

Date: May 16, 2025

To,
Sr. General Manager
Listing Department
BSE Limited
Phiroze Jeejeebhoy Towers
Dalal Street
Mumbai – 400 001

BSE Scrip Code: 544319

To,
Sr. General Manager
Listing Department
National Stock Exchange of India Limited
Exchange Plaza, C-1, Block G
Bandra Kurla Complex
Bandra (E), Mumbai – 400 051

NSE Symbol: SENORES

Sub.: Intimation under Regulations 30 of the Securities and Exchange Board of India (Listing Obligations and Disclosure Requirements) Regulations, 2015 - Investor Presentation

Dear Sir/Madam,

In terms of Regulation 30 of Securities and Exchange Board of India (Listing Obligations and Disclosure Requirements) Regulations, 2015 and our intimation dated May 12, 2025 regarding the Earnings Conference Call, we enclose herewith the Investor Presentation for Q4 and FY25.

Please be informed that the Investor Presentation for Q4 and FY25 enclosed herewith shall supersede the Investor Presentation submitted earlier today.

The aforesaid information is also being hosted on the Company's website at www.senorespharma.com.

You are requested to take the same on record.

Thanking you.

For Senores Pharmaceuticals Limited

Vinay Kumar Mishra
Company Secretary and Compliance Officer
ICSI Membership No.: F11464

Enclosures: As above

Senores Pharmaceuticals Limited

1101 to 1103, 11th Floor, South Tower, One42, Opp. Jayantilal Park,
Ambali Bopal Road, Ahmedabad-380054, Gujarat, India

P: +91 79 2999 9857 | E: info@senorespharma.com
W: www.senorespharma.com | CIN No.: L24290GJ2017PLC100263

Investor Presentation

May 2025



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This Presentation contains data and statistics from certain reports and the F&S Report, which is subject to the following disclaimer Frost & Sullivan has taken due care and caution in preparing this report (“Overview of the Global Pharma Market” Report) based on the information obtained by Frost & Sullivan from sources which it considers reliable (“Data”). This Report is not a recommendation to invest / disinvest in any entity covered in the Report and no part of this Report should be construed as an expert advice or investment advice or any form of investment banking within the meaning of any law or regulation. Without limiting the generality of the foregoing, nothing in the Report is to be construed as Frost & Sullivan providing or intending to provide any services in jurisdictions where Frost & Sullivan does not have the necessary permission and/or registration to carry out its business activities in this regard. Senores Pharmaceuticals Limited will be responsible for ensuring compliances and consequences of non-compliances for use of the (“Overview of the Global Pharma Market”) Report or part thereof outside India. No part of this Frost & Sullivan Report may be published/reproduced in any form without Frost & Sullivan’s prior written approval.



Financial Highlights Q4 and FY25

Key Operational Highlights

Regulated Markets

22

Owned Commercial Products

28

Pipeline CGT Opportunity Products

61

Approved ANDA Products

22

CDMO/CMO Commercial Products

51

Pipeline Products

69

CDMO/CMO Pipeline Products

Emerging Markets

285

Approved Products

636

Products Under Registration

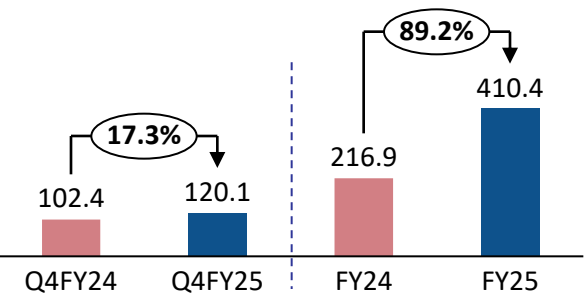
40+

Countries Present

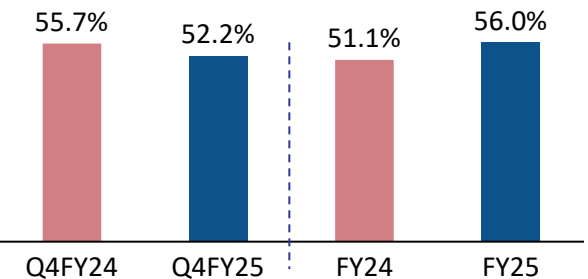
Consolidated Financial Highlights – Q4 & FY25

Income up by **91%**
Y-o-Y for FY25

Income (INR Crs)*

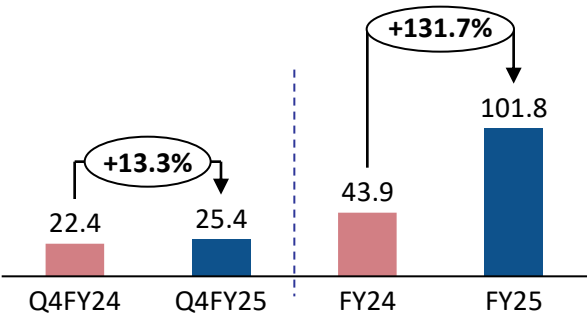


Gross Margin (%)

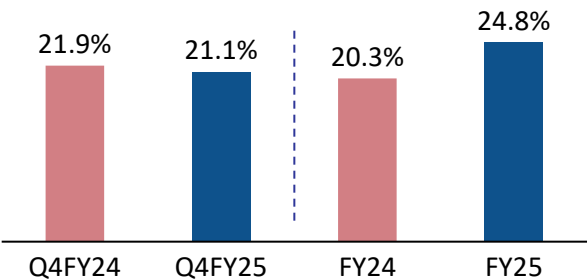


EBIDTA up by **145%**
Y-o-Y for FY25

EBITDA (INR Crs)

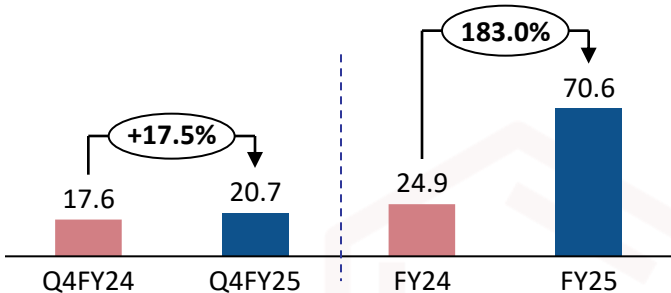


EBITDA Margin (%)

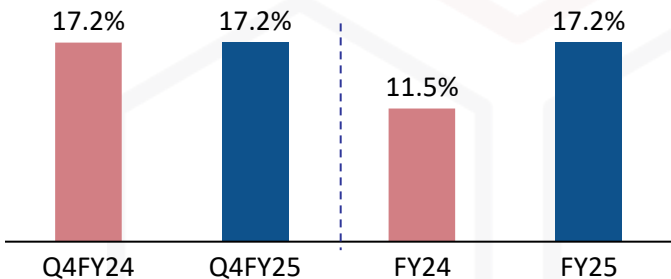


PBT up by **183%**
Y-o-Y for FY25

PBT (INR Crs)

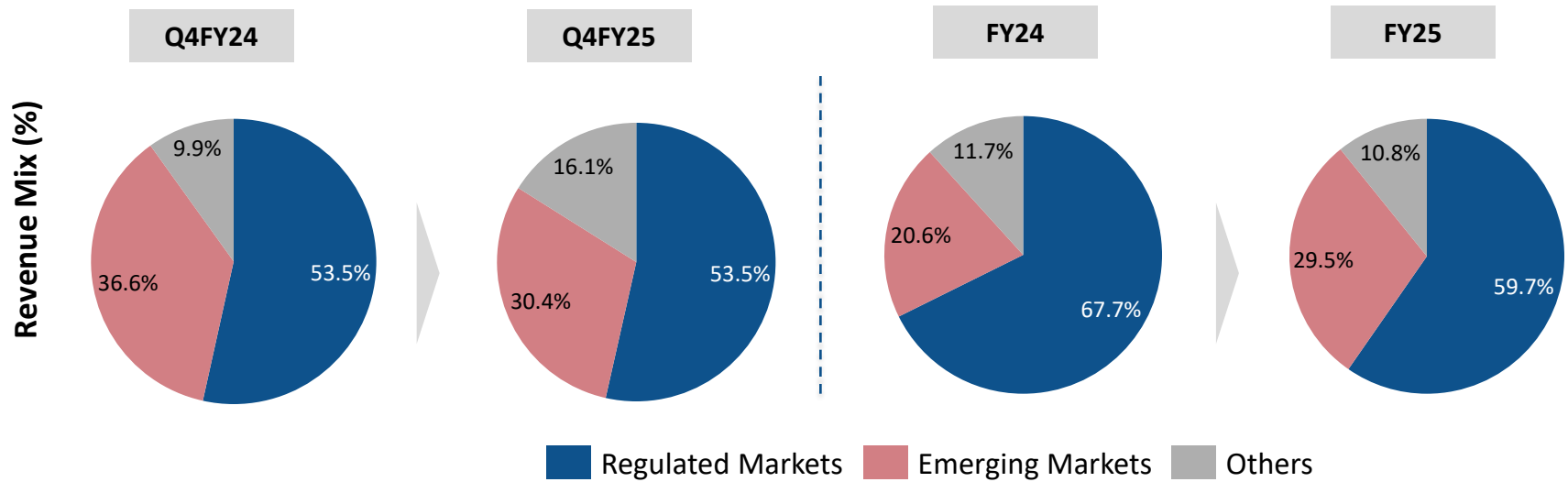
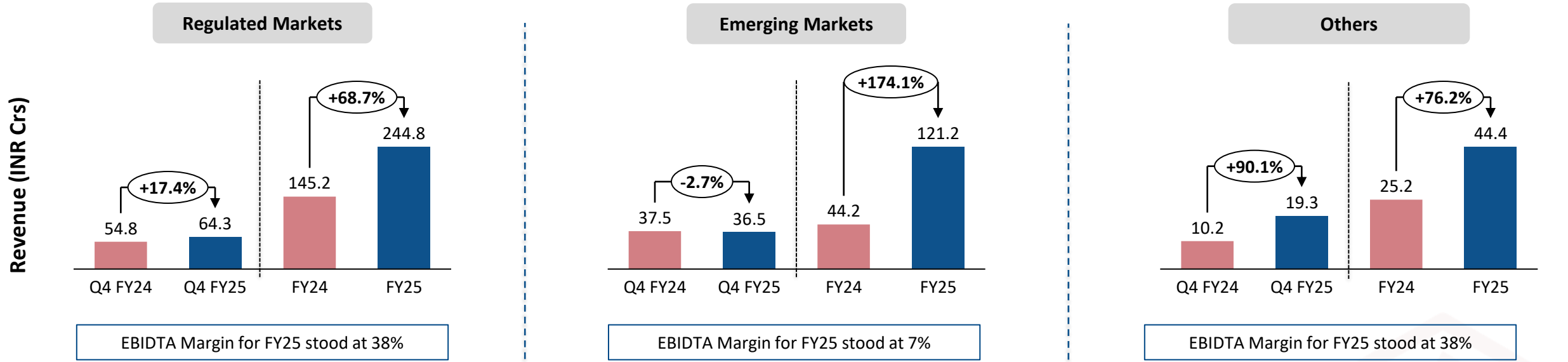


PBT Margin (%)



* Income includes Revenue from Operations and Other Operating Income.
Note: FY24 includes financials of Ratnatris Pharmaceuticals from 14th December 2023 to 31st March 2024

Business Segment Financial Highlights – Q4 & FY25



- **Regulated Markets**, contribute to **60% of total revenue** for FY25 and will continue to remain the **key focus market** going forward.
- Business model in the **Emerging Markets** is beginning to **take shape**. Share of Emerging markets has increased to **~30%** for FY25 and this segment is expected to grow steadily from here on

Note: FY24 includes financials of Ratnatris Pharmaceuticals from 14th December 2023 to 31st March 2024.



Swapnil Shah
Managing Director

Commenting on the Q4 & FY25 performance of the company Mr. Swapnil Shah, Managing Director for Senores Pharmaceuticals Limited said,

*“We are pleased to report strong performance for FY25, driven by our strategy of developing niche products for Regulated Markets and expanding our CDMO/CMO operations. At the same time, we have continued to grow our presence and product portfolio in Emerging Markets. Our **Total Income** and **Profit before Tax** for FY25 have increased by **91% & 183% respectively** on year-on-year basis, and we are optimistic about sustaining this momentum going forward.*

We continued to make steady progress on our stated strategies. During the quarter, we expanded our portfolio in the Regulated Markets through acquisition of 14 ANDA’s from Dr. Reddy’s and 1 ANDA from Breckenridge Pharmaceuticals. A large part of this acquired ANDA basket has considerable government contract opportunities. Further, our CDMO/CMO segment is gaining traction and continued to scale up well. We have launched 1 new product in the CDMO/CMO segment during the quarter and have 69 products in pipeline.

Our shift in product portfolio and go-to-market strategies in the Emerging Markets business is beginning to give the desired results, which can be seen in the improvement in margin during the quarter. We have registered 48 new products across the Emerging Markets during Q4FY25, taking the total portfolio to 285 registered products.

Going forward, although the uncertainties due to the tariff situation in USA will have to be closely monitored, we believe we are well placed to navigate the same, given our USFDA-approved manufacturing facility in USA.

We will continue to focus on three key pillars going forward – Expansion of ANDA portfolio in Regulated Markets; Steady scale-up of CDMO/CMO segment in the Regulated Markets; and Portfolio expansion and profitability improvement in the Emerging Markets.”

Consolidated Profit & Loss Account – Q4 & FY25

Profit and Loss (INR Crs)	Q4 FY25	Q4 FY24	YoY	FY25	FY24*	YoY
Revenue	108.7	97.4	11.6%	390.8	209.0	87.0%
Other Operating Income	11.5	5.0		19.5	7.9	
Total Income	120.2	102.4	17.3%	410.4	216.9	89.2%
Total COGS	57.5	45.4		180.7	106.1	
Gross Profit	62.7	57.0	9.9%	229.7	110.7	107.4%
Gross Margin (%)	52.2%	55.7%	-350 bps	56.0%	51.1%	490 bps
Employee Cost	14.6	18.2		60.4	35.5	
Other Expenses	22.7	16.4		67.5	31.3	
EBITDA	25.4	22.4	13.4%	101.8	43.9	131.7%
EBITDA Margin (%)	21.1%	21.9%	-70 bps	24.8%	19.4%	450 bps
Depreciation	5.2	3.7		16.8	10.0	
Other Income	5.8	2.0		7.2	0.5	
EBIT	26.0	20.6	25.9%	92.1	34.4	167.9%
Finance Cost	5.3	3.0		21.6	9.4	
Profit before Tax	20.7	17.6	17.4%	70.6	24.9	183.0%
PBT Margin (%)	17.2%	17.2%	0 bps	17.2%	11.5%	570 bps
Tax + Minority Interest	2.8	1.7		12.0	-7.8	
Profit after Tax and Minority Interest	17.8	15.9	11.9%	58.6	32.7	79.1%
Profit after Tax and Minority Interest Margin (%)	14.8%	15.6%	-70 bps	14.3%	15.1%	-80 bps
EPS (INR)	3.8	8.0		16.1	12.2	

*FY24 includes financials of Ratnatris Pharmaceuticals from 14th December 2023 to 31st March 2024

Other Financial Highlights – Q4 & FY25

Particulars (Rs Crs)	Q4FY25	Q4FY24	FY25	FY24
A: Other Operating Income				
Product related service Income	4.91	3.78	7.68	4.13
Miscellaneous - Other Operating Income	7.24	2.03	7.94	2.72
Gain on Foreign Exchange Fluctuation (Net)	-0.67	0.89	3.92	1.07
Total Other Operating Income	11.48	6.70	19.54	7.92
B: Interest Income	5.82	0.28	7.15	0.46
Total Other Income	17.30	6.98	26.69	8.38

Capex Plans:	In Cr
Capex Incurred during FY 2025	157.5
Capex planned for next year	251.0

Working Capital Days	FY25	FY24
Inventory	43	69
Trade Receivables	108	228
Trade Payables	182	435

Consolidated Balance Sheet

ASSETS (INR Crs)	Mar-25	Mar-24
Non-Current Assets		
Property, Plant and Equipment	198.9	152.2
Capital Work-in-Progress	44.2	17.8
Goodwill	38.2	38.2
Intangible Assets	54.3	35.9
Right of Use Assets	9.4	9.1
Intangible Assets under Development	128.3	79.3
Non-Current Financial Assets	5.2	20.5
Deferred Tax Assets (Net)	18.7	15.0
Other Non-Current Assets	7.4	3.0
Total Non-Current Assets	504.4	371.0
Current Assets		
Inventories	56.6	37.4
Trade Receivables	123.9	112.0
Cash & Cash Equivalents	105.4	7.7
Bank Balances Other Than Cash & Cash Equivalents	280.1	5.4
Loans	0.0	0.3
Other Current Financial Assets	117.0	66.2
Other current assets	39.5	22.0
Total Current Assets	722.4	250.9
Total ASSETS	1,226.9	621.9

EQUITY AND LIABILITIES (INR Crs)	Mar-25	Mar-24
Equity		
Equity Share Capital	46.1	30.5
Other Equity	740.1	173.8
Total Shareholder's Fund	786.2	204.3
Non-Controlling Interest	26.1	27.4
Total Equity	812.2	231.7
Non-Current Liabilities		
Borrowings	162.5	133.7
Lease Liabilities	8.2	7.8
Provisions	3.3	1.2
Total Non-Current Liabilities	173.9	142.7
Current Liabilities		
Borrowings	142.3	114.7
Lease Liabilities	1.8	1.5
Trade Payables	67.2	113.0
Other Current Liabilities	6.0	5.2
Other Current Financial Liabilities	7.8	4.6
Provisions	2.1	1.4
Current Tax Liabilities (net)	13.5	7.1
Total Current Liabilities	240.7	247.5
Total EQUITY AND LIABILITIES	1,226.9	621.9

Consolidated Abridged Cash Flow Statement

Cash Flow Statement (INR Mn)	FY25	FY24
Profit/(Loss) Before Tax	70.6	24.9
Adjustments for Non-Cash and Non-Operational Expenses / (Incomes)	27.1	17.9
Operating Profit / (Loss) Before Working Capital Changes	97.7	42.8
Changes in Working Capital	-134.4	-66.1
Cash from Operations	-36.7	-23.2
Income Tax (Paid) / Refunded	-9.2	-2.5
Net Cash Flow from Operating Activities (A)	-45.9	-25.7
Net Cash Flow from Investing Activities (B)	-429.5	-54.2
Net Cash Flow from Financing Activities (C)	573.1	87.0
Net Change in Cash & Cash Equivalents during the Year (A+B+C)	97.7	7.0
Cash & Cash Equivalents at the Beginning of the Period	7.6	0.1
Cash & Cash Equivalents at the End of the Period	105.4	7.6

Note: FY24 includes financials of Ratnatris Pharmaceuticals from 14th December 2023 to 31st March 2024

Update on Utilization of Funds from IPO

Sr. No	Object as per Offer Document	Amount as proposed in the Offer Document	Amount utilized as on March 31, 2025	Amount Unutilized as on March 31, 2025	Remarks
1	Investment in one of our Subsidiaries, Havix, to fund capital expenditure requirements for setting up a manufacturing facility for the production of sterile injections in our Atlanta Facility	107.0	-	107.0	Out of the unutilized amount Rs 130 crores parked as Fixed Deposit with HDFC Bank, Rs 209 crores parked as Fixed Deposit with ICICI Bank and balance held in HDFC bank account
2	Re-payment/pre-payment, in full or in part, of certain borrowings availed by our Company	73.5	43.6	29.9	
3	Investment in our Subsidiary, namely, Havix, for re-payment/pre-payment in full or in part, of certain borrowings availed by such Subsidiaries	20.2	3.9	16.3	
4	Funding the working capital requirements of our Company	43.3	20.2	23.1	
5	Investment in our Subsidiaries, namely Senores Pharmaceuticals Inc. and Ratnatris to fund their working capital requirements	59.5	22.4	37.1	
6	Funding inorganic growth through acquisition and other strategic initiatives and general corporate purposes	154.4	35.7	118.7	
7	Offer expenses	42.2	31.0	11.2	
	Total	500.0	156.8	343.3	

About Senores Pharmaceuticals Limited



Brief Overview

Global Research driven **Formulation Focused** company engaged in **developing & manufacturing** of pharmaceutical products predominantly for the **Regulated Markets of the US, Canada & UK** across various **therapeutic areas**



Niche Product Identification

Company focusses on **identification**, **development** and **commercialization** of products



R&D Capabilities

Identification, **development & manufacturing** of diverse range of **specialty**, **underpenetrated & complex** pharmaceutical products across therapeutic areas & dosage forms



Regulated Market Approved Facility

1 USFDA & DEA approved **formulation facility** in the **US**

Partnerships

Partnerships with leading pharmaceutical companies across Regulated as well as Emerging Markets

At a Glance



Business Segments



Regulated Markets Business

Primarily serves the **US, Canada** and **United Kingdom**

USFDA approved **facility in Atlanta, US**

22 Commercialized Products

61 Approved ANDA Products

51 Pipeline Products

28 Pipeline CGT Opportunity Products



Emerging Markets Business

Develop & manufacture pharmaceutical products for the Emerging Markets

WHO-GMP approved **facility at Chhatral**

Present in **40+ Countries**



Product registration for **285 Products**

Approval for manufacturing facility from regulatory bodies of **10 countries**

Product applications filed for **636 Products**

API Business

Manufacture APIs and caters to the **domestic market** and **SAARC countries**

Naroda facility compliant with Indian GMP

Commercialized **16 APIs**

Critical Care Injectables Business

Supply of critical care injectables across **India** to various **hospitals** through the distributors

Field presence with **47 employees**

Launched **56 Products**

Key Events & Milestones



Incorporated as “**Senores Pharmaceuticals Pvt. Ltd.**” in India.

2017



Incorporated a wholly owned subsidiary, **Senores Pharmaceuticals Inc** in the US

2021



Entered into a new segment of API with acquisition of **Ratnagene Lifescience Pvt. Ltd.**¹

2023

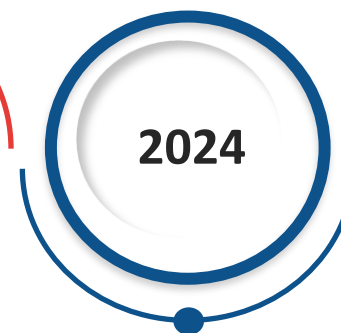


Acquired majority stake in Havix, enhancing the ability to serve Regulated Markets through **USFDA approved facility in US**



Consolidated presence in emerging markets by acquiring shares in **Ratnatris Pharmaceuticals Pvt. Ltd.**

2023



Successful Listing on Stock Exchanges in India on BSE & NSE

2024



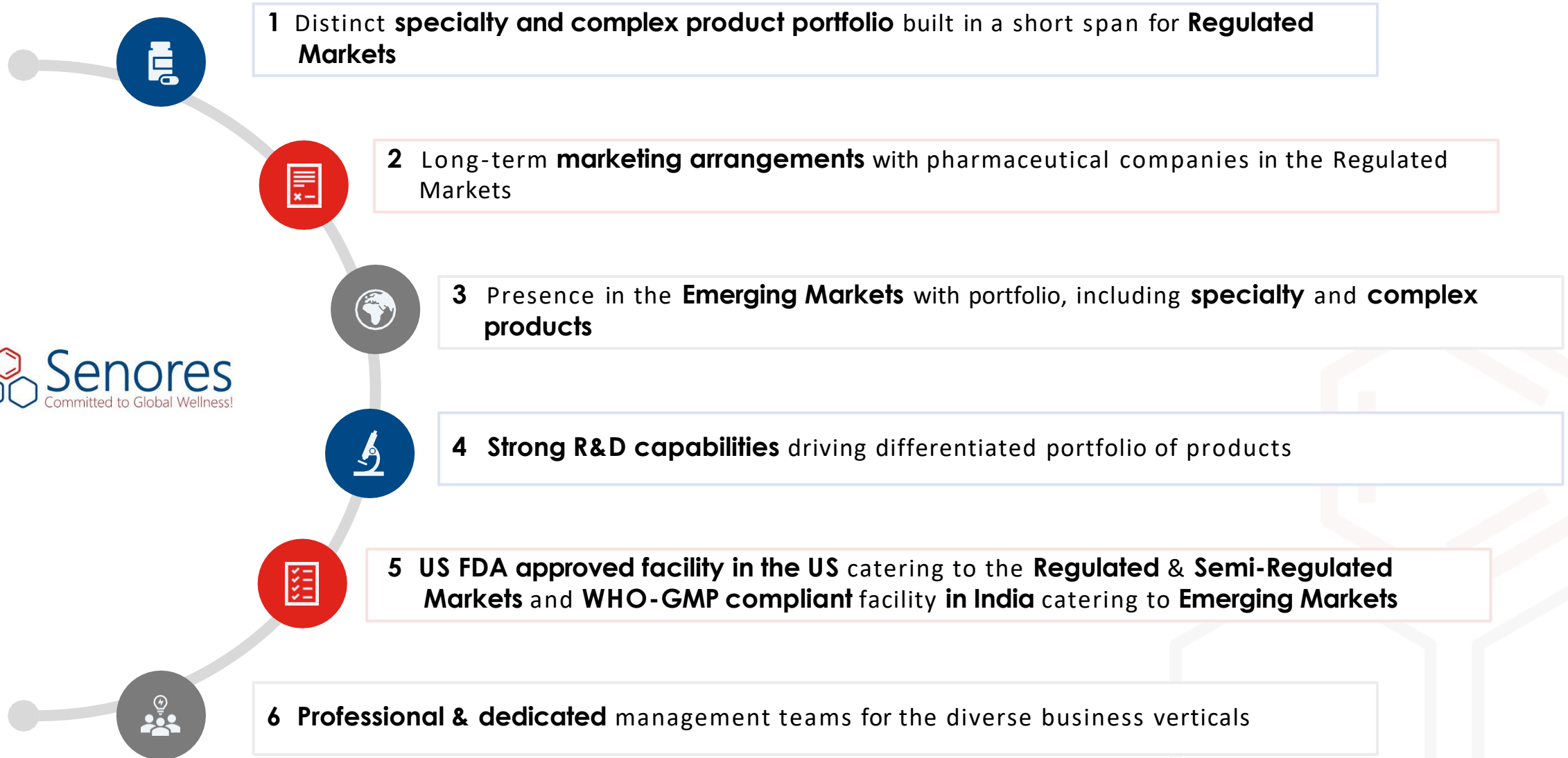
Commenced manufacturing activities at its **greenfield API plant** located at Gujarat

2025

Key Strengths

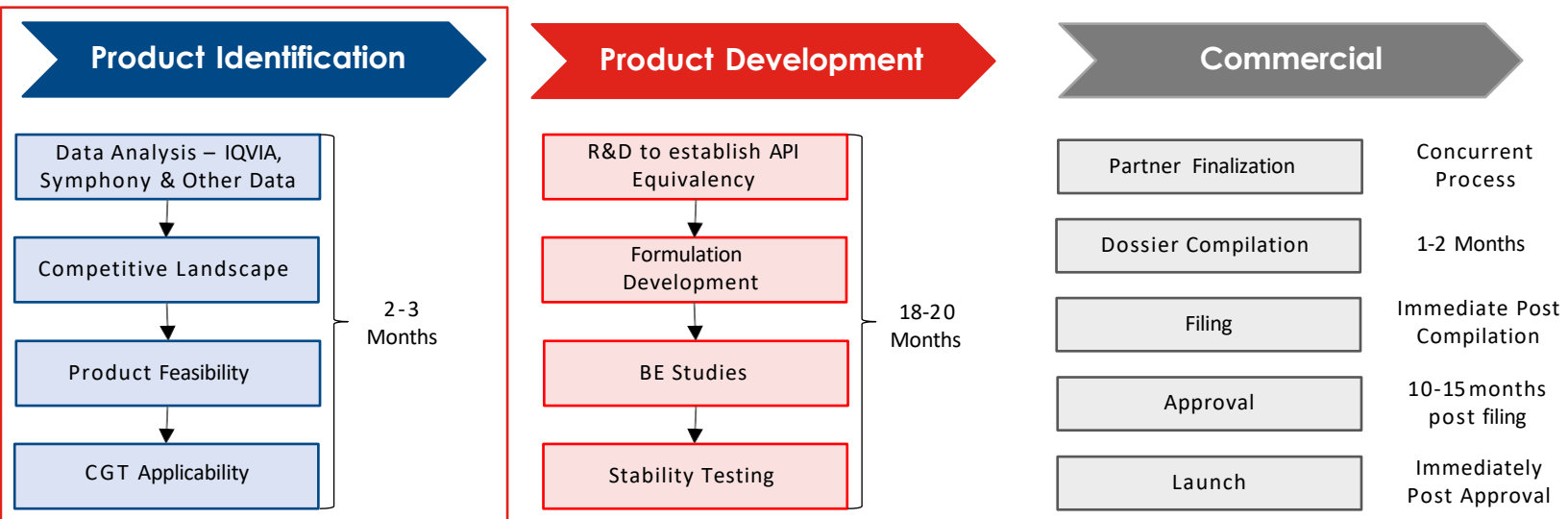


Key Strengths of the Company



Distinct Specialty & Complex Product Portfolio Built in a Short Span for Regulated Markets

Demonstrated Capability to propel Products from Conception to Commercialization



ANDA Products

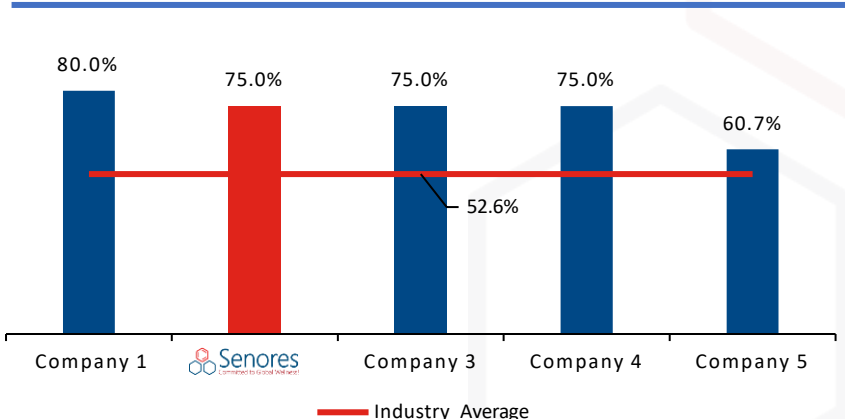


2nd Highest in terms of proportion of CGT Exclusivity amongst Peers

- CGT is granted when FDA determines there is **inadequate generic competition**
- Exclusivity Period for **180 days**
 - Competing Generic Versions of the drug **cannot be marketed**
- Allows to **establish a foothold** in the market
- Opportunity** for **Market Penetration & Revenue Growth**
- Lower price erosion** due to **lesser competition**

Source: F&S Report
Note: All the numbers are as of 30th September, 2024

Proportion of Ingredients with CGT Exclusivity



Presence in Regulated Markets through Long-term marketing arrangements

MARKETED PRODUCTS

Business Model

Identify, Develop & Commercialize specialty & complex niche products in mid-market range

Revenue Model

In-Licensing Fee

Transfer Pricing

Profit Sharing

- **Long-term Marketing & Distribution Agreements**
- Strength lies in taking a product from **conceptualization to commercialization**
- **22 Commercialized** products
- **61 Approved ANDA Products**

CDMO / CMO

Business Model

Customized formulation, development & manufacturing capabilities for customers

Revenue Model

Tech Transfer/
Developmental Cost

Transfer Pricing

Service Income

- **Partner with CDMO customers early** in the drug development process
- **Recurring Revenue with Steady & Predictable Cash Flows**
- Contracts for more than **40** products in **US, Canada, UK, South Africa, UAE, Israel, Denmark, Saudi Arabia & Vietnam.**
- Eligible for manufacturing formulations having **controlled substances**
- **Complied for catering to government supplies in the US**

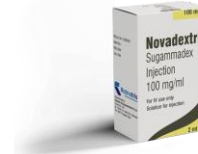
Pharmaceutical companies increasingly favor partnering with one-stop-shop solution providers that seamlessly integrate both development and manufacturing services

Presence in the Emerging Markets with Portfolio Including Specialty and Complex Products

Special Focus

Niche & Specialty patented products in the US with early launch opportunity in Emerging Markets

Sugammadex



Ferric Carboxymaltose



Tofacitinib



Sacubitril + Valsartan



Apixaban



Eltrombopag Olamine



Diverse Business Models

P2P Model

Distributor Model

CDMO Model

Own Brands²

Top Emerging Markets¹

Philippines



Peru



Ghana



Nigeria



Myanmar



Guatemala



Kenya



40+

Countries Marketed

921

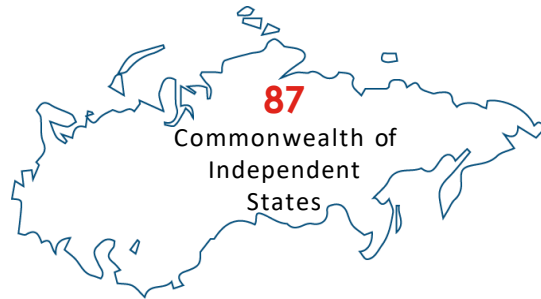
Total Products

285

Product Registrations

636

Product Applications Filed



73 | 14
Registered | Filed



72 | 331
Registered | Filed



26 | 112
Registered | Filed



113 | 174
Registered | Filed

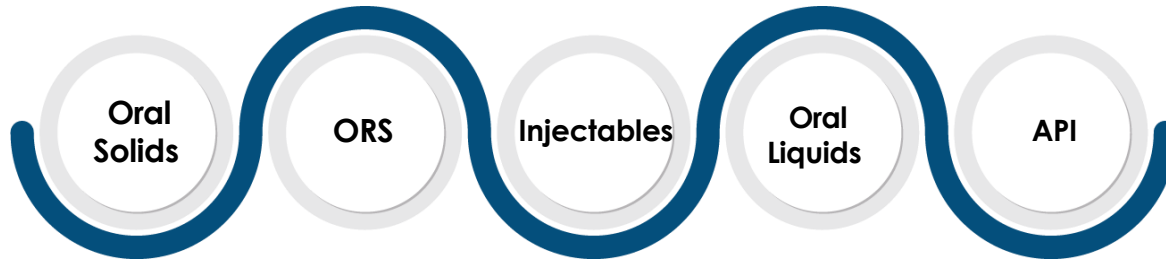


1 | 5
Registered | Filed

Partners to Global Pharmaceutical Companies in Regulated & Emerging Markets



Strong R&D Capabilities Driving Differentiated Product Portfolio



R&D Site 1 - USA

- In-House R&D in US for Controlled Substances



Identify

- Internal research
- Information available on subscribed databases



R&D Site 2 - India

- Helps in dossier preparation
- Submission of ANDA applications



Team

- **64** people
- 2 members having doctoral qualifications



R&D Site 3 - India

Consolidating R&D activities at Ahmedabad. R&D Centre spread around 11,750 sq. ft.



Margin

The focus on R&D has allowed the company to maintain high profit margins compared to it's industry peers



Accredited Manufacturing Facilities

Atlanta Facility

Facility

2
Manufacturing Lines


185,300 Sq. Ft.
Total Area

OSD

1.2 Bn units
Annual Installed Capacity




Strong Regulatory Track Record




USFDA Approved

4 USFDA, received no Form 483 & 8 Customer Audits



DEA Compliant

Eligible for Manufacturing Controlled Substances



BAA Compliant

Ability to caters to government supplies

Ahmedabad, Chhatral Facility



35,205 Sq. Mtr.
Total Area

12
Manufacturing Lines

4
Dosage Forms

General Oral Dosage

1397.96 Mn
Annual Installed Capacity¹

Injectables

49.92 Mn
Annual Installed Capacity¹

Beta Lactum Orals

511.68 Mn
Annual Installed Capacity¹

Facility Approvals

KUWAIT MOH	CAMBODIA MOH	SRI LANKA NMRA	IVORY COAST DPM	KENYA PPB
PHILIPPINES FDA	NIGERIA NAFDAC	PERU DIGEMID	ZAMBIA MOH	LIBERIA MOH

Corporate Structure



Group holding company based in India
Houses Domestic Critical Care Injectables business & API business



Senores Pharmaceuticals Inc.
Marketing cum ANDA
Holding Company

US based company holding
ANDAs and mutually
exclusive marketing
relationships with customers

100% Subsidiary



Havix Group Inc
Caters to Regulated Market

Engaged in the Contract
Development & Manufacturing
of Generic Pharmaceutical
Formulations for the US, Canada

67.77%¹ Subsidiary



Ratnatris Pharmaceuticals Pvt. Ltd.
Caters to Emerging Market

Development, Manufacturing
and Marketing of Generic
Pharmaceutical Formulations

69% Subsidiary

Note 1 : 49.91% held by the holding company & 16.66% held by it's wholly owned subsidiary company named Senores Pharmaceuticals Inc

Professional and Dedicated Management Team



Swapnil Jatinbhai Shah
Promoter & Managing Director

- 17+ years of experience in the pharma sector
- Part of the core management team & leads overall company functions: product portfolio, corporate strategy, business development, strategic management



Sanjay Shaileshbhai Majmudar
Chairman & Non-Executive, Non-Independent Director

- 40+ years of experience
- Director on the board of AIA Engineering Ltd., Ashima Ltd., M&B Engineering Ltd. & Welcast Steels Ltd.



Deval Rajnikant Shah
Whole-Time Director & CFO

- 40+ years of experience in Pharma, Engineering & Chartered Accountancy
- Previously associated with SAI Consulting Engineers Pvt. Ltd. as CFO



Chetan Bipinchandra Shah
Whole-Time Director & COO

- 25+ years of experience in Pharma
- Previously associated with Torrent Pharma & Cadila Pharma. Also associated with Reliance Retail Ltd., Reliance Fresh Ltd. & Reliance Corporate IT Park Ltd.



Jitendra Babulal Sanghvi
Non-Executive, Non-Independent Director

- 15+ years of experience
- Director of Ratnaris Pharma since Aug 11, 2009
- Conferred with "Young Pharma Entrepreneur of the Year" award in 2013



Ashokkumar Vijaysinh Barot
Promoter, Non-Executive, Non-Independent Director

- 21+ years of Pharmaceutical experience
- Non-Executive Director on the board of Di-Cal Pharma Pvt. Ltd. Since Nov 6, 2008. Registered pharmacist with the state pharmacy council of Gujarat



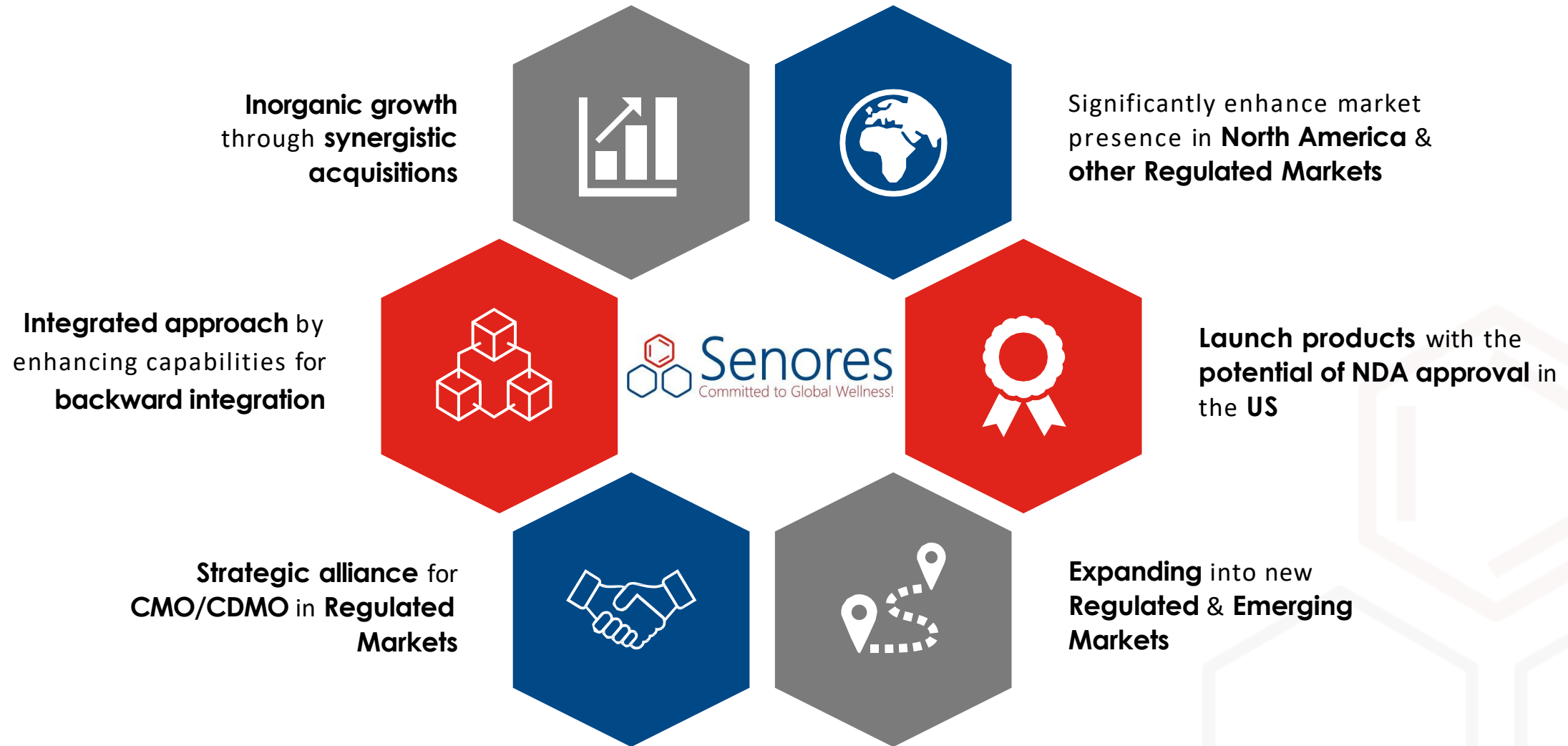
Deepak Jain
Vice President Regulatory Affairs

- 11+ years of Pharmaceutical experience
- Previously Associated with Cadila Healthcare Ltd.. As Deputy General Manager

Growth Strategies

The background features a dark blue gradient on the left and a lighter blue gradient on the right, separated by a diagonal line. A thick red stripe and a thinner white stripe run parallel to this diagonal line. Faint, light blue network patterns of interconnected dots and lines are visible in the background.

Key Strategies

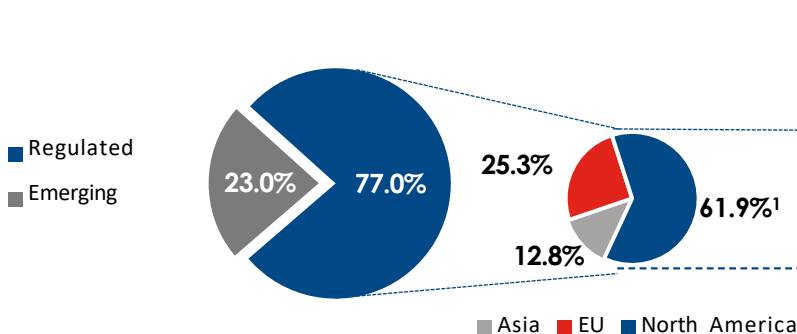


Enhance Market Presence of Marketed Products in North America and Other Regulated Markets

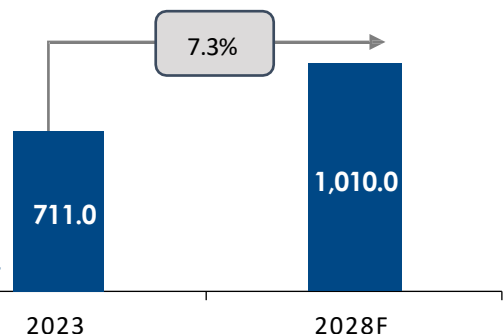
US accounts for 43% of the global pharma market, 56% of the regulated market & 91% of North America Market

Persistent drug shortage calls for an increased supply of generic drugs

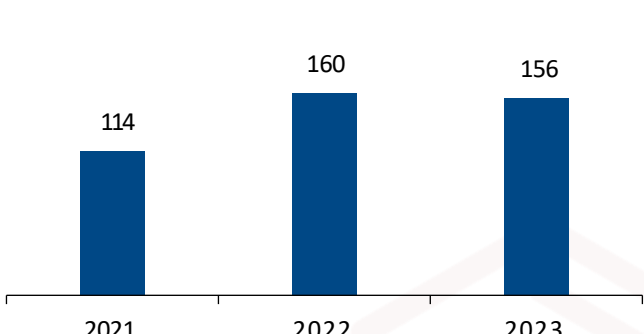
Global pharma market by classification (2023, USD bn)



US pharma market size (USD bn)



Number of new drug shortages



Enhance market presence in North America along with the partners

✓ Launch generic products with the potential of NDA approval in the US

✓ Set up niche sterile injectables manufacturing facility in the US

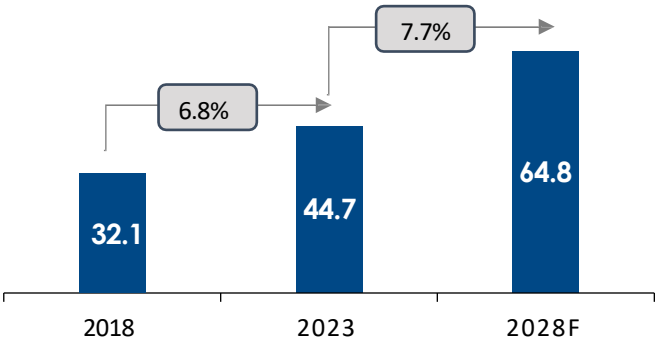
✓ Inorganic growth through synergistic acquisitions

Source: F&S Report
Note: 1: Pharma market value for North America is USD 778.8 bn, Europe is USD 318.5 bn & Asia is USD 161.3 bn

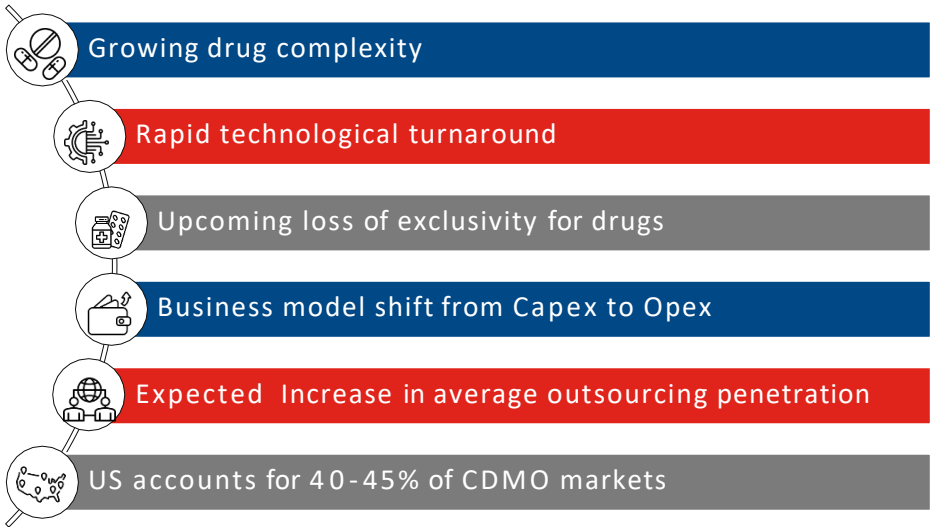
Entering into Strategic Alliance for CMO/CDMO in Regulated Markets

CDMO market growing faster than the pharma market which is growing at 7.7%

US CDMO/CMO market (USD bn)



Growth propellers



Strategic alliance for CMO/CDMO in Regulated Markets

The dependence on CDMO/CMO's has increased as they offer



Appended Manufacturing Capacity



Access to new Markets



Mitigate investments, production & supply risk



Bring Technological Overhaul



Eligible for Manufacturing Controlled Substances



Complied to cater to Government Supplies

Offering benefits such as

Focus on core competencies

Flexibility & Scalability

Early to Market advantage

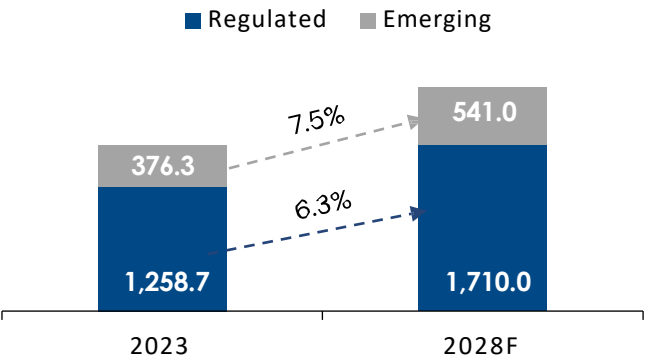
Cost Advantages

Access to global talent & advanced technologies

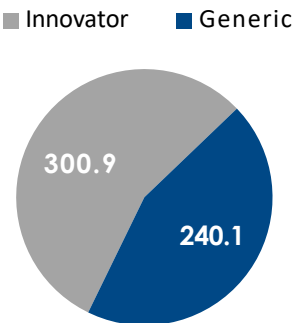
Expanding into New Regulated & Emerging Markets

Regulated & Emerging markets witnessing strong growth

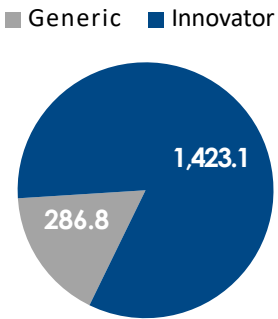
Global pharma market by classification (USD bn)



Emerging pharma market by Product (2028F)



Regulated pharma market by Product (2028F)



Expanding into New Regulated & Emerging Markets



Focus on niche and complex range of products with higher margin profile



Leverage presence in Regulated Markets, to increase reach in Semi Regulated Markets



Tapping into new mid-tier markets such as Brazil, Australia & New Zealand



Leveraging knowledge of Emerging markets to manufacture more products by setting up new facilities



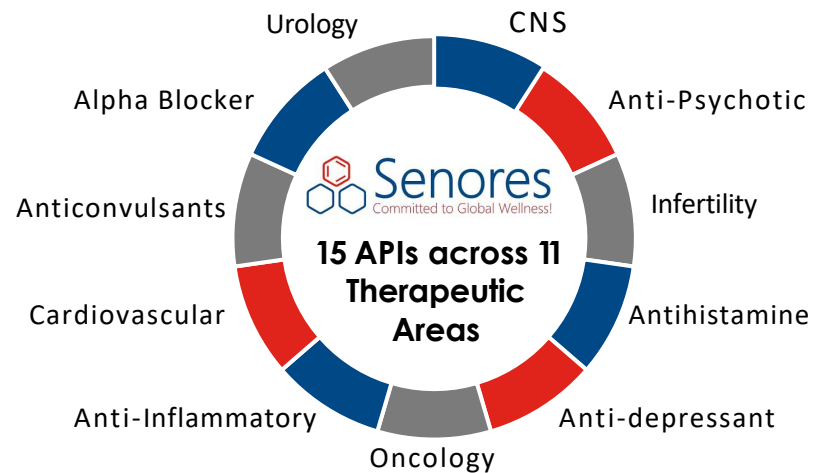
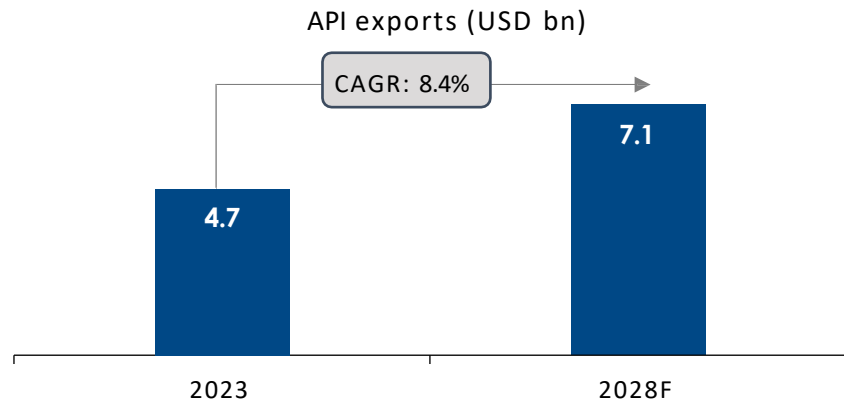
Opportunities for registering & marketing value added niche formulations in Emerging Markets: Philippines, Uzbekistan, Peru, Ghana, Tanzania, Kenya, Libya and Guatemala



Multiple marketing and distribution models to enhance presence in Emerging market countries

Enhancing Capabilities for Backward Integration

Indian API export market to grow at a CAGR of 8.4%



Pursuing an integrated approach for greater backward integration

Proposed Expansion Plans

Setup Greenfield unit - Increase API manufacturing capacity from 25 MTPA to 169 MTPA

Manufacture API for Regulated and Semi-Regulated markets

Benefits of API manufacturing capabilities



Supply Chain Control



Cost Savings



Quality Assurance



Flexibility & Innovation



Reduced Time to Market



Competitive Advantage



Diversified revenue stream & business resilience

Backward integration into API manufacturing reduces reliance on third-party vendors & enhances market competitiveness

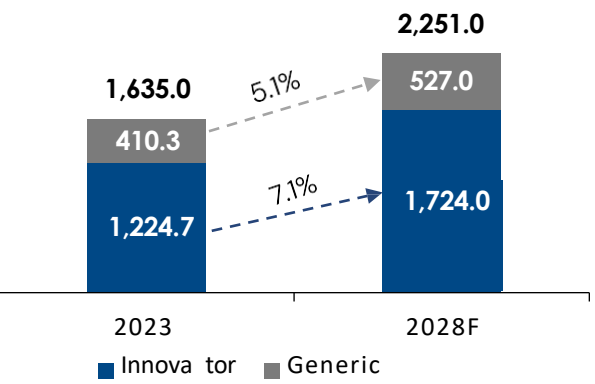
Industry Overview



Global Pharma Market

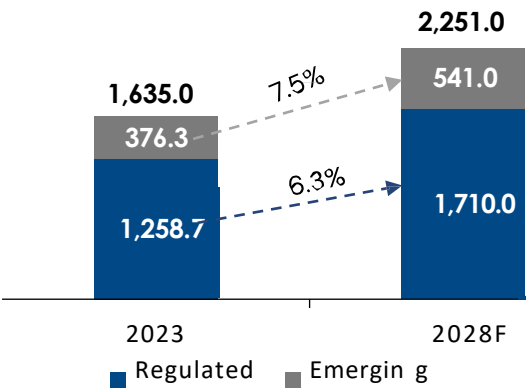
Global pharma market is expected to grow at 6.6% from 2023-28...

Global pharma market by product type (USD bn)



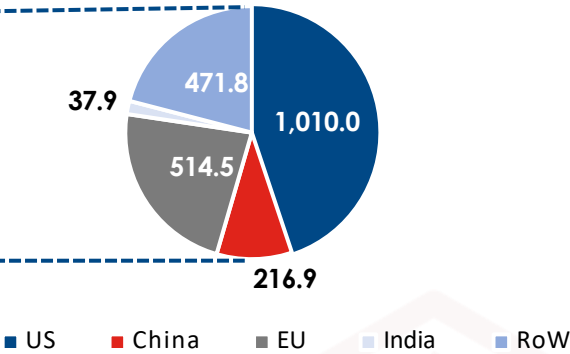
...regulated market will continue to dominate the global pharma market

Global pharma market by region (USD bn)



US continues to exert influence on Global pharma market

Share in global pharma market by region (2028F)



Aging Population

Population of people over 60 expected to double and reach ~2.1 Bn by 2050, increasing age related conditions

Increasing incidences of chronic diseases

~ 1 in 3 adults have multiple chronic conditions, requiring lifelong use of drugs, with costs estimated to reach USD 47 tn by 2030



Investments in R&D

R&D investments drive market growth by discovering breakthrough treatments for prevalent and emerging diseases by expanding therapeutic segments

Consumer awareness

Post-COVID-19, the OTC segment grew massively due to heightened consumer awareness about health, wellness, and preventive care

Demand from developing nations

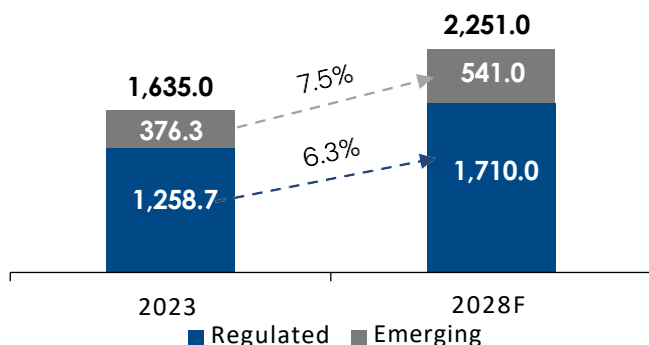
Rising instances of chronic conditions and persistent burden of infectious diseases drive dual demand

Regulated Pharma Market

Regulated markets to dominate the global pharmaceutical sector, driven by their access to a growing innovative drug market & a thriving generics market

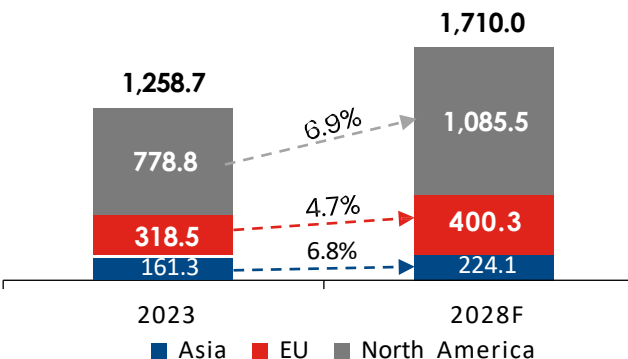
Regulated markets constitute 77% of the global pharmaceutical sector (2023)

Global pharma market by classification (USD bn)



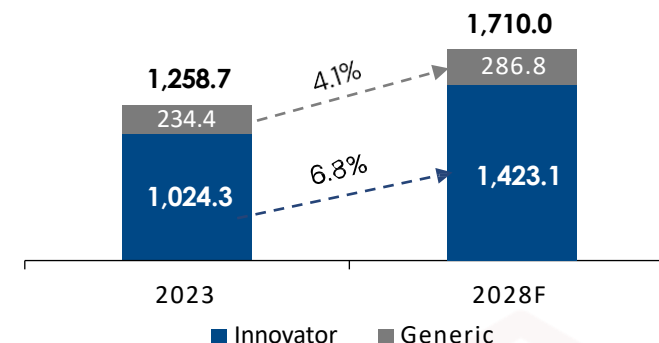
North America accounts for 62% of regulated pharma market (2023)

Regulated pharma market by region (USD bn)



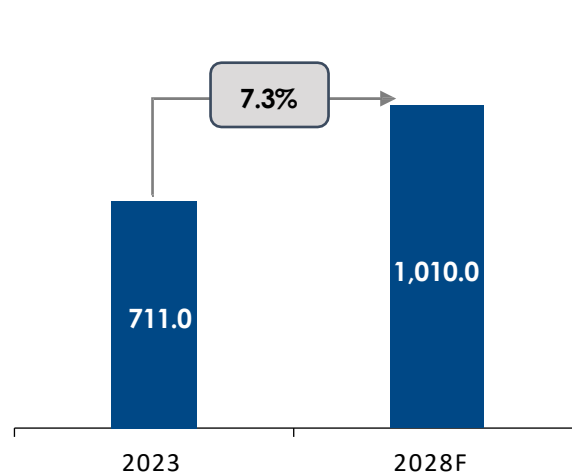
New generics and patent expirations to drive drug market growth

Regulated pharma market by product type (USD bn)



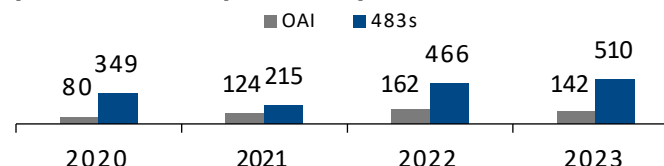
US accounts for 43% of the global pharma market, 56% of the regulated market and 91% of North American market (2023)

US Pharma Market size (USD bn)

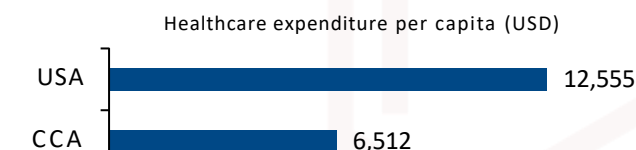


Characteristics of US healthcare market

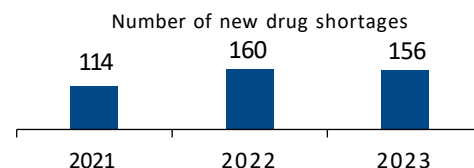
High quality standards impose entry barriers



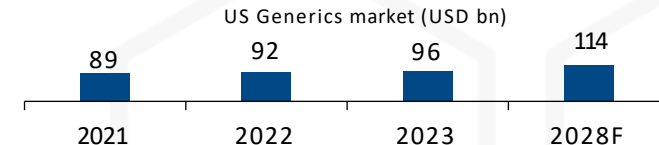
Very high healthcare expenditure per capita



Persistent drug shortage calls for an increased supply of generic drugs



Increasing generic adoption to navigate high healthcare costs



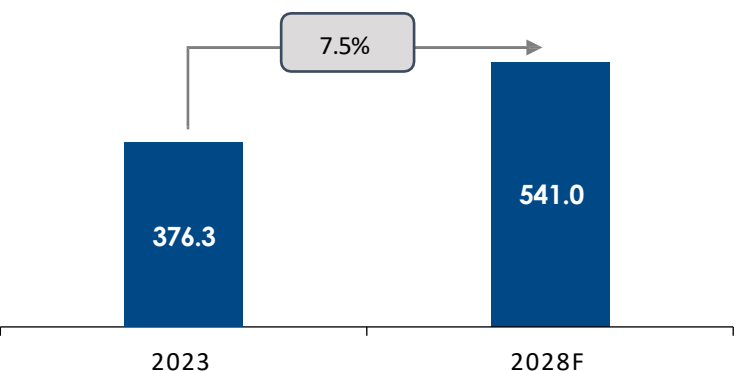
Growing drug demand with a simultaneous need to control costs has increased import dependence, particularly from India

Emerging Pharma Market

Emerging pharma market

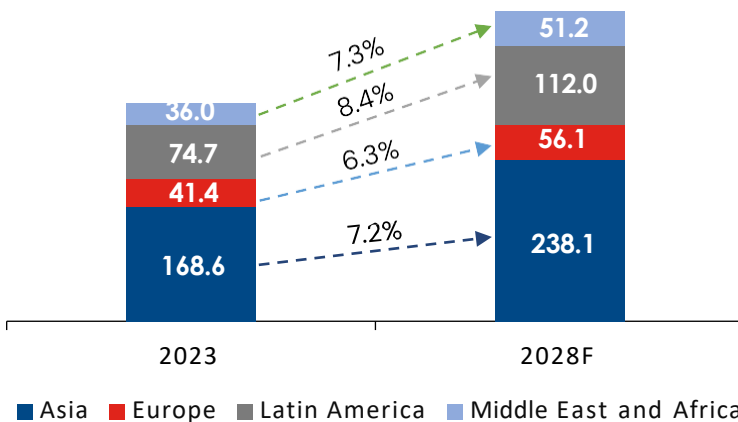
Emerging markets to outpace developed ones...

Emerging pharma market size (USD bn)



...driven by population growth, expanding disease burden, & investments in healthcare & infrastructure

Emerging pharma market size by region (USD bn)

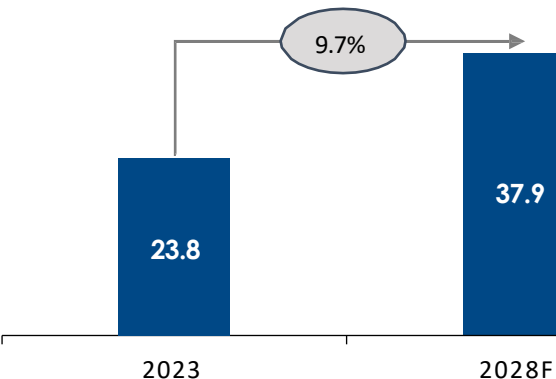


Source: F&S Report

Overview of India pharma market

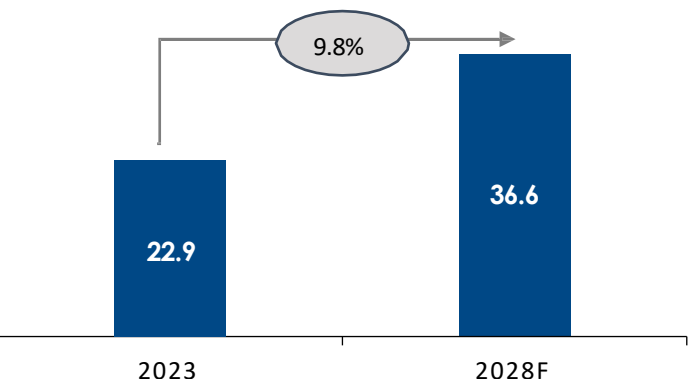
IPM is amongst the fastest growing in the world...

Indian pharma market size (USD bn)



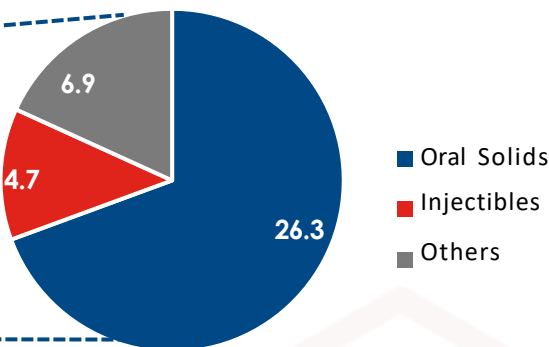
...dominated by generic drugs constituting ~96% of drug consumption

Indian generic pharma market size (USD bn)



Oral solids will dominate IPM

Indian pharma market by dosage (2028F, USD bn)



Growth Drivers

- ✓ Insurance penetration
- ✓ Increase in chronic patient population
- ✓ Availability of affordable & innovative generics
- ✓ Drug access focused government schemes
- ✓ Growth in hospital business segment

SENORES PHARMACEUTICALS LIMITED

CONTACT US



Senores Pharmaceuticals Limited

CIN: L24290GJ2017PLC100263

Mr. Deval Shah – Whole Time Director & CFO

investors@senorespharma.com



Strategic Growth Advisors Private Limited

CIN: U74140MH2010PTC204285

Mr. Sagar Shroff / Mr. Tanay Shah
sagar.shroff@sgapl.net / tanay.shah@sgapl.net
+91 98205 19303 / +91 98333 91899