

June 8, 2018

BSE Limited

Floor 25, P. J. Towers Dalal Street, Mumbai - 400 001 National Stock Exchange of India Limited Exchange Plaza Bandra Kurla Complex Bandra (E)

Mumbai - 400 051

Dear Sirs,

Sub.: Intimation of Schedule of Investor/Analyst meetings on June 11, 2018

Pursuant to the provisions of Regulation 30 of the Securities and Exchange Board of India (Listing Obligations and Disclosure Requirements) Regulations, 2015, we enclose the details of investor/analyst meetings scheduled on June 11, 2018.

The schedule may undergo change due to exigencies on the part of Investors/Analysts/ Company.

We also enclose the presentation to be used during the meetings.

This is for your kind information and records.

Thanking you,

Yours faithfully, For Jubilant Life Sciences Limited

Rajiv Shah Company Secretary

Encl.: as above

A Jubilant Bhartia Company



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CIN: L24116UP1978PLC004624

Investor/Analyst meetings scheduled on 11th June, 2018

Meeting Date	Type of Meeting	Name of fund/firm	Location
Monday, 11 th June 2018	One on One	 HDFC Mutual Fund (Credit) Axis Mutual Fund (Fixed Income) UTI Mutual Fund (Fixed Income) ICICI Prudential Mutual Fund (Credit) SBI Life Insurance (Fixed Income) Yes Bank (Debt Capital Markets) 	Mumbai





Jubilant Life Sciences Limited

June 2018

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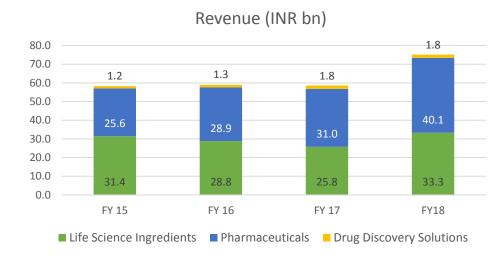
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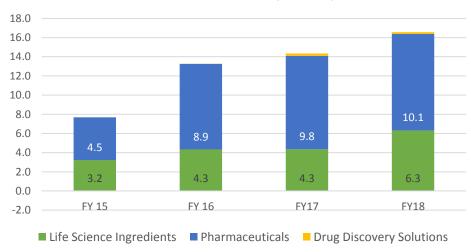
- 1. The numbers for the quarter have been reclassified and regrouped wherever necessary
- 2. Closing Exchange Rate for USD 1 at Rs. 66.25 as on March 31'16, Rs. 64.85 as on March 31'17, Rs. 63.88 as on December 31'17 & Rs. 65.17 as on March 31'18
- The Exchange fluctuation on long term forex loans in Indian books have been amortized over the tenure of the loan period as recommended under Indian Accounting Standards
- 4. Financial numbers upto FY 2015 are as per erstwhile Indian Generally Accepted Accounting Principles (I-GAAP), whereas for FY 2016 onwards, it is as per Indian Accounting Standards

Jubilant Life Sciences At a Glance

- Global integrated pharma and life sciences solutions provider with a track record of about 40 years
- Strategic presence in Injectables with USFDA approved Manufacturing facilities in North America
- Strong positions in products across niche businesses such as Radiopharmaceuticals/Allergy Therapy Products
- Expertise in Chemistry and manufacturing spans across about 4 decades of experience
- 4 USFDA approved manufacturing facilities in North America and 2 USFDA approved manufacturing facilities in India
- 5 state-of-the-art Life Sciences Ingredients manufacturing facilities in India
- Employs around 7,600 people globally, including around 2,200 in North America and around 1,100 dedicated to R&D



Business EBITDA (INR bn)





Jubilant Life Sciences Overview

Pharmaceuticals

Specialty Injectables

- √ Radiopharmaceuticals
- ✓ CMO of Sterile Injectables
- ✓ Allergy Therapy Products

Generics

- ✓ Solid Dosage Formulations
- ✓ Active Pharmaceutical Ingredients
- ✓ India Branded Pharmaceuticals

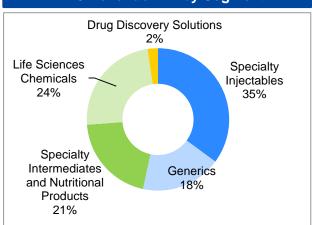
Life Science Ingredients

Specialty Intermediates and Nutritional Products

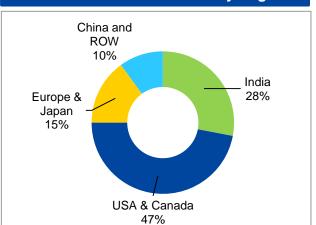
Life Sciences Chemicals

Drug Discovery Solutions

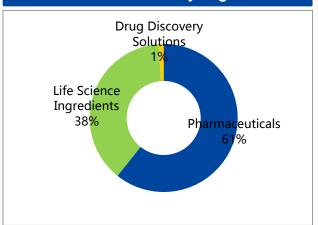
FY18 Revenue Mix by Segment



FY18 Revenue Contribution by Region



FY18 EBITDA Mix by Segment





Key Credit Strengths

Global Competitive Edge Due to Low Cost and Vertical Integration Strong Leadership in Key Products and Markets 3 De-risked Business Model With Diverse Sources of Revenue 4 Strong Pipeline of Products with Deep R&D Capabilities 5 High-Quality, World -Class, Low Cost Manufacturing Footprint 6 Experienced Management team with high standards of corporate governance



Global Competitive Edge Due to Low Cost and Vertical Integration

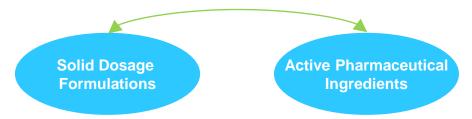
Vertical integration across the value chain enables cost competitive advantage resulting in higher margins

Pharmaceuticals

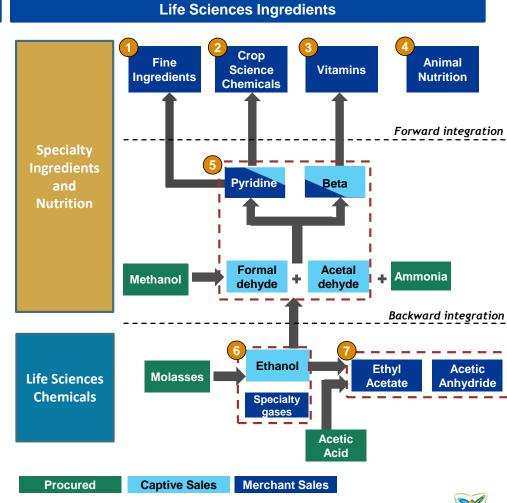
- Presence across the spectrum offering products/services and delivering innovative solutions for customers
- Specialty Injectables



 APIs from the manufacturing facilities are used for dosage formulations under generic business



 Majority of commercial solid dosage formulations are based on in-house APIs





Leadership Position in Key Products...

Pharmaceuticals Pharmaceutical						
Specialty Injectables	 Leader in Radiopharmaceutical products such as MAA, DTPA and I-131 Ranked in top 5 Contract Manufacturers in North America for sterile injectables Leading supplier in North America for Allergy Therapy Products 					
Generics	 US leadership positions in certain Solid Dosage Formulations products in CVS (Terazosin), CNS (Prochlorperazine) and Methylprednisolone Global leadership positions in certain key API products in CVS (Valsartan, Irbesartan) and CNS (Oxcarbazepine, Carbamazepine, Tramadol, Donepezil) 					

Life Sciences Ingredients

Speciality Intermediates and Nutritional Products	 Global leadership position in Pyridines & derivatives Second largest producer globally in Vitamin B3
Life Science Chemicals	 Globally #4 in Acetic Anhydride (Merchant Sales) Globally #7 in Ethyl Acetate

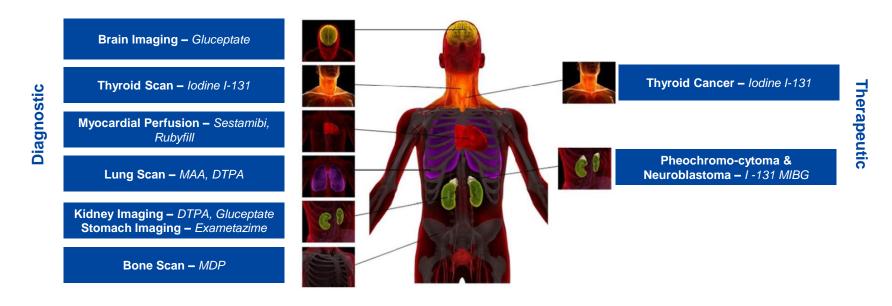




...With Strong Position in Radiopharma & Allergy Therapy Products Business

Radiopharmaceuticals

- Wide array of therapeutic and diagnostic products, with market leadership and limited competition across key products in North America
- Demonstrated strong growth in Radiopharmaceuticals business
 - Future growth to be driven by a strong product pipeline
 - Ruby-fill, used for heart imaging, is expected to be a key growth driver



Allergy Therapy Products

- Strong market position in Allergy Therapy Products in North America
 - Among top 3 in the US for allergy immunotherapy with dominant market penetration
- US Market leader for manufacture of Stinging Insect Venom Immunotherapy Products





De-risked Business Model With Diverse Sources of Revenue

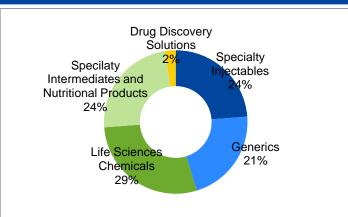
De-risked business model

- Presence in niche Specialty businesses that have high barriers to entry
- Forward integration in Pyridine related derivatives to ensure higher utilization and profitability
- Diverse end-use industry segmentation in Life Science Ingredients with focus on Pharmaceutical and Agrochemical industries

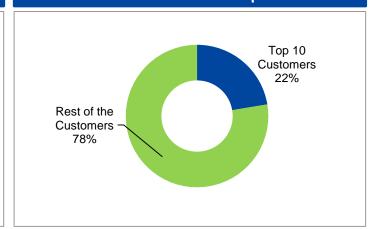
No Concentration Risk

- Business: Life Science Chemicals is the largest segment and accounts for 29% of FY18 Revenue
- Customers: Top 10 customers account for 22% of FY18 Revenue
- Products: Top 10 products account for 48% of FY18 Revenue
- Geography: 72% of FY18 Revenue from international markets with products & services reaching over 100 countries

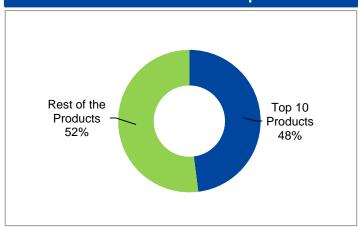
FY18 Revenue Mix by Segment



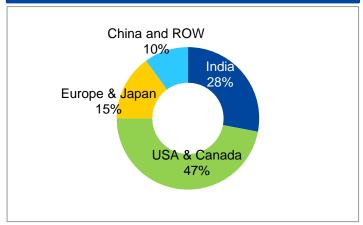
FY18 Revenue Contribution of Top 10 Customers



FY18 Revenue Contribution of Top 10 Products



FY18 Revenue Contribution by Region

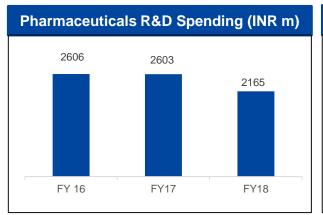


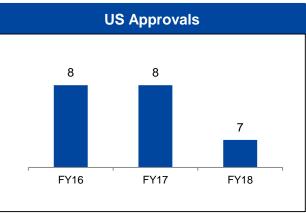




Strong Pipeline of Products With Deep R&D Capabilities

- Strong R&D capabilities demonstrated by complex and niche product filings in formulations and API
- ▶ Strong R&D support with a dedicated workforce of around 1,100 R&D professionals





Product pipeline as on March 31, 2018							
	Dosage (Orals)			Steriles			
Region	Total Approval Pending		Total Filings	Approval	Pending		
US	94	59	35	12 ⁽¹⁾	10 ⁽²⁾	2	
Canada	22	22	1	13	13	-	
Europe	103	95	8	10	10	-	
ROW	710	504	206	44	40	4	
Total	929	680	249	79 ⁽³⁾	73	6	

94 filings + 4 ANDAs for dosage (sterile)

35 ANDAs pending (oral solid) approvals + 2 pending ANDA approval for dosage (sterile)

Total 10 ANDAs filed in FY18

Total 7 ANDA approvals
received from USFDA in FY18

75 Radio Pharma filings (includes 8 filings in US / 8 approved)

41 commercial APIs
93 US DMFs filed



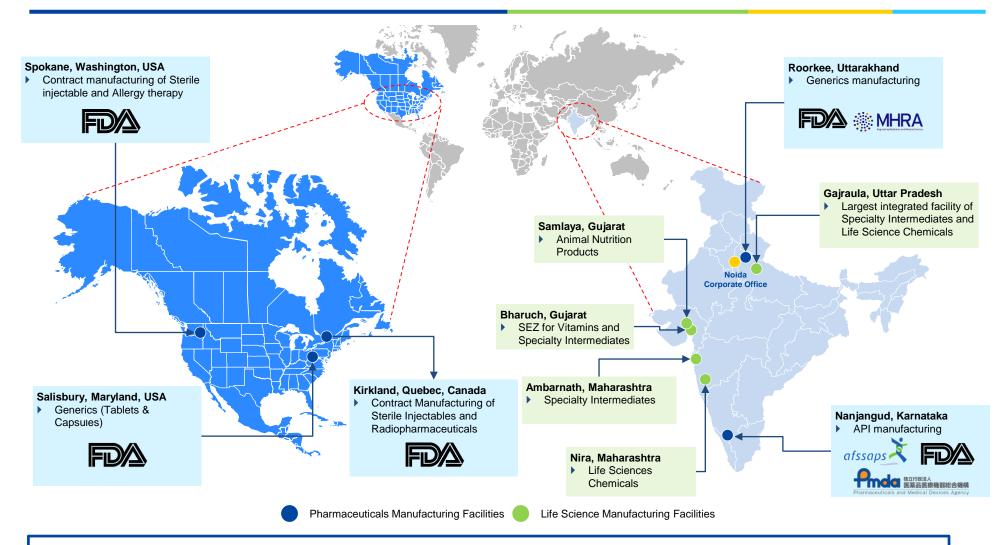
¹⁾ Includes 4 ANDA filings for dosage (sterile) and 8 radiopharma US filings

⁽²⁾ Includes 2 ANDA filings for dosage (sterile) and 8 radiopharma US filings

⁽³⁾ Includes 4 ANDA filings for dosage (sterile) and 75 radiopharma filings



High-Quality, World-Class, Low Cost Manufacturing Footprint



4 USFDA approved manufacturing facilities in North America and 2 USFDA approved manufacturing facilities in India 5 state-of-the-art Life Science Ingredients manufacturing facilities in India





Experienced Management team with high standards of corporate governance



Shyam S Bhartia Chairman 39 industry years in pharma, specialty chemicals, foods, oil and gas, aerospace and IT



Hari S Bhartia
Co-Chairman & Managing Director
33 industry years in pharma, specialty
chemicals, foods, oil and gas, aerospace
and IT



R Sankaraiah Executive Director – Finance 32 years Exp.



Dr. Raju MistryChief Human
Resources Officer
25 years Exp.



Dr. Ashutosh Agarwal *Chief Scientific Officer* 35 years Exp.



Siddhartha Pahwa Chief of Supply Chair 21 years Exp.



Ajay Khanna Chief - Strategic & Public Affairs 37 years Exp.

Pharmaceuticals

✓ IT

✓ HR



Finance

Operations

Pramod Yadav, CEO - Jubilant Pharma 29 years of Industry Experience

Life Sciences Ingredients



Rajesh Srivastava, whole-time Director and CEO – Life Sciences Ingredients
29 years of Industry Experience

Drug Discovery Solutions*



Marcel J Velterop, President -Drug Discovery Solutions 28 years of Industry Experience

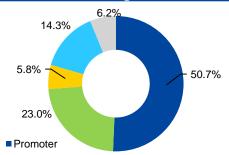
- ✓ Radiopharmaceuticals
- ✓ CMO of Sterile Injectables
- ✓ Allergy Therapy Products
- ✓ Solid Dosage Formulations
- ✓ Active Pharma Ingredients
- India Branded Pharmaceuticals

- ✓ Advanced Intermediates
- ✓ Fine Ingredients
- ✓ Crop Science Ingredients
- ✓ Nutritional Products (Human and Animal)

✓ IT

- ✓ Life Science Chemicals
- ✓ Ethanol
- ✓ Finance
- ✓ Operations
 ✓ HR
- ✓ Quality
 ✓ EHS

Shareholding Structure



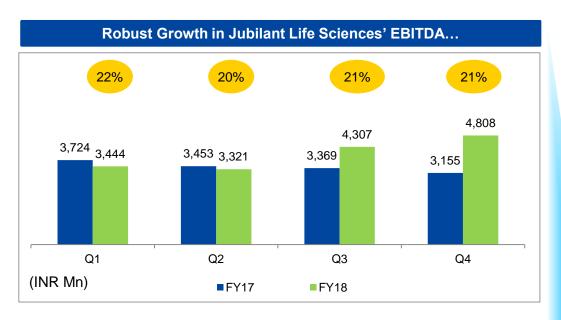
- Foreign Institutional Investors
- Financial Institutions / MFs / Banks
- ■NRI and Indian Public
- Others

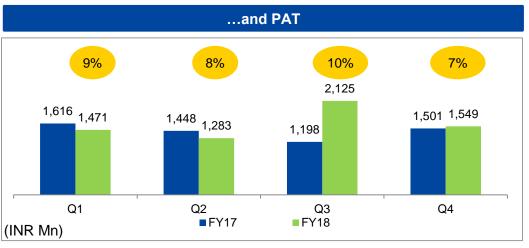
As of 31st Mar, 2018

Corporate Functions

Businesses

Performance Back on Track With Business Related Concerns Resolved





Successful Resolution of Warning Letters at Montreal and Spokane

Montreal CMO Facility

- In Feb 2013, Jubilant's manufacturing facility Jubilant HollisterStier (JHS) located at Kirkland, Quebec, Canada received a Warning Letter from the FDA
- The facility engages in contract manufacturing of injectables for large pharma companies and also for the Radiopharma business
- It was re-inspected in Sep 2013 and all issues related to the facility were successfully resolved in Feb 2014 with the warning letter closed out in Sep 2014
- Successful inspection of the facility by USFDA in December 2015 and December 2016

Spokane CMO Facility

- In Dec 2013, Jubilant's manufacturing facility JHS located at Spokane, Washington State, US received a Warning Letter from the FDA
- During H1 FY14, the contract manufacturing operations at Spokane contributed 7% to consolidated sales and 4% to consolidated EBITDA.
- ▶ The facility was re-inspected by FDA in Apr 2014 and Dec 2014
- In Jun 2015, Spokane facility was upgraded to the status of Voluntary Action Indicated (VAI) and ramp up of operations is underway
- Successful inspection of the facility by USFDA in July 2015, September 2016, November 2016 and September 2017





Pharmaceuticals Segment

Overview

	Specialty Injectables
	opoliarly injustables
Radio Pharma	 One of the leading US player involved in development, manufacturing, marketing and distribution of radiopharmaceutical products Leadership position in some of the radiopharmaceutical products with high profitability
	 Strong portfolio of differentiated products including RUBY-FILL[®] and I-131 MIBG
СМО	 Fully integrated contract manufacturer for innovator pharma companies with healthy order book Operating from 2 facilities at Spokane, USA and Montreal, Canada Broad range of capabilities including sterile liquids and lyophilized products, OCLs, biologics etc.
Allergy Therapy Products	 Provides allergy antigens, skin testing devices, and custom patient prescriptions in allergy immunotherapy area One of the top players in the US market Strong brand recall with ~100 years of experience

Generics

1

Solid Dosage Formulations

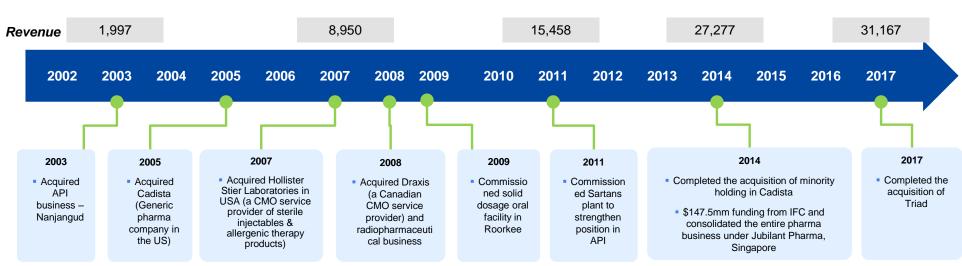
- US focused formulations player with a growing presence in Japan, Australia and emerging markets
- Focus on low competition generics
- Front-end presence in US via 100% subsidiary Cadista
- Leveraging low cost R&D out of India with strong pipeline of products

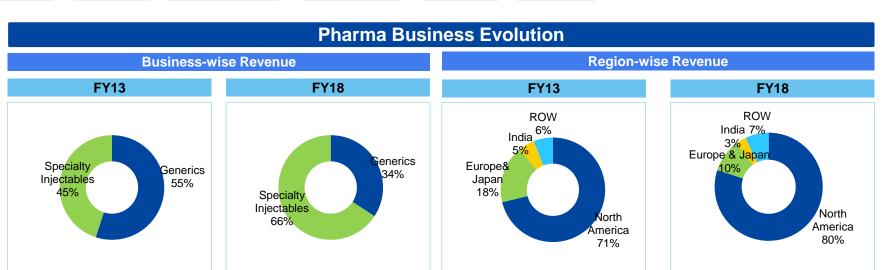
2

- Active
 Pharmaceutical
 Ingredients
- Focus on cost competitiveness and regulated markets leading to superior margins in industry
- Vertically integrated operations with formulations business
- Well positioned in some of the key products in chosen therapeutic areas of CNS, CVS and anti-infectives



Value Creation in Pharmaceutical Businesses







Overview

Pharmaceuticals

Generics

Revenue (INR m)

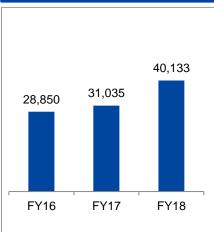
Overview

▶ Radiopharmaceuticals: North America market leadership in nuclear imaging products

Specialty Injectables

- ▶ Contract Manufacturing (CMO) of sterile injectables for large pharmaceuticals and biotech companies
- ▶ Allergy Therapy Products: Focus on diagnostic allergenic extracts with 53% US Allergy market penetration

- ▶ Solid Dosage Formulations: Strong pipeline of 94 ANDAs filed and 35 ANDAs pending approvals as on March 31, 2018
- APIs: Vertically integrated operations with API manufacturing for captive consumption as well as external sales
- Indian Branded Pharmaceuticals: Recent foray with the launch of Cardiovascular and Diabetic division



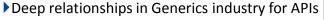
FY18 Revenue Contribution to Company





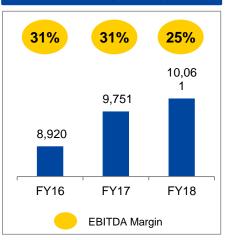
Key Highlights

- ▶ Reported YoY growth of 59% for FY18
- Expect to launch new differentiated products going forward
- Healthy order book position in the CMO business



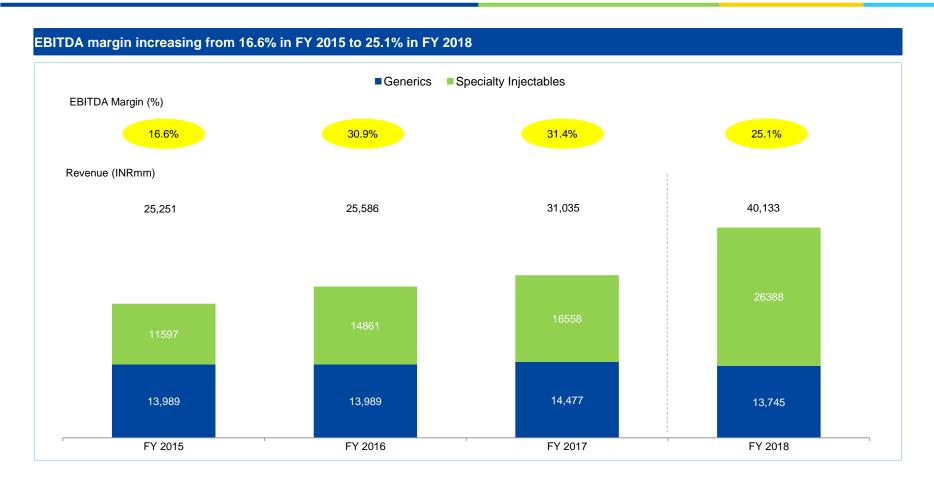
Front-end presence in US via 100% subsidiary Cadista







Pharma business has shown Strong Profitability



- Strong EBITDA growth in FY2018 due to:
 - Improved realization in RadioPharma
 - Revival of CMO of Sterile Injectables business



We have a unique US led business model which makes us standout amongst Indian pharma companies

Strength in North America

~79% of Pharma Revenues from US and Canada

- Leadership positions in products across niche businesses such as Radiopharmaceuticals/Allergy Therapy Products
- Amongst the rapidly growing and profitable Gx players
- Pharma has about 70% of asset base in North America

Strong Linkages b/w Businesses

Vertical integration gives cost advantage resulting in high margins

- APIs from the manufacturing facilities are used for dosage formulations under Gx business
- Gx R&D capabilities supporting Gx product launches of DI and Allergy
- Gx leveraging CMO for new products: Injectables and oral liquids

Diversified Low Risk Business Model Market leadership in stable, niche specialty segments with high entry

Diversified customer base, product lines and product sourcing minimizes concentration risk

barriers

Deep R&D Capabilities

Continuous innovation for new products

- Complex and niche product filings in formulations and API segments
- including 12 injectables filings in the US
- 94 ANDAs in Oral Solids filed and 59 approvals in the US as on March 31, 2018

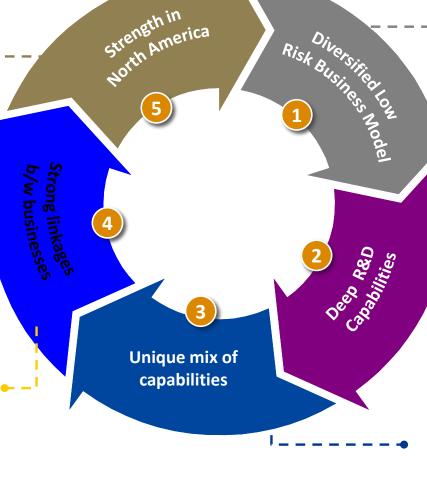
Unique mix of capabilities

Technology: Capabilities in manufacturing of API, oral solids as well as sterile products **Sales:**

North America: Relationship with retailers/ wholesalers, doctors, Radio-pharmacies & Large Pharma companies

RoW

Strong B2B partnerships in 30+ markets Front-end presence in Europe





Key Growth Drivers

New launches including differentiated products in niche businesses Strong order book in CMO business – expect good ramp up of operations and higher capacity utilization in the coming quarters Target deeper penetration in North America and accelerating growth to become a leading Allergy Therapy **Specialty** products Company in the US **Injectables** New targets, research opportunities and more collaborative programs in discovery services Strategic positioning in sterile injectables with limited competition **Pharmaceuticals** Expected shortage of quality manufacturing sites for injectables in North America given increased compliance and regulatory focus Portfolio of 94 ANDAs filed with 35 pending approvals as on March 31, 2018 Received 7 ANDA approvals in FY18 – launch of approved ANDAs to drive near term growth • Geographic expansion across Europe, Japan and Emerging Markets in Solid Dosage Formulations – portfolio of 835 filings and 621 approvals ex US as on March 31, 2018 **Generics** Deep relationships with customers ▶93 US DMFs, 42 CEPs in Europe and 39 Canadian DMFs as on March 31, 2018 Expected filings include specialized segments such as injectables and ophthalmics







Life Science Ingredients

Overview

Life Sciences Ingredients

Speciality Intermediates & Nutritional Products

▶ Specialty Intermediates

- •Global leadership in Pyridines, Picolines and derivatives
- Competitive advantage on account of lower cost due to vertical integration and process efficiencies

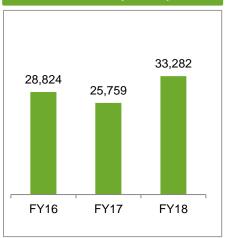
Nutritional Products

- •Key products: Vitamins B3 and B4
- Second largest producer globally in Vitamin B3

Life Sciences Chemicals

- Leadership in domestic market and significant global presence
- Key products are acetyls which include Ethyl Acetate and Acetic Anhydride
- Leadership positions in Acetic Anhydride (globally 4th largest in merchant sales) and Ethyl Acetate (globally 7th largest)

Revenue (INR m)

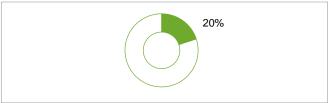


FY18 Revenue Contribution to Company

Key

Highlights

Overview



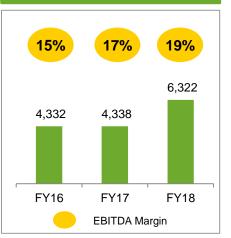
▶ Improved performance in Fine Ingredients aided by improvement in pricing and volumes

- Anti-Dumping Duty on Pyridines reduced from 24.6% to 17.6%
- Forward integration into Fine Ingredients, Crop Science Ingredients and Nutritional Products for higher value realization



- ▶ New contracts have been signed across applications
- ▶ Large capacities of Ethyl Acetate and Acetic Anhydride
- Manufacture using eco-friendly green route compared with crude used by competitors

EBITDA (INR m)





Key Growth Drivers

Specialty Intermediates and Nutritional Products

- Conversion of existing product specific plants to multi-purpose plants to meet enhanced customer demand requirements
- New Product launches in Specialty Intermediate plants
- Increased supply of Life Sciences intermediates for Pharma related applications
- Increased focus on new, non-traditional geographies for Pyridine
- Increased demand for higher value-added products in Fine Ingredients and Nutritional Products

Life Sciences Chemicals

- ▶ Higher capacity utilization to aid growth for Acetic Anhydride
- Expanding geographic reach into Key Markets such as US and Europe
- Cost advantages:
 - Efficient process
 - Volume advantage in procurement
 - •Multimodal transportation and reverse logistics to optimize costs

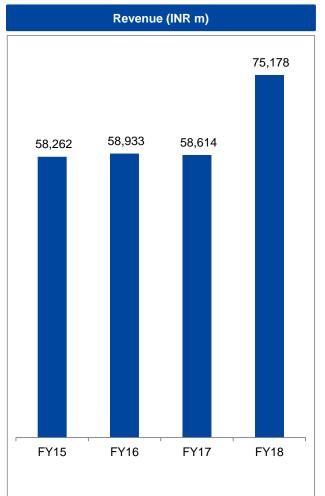


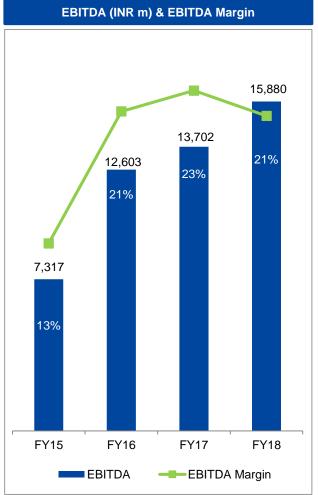


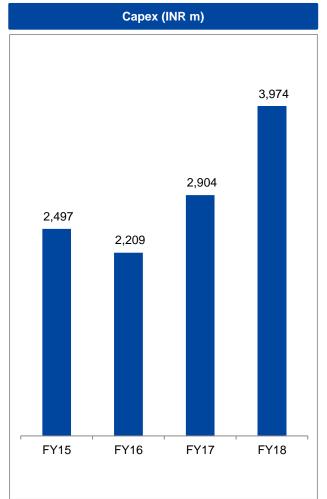


Financial Overview

Financial Overview

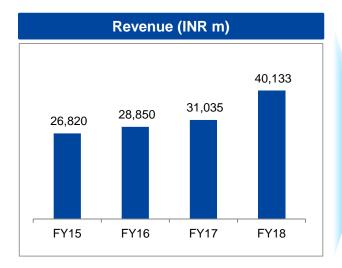


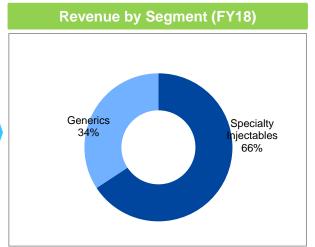


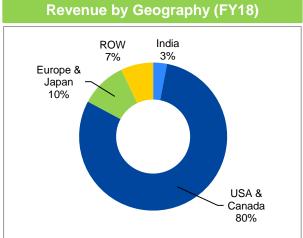


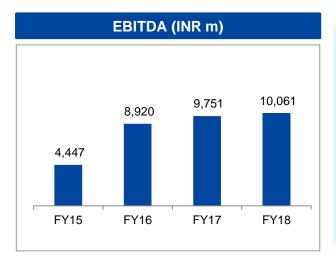


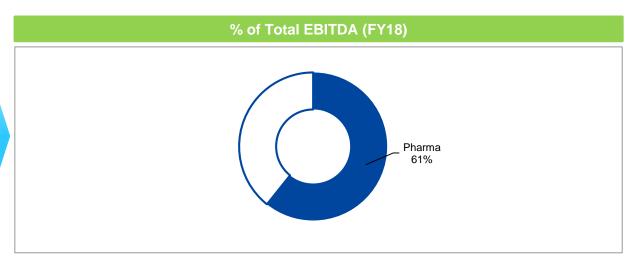
Pharma Business: Overview





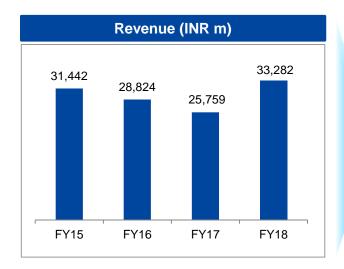


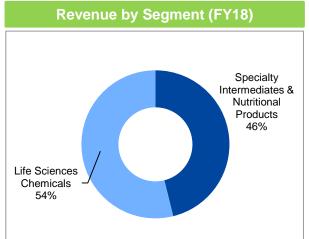


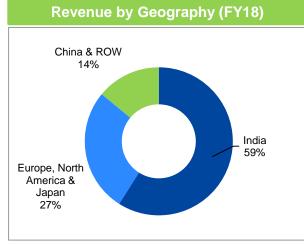


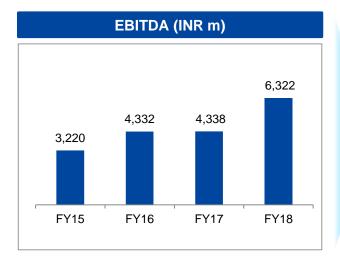


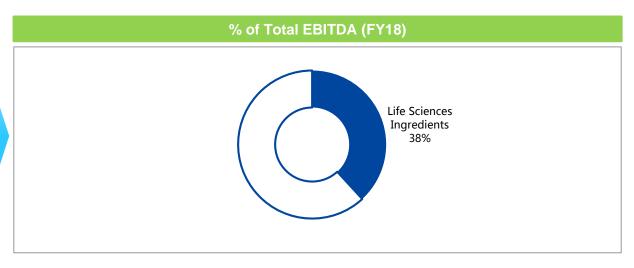
Life Sciences Ingredients: Overview





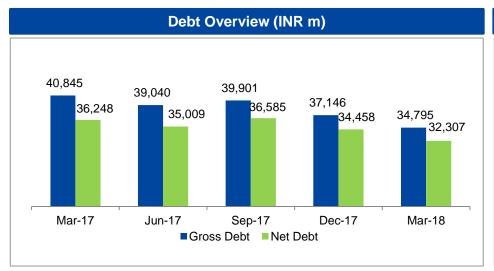


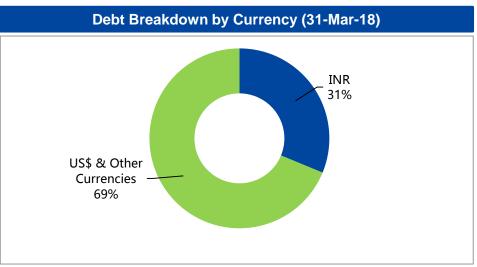


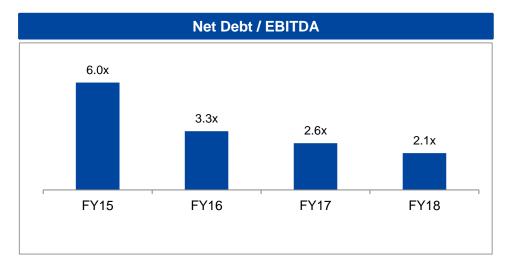




Strong Balance Sheet







- Reorganized debt portfolio in such a way that free cash flows would be adequate for scheduled loan repayments
- Prepaid all rupee loans scheduled for repayment during next two years
- Net Debt reduction of Rs. 4,059 Million in FY18 after Net Debt reduction of Rs. 5,056 Million in FY17



Q4'FY18 and FY18 Results Analysis



Income Statement – Q4'FY18

Particulars	Q4'FY17	Q3'FY18	Q4'FY18	YoY Growth	QoQ Growth
			(%)		
Total Revenue from Operations (Net of Excise)	1,598	2,068	2,252	41%	9%
Pharmaceuticals	804	1,101	1,238	54%	12%
Life Science Ingredients	743	922	968	30%	5%
Drug Discovery Solutions	51	45	47	-8%	5%
Total Expenditure	1,293	1,651	1,794	39%	9%
Other Income	11	3	23	118%	611%
EBITDA from Operations					
Pharmaceuticals	216	245	298	38%	22%
Life Science Ingredients	116	206	187	62%	-9%
Drug Discovery Solutions	6	3	10	85%	230%
Corporate (Expenses)/Income	(22)	(23)	(15)		
Total	316	431	481	52%	12%
One time expenses	0	(11)	0		
Reported EBITDA	316	420	481	52%	14%
Depreciation and Amortization	75	82	182		
Normal	75	82	91	21%	11%
One-time	0	0	91		
Finance Cost	80	77	72	-10%	-6%
Profit before Tax	160	261	227	42%	-13%
Tax Expenses (Net)	11	48	74	570%	53%
Minority Interest	(1)	0	(2)		
Net Profit After Tax and Minority Interest	150	213	155	3%	-27%
Normalised Profit after Tax	150	223	226	51%	1%
Earnings Per Share - Face Value Re. 1 (Rs.)	9.63	13.64	9.94		
Normalised Earnings Per Share	9.63	14.33	14.53		
	(%)			(bps)	
EBITDA Margins from Operations	19.7%	20.8%	21.3%	161	52
Pharmaceuticals	26.9%	22.2%	24.1%	(278)	189
Life Science Ingredients	15.6%	22.3%	19.3%	378	(299)
Drug Discovery Solutions	10.9%	7.0%	22.0%	1,112	1,504 JU
Net Margins	9.4% 30	10.3%	6.9%	(252)	(340) LIF

Financial Highlights – Q4'FY18

- Consolidated revenue at Rs. 2,252 Crore, up 41% YoY and 9% QoQ
 - Pharmaceuticals revenue at Rs. 1,238 Crore, contributing 55% to revenue, up 54% YoY and 12% QoQ
 - Life Science Ingredients revenue at Rs. 968 Crore, contributing 43% to revenue, up 30% YoY and 5% QoQ
 - Drug Discovery Solutions revenue at Rs. 47 Crore, contributing 2% to revenue, 5% QoQ
 - International revenues at Rs. 1,623 Crore, contributing 72% to revenue; growing 41% YoY
- EBITDA from Operations of Rs. 481 Crore, up 52% YoY and 12% QoQ with margins of 21.3%
 - Pharmaceuticals EBITDA of Rs. 298 Crore, up 38% YoY and 22% QoQ with margins of 24.1%
 - Life Science Ingredients EBITDA at Rs. 187 Crore; growth of 62% YoY and margins at 19.3%
 - Drug Discovery Solutions EBITDA at Rs. 10 Crore, growth of 85% YoY and 230% QoQ; margins at 22%
- Depreciation and amortization of Rs. 182 Crore, including Rs. 91 Crore one-time charge of product development expenses due to rationalization of product portfolio to reflect the current market conditions prevailing in the global generic markets, US in particular. As a matter of prudence, the company has chosen to take a charge in the P&L consistent with IND-AS which has no cash impact during the quarter
- Finance costs lower 10% YoY and 6% QoQ at Rs. 72 Crore. Finance costs include borrowing costs of Rs. 51 Crore and non-cash charge on Stock Settlement Instrument of Rs. 21 Crore
- Reported PAT at Rs. 155 Crore, with Net Margins at 6.9% and EPS of Rs. 9.94 for Re. 1 FV; Normalized PAT after adjusting for one-time charges stood at Rs. 226 Crore, up 51% YoY, with Normalized EPS of Rs. 14.53
- Net Debt reduction of Rs. 263 Crore after incurring capital Expenditure of Rs. 141 Crore



Income Statement – FY18

Particulars	FY17	FY18	YoY Growth
	(Rs Crs)	(%)	
Total Revenue from Operations (Net of Excise)	5,861	7,518	28%
Pharmaceuticals	3,104	4,013	29%
Life Science Ingredients	2,576	3,328	29%
Drug Discovery Solutions	182	176	-3%
Total Expenditure	4,516	5,999	33%
Other Income	25	40	61%
EBITDA from Operations			
Pharmaceuticals	975	1,006	3%
Life Science Ingredients	434	632	46%
Drug Discovery Solutions	26	19	-28%
Corporate (Expenses) /Income	(65)	(69)	
Total	1,370	1,588	16%
One time expenses - Acquisition related	0	(30)	
Reported EBITDA	1,370	1,558	14%
Depreciation and Amortization	291	415	
Normal	291	324	11%
One-time	0	91	
Finance Cost	341	284	-17%
Profit before Tax	738	859	16%
Tax Expenses (Net)	163	225	38%
Minority Interest	(1)	(8)	
Net Profit After Tax and Minority Interest	576	643	12%
Normalised Profit after Tax	576	744	29%
Earnings Per Share - Face Value Re. 1 (Rs.)	36.93	41.25	
Normalised Earnings Per Share	36.93	47.77	
	(%)	(%)	
EBITDA Margins from Operations	23.4%	21.1%	(225)
Pharmaceuticals	31.4%	25.1%	(635)
Life Science Ingredients	16.8%	19.0%	215
Drug Discovery Solutions	14.2%	10.5%	(371)
Net Margins	9.8%	8.6%	(127) [UBILA I

Financial Highlights – FY18

- Consolidated revenue at Rs. 7,518 Crore, up 28% YoY
 - Pharmaceuticals revenue at Rs. 4,013 Crore, up 29% YoY, contributing 53% to the revenues
 - LSI revenue at Rs. 3,328 Crore, up 29% YoY, contributing 44% to the revenues
 - Drug Discovery Solutions revenue at Rs. 176 Crore, contributing 2% to the revenues
 - International revenues at Rs. 5,417 Crore, contributing 72% to the revenues; growing 28% YoY
- EBITDA from Operations of Rs. 1,588 Crore, up 16% with margins of 21.1%
 - Pharmaceuticals EBITDA of Rs. 1,006 Crore, up 3% YoY with margins of 25.1%
 - Highest-ever Life Science Ingredients EBITDA at Rs. 632 Crore, up 46% YoY as compared to Rs. 434 Crore in FY17; margins at 19%, up from 16.8% in FY17
 - Drug Discovery Solutions segment EBITDA at Rs. 19 Crore
- Depreciation and amortization of Rs. 415 Crore, including Rs. 91 Crore one-time charge of product development expenses due to rationalization of product portfolio to reflect the current market conditions prevailing in the global generic markets, US in particular. As a matter of prudence, the company has chosen to take a charge in the P&L consistent with IND-AS which has no cash impact during the year
- Finance costs lower 17% YoY at Rs. 284 Crore. Finance costs include borrowing costs of Rs. 220 Crore and non-cash charge on Stock Settlement Instrument of Rs. 64 Crore
- Reported PAT at Rs. 643 Crore, growth of 12% YoY with Net Margins at 8.6% and EPS of Rs. 41.25 for Re. 1 FV; Normalized PAT after adjusting for one-time charge on account of product rationalization and acquisition costs at Rs. 744 Crore, up 29% YoY; Normalized EPS of Rs. 47.77
 - Net Debt reduction of Rs. 406 Crore after Capital Expenditure of Rs. 397 Crore and Triad acquisition



Segmental Revenue Analysis – FY18

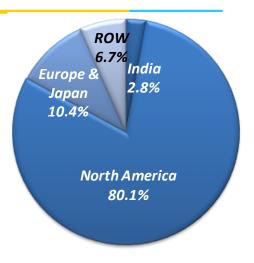
Segmental Revenue Analysis	Revenue (Rs. Crs.)		Revenue Mix (%)	YoY Growth %	
	FY17	FY18	1411X (70)	CIOWEII /0	
Pharmaceuticals	3,104	4,013	53%	29%	
Generics	1,448	1,375	18%	-5%	
Specialty Injectables	1,656	2,639	35%	59%	
Life Science Ingredients	2,576	3,328	44%	29%	
Specialty Intermediates and Nutritional Products	1,365	1,535	20%	12%	
Specialty Intermediates	905	970	13%	7%	
Nutritional Products	460	565	8%	23%	
Life Science Chemicals	1,211	1,793	24%	48%	
Drug Discovery Solutions	182	176	2%	-3%	
Income from Operations (excl. IDTs)	5,861	7,518	100%	28%	
Inter Divisional Sales (IDTs)	529	525	7%	-1%	

- Company reports highest-ever revenue of Rs. 7,518 Crore, driven by improvement in Pharmaceuticals and Life Science Ingredients
- Pharmaceuticals segment reports revenue of Rs. 4,013 Crore, 29% YoY growth on account of healthy organic growth and acquisition of Triad, which contributed Rs. 767 Crore to Revenues
- LSI segment posts income of Rs 3,328 Crore, up 29% YoY and contributes 44% to total revenue driven by both volume and price



Pharmaceuticals Segment Highlights

Geo-wise Revenue (Rs crs)	Q4'FY17	Q3'FY18	Q4'FY18	Mix %	YoY %	QoQ %
International	767	1,067	1,203	97%	57%	13%
North America	580	915	992	80%	71%	8%
Europe & Japan	130	85	129	10%	-1%	51%
ROW	57	66	82	7%	44%	24%
India	37	34	35	3%	-6%	3%
Income from Operations	804	1,101	1,238	100%	54%	12%



Geo-wise Revenue (Rs crs)	FY17	FY18	Mix %	YoY %
International	2,923	3,878	97%	33%
North America	2,150	3,189	79%	48%
Europe & Japan	466	402	10%	-14%
ROW	307	286	7%	-7%
India	181	135	3%	-25%
Income from Operations	3,104	4,013	100%	29%





Pharmaceuticals Segment Highlights

- Pharmaceuticals revenue at Rs. 4,013 Crore, up 29% YoY, contributing 53% to the revenues
- Specialty Injectables reported revenues of Rs. 2,639 Crore, showing robust growth of 59% YoY,
 contributing 66% to Pharma segment sales as compared to 53% in FY17
- Revenues in Generics business stood at Rs. 1,375 Crore, contributing 34% to the segment sales
 - Successful USFDA inspections of our CMO Spokane, Radiopharmaceuticals and APIs facilities
 - Steady ramp-up of Rubyfill installations in the US market
 - Received approval from Health Canada for Rubyfill Elution System
 - Drax Exametazime approval received and product launched in the USA
 - Received USFDA approval for new pulmonary indication in DraxImage DTPA
 - Ramp up of CMO business underway with order book of US\$ 702 Million and addition of four new customers
 - Signed long term contract for supply of animal health product



Regulatory Status

Regulatory Agency	Cadista USA	Roorkee India	CMO / Allergy Spokane	CMO Montreal	JDI Montreal Canada	Nanjangud India	•	All sites successfully inspected and fully compliant with USFDA
(USA)	Mar 2017	Mar 2017	Sep 2017	Dec 2016	Sep 2017	Oct 2017		regulations
Health Canada (Canada)				Oct 2017	Apr 2016	Oct 2017	•	Use the experience from multiple Agency
(Japan)		Dec 2015	Feb 2017			May 2016		inspections to enhance compliance status of all
(India SLA / CDSCO)		Sep 2015				Sep 2016		sites
(Brazil)				May – June 2016		Mar 2015	•	World class quality control practices
TC. Sağlık Bakanlığı (Turkey)			Mar 2015				•	Global quality control
Cofepris Contain Fabrica on 1- Prescribe (Mexico)						Aug 2015		function reporting to the Corporate Board

Portfolio of R&D products – Filings and Approvals

Product pipeline as on March 31, 2018										
Dosage (Orals)			Steriles							
Region	Total Filings	Approval	proval Pending		Approval	Pending				
US	94	59	35	12	10	2				
Canada	22	22	-	13	13	-				
Europe	103	95	8	10	10	-				
ROW	710	504	206	44	40	4				
Total	929	680	249	79	73	6				

We have a total of 1,008 filings across geographies

- 929 filings in Dosage (Orals)
- > 79 filings in sterile (injectables & ophthalmics)
- ➤ 680 Dosage (Orals) and 73 injectables have been approved
- 255 filings (249 Dosage (Orals) and 6 sterile products) are pending approval



Portfolio of R&D products – Filings and Approvals

Dosage (Orals)

- Filed 94 ANDAs in the US
 - i. 59 ANDAs have been approved and 35 ANDAs are pending approval
 - ii. Filed 10 ANDAs in FY18 (in-house) + 1 Inlicensed product
- Made 835 filings in ROW (ex-US) markets including Canada, Europe and Japan
 - i. 621 filings have been approved and 214 filings are pending approval
- In-licensing of two products in the US market, 1 is approved

Injectable and Others

- Total 4 ANDAs filed and approvals for 2 have been received
 - 1 ANDA for injectable was filed in FY18

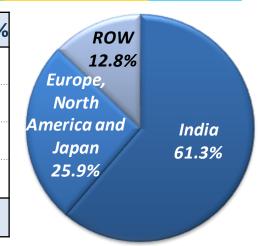
Radiopharmaceuticals

- Filing status as on March 31, 2018:
 - 8 approved registrations in the US
 - 13 registrations in Canada which are all approved
 - 10 registrations in Europe of which are all approved
 - In ROW, we have a total of 44 registrations/licenses, of which 4 are pending for approval
- There are 5 products under development



LSI Segment Highlights

Geo-wise Revenue (Rs crs)	Q4'FY17	Q3'FY18	Q4'FY18	Mix %	YoY %	QoQ %
International	335	371	374	39%	12%	1%
Europe, North America and Japan	227	243	250	26%	10%	3%
ROW	108	128	124	13%	15%	-3%
India	408	551	593	61%	45%	8%
Income from Operations	743	922	968	100%	30%	5%



Geo-wise Revenue (Rs crs)	FY17	FY18	Mix %	YoY %
International	1,146	1,366	41%	19%
Europe, North America and Japan	802	884	27%	10%
ROW	343	482	14%	41%
India	1,430	1,962	59%	37%
Income from Operations	2,576	3,328	100%	29%

ROW, 14.5%
Europe,
North India,
America and
Japan, 26.6%

Witnessed growth across all key geographies



LSI Segment Highlights

FY18 revenue at Rs. 3,328 Crore, up 29% YoY, contributing 44% to the revenues

Specialty Intermediates:

- Revenues at Rs. 970 Crore, up 7% YoY
- Launched 7 products during the year and 6 more to be launched in FY19
- Commissioned cGMP Plant in Bharuch for supplies to pharma and agro-intermediates
- Some De-bottlenecking initiatives completed and underway to meet additional demand

Life Science Chemicals:

- Revenues at Rs. 1,793 Crore, up 48% YoY
- The capacity enhancement project of Acetic Anhydride is progressing as per plan to achieve target completion in FY'19. This will provide additional revenue of Rs. 300 Crore at full capacity
- 4th largest ethanol supplier in the government blending program. Won a contract to generate revenues of Rs. 300 Crore from Dec'17 to Nov'18
- Higher availability of molasses with lower prices due to good sugar production

Nutritional Products:

Revenues at Rs. 565 Crore, up 23% YoY



Drug Discovery Solutions Segment Highlights

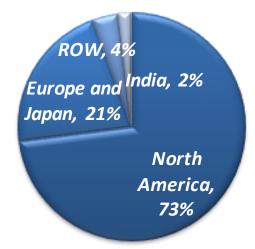
Revenue at Rs. 176 Crore during FY18

- Pipeline of Integrated Drug Discovery Projects, functional projects & FTE business continues to be strong
- Increased effort & focus on in-house proprietary research for out-licensing molecules or onboarding new projects

Geo-wise Revenue (Rs crs)	Q4'FY17	Q3'FY18	Q4'FY18	Mix %	YoY %	QoQ %
International	49	43	46	98%	-5%	6%
North America	37	27	39	83%	4%	44%
Europe & Japan	11	16	3	7%	-68%	-78%
ROW	1	1	4	8%	625%	345%
India	2	1	1	2%	-67%	-29%
Income from Operations	51	45	47	100%	-8%	5%

ROW 7.9% Europe & India Japan 7.4%	
North America 82.9%	

Geo-wise Revenue (Rs crs)	FY17	FY18	Mix %	YoY %
International	178	173	98%	-3%
North America	131	129	73%	-2%
Europe and Japan	44	38	21%	-14%
ROW	3	7	4%	112%
India	4	3	2%	-7%
Income from Operations	182	176	100%	-3%



Expenditure Analysis – Q4'FY18

Expenses (Rs Crs)	Q4'FY17	% of Sales	Q3'FY18	% of Sales	Q4'FY18	% of Sales	YoY Growth %	QoQ Growth %
Material Cost and Change in Inventory	548	34%	707	34%	817	36%	49%	16%
Purchases of stock-in-trade	42	3%	7 9	4%	73	3%	73%	-8%
Employee benefits expense	324	20%	423	20%	454	20%	40%	7%
Power and fuel expense	93	6%	120	6%	112	5%	20%	-7%
Others	285	18%	322	16%	338	15%	18%	5%
Total Expenses (Excluding Excise Duty)	1,293	81%	1,651	80%	1,794	80%	39%	9%
Depreciation and Amortization	75	5%	82	4%	182	8%	142%	122%
Total Finance Costs	80	5%	77	4%	72	3%	-10%	-6%
Borrowing costs	54	3%	57	3%	51	2%	-5%	-10%
Stock Settlement Instrument Charge	26	2%	20	1%	21	1%	-19%	5%

- Material Costs as percentage of sales increase due to increase in revenues in LSI
- Power & Fuel as percentage of sales stable
- Employee benefits expenses and Other Expenses higher on account of acquisition of Triad in addition to annual increase in Employee Benefits
- Total Finance costs lower 10% YoY. Borrowing costs lower by 5% YoY. Stock settlement instrument is a non-cash debit to P&L, on account of convertible instrument issued to IFC of US\$ 58.2 Million as a mandatory conversion option at IPO of JPL

Expenditure Analysis – FY18

Expenses (Rs Crs)	FY17	% of Sales	FY18	% of Sales	YoY Growth %
Material Cost and Change in Inventory	1,814	31%	2,626	35%	45%
Purchases of stock-in-trade	185	3%	243	3%	31%
Employee benefits expense	1,231	21%	1,556	21%	26%
Power and fuel expense	334	6%	425	6%	27%
Others	952	16%	1,150	15%	21%
Total Expenses (Excluding Excise Duty)	4,516	77%	5,999	80%	33%
Depreciation and Amortization	291	5%	415	6%	42%
Total Finance Costs	341	6%	284	4%	-17%
Borrowing costs	287	5%	220	3%	-23%
Stock Settlement Instrument Charge	54	1%	64	1%	19%

- Material Costs as percentage of sales increase due to increase in revenues in LSI
- Power & Fuel as percentage of sales stable
- Employee benefits expenses and Other Expenses stable as percentage to sales
- Total Finance costs lower 17% YoY. Borrowing costs lower by 23% YoY. Stock settlement instrument is a non-cash debit to P&L, on account of convertible instrument issued to IFC of US\$ 58.2 Million as a mandatory conversion option at IPO of JPL



Debt Profile

Particulars	31-Mar-17	31-Dec-17	31-Mar-18
Foreign Currency Loans	(\$ Mn)	(\$ Mn)	(\$ Mn)
Standalone	19	0	9
Subsidiaries (incl. Stock Settlement Instrument)	407	374	358
Total	426	374	367
Rupee Loans	(Rs. Crs)	(Rs. Crs)	(Rs. Crs)
Standalone	1,241	1,227	1,007
Subsidiaries	82	101	7 9
Total	1,323	1,327	1,086
Gross Debt	(Rs. Crs)	(Rs. Crs)	(Rs. Crs)
Standalone	1,361	1,227	1,066
Subsidiaries	2,723	2,488	2,414
Total	4,084	3,715	3,480
Cash & Equivalent	460	269	249
Net Debt	3,625	3,446	3,231
Change in debt on account of exchange rate difference from 31-March, 2017	0	36	-12
Net Debt - Adjusted for foreign exchange difference	3,625	3,482	3,219
Reduction in Net Debt from March,2017 Adjusted for Exchange Diff.		-143	-406
Closing Exchange Rate (Rs./USD)	64.85	63.88	65.17

- Net debt at Rs. 3,219 Crore compared to Rs. 3,625 Crore on 31-Mar-17 on constant currency basis
 - Debt reduction of Rs. 263 Crore during the quarter and Rs. 406 Crore for FY18
- Average blended interest rate for FY18 at 5.97% pa Re loans @ 8.16%, \$ loans @ 4.71%
- Stock Settlement Instrument (IFC) of US\$ 58.2 Million at 10% per annum discount to Jubilant Pharital Limited IPO price

Outlook

- ➤ We see a clear roadmap for growth in our businesses both in revenues and profitability in FY19, with H2'FY19 expected to be better than H1'FY19. The growth drivers for each of the business are as follows:
 - Specialty Injectables:
 - Growth from new products and execution of existing contracts in Radiopharma
 - Full year impact of Triad business in our operations with break-even profitability
 - Healthy order book and new customer additions, supported by higher production and new capacities, to deliver better results in CMO
 - Higher sales of existing products and new capacities in Allergy Therapy products
 - Generics: Higher volumes from new product launches and new markets
 - Life Science Ingredients: Better demand for existing products, new capacities from ongoing investments and de-bottlenecking initiatives, and launch of new products to augment growth
- ➤To meet the increased demand in our businesses, we plan to invest about Rs. 550 Crore in capital expenditure in FY19. In addition, we plan to invest Rs. 300 Crore in R&D during the year, including Rs. 150 Crore in Product Development expenditure. We will continue our efforts to strengthen balance sheet by reducing debt and improving financial ratios



Our Vision & Promise

OUR VISION

To acquire and maintain global leadership position in chosen areas of businesses
 To continuously create new opportunities for growth in our strategic businesses
 To be among the top 10 most admired companies to work for
 To continuously achieve a return on invested capital of at least 10 points higher than the cost of capital

OUR PROMISE

Caring, Sharing, Growing

We will, with utmost care for the environment and society, continue to enhance value for our customers by providing innovative products and economically efficient solutions; and for our stakeholders through growth, cost effectiveness and wise investment of resources

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