

Natco Pharma Limited

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June 3, 2017

Corporate Relationship Department The Bombay Stock Exchange Limited Dalal Sreet, Fort Mumbai 400 001 Manager – Listing M/s. National Stock Exchange of India Ltd "Exchange Plaza", Bandra – Kurla Complex Bandra (E) <u>Mumbai 400 051</u>

Scrip Code: 524816 Scrip Code: NATCOPHARM

Dear Sir

Please find enclose herewith the corporate presentation.

Thanking you

For NATCO Pharma Limited

M. Adinarayana

Company Secretary &

Vice President (Legal & Corp. Affairs)



CORPORATE PRESENTATION

June 2017



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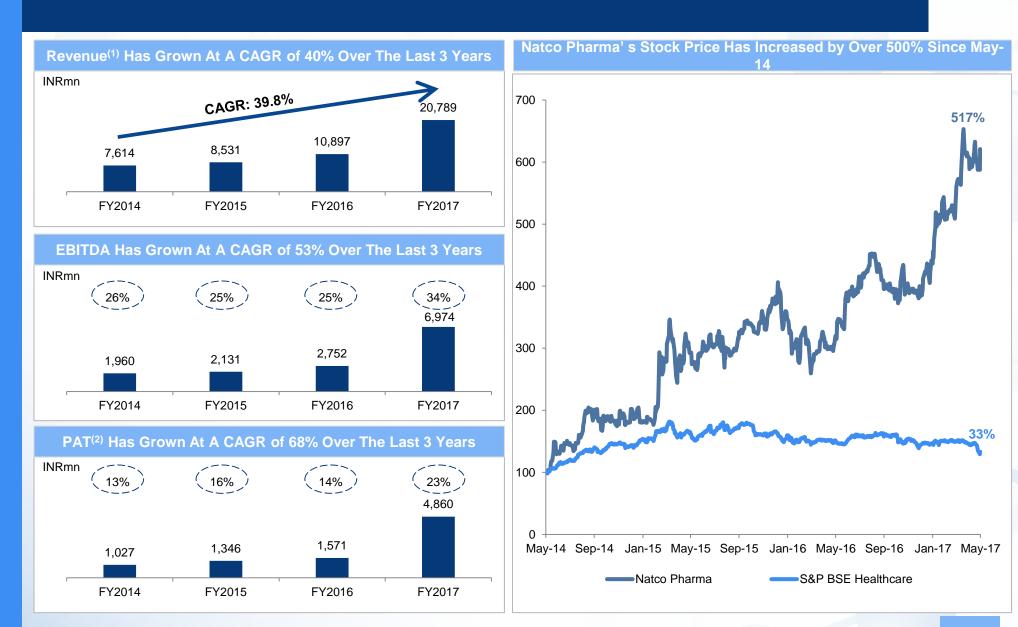
Natco Pharma at a Glance



- Vertically integrated pharmaceutical company with well diversified businesses across geographies India, US and RoW
- Leading position in the domestic Oncology and Gastro Hepatology segments
 - Portfolio of market leading brands catering to various oncology diseases including breast, brain, bone, lung and ovarian cancer
 - Amongst the first company's in India to launch the generic version of Gilead's Sovaldi (Sofosbuvir) and its combinations under its brands Hepcinat and Hepcinat LP for the treatment of Hep-C ('Hepatitis C') in India
- Focused on complex generics for the US markets with a portfolio of 43 niche ANDA filings including 20 Para IV filings⁽¹⁾
 - First Company to launch the generic version of Roche's Tamiflu in partnership with Alvogen
- Strong impetus on R&D with about 350 employees dedicated to R&D
- Reported revenues⁽²⁾ of INR 20,789mn for the financial year ended 31st March 2017
- Listed on the Bombay Stock Exchange and National Stock Exchange with a market capitalization⁽³⁾ of USD 2.5bn
- Incorporated in 1981 and headquartered in Hyderabad with over 4,000 employees across all locations

Track Record of Consistent Growth and Value Creation





March 2014 are as per Indian GAAP Margins

Represents consolidated gross revenue and includes other income Represents PAT after minority interest

Key Business Segments





	Formu	lations	API	Othors	
	Domestic International (Domes		(Domestic & Exports)	Others	
Overview	 Leading player in the domestic oncology and Gastro Hepatology segment with a portfolio of market leading brands Recent foray into the Cardiology and Diabetology segments Specialist sales force of over 350 personnel and over 500 distributors 	 Focused on complex generics for the US Front end partnerships with leading global generic pharma companies 43 niche ANDA filings including 20 Para IV filings Emerging presence in Asia, Europe, Canada, Brazil and other RoW markets 	 Strategically important division Vertical integration for its FDF portfolio Filed 37 DMFs in US with over 10 products under development Exports focused on the US, Europe and Brazil 	 Operations in Brazil, Canada, Singapore and Australia Selective contract manufacturing business and other operating income 	
FY17 Revenue (INRmn)	8,810 *	8,370	1,838	1,771	
FY17 Revenue Contribution	42%	40%	9%	9%	
Growth FY17 over FY16	39%	262%	13%	9%	

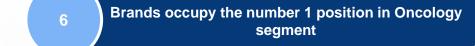
Strong Growth in Domestic Formulations Business





INRmn



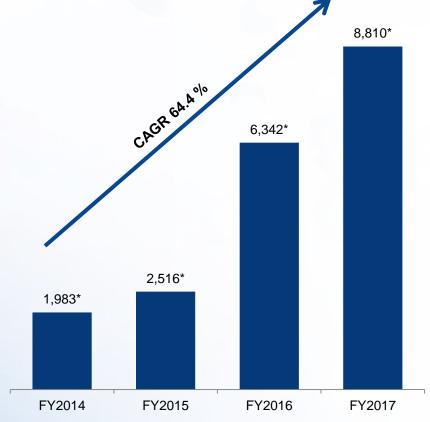








Among the first companies to launch generic Hep-C range of products



Leading Position in Domestic Oncology Segment



Oncology Division Overview

- Entered the segment with launch of generic version of Imatinib in 2003
- Portfolio of market leading brands 6 brands occupy the number 1 position
- Progressively widened its oncology product range from 6 in 2003-04 to
 28*
- Sales and marketing of the product is supported by over 75 sales representatives and strategically located logistics network of over 500 distributors
- #1 Domestic Company in Oncology

Oncology Portfolio

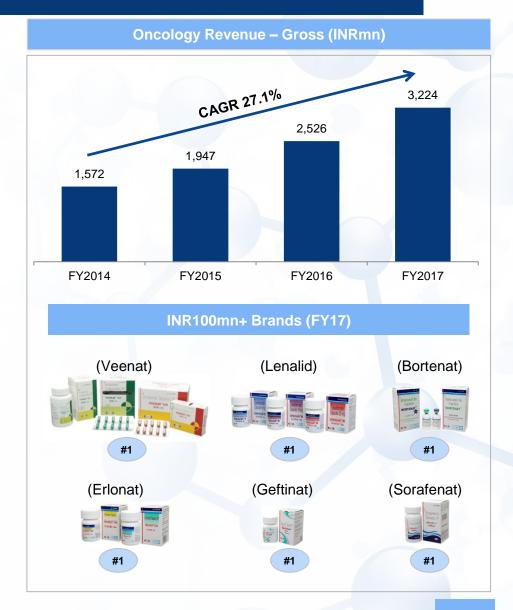
Hematology

Solid Tumors

of Active Brands*







Gastro Hepatology – Leading Market Position in Hep-C

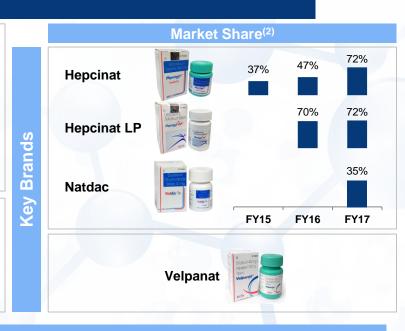


Market Leading Hep-C **Franchise**

- First Company to launch generic Sofosbuvir and its combinations for the treatment of Hep-C in India & Nepal under its brand Hepcinat & Hepcinat LP
- Non-exclusive licensing agreement with Gilead Sciences for 101 countries including India reaching a target population of 103 million people
- Launched generic Daclatasvir in India under its brand Natdac
- Market leading positions across the Hep-C class of drugs in India
- Sales and marketing of the product is supported by over 120 sales representatives

Extending the Hep-C **Franchise**

- Launched an oral fixed-dose combination of Sofosbuvir and Velapatasvir under its brand Velpanat
- Medicine is highly efficacious across all genotypes 1 to 6
- Limited competition with only one other competitor in the domestic market
- Aggressively foraying into RoW markets



Expanding Into Emerging Markets Of Asia And Africa

countries



Demonstrating Strong Revenue Growth (INRmn)



Expanding Domestic Presence with Launch of New CnD Division



Cardiology and Diabetology

- Launched Cardiology and Diabetology (CnD) division in early 2017
- Focus will be on niche molecules with high barriers to entry



US Market - Focus on Complex Generics



US FDF product portfolio is predominantly focused on high-barrier-to-entry products that are typically characterised by one or more of the following:

- Intricate chemistry
- Challenging delivery mechanism
- Difficult or complex manufacturing process
- May face complex legal and regulatory challenges

	Overview of Key Filings					
	Key Brand	Molecule	Therapeutic Segment / Indication	Dosage Form	Para IV Para III	Market Size (US\$mn)#
	Copaxone 20&40mg	Glatiramer 20&40mg	Multiple Sclerosis	PFS	✓	4,138.4
	Tamiflu	Oseltamivir Suspension	Influenza Infection	Suspension	√	213.1
ne	Vidaza	Azacitidine	Myelodysplastic syndrome	Injection	✓	203.1
Pipeline	Fosrenol	Lanthanum Carbonate	End stage renal disease	Tablets	√	119.0
	Revlimid*	Lenalidomide	Multiple Myloma	Capsules	✓	4,416.9
	Tracleer*	Bosentan	Hypertension	Tablets	✓	367.5
	Nexavar*	Sorafenib	Liver, Kidney Cancer	Tablets	✓	330.7

- Pipeline of niche and complex generics products in US
- 43 niche ANDA filings including 20 Para IV filings⁽¹⁾
- 22 approved ANDAs (1 yet to be launched)

De-risked Business Model through Partnerships with Global Pharmaceutical Players



Mitigation Strategy

US Market reach and Regulatory Challenges

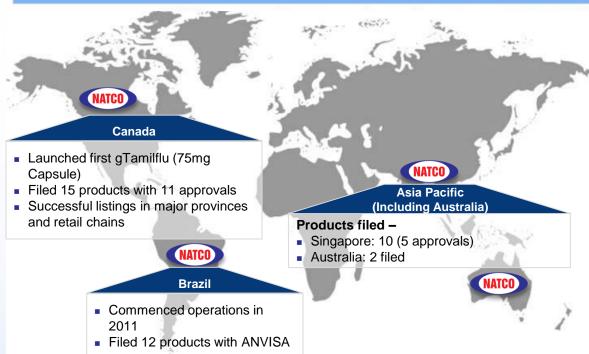
- Adopted and successfully implemented partnership strategy for international formulations product
 - Has product specific partnerships with global generic players at different stages of a potential ANDA filing
 - Entered into de-risked arrangements with marketing partner with the partner responsible for the litigation and regulatory process to secure the ANDA approval
 - Profit sharing arrangements with the front end partners to ensure Natco participates in the up-side. Recent success with the launch of generic Tamiflu capsules in partnership with Alvogen & Doxorubicin Liposomal Injection with Dr. Reddy's Laboratories Ltd.



Expanding RoW Presence



RoW growth to be driven by Hep-C franchise expansion and continued business in LatAm and other Asian countries



Hep-C driven markets

- Filed in over 40 countries
- Received approvals and import permits for over 10 countries

Europe

- Sell our products in Eastern Europe, UK and Germany
- 4 approvals
- Distribution arrangements with our business partner

Venezuela

Sell our FDF products (oncology) to third parties

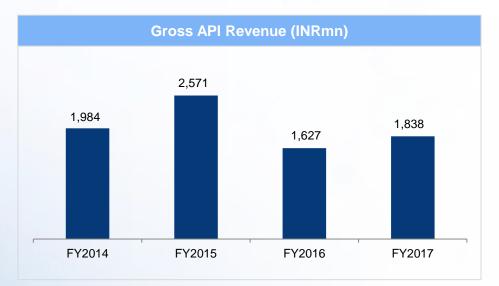
Strong In-House API Development with Vertical Integration for Key Formulation Products

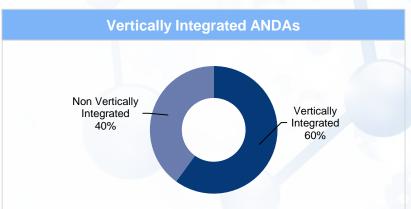


- Strategically important business develops APIs primarily for captive consumption of its FDF portfolio as well as third party sales
- Portfolio of 37 US DMFs with over 10 products under development (as of 31-March-2017)
- Focuses on complex molecules in oncology and CNS segments
 - Other therapeutic areas of focus includes Anti-asthmatic, Anti-depressant, Anti-migraine, Anti-osteoporosis and G I Disorders
- Exports are focused on the US, EU, Canada, Latin America and South-East Asia
- Vertical integration for several APIs a key competitive advantage

API Strengths

- Complex multi-step synthesis & scale-up
- Semi-synthetic fusion technologies
 - Fermentation / Biotech / Synthetic / Separation technologies
- Containment / High potency APIs
- Peptide (Solid phase) pharmaceuticals





Expansion plans to significantly augment API manufacturing capacity

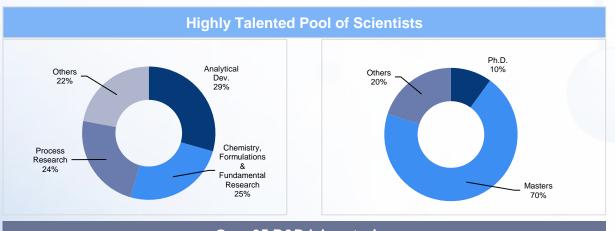
Strong Research & Development Capabilities



Strong R&D capabilities demonstrated by its complex and niche product filings in formulations and API segments

 Two well equipped research facilities with capabilities across synthetic chemistry, biotech & fermentation, nano pharmaceuticals, new drug discovery & cell biology

R&D Expense (INRmn) and as % of Standalone Revenue 6.4% 7.1% 6.7% 6.1% 1,235 FY2014 FY2015 FY2016 FY2017



22 ANDAs Approved
(1 vet to be launched)

20 Para IV Filings⁽¹⁾

37 US DMFs Filed(1)

Over 10 API products Under Development



189 International Patents Filed⁽¹⁾

133 International Patents
Granted

185 Indian Patents Filed(1)

83 Indian Patents Granted

Commitment to Manufacturing Excellence with a Culture of Quality and Compliance



cility

International Markets Formulations

Kothur Facility

Visakhapatnam



Capability

Regulatory

Approvals

- Tablets, Capsules, Pellets, Injectables
- GMP, USFDA, German Health Authority, ANVISA
- Other Highlights
- US FDA January 2017

- Cytotoxic & other Oral Solid Dosages
- na
- Targeted towards US & other International regulated markets
- Located in a Special
 Economic Zone (SEZ)

API Manufacturing Facilities

Mekaguda Facility	Chennai Fa
 GMP, USFDA, German Health 	

Key Regulatory Approvals

Last Audit

US FDA – January 2015 (Last approval)

(Mexico)

- GMP
- US FDA August 2016 (Last approval)

Guwahati Facility

Expansion plans to significantly augment API manufacturing capacity

Authority, PMDA (Japan), Cofepris

Focus on renewable energy to reduce manufacturing cost

Domestic Market Formulations

Nagarjuna Sagar Facility



Ampoules, Vials, Lyophilized vials,

Parenterals, Sterile Dry Powders

Dehradun Unit 6 Facility



• Tablets, Capsules, Injectables

Dehradun Unit 7 Facility



Key Regulatory Approvals

Capability

GMP

GMP

 GMP, Public Health Service of the Netherlands (EU GMP

Tablets, Capsules

GMP Compliant Facility

Tablets, Capsules



Natco's Near and Long-Term Goals



Domestic Branded Formulations

- Maintain leadership position in Oncology and Gastro Hepatology segment
- Launch 8- 10 new products
- Traction in the newly launched CnD division
- Extension of the Hep-C franchise via the launch of Velpanat
- Target 20% plus YoY growth

Complex Generics

- Near term approval for some key molecules –
 TAD for generic Copaxone in June 2017
- Also anticipating approval for Lanthanum Carbonate, Azacitidine, Bosentan
- Extension of the Tamiflu franchise via launch of the suspension version
- Profit share component via the launch of the Doxorubicin Liposomal injection

Enter new attractive segments

Key molecules currently filed in the market

2020+

2017

Experienced Management





Mr. V.C Nannapaneni Chairman and Director



Over 42 years of experience in the Pharmaceutical Industry



Mr. Rajeev Nannapaneni Vice Chairman & CEO

Holds bachelors degree in Quantitative Economics and History from Tufts University, Boston, USA

Holds wide experience and exposure in General Management and Product Development



Dr. A.K.S Bhujanga Rao President (R&D and Technical)

- Awarded Ph.D.in Synthetic Organic Chemistry from the Indian Institute of Science (IISc), Bangalore
- Wide expertise in technology transfer to commercial scale, quality control regulatory affairs and Patents



Dr. Linga RaoPresident (Technical Affairs)

- Holds Masters degree in Science (Applied Chemistry) & Ph.D in Chemistry from JNTU, Hyderabad
- Over 35 years of experience in the pharmaceutical industry and has been working with Natco for over 21 years



Mr. P.S.R.K Prasad Executive Vice President

- Holds B.E. Mech. Engg. from Andhra University, Visakhapatnam
- Responsible for looking after the general administration, engineering, regulatory, training, environmental matters, safety, health, production and maintenance activities of the Company



M. AdinarayanaCompany Secretary &
VP-Legal & Corporate Affairs

- Bachelors in Commerce and Bachelors in Law from Andhra University, Fellow Member of Institute of Company Secretaries
 of India
- 22+ years of experience within the Company in legal, secretarial and patent litigation areas



Mr. S.V.V.N.Appa Rao

- Over 25 years of experience including 20 years within the Company covering areas of accounting, financial controller, treasury
- Responsible for finance and treasury functions at the Company

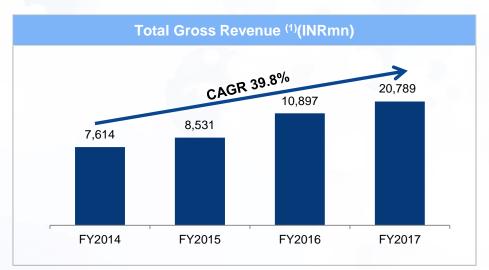


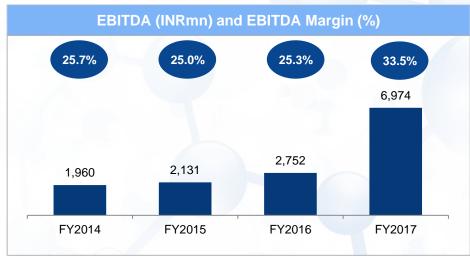
Mr. Rajesh Chebiyam Vice President - Business Development & Corp Support

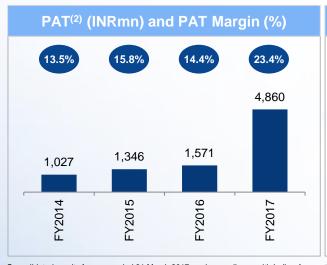
- Holds MBA from Babson College (USA) and Masters degree in Chemical Engineering from University of Rhode Island
- 20+ years of experience across supply chain, operations, business development, sales and strategy

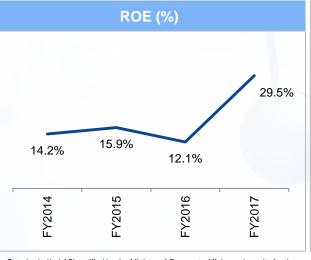
Demonstrated Track Record of Topline and Earnings Growth

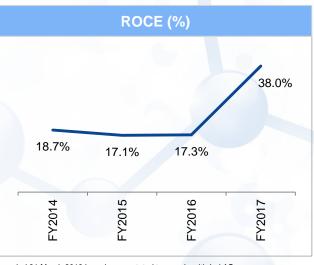












Consolidated results for year ended 31 March 2017 are in compliance with Indian Accounting Standards (Ind AS) notified by the Ministry of Corporate Affairs and results for the year ended 31 March 2016 have been restated to comply with Ind AS. Results for the financial years ended 31 March 2015 and 31 March 2014 are as per Indian GAAP

⁽¹⁾ Represents consolidated gross revenue and includes other income

⁽²⁾ Represents PAT after minority interest

Historical Financials



Consolidated Profit & Loss Statement (INRmn)

Particulars	31-Mar-17	31-Mar-16
Revenue from operations (gross)	20,650	10,804
Less : Excise duty		
Revenue from operations (net)	20,650	10,804
Other income	139	94
Total revenue	20,789	10,897
Expenses		
Cost of material consumed	5,208	3,037
Purchase of stock in trade	971	152
Change in Inventory	(188)	(483)
Employee benefits	2,432	1,798
Finance costs	185	229
Depreciation	544	508
Other expenses	5,393	3,641
Prior period expenses	0	C
Total expenses	14,545	8,882
Profit before exceptional items and tax	6,244	2,015
Exceptional item	0	C
Profit before tax	6,244	2,015
Current Tax	1,393	441
Deferred Tax Benefit	1	38
PAT (Before Minority interest)	4,849	1,536
Profit from discontinued operations, net of tax		22
Minority Interest	(11)	(13)
PAT (After Minority interest)	4,860	1,571

Consolidated Balance Sheet (INRmn)

Particulars

Particulars	31-Mar-17	31-Mar-16
Equity share capital	349	348
Other equity	16,144	12,609
Non-controlling interest	40	49
Total of Equity	16,533	13,007
Financial Liabilities		
Borrowings	-	-
Other financial liabilities	8	8
Employee benefit obligations	219	125
Deferred tax liabilities	150	146
Total Non-current liabilities	377	279
Financial liabilities		
Borrowings	2,216	984
Trade payables	2,627	2,755
Other financial liabilities	1,014	815
Current tax liabilities, net	133	34
Other current liabilities	258	327
Employee benefit obligations	18	15
Total Non-current liabilities	6,265	4,929
Total equity and liabilities	23,175	18,215
Property, plant and equipment	8,272	7,046
Capital work-in-progress	3,362	2,118
Other intangible assets	58	55
Investments	1	1
Other financial assets	131	106
Other non-current assets	478	521
Total Non-current assets	12,302	9,847
Inventories	3,489	3,573
Financial Assets		
Investments	321	221
Trade receivables	4,751	2,616
Cash and cash equivalents	235	242
Other bank balances	123	210
Loans	34	28
Other financial assets	752	770
Income tax asset	-	34
Other current assets	1,166	676
Total current assets	10,873	8,368
Total assets	23,175	18,215

Historical Financials (Cont'd)



Segment Breakdown (INRmn)

Revenue Division (INRmn)	31-Mar-17
API, Domestic	512
API, Exports	1,325
API Gross Revenue	1,838
Formulations, Exports	3,947
Profit sharing Income	4,423
Formulations Onco	3,224
Formulations, Brand Pharma Non Onco	4,802
Formulations, 3rd party, & miscel	784
Formulations Gross Revenue	17,180
Service Income-Exports	(89)
Service Income-Domestic	0
Service Income	(89)
Other Operating & Non-Operating Income	1,208
Adjustments as per IndAS	21
Stand-Alone Total Gross Revenue	20,159
Total Revenue, all subsidiaries	630
TOTAL Gross Revenue	20,789