margin (%)	34%	33%	48%	34%	41%

Key Priorities – Allergy Immunotherapy

- #2 player in US Sub-Cutaneous Immunotherapy market (Venom and Non-Venom) of >\$200M. High barriers to entry as products are branded biologicals with regulatory approvals grandfathered in
 - Further strengthen the prescriber base for Venom immunotherapy in the US through continuous brand building. Sole supplier of venom in US
 - Focus on increasing market share in Non Venom Allergenic extracts (e.g., Dog, Cat, Mite allergy) and Skin Testing Devices in US. #2 player in US
 - Gain market share in Europe and other non-US markets across Venom product category
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-



Financial Highlights – CDMO Sterile Injectables (Rs Crs)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	299	254	301	562	555
% of Company Revenue	19%	16%	18%	18%	17%
EBITDA	71	41	56	203	97
EBITDA Margin (%)	24%	16%	19%	36%	17%

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Adjusted Revenue	277	254	301	470	555
Adjusted EBITDA	51	41	56	116	97
Adjusted EBITDA Margin	18%	16%	19%	25%	17%

Adjusted Revenue and EBITDA excludes the one-off COVID related business

Key Priorities – CDMO Sterile Injectables

- Global Fill and Finish Sterile Injectable markets of USD 13Bn, with double digit growth rate projected over next 5 years
- Focus is on-time and at-cost execution of USD 370Mn capacity expansion in Spokane and Montreal, to double the CMO capacity over next 5+ years in a phased manner
- Cooperative agreement with US Govt. for USD 149.6 Mn and concessional loan from Canadian Govt. for ~USD 48 Mn
- Leverage differentiated technical know-how to further build scale, e.g., Hormones, Ophthalmic, and Vaccines etc.
- CMO Montreal facility received OAI from the US FDA in May 2023. Engaging with the US FDA to address its observations and resolve the OAI status at the facility





Financial Highlights – Generics (Rs Crs)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	161	202	172	340	375
% of Company Revenue	10%	13%	10%	11%	24%
EBITDA	-82	-21	-50	-155	-71
EBITDA Margin (%)	(51%)	(10%)	(29%)	(46%)	(19%)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Adjusted Revenue	165	202	172	343	375
Adjusted EBITDA	-69	-21	-46	-143	-67
Adjusted EBITDA Margin	(42%)	(10%)	(27%)	(42%)	(18%)

Adjusted Revenue and EBITDA excludes the Remdesivir business and one-time discount to customers in US market due to pricing pressure and shelf stock adjustment in certain products

Key Priorities – Generics

- Continue quality improvement initiatives and engagement with the US FDA for resolution of Import Alert at the Roorkee facility.
- Salisbury site is compliant with US FDA. Roorkee site is compliant with other key non-US markets, e.g., MHRA, Japan, South Africa, and Canada.
- Focus on implementation of Rs 150 Cr cost optimization opportunities. Benefits have started getting reflecting in performance from Q1'FY24 onwards
- Re-prioritise geography-mix to accelerate growth in branded markets such as India and select International markets
- Continue to strengthen leadership position in select products across markets





Financial Highlights – Drug Discovery Services (Rs Crs)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	150	103	115	268	218
% of Company Revenue	9%	7%	7%	9%	14%
EBITDA	54	22	26	93	47
EBITDA Margin (%)	36%	21%	22%	35%	22%

Key Priorities – Drug Discovery Services

- Leverage state of the art infrastructure and differentiated technical know-how, e.g., Integrated Drug Discovery, DMPK to drive new customer acquisitions in drug discovery
- Continue to invest in capabilities for improving productivity, speeding up time to market and lowering cost of innovation
- Further strengthen the CDMO contract pipeline within existing and new technologies



Financial Highlights – CDMO - API (Rs Crs)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	170	177	165	332	341
% of Company Revenue	11%	11%	10%	11%	22%
EBITDA	14	13	15	21	28
EBITDA Margin (%)	8%	7%	9%	6%	8%

Key Priorities – CDMO - API

- In March 2023, the API plant at Nanjangud saw reversal of OAI status to compliant VAI status from USFDA, driven by Quality Improvement Initiatives at site.
- Going forward, focus is to drive higher capacity utilization including through launch of new products and by acquiring new customers globally
- Operations transformation program underway to increase productivity while lowering costs. Benefits will start becoming visible from H2'FY24



Key Priorities – Proprietary Novel Drugs

- Clinical stage precision therapeutics business advancing potent and selective small molecules to address unmet medical needs in oncology and autoimmune diseases
- Wholly owned assets; opportunities to explore institutional funding, as well as maximize partnerships to get non-dilutive funding
- Emphasis on cost optimized operating model with a focus on value creation
- Business' most advanced program - first in class dual inhibitor of LSD1/HDAC6 is undergoing Phase I/II clinical trials
- Another program – PRMT5 Brain penetrant has received IND approval
- LSD1/HDAC6 and PRMT5 have the potential to address high unmet medical needs globally with multi-billion-dollar market size

Segment Financial Results - Consolidated (Rs Crs)

Segment Revenue	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24	FY23
1. Radiopharma	658	691	741	1,250	1,432	2,552
Radiopharmaceuticals	248	204	251	444	455	872
Radiopharmacies	410	487	490	806	977	1,681
2. Allergy Immunotherapy	156	151	179	286	330	603
3. CDMO Sterile Injectables	299	254	301	562	555	1,155
4. Generics	161	202	172	340	375	762
5. CRDMO	320	280	279	600	559	1,185
Drug Discovery Services	150	103	115	268	218	522
CDMO - API	170	177	165	332	341	662
6. Proprietary Novel Drugs	0	0	0	4	0	4
Unallocable Corporate Income	6	9	8	10	17	22
Total Revenue	1,600	1,587	1,680	3,051	3,267	6,282

Segment EBITDA	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24	FY23	FY23
	Margin	Margin	Margin	Margin	Margin	Margin	Margin
1. Radiopharma	146	95	138	219	233	378	15%
Radiopharmaceuticals	163	66%	93	46%	132	53%	465
Radiopharmacies	(17)	(4%)	2	0%	6	1%	(87)
2. Allergy Immunotherapy	53	34%	50	33%	86	48%	205
3. CDMO Sterile Injectables	71	24%	41	16%	56	19%	345
4. Generics	(82)	(51%)	(21)	(10%)	(50)	(29%)	(230)
5. CRDMO	68	21%	35	12%	41	15%	199
Drug Discovery Services	54	36%	22	21%	26	22%	164
CDMO - API	14	8%	13	7%	15	9%	35
6. Proprietary Novel Drugs	(10)	(10)	(8)	(17)	(18)	(35)	17%
Unallocable Corporate (Expenses) / Income	(14)	(12)	(11)	(25)	(23)	(48)	5%
Total EBITDA (Reported)	232	14%	178	11%	252	15%	815
				436	14%	430	13%



About Jubilant Pharmova Limited

Jubilant Pharmova Limited (formerly Jubilant Life Sciences Limited) is a company with global presence that is involved in Radiopharma, Allergy Immunotherapy, CDMO Sterile Injectables, Contract Research Development and Manufacturing Organisation (CRDMO), Generics and Proprietary Novel Drugs businesses.

In the Radiopharma business, the Company is involved in manufacturing and supply of Radiopharmaceuticals with a network of 46 radio-pharmacies in the US. The Company's Allergy Immunotherapy business is involved in the manufacturing and supply of allergic extracts and venom products in the US and in some other markets such as Canada, Europe and Australia. Jubilant through its CDMO Sterile Injectables business offers manufacturing services including sterile fill and finish injectables (both liquid and lyophilization), full-service ophthalmic offer (liquids, ointments & creams) and ampoules.

The CRDMO business of the Company includes the Drug Discovery Services business that provides contract research and development services through two world class research centers in Bangalore and Noida in India and the CDMO-API business that is involved in the manufacturing of Active Pharmaceutical Ingredients. Jubilant Therapeutics is involved in Proprietary Novel Drugs business and is an innovative biopharmaceutical company developing breakthrough therapies in the area of oncology and auto-immune disorders. The company operates six manufacturing facilities that cater to all the regulated market including USA, Europe and other geographies. Jubilant Pharmova Limited has a team of over 5,500 multicultural people across the globe. The Company is well recognized as a 'Partner of Choice' by leading pharmaceuticals companies globally.

For more information, please contact:

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Disclaimer

Statements in this document relating to future status, events, or circumstances, including but not limited to statements about plans and objectives, the progress and results of research and development, potential product characteristics and uses, product sales potential and target dates for product launch are forward-looking statements based on estimates and the anticipated effects of future events on current and developing circumstances. Such statements are subject to numerous risks and uncertainties and are not necessarily predictive of future results. Actual results may differ materially from those anticipated in the forward-looking statements. Jubilant Pharmova may, from time to time, make additional written and oral forward looking statements, including statements contained in the company's filings with the regulatory bodies and its reports to shareholders. The company assumes no obligation to update forward-looking statements to reflect actual results, changed assumptions or other factors.



Financial Results

Quarter Ended September 30, 2023

October 2023

Jubilant Pharmova is uniquely positioned to create sustained Shareholder Value



An integrated global pharmaceuticals and contract research company



Over 5,500 people globally, including over 2,100 in North America



6 manufacturing facilities catering to regulated markets including USA, Europe and other geographies



Strong position in Radiopharmaceuticals, Allergy Immunotherapy and CDMO Sterile Injectables



One of the leading and growing India based Contract Research and Development companies



Proprietary business has strong portfolio of programs in oncology and auto immune disorders



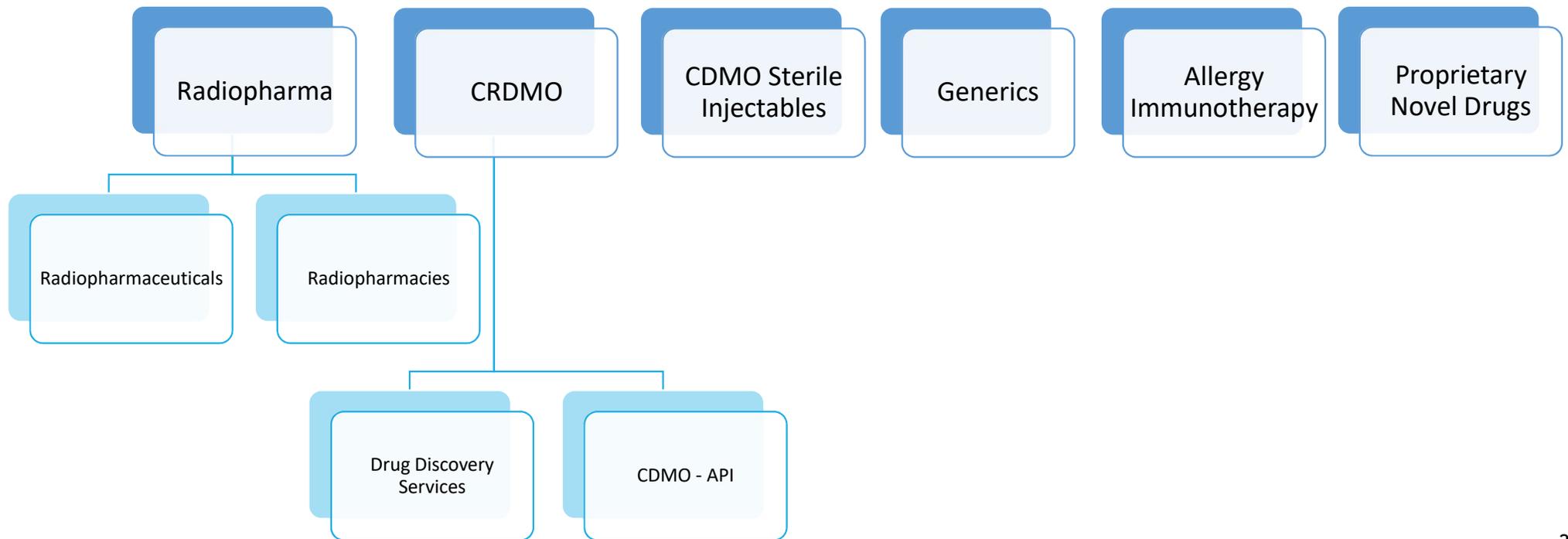
Drug Discovery services through two world-class centers in Bengaluru and Greater Noida



FY23 Revenue ~Rs 6,300 Crs (~US\$ 783 million)

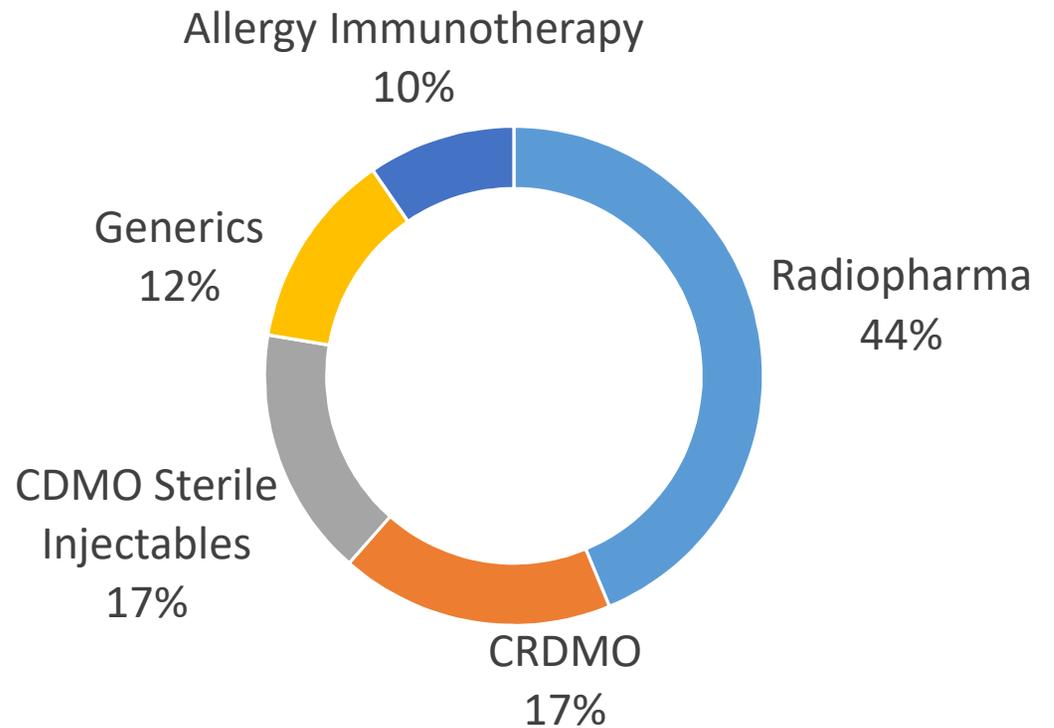
Business Structure

Jubilant Pharmova Limited





H1'FY24 – Segment Wise Revenue Split



Business Snapshot

- #3 radiopharmaceutical manufacturer in the US
- Manufacturing facility based in Montreal Canada
- # 2 network in the US with 46 radiopharmacies

Radiopharma

- Leading contract manufacturer for Sterile Injectables
- Differentiated technologies, viz. hormonal steroids, vaccines
- Manufacturing facilities in Spokane and Montreal

CDMO - Sterile Injectables

- Manufacturing facilities at Roorkee, India and Salisbury, US
- Focus on quality leadership and compliances
- Market leadership in select products in US and branded markets

Generics

- Manufacturing facility at Nanjangud, India
- Over 50% of API sales are to regulated markets
- Strong market share in CNS / CVS products globally

CDMO - API

- #2 player in the US allergenic extract market
- Sole supplier of venom in the US
- Manufacturing facility at Spokane, WA (USA)

Allergy Immunotherapy

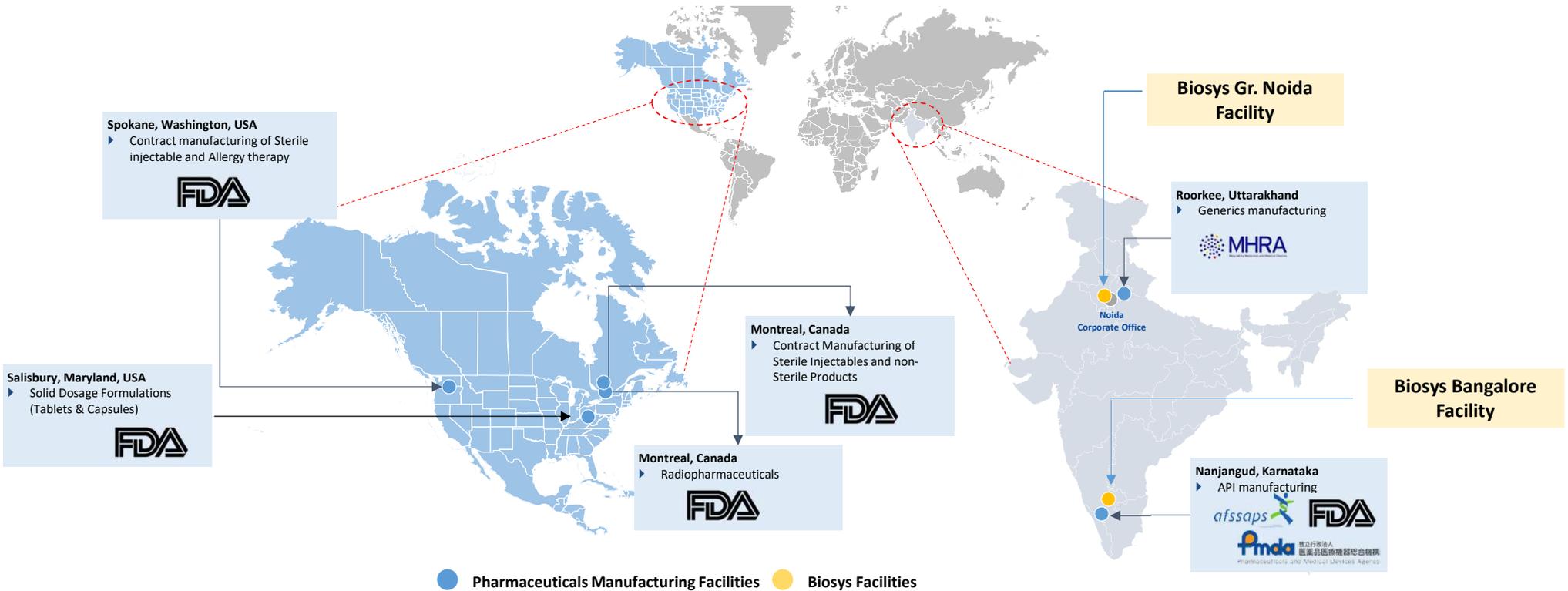
- Fully integrated Drug Discovery services provider
- Facilities in Greater Noida and Bengaluru
- Provides Drug Discovery and CDMO services to global innovators

Drug Discovery Services

- High potential programs in the area of oncology and autoimmune disorders
- Lead program LSD1/HDAC6 inhibitor has successfully started Phase I trials
- IND approval for second program, JBI-778, an Oral, Brain Penetrant PRMT5 Inhibitor

Proprietary Novel Drugs

High-Quality, World-Class Manufacturing Footprint and Operational Facilities



- 6 manufacturing facilities catering to the regulated markets, including USA, Europe and other geographies.
- Contract research and development services through 2 world-class research centers in Bengaluru and Greater Noida in India.

Financial Highlights

Q2 & H1'FY24



Figures in Rs Crs

Particulars	Q2'FY23	Q1'FY24	Q2'FY24
Total Revenue	1,600	1,587	1,680
Adjusted Revenue¹	1,582	1,587	1,704
Reported EBITDA	232	178	252
	14.5%	11.2%	15.0%
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	14.2%	11.2%	16.9%
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Profit Before Tax	36	25	98
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Reported PAT	5	6	62
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- Exceptional cost of Rs 57 Crore in Q2'FY23 included Rs 48 Crore of foreclosure charges related to bond repayment and balance due to write-off of capitalized debt origination costs

1. Adjustments include non-recurring / one-off revenues related to Remdesivir, Covid related revenues in CDMO Sterile Injectables business and one-time discount to customers of Generics business in US market due to pricing pressure and shelf stock adjustment in certain products
2. *Adjusted Profit Before Tax / PAT is after adjusting for exceptional items and adjustments factored in EBITDA*

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Figures in Rs Crs

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Total Revenue	3,051	3,267
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Reported PAT	52	68
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2. *Adjusted Profit Before Tax / PAT is after adjusting for exceptional items and adjustments factored in EBITDA*

Financial Results Overview H1'FY24 - Consolidated

Key Ratios H1'FY24 - Consolidated

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Net Debt (Constant Currency)	2,426	2,475
Net Debt / EBITDA	2.98	3.06
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1. EBITDA is on TTM basis
2. Exchange rate: 1USD = INR 83.05 as on Sep 30, 2023 and 1USD = INR 82.17 as on Mar 31, 2023

Financial Highlights

Radiopharmaceuticals

Figures in Rs Crs

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	248	204	251	444	455
% of Company Revenue	16%	13%	15%	15%	14%
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Key Priorities

Radiopharmaceuticals

- Maintain leadership position in stable high margin core portfolio in North America, e.g., lung functional imaging and thyroid targeted radiotherapeutics
- Innovation leader in PET cardiac imaging through proprietary RUBY-FILL (best in class cardiac imaging product). Further accelerate Ruby-Fill installs in US and other global markets.
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 - Peak potential market size for MIBG is around USD 240 Mn
- Continue launch of high-growth innovation products. Launched Technetium Mertiatide Injection in Q1'FY24. One more launch planned in FY24 subject to regulatory approvals



**Ruby-fill Elution
System with
Generator**





Financial Highlights Radiopharmacies

Figures in Rs Crs

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Revenue	410	487	490	806	977
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EBITDA Margin (%)	(4%)	0%	1%	(5%)	1%

Key Priorities

Radiopharmacies

- *Accelerate sales of high growth new products, e.g., Ga-PSMA, and to further gain market share in existing SPECT products*
- *Maintain current momentum of strong growth in 3rd party sales*
- *Leverage existing cyclotrons to capture share of PET product growth*
- *Additionally, explore opportunity to further expand presence into PET radiopharmacies, due to strong demand of PET products, such as PET-PSMA*
- *Continue to enhance operational and procurement efficiencies leading to improvement in financial performance in FY24*



Financial Highlights

Allergy Immunotherapy

Figures in Rs Crs

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	156	151	179	286	330
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Key Priorities

Allergy Immunotherapy

- *#2 player in US Sub-Cutaneous Immunotherapy market (Venom and Non-Venom) of >\$200M. High barriers to entry as products are branded biologicals with regulatory approvals grandfathered in*
- *Further strengthen the prescriber base for Venom immunotherapy in the US through continuous brand building. Sole supplier of venom in US*
- *Focus on increasing market share in Non Venom Allergenic extracts (e.g., Dog, Cat, Mite allergy) and Skin Testing Devices in US. #2 player in US*
- *Gain market share in Europe and other non-US markets across Venom product category*

Financial Highlights

CDMO Sterile Injectables

Figures in Rs Crs

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Revenue	299	254	301	562	555
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EBITDA Margin (%)	24%	16%	19%	36%	17%
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- Focus is on-time and at-cost execution of USD 370Mn capacity expansion in Spokane and Montreal, to double the CMO capacity over next 5+ years in a phased manner
- Cooperative agreement with US Govt. for USD 149.6 Mn and concessional loan from Canadian Govt. for ~USD 48 Mn
- Leverage differentiated technical know-how to further build scale, e.g., Hormones, Ophthalmic, Vaccines etc.
- CMO Montreal facility received OAI from the US FDA in May 2023. Engaging with the US FDA to address its observations and resolve the OAI status at the facility

Financial Highlights Generics

Figures in Rs Crs

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Revenue	161	202	172	340	375
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EBITDA Margin (%)	(51%)	(10%)	(29%)	(46%)	(19%)

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Adjusted Revenue	165	202	196	343	399
Adjusted EBITDA	(69)	(21)	(13)	(143)	(34)
Adjusted EBITDA Margin	(42%)	(10%)	(7%)	(42%)	(9%)

- Adjusted Revenue and EBITDA excludes the Remdesivir business and one-time discount to customers in US market due to pricing pressure and shelf stock adjustment in certain products

Key Priorities

Generics

- Continue quality improvement initiatives and engagement with the US FDA for resolution of Import Alert at the Roorkee facility.
- Salisbury site is compliant with US FDA. Roorkee site is compliant with other key non-US markets, e.g., MHRA, Japan, South Africa, Canada.
- Focus on implementation of Rs 150 Cr cost optimization opportunities. Benefits have started getting reflecting in performance from Q1'FY24 onwards
- Re-prioritise geography-mix to accelerate growth in branded markets such as India and select International markets
- Continue to strengthen leadership position in select products across markets

Financial Highlights

Drug Discovery Services

Figures in Rs Crs

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	150	103	115	268	218
% of Company Revenue	9%	7%	7%	9%	7%
EBITDA	54	22	26	93	47
EBITDA Margin (%)	36%	21%	22%	35%	22%



Key Priorities

Drug Discovery Services

- Leverage state of the art infrastructure and differentiated technical know-how, e.g., Integrated Drug Discovery, DMPK to drive new customer acquisitions in drug discovery
- Continue to invest in capabilities for improving productivity, speeding up time to market and lowering cost of innovation
- Further strengthen the CDMO contract pipeline within existing and new technologies



Financial Highlights

CDMO - API

Figures in Rs Crs

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	170	177	165	332	341
% of Company Revenue	11%	11%	10%	11%	10%
EBITDA	14	13	15	21	28
EBITDA Margin (%)	8%	7%	9%	6%	8%



Key Priorities

CDMO - API

- In March 2023, the API plant at Nanjangud saw reversal of OAI status to compliant VAI status from USFDA, driven by Quality Improvement Initiatives at site.
- Going forward, focus is to drive higher capacity utilization including through launch of new products and by acquiring new customers globally
- Operations transformation program underway to increase productivity while lowering costs. Benefits will start becoming visible from H2'FY24.





Financial Highlights Proprietary Novel Drugs

Figures in Rs Crs

Particulars	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24
Revenue	0	0	0	4	0
EBITDA	(10)	(10)	(8)	(17)	(18)

Key Highlights & Priorities

Proprietary Novel Drugs

- Clinical stage precision therapeutics business advancing potent and selective small molecules to address unmet medical needs in oncology and autoimmune diseases
- Wholly owned assets; opportunities to explore institutional funding, as well as maximize partnerships to get non-dilutive funding
- Emphasis on cost optimized operating model with a focus on value creation
- Business' most advanced program - first in class dual inhibitor of LSD1/HDAC6 is undergoing Phase I/II clinical trials
- Another program – PRMT5 Brain penetrant has received IND approval
- LSD1/HDAC6 and PRMT5 have the potential to address high unmet medical needs globally with multi-billion-dollar market size

Business Enablers



Sustainability continues to be an important focus area for us

Participated in S&P DJSI Assessment:

- Achieved **94 percentile** in the Global Pharmaceutical Industry
- Among the **top 6% companies globally**

- Received Gold Rating
- Achieved **92 percentile (Score 67/100)**

CLIMATE LEADERS 2022

CLIMATE LEADERS Asia-Pacific 2022

Climbed from 23rd to **6th position** on **Responsible Business Ranking** by the joint ET-Future scope 8th Sustainability Index Report

CDP DISCLOSER 2022	Climate	B
	Water	B
	Supply Chain	A-



- 2001**: ISO 14000 Certification
- 2002**: Sustainability Policy Adopted
- 2003**: Sustainability Report Released
- 2005**: Became GRI Organization Stakeholder Member
- 2008**: Jubilant Bhartia Foundation CSR Wing Launched
- 2009**: Climate Change Mitigation and Green Supply Chain Policy
- 2010**: Became UNGC Signatory and Participation in CDP
- 2013**: 1st EvoVadis Review conducted
- 2015**: SoFI Sustainability Software Launched
- 2019**: Sustainability Goals created aligned with UNSDG
- 2021**: Dow Jones Sustainability Index (DJSI)



Annexure

Segment Financial Results Overview | Consolidated

Figures in Rs Crs 

Segment Revenue	Q2'FY23	Q1'FY24	Q2'FY24	H1'FY23	H1'FY24	FY23
1. Radiopharma	658	691	741	1,250	1,432	2,552
Radiopharmaceuticals	248	204	251	444	455	872
Radiopharmacies	410	487	490	806	977	1,681
2. Allergy Immunotherapy	156	151	179	286	330	603
3. CDMO Sterile Injectables	299	254	301	562	555	1,155
4. Generics	161	202	172	340	375	762
5. CRDMO	320	280	279	600	559	1,185
Drug Discovery Services	150	103	115	268	218	522
CDMO - API	170	177	165	332	341	662
6. Proprietary Novel Drugs	0	0	0	4	0	4
Unallocable Corporate Income	6	9	8	10	17	22
Total Revenue	1,600	1,587	1,680	3,051	3,267	6,282

Segment EBITDA	Q2'FY23	Q2'FY23 Margin	Q1'FY24	Q1'FY24 Margin	Q2'FY24	Q2'FY24 Margin	H1'FY23	H1'FY23 Margin	H1'FY24	H1'FY24 Margin	FY23	FY23 Margin
1. Radiopharma	146	22%	95	14%	138	19%	219	18%	233	16%	378	15%
Radiopharmaceuticals	163	66%	93	46%	132	53%	256	58%	226	50%	465	53%
Radiopharmacies	(17)	(4%)	2	0%	6	1%	(38)	(5%)	8	1%	(87)	(5%)
2. Allergy Immunotherapy	53	34%	50	33%	86	48%	97	34%	136	41%	205	34%
3. CDMO Sterile Injectables	71	24%	41	16%	56	19%	203	36%	97	17%	345	30%
4. Generics	(82)	(51%)	(21)	(10%)	(50)	(29%)	(155)	(46%)	(71)	(19%)	(230)	(30%)
5. CRDMO	68	21%	35	12%	41	15%	114	19%	76	13%	199	17%
Drug Discovery Services	54	36%	22	21%	26	22%	93	35%	47	22%	164	31%
CDMO - API	14	8%	13	7%	15	9%	21	6%	28	8%	35	5%
6. Proprietary Novel Drugs	(10)		(10)		(8)		(17)		(18)		(35)	
Unallocable Corporate (Expenses) / Income	(14)		(12)		(11)		(25)		(23)		(48)	
Total EBITDA (Reported)	232	14%	178	11%	252	15%	436	14%	430	13%	815	13%

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