

Natco Pharma Limited Regd. Off. : NATCO HOUSE', Road No. 2, Banjara Hills, Hyderabad - 500034.

Telangana, INDIA. Tel : +91 40 23547532, Fax : +91 40 23548243 CIN : L24230TG1981PLC003201, www.natcopharma.co.in

August 9, 2018

Corporate Relationship Department M/s. BSE Ltd Dalal Sreet, Fort <u>Mumbai 400 001</u> 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 enclosed herewith the Investor Presentation for the Quarter ended June 30, 2018.

Thanking You.

For NATCO Pharma Limited

MANarayana

M. Adinarayana Company Secretary & Vice President (Legal & Corp. Affairs)



INVESTOR PRESENTATION

Q1 FY19, August 2018

Disclaimer / Important Disclosure

NATCO

THIS PRESENTATION (PRESENTATION) IS NOT AN OFFER TO SELL ANY SECURITIES OR A SOLICITATION TO BUY ANY SECURITIES OF NATCO PHARMA LIMITED OR ITS SUBSIDIARIES OR JOINT VENTURES (TOGETHER, THE "COMPANY").

The material that follows is a Presentation of general background information about the Company's activities as at the date of the Presentation or as otherwise indicated. It is information given in summary form and does not purport to be complete and it cannot be guaranteed that such information is true and accurate. This Presentation has been prepared by and is the sole responsibility of the Company. By accessing this Presentation, you are agreeing to be bound by the trading restrictions. It is for general information purposes only and should not be considered as a recommendation that any investor should subscribe / purchase the Company shares.

This Presentation includes statements that are, or may be deemed to be, "forward-looking statements". These forward-looking statements can be identified by the use of forward-looking terminology, including the terms "believes", "estimates", "anticipates", "projects", "expects", "intends", "may", "will", "seeks" or "should" or, in each case, their negative or other variations or comparable terminology, or by discussions of strategy, plans, aims, objectives, goals, future events or intentions. These forward-looking statements include all matters that are not historical facts. They appear in a number of places throughout this Presentation and include statements regarding the Company's intentions, beliefs or current expectations concerning, amongst other things, its results or operations, financial condition, liquidity, prospects, growth, strategies and the industry in which the Company operates.

By their nature, forward-looking statements involve risks and uncertainties because they relate to events and depend on circumstances that may or may not occur in the future. Forward-looking statements are not guarantees of future performance including those relating to general business plans and strategy of the Company, its future outlook and growth prospects, and future developments in its businesses and its competitive and regulatory environment. No representation, warranty or undertaking, express or implied, is made or assurance given that such statements, views, projections or forecasts, if any, are correct or that the objectives of the Company will be achieved. There are some important factors that could cause material differences to Company's actual results. These include (i) our ability to successfully implement our strategy (ii) our growth and expansion plans (iii) changes in regulatory norms applicable to the Company (iv) technological changes (v) investment and business income (vi) cash flow projections etc. (vii) exposure to market as well as other risks.

The Company, as such, makes no representation or warranty, express or implied, as to, and does not accept any responsibility or liability with respect to, the fairness, accuracy, completeness or correctness of any information or opinions contained herein. The information contained in this Presentation, unless otherwise specified is only current as of the date of this Presentation. The Company assumes no responsibility to publicly amend, modify or revise any forward looking statements, on the basis of any subsequent development, information or events, or otherwise. Unless otherwise stated in this Presentation, the information contained herein is based on management information and estimates.

Any opinions expressed in this presentation are subject to change without notice. The presentation should not be construed as legal, tax, investment or other advice. None of the Company or any of its affiliates, advisers or representatives accepts any liability whatsoever for any loss howsoever arising from any information presented or contained in this presentation. The information contained in this presentation has not been independently verified. Furthermore, no person is authorized to give any information or make any representation which is not contained in, or is inconsistent with, this presentation. Any such extraneous or inconsistent information or representation, if given or made, should not be relied upon as having been authorized by or on behalf of the Company. Further, past performance is not necessarily indicative of future results.

This document is just a Presentation for information purposes and private circulation only and is not intended to be a "prospectus" or "offer document" or a "private placement offer letter" (as defined or referred to, as the case may be, under the Companies Act, 2013). It is clarified that this Presentation is not intended to be a document offering for subscription or sale of any securities or inviting offers from the Indian public (including any section thereof) or from persons residing in any other jurisdiction including the United States for the subscription to or sale of any securities including the equity shares of the Company or any of its subsidiaries. No part of it should form the basis of or be relied upon in connection with any investment decision or any contract or commitment to purchase or subscribe for any securities. None of the Company's securities may be offered or sold in the United States without registration under the U.S. Securities Act of 1933, as amended, except pursuant to an exemption from registration there from.

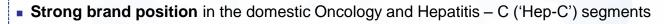
This document has not been and will not be reviewed or approved by a regulatory authority in India or by any stock exchange in India. This presentation is confidential and this presentation or any part thereof should not be used or relied upon by any other party or for any other purpose and should not be copied, reproduced, recirculated, redistributed, published in any media, website or otherwise, in any form or manner, in part or as a whole, without the express consent in writing of the Company. Any unauthorized use, disclosure or public dissemination of information contained herein is prohibited. The distribution of this presentation in certain jurisdictions may be restricted by law. Accordingly, any persons in possession of the aforesaid should inform themselves about and observe any such restrictions.

Natco Pharma at a Glance





Vertically integrated pharmaceutical company with presence across geographies – India, US and RoW



- Portfolio of brands catering to various oncology diseases including breast, brain, bone, lung and ovarian cancer
- Launched the generic version of Gilead's Sovaldi (Sofosbuvir) and its combinations for the treatment of Hep-C in India



Focused on complex generics for the US markets with niche Para IV and Para III filings



Strong focus on R&D with over 400 employees dedicated to R&D⁽³⁾



Total revenues⁽¹⁾ of INR 22,424 mn for the financial year ended 31st March 2018



• Listed on the BSE and NSE with a market capitalization⁽²⁾ of **USD 2.14bn**



Incorporated in 1981 and headquartered in Hyderabad with over 4,830 employees across all locations⁽³⁾

Track Record of Consistent Growth



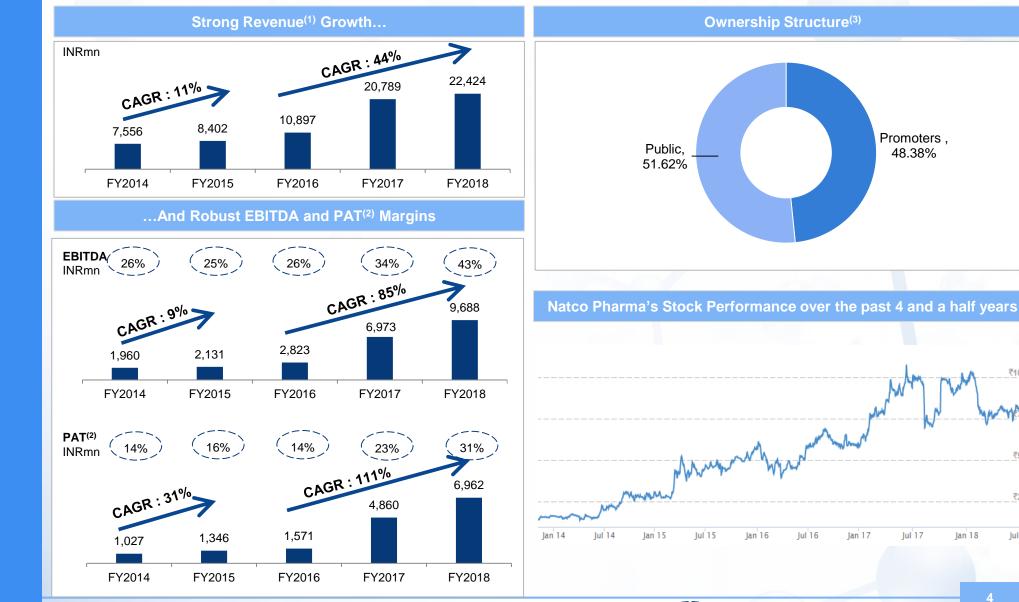
Promoters,

48.38%

Jul 17

#

Margins



FY2014 and FY2015 numbers have been prepared under IGAAP, whereas FY2016, FY2017 and FY2018 numbers have been prepared under Ind AS Source: BSE, as of 31 March, 2018 (3) Represents consolidated gross revenue and includes other income (1)

Represents PAT after minority interest (2)

Jan 18

₹1000

₹500

₹250

Jul 18

Key Business Segments





	Formu	llations	API	Others	
	Domestic	International	(Domestic & Exports)	Others	
Overview	 Strong brand position in the domestic oncology and Hepatitis-C segments Recent foray into the Cardiology and Diabetology segments Specialist sales force of over 350 personnel and over 400 distributors Fall in FY18 revenue due to decline in HepC market size. Expect growth with target launch of 8-10 molecules per year 	 Focused on complex generics for the US Front end partnerships with leading global generic pharma companies Niche Para IV and Para III filings Emerging presence in Asia, Europe and developing markets 	 Strategically important division Vertical integration for its Finished Dosage Formulation ('FDF') portfolio Filed 42 DMFs in the US with niche products under development Exports focused on Europe and emerging markets 	 Operations in Brazil, Canada, Singapore and Australia Selective contract manufacturing business and other operating income 	
FY18 Revenue (INRmn)	7,202*	10,418	2,854	1,950	
FY18 Revenue Contribution	32%	47%	13%	9%	
Growth FY18 over FY17	- 18%	25%	55%	10%	

* Includes third party sales

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

Key Products in Pipeline

	Key Brand	Molecule	Therapeutic Segment / Indication	Para IV
	Gilenya	Fingolimod	Multiple Sclerosis	\checkmark
	Treanda	Bendamustine	Cancer, CLL	\checkmark
	Nexavar	Sorafenib	Liver, Kidney Cancer	✓
	Tracleer	Bosentan	Hypertension	Para III
ed	Revlimid ⁽¹⁾	Lenalidomide	Multiple Myeloma	✓
Be Launched	Afinitor	Everolimus (higher strength)	Kidney Cancer	\checkmark
	Zytiga	Abiraterone	Prostate Cancer	\checkmark
٩	Tarceva	Erlotinib	NSCLC, Pancreatic Cancer	~
	Kyprolis	Carfilzomib	Multiple Myeloma	✓
	Aubagio	Teriflunomide	Multiple Sclerosis	✓
	Eliquis	Apixaban	Anticoagulant	✓
	Pomalyst	Pomalidomide	Multiple Myeloma	\checkmark
	Sovaldi	Sofosbuvir	Anti-Viral / Hep C	✓

Low Risk Business Model through Partnerships with Global Pharmaceutical Players

- Adopted and successfully implemented partnership strategy for international formulation products
 - Has product specific partnerships with global generic players at different stages of a potential ANDA filing
 - Low risk business model:
 - Marketing partner typically responsible for the litigation and regulatory process to secure the ANDA approval
 - Multi-site approvals
 - Multi-sourcing arrangements
 - Profit sharing arrangements with the front end

partners.

- Pipeline of niche and complex generics products in US
- 29 approved ANDAs⁽²⁾
- 16 Para IVs yet to be launched ⁽²⁾

(2) As of March 31, 2018. Approval received either by Natco or its marketing partner

Strong Growth in Domestic Formulations Business



Therapeutic

Domestic Formulation Sales⁽¹⁾: Market Leading Growth



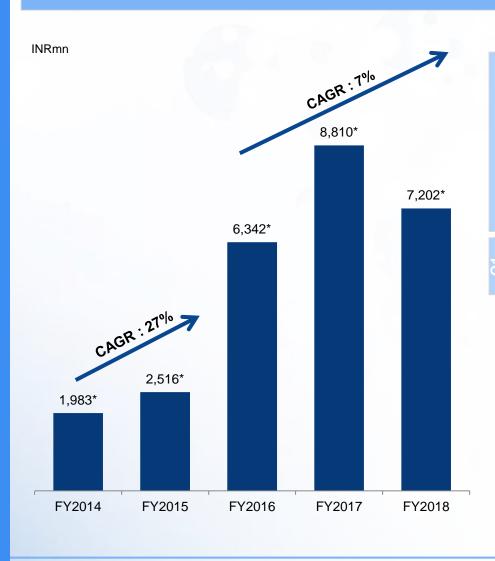
Dosage Form

Molecule

Brand

6

350-



				Segment
	Alphalan	Melphalan	Injection	Oncology
18	Carfilnat	Carfilzomib	Injection	Oncology
	Pomalid	Pomalidomide	Capsules	Oncology
	Tafnat	Tenofovir Alafenamide	Tablets	Speciality Pharma
F	Velpanat	Sofosbuvir + Velpatasvir	Tablets	Speciality Pharma
	Arganat	Argatroban	Injection	Cardiology
	Dabigat	Dabigatran Etexilate Mesylate	Capsules	Cardiology
- 6	Denopsy	Teriflunomide	Tablets	Speciality Pharma
З Ш	Posanat	Posaconazole	Injection	Speciality Pharma

Strong position in Oncology and Hepatitis-C domains

Brands in excess of INR 100mn+ sales in Oncology segment

> Brands occupy the #1 position in Hep-C segment

Sales force in India across Oncology, Gastro Hepatology, Cardiology and Diabetology

4

FY2014 and FY2015 numbers have been prepared under IGAAP, whereas FY2016, FY2017 and FY2018 numbers have been prepared under Ind AS

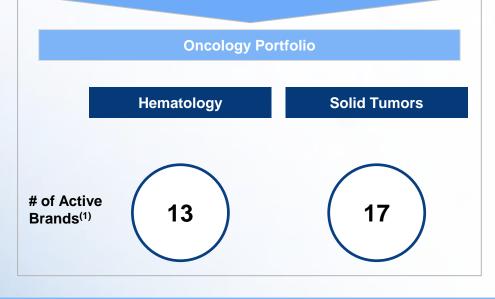
(1) Represents gross revenue
 * Includes third party sales

Strong Market Position in Domestic Oncology Segment

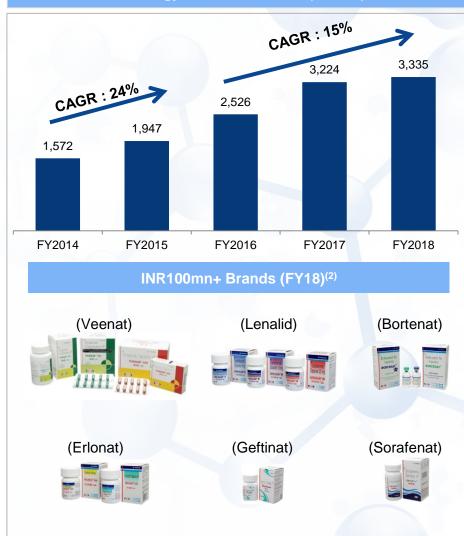


Oncology Division Overview

- Entered the segment with launch of generic version of Imatinib in 2003
- Portfolio of well recognized brands 6 brands with INR 100mn+ sales in the oncology segment
- Progressively widened its oncology product range from 6 in 2003-04 to 30⁽¹⁾
- Sales and marketing of the product is supported by approximately 70 sales representatives and strategically located logistics network of distributors



Oncology Revenue – Gross (INRmn)



FY2014 and FY2015 numbers have been prepared under IGAAP, whereas FY2016, FY2017 and FY2018 numbers have been prepared under Ind AS (1) As on 31 March, 2018 (2) Source: Company

Pharma Specialties



Gastro Hepatology – Leading Market Position in Hep-C Portfolio	 Launched 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 105 countries including India Launched generic Daclatasvir in India under its brand Natdac and an oral fixed-dose combination of Sofosbuvir and Velapatasvir under its brand Velpanat Market leading positions across the Hep-C class of drugs in India Sales and marketing of the product is supported by approximately 120 sales representatives
the Hep-C first t	Ached Hepcinat Plus, a generic fixed dose bination of Sofosbuvir and Daclatasvir for the time in India ying into RoW markets Emerging portfolio of Hep B - Launched Tenofovir Alafenamide tablets under its brand Tafnat as an extension to existing Hep B portfolio of Tenofovir (Teravir) & Entecavir (X-Vir) tablets

Expanding Into Emerging Markets Of Asia And Africa

- Import Permits & Approvals Received
- Access via Agreement with Gilead

Expanding Domestic Presence with CnD Division



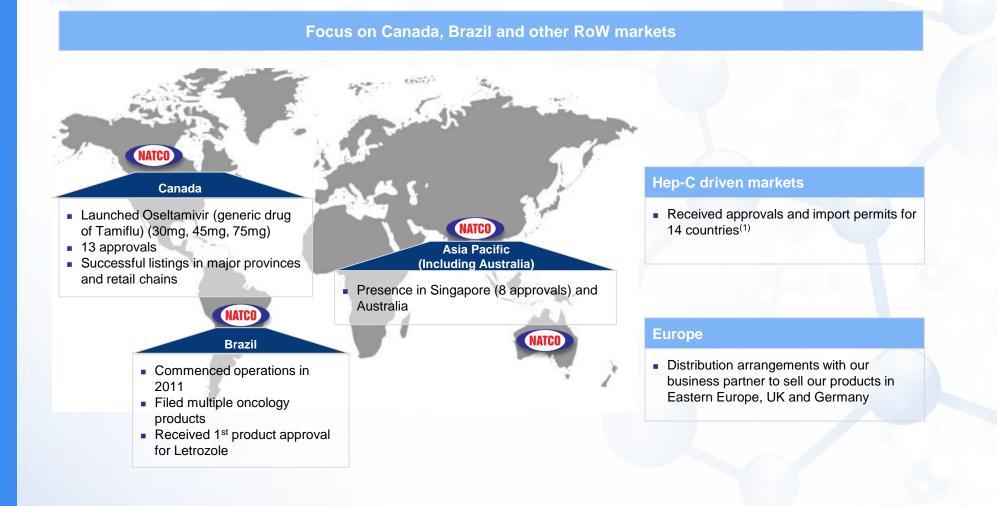


- Launched Cardiology and Diabetology (CnD) division in early 2017
- First to launch Argatroban injection and Dabigatran in India for treatment of patients with thrombosis syndrome
- Focus will be on niche molecules with high barriers to entry



Expanding RoW Presence





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 42 US DMFs⁽¹⁾ with with niche products under development
- Focuses on complex molecules in oncology and CNS segments
 - Other therapeutic areas of focus includes Anti-asthmatic, Antidepressant, 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

	Mekaguda Facility	Chennai Facility
Chemistry Skills	 Complex chemistry peptides 	 Cytotoxic API's and Biotechnology based products Synthetic chemistry
Key Regulatory Approvals	 GMP, USFDA, German Health Authority, PMDA (Japan), Cofepris (Mexico) 	 GMP, USFDA
Last US FDA Audit	 US FDA audit with Zero observations completed in February 2018 	 US FDA audit – EIR Received August 2016

Expansion plans to augment API manufacturing capacity

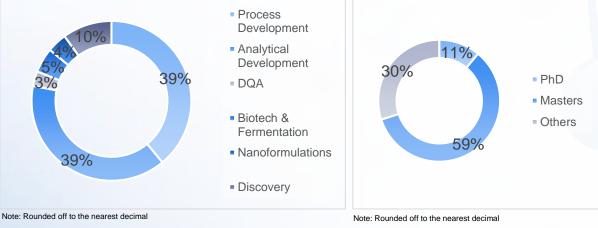
Research & Development Capabilities



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

Two 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 6.7% 7.8% 7.1% 6.0% 6.4% 1,665 1,216 703 517 407 FY2018 FY2014 FY2015 FY2016 FY2017 Talented Pool of Scientists (Total no: 271)⁽¹⁾



Over 35 R&D laboratories in 2 research facilities

FY2014 and FY2015 numbers have been prepared under IGAAP, whereas FY2016 and FY2017 numbers have been prepared under Ind AS As of March 31, 2018.

As of March 31, 2018. Approval received either by Natco or its marketing partner (2)



#

% of standalone revenues

Several International and Indian patents filed and granted

29 ANDAs Approved⁽²⁾

16 Para IVs to be Launched⁽²⁾

42 US DMFs Filed⁽²⁾

13

Commitment to Manufacturing Excellence with a Culture of Quality and Compliance



	International Markets Formulations						
	Kothur	Facility	Visakhapatnam Facility				
Capability	Tablets, Capsules, Pellets, Injectables		Cytotoxic & other Oral Solid Dosages				
Key Regulatory Approvals	 GMP, USFDA, German Health Authority, 	ANVISA	• na				
Other Highlights	 US FDA audit – EIR Received July 2017 		 Targeted towards US & other International regulated markets Located in a Special Economic Zone (SEZ) Expect to be ready in 2018 				
		Domestic Market Forr	nulations				
	Nagarjuna Sagar Facility	Dehradun Unit 6 Facility	Dehradun Unit 7 Facility	Guwahati Facility			
		50					

	And a state of the	2 Kanna Al		
Capability	 Ampoules, Vials, Lyophilized vials, Parenterals, Sterile Dry Powders 	 Tablets, Capsules, Injectables 	 Tablets, Capsules 	 Tablets, Capsules
Key Regulatory Approvals	GMP	GMP	 GMP, Public Health Service of the Netherlands (EU GMP) 	 GMP Compliant Facility

DESIDE FULLING

Experienced Management



	Mr. V.C Nannapaneni Chairman and Managing Director	 Holds Masters degree in Pharmaceutical Administration from the Long Island University, USA Over 4 decades of experience in the pharmaceutical industry
R.	Mr. Rajeev Nannapaneni Vice Chairman & CEO	 Holds bachelors degree in Quantitative Economics and History from Tufts University, Boston, USA Has 15 years of experience in the pharmaceutical industry
	Mr. P.S.R.K Prasad Executive Vice President (Corp. Engineering Services)	 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
Q	Dr. Linga Rao President (Technical Affairs)	 Holds Masters degree in Science (Applied Chemistry) & Ph.D in Chemistry from JNTU, Hyderabad Over 4 decades of experience in the pharmaceutical industry and has been working with Natco for over 23 years
	M. Adinarayana Company Secretary & VP-Legal & Corporate Affairs	 Bachelors in Commerce and Bachelors in Law from Andhra University, Fellow Member of Institute of Company Secretaries of Inda 24+ years of experience within the Company in legal, secretarial and patent litigation areas
	Mr. S.V.V.N.Appa Rao CFO	 Over 27 years of experience including 22 years within the Company covering areas of accounting, financial controller, treasury Responsible for finance and treasury functions at the Company
	Dr. Pulla Reddy M Executive Vice President - R&D	 Holds Masters in Science (Chemistry) and Ph.D in Chemistry, both from University of Hyderabad. Did postdoctoral research for 2.5 years at University of Zurich, Switzerland 24 years experience at Natco with key role in developing novel commercially viable processes for over 100 APIs and intermediates
R	Dr. Rami Reddy B Director - Formulations	 Holds M. Pharm and Ph.D. (Pharmaceutics) degree from Nagpur University 32 years of experience in the Pharmaceutical Formulation industry. Responsible for Formulation plant operations, Product development and Regulatory compliance
	Mr. Rajesh Chebiyam Vice President - Acquisitions, Institutional Investor Mgmt. & Corporate Communications	 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

Natco's Near and Long-Term Goals



Domestic Branded Formulations	าร
-------------------------------	----

Near-term Strategies

Long-term

Strategies

- Maintain leadership position in Oncology and Hepatitis-C segment
- Intensify the focus of CnD pipeline for niche launches
- Launch 8-10 new products

- Enter new attractive segments
- Growth through inorganic strategies

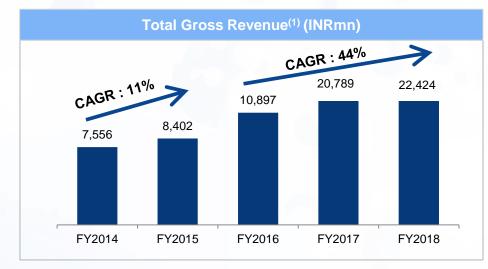
- **Complex Generics & Export Markets**
 - Focus on growth in key subsidiaries of Canada & Brazil
 - Intensify regulatory filings rate in RoW markets led by Hep-C portfolio

- Focus on a select few high-potential filings, predominantly differentiated products through either Novel Drug Delivery Systems (NDDS) or complex chemistries
- Strategic alliances in RoW markets for further growth

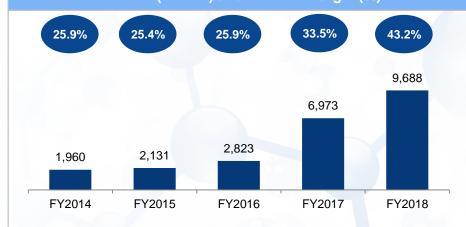
Demonstrated Track Record of Topline and Earnings Growth

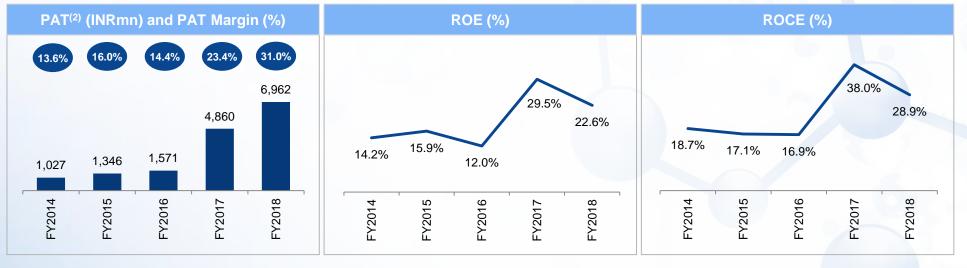


FY2014 and FY2015 numbers have been prepared under IGAAP, whereas FY2016 and FY2017 numbers have been prepared under Ind AS



EBITDA (INRmn) and EBITDA Margin (%)





(1) Represents consolidated gross revenue and includes other income

(2) Represents PAT after minority interest

Historical Financials



Consolidated Profit & Loss Statement (INRmn)

Particulars	31-March-2018	31-March-2017
Revenue		
Revenue from operations (Refer note 5)	22,020	20,650
Other income	404	139
Fotal revenues	22,424	20,789
Expenses		
Cost of materials consumed	3,905	5,208
Excise duty (Refer note 5)	172	448
Purchases of stock-in-trade	459	971
Changes in inventories of finished goods, stock-in -trade and work-in- progress	(253)	(188)
Employee benefits expense	3,256	2,432
Finance costs	154	185
Depreciation and amortisation expense	662	544
Other expenses	5,197	4,945
Total expenses	13,552	14,545
Profit before tax	8,872	6,244
Tax expense / (credit)		
Current tax	2,199	1,354
Deferred tax	-	1
Minimum alternative tax credit	(279)	-
Tax for earlier years		40
Profit after tax	6,952	4,849
Other comprehensive income (net of taxes)		
Items that will not be reclassified to profit or loss		
Re-measurement gains/(losses) on defined benefit plans	(42)	(50)
Net (loss)/gain on FVTOCI equity securities	(2)	28
Items that will be reclassified to profit or loss		
Exchange differences on translation of foreign operations	(8)	(12)
Total comprehensive income	6,900	4,815
Profit attributable to:		
Owners of the parent	6,962	4,860
Non-controlling interests	(10)	(11)
Total comprehensive income attributable to:		
Owners of the parent	6,910	4,826
Non-controlling interests	(10)	(11)
Paid-up equity share capital of ₹2 each	369	349
Other equity (Revaluation reserve ₹Nil)	30,353	16,144
Earnings per share (non-annualised)		
Basic (in ₹)	39.26	27.78
Diluted (in ₹)	39.13	27.75

Particulars	31-March-2018	31-March-201
ASSETS		
Non-current assets		
(a) Property, plant and equipment	10,127	8.272
(b) Capital work-in-progress	4,800	3,363
(c) Intangible assets	59	58
(d) Financial assets		
Investments	81	1
Other financial assets	150	131
(e) Current-tax assets (net)	18	-
(f) Other non-current assets	609	478
	15,844	12,303
Current assets	10,011	12,000
(a) Inventories	4,384	3,489
(b) Financial Assets	4,304	3,403
Investments	684	321
Trade receivables	6,375	4,752
Cash and cash equivalents	217	4,752
		123
Bank balances other than cash and cash equivalents	1,620 45	
Loans		35
Other financial assets	6,142	752
(c) Other current assets	1,840	1,166
	21,307	10,873
Total assets	37,151	23,176
EQUITY AND LIABILITIES		
Equity		
(a) Equity share capital	369	349
(b) Other equity	30,353	16,144
Equity attributable to owners	30,722	16,493
Non-controlling interest	38	41
Total of Equity	30,760	16,534
Liabilities		
Non-current liabilities		
a) Financial liabilities		
Other financial liabilities	8	8
(b) Provision for employee benefits	324	219
(c) Deferred tax liabilities (net)	139	150
	471	377
Current liabilities	71.1	
(a) Financial liabilities		
Borrowings	1,732	2,216
Trade payables	2,691	
· · · · · · · · · · · · · · · · · · ·	· · · · · · · · · · · · · · · · · · ·	2,627
Other financial liabilities	1,024	1,014
(b) Other current liabilities	310	257
(c) Provision for employee benefits	137	18
(d) Current-tax liabilities (net)	26	133
	5,920	6,265
Total equity and liabilities	37.151	23,176
· · ·		,

FY2014 and FY2015 numbers have been prepared under IGAAP, whereas FY2016 and FY2017 numbers have been prepared under Ind AS

Historical Financials (contd.)



Segmental Breakdown (INR Mn)						
Revenue Division	Q1 FY19	FY18	Q4FY18	Q1 FY18	FY17	
API, Domestic	64.5	355.3	82.0	67.2	512.2	
API, Exports	608.2	2,498.6	514.5	795.9	1,325.5	
API Gross Revenue	672.7	2,853.9	596.5	863.1	1,837.7	
Formulations, Exports	262.9	2,050.7	590.2	436.1	3,947.4	
Income from Profit Sharing/Service Income	2,103.5	8,367.9	4,332.0	901.8	4,328.2	
Formulations Onco (including CnD)	929.5	3,380.6	828.2	731.0	3,224.3	
Formulations, Brand Pharma Non-Onco	741.9	3,103.5	624.2	844.7	4,801.6	
Formulations, 3rd party, & miscel	241.7	718.2	133.0	246.2	784.6	
Formulations Gross Revenue	4,279.6	17,620.9	6,507.6	3,159.8	17,086.1	
Other operating and non- operating income	561.7	1,003.9	391.5	311.9	1,236.0	
Stand-Alone Total Net Revenue	5,514.0	21,478.7	7,495.7	4,334.8	20,159.8	
Total Revenue, all subsidiaries	231.0	945.3	383.0	152.2	630.0	
Consolidated Total Net Revenue	5,745.0	22,424.0	7,878.7	4,487.0	20789.8	

	Q1 FY19	FY18	Q4 FY18 Q1 FY18		FY17
Total Revenues	5,745	22,424	7,879	4,487	20,789
EBITDA	2,535	9,688	4,034	1401	6,973
EBITDA Margin (%)	44.1%	43.2%	51.2%	31.2%	33.5%
PAT (after minority interest)	1,816	6,962	2,999	940	4,860
PAT Margin (%)	31.6%	31.0%	38.1%	20.9%	23.4%