

**Date:** January 23, 2025

To,
Sr. General Manager
Listing Department
BSE Limited

Phiroze Jeejeebhoy Towers Dalal Street Mumbai – 400001

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 January 10, 2025 regarding the Earnings Conference Call, we enclose herewith the Investor Presentation for Q3 and 9M FY25.

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

W: www.senorespharma.com | CIN No.: U24290GJ2017PLC100263

# Investor Presentation JANUARY 2025



# Safe Harbor



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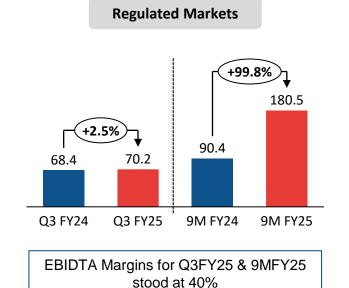
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# Financial Highlights Q3 and 9M FY25

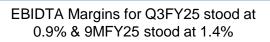
# Q3 & 9M FY25 Business Segment Financial Highlights\*



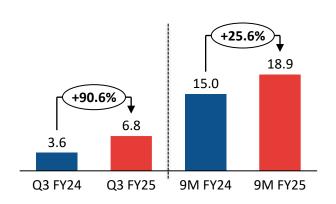




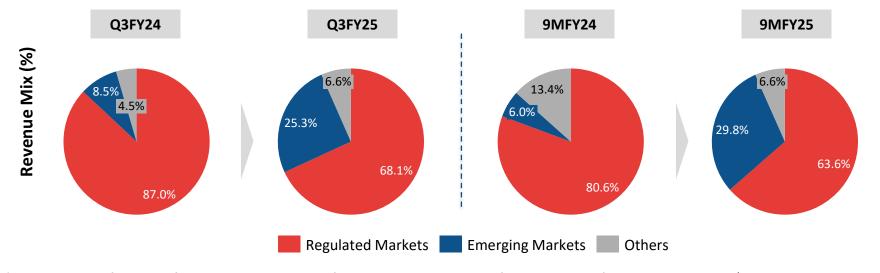
# Emerging Markets +1,164.8% 84.6 84.6 Q3 FY24 Q3 FY25 9M FY24 9M FY25



### Others (API, Injectables)



EBIDTA Margins for Q3FY25 stood at 50% & 9MFY25 stood at 31%



- Regulated Markets, contribute >63% of total revenue for 9M 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 9M FY25 and this segment is expected to grow steadily from here on

# **Q3 FY25 Key Operational Highlights**



**Regulated Markets** 

22

**Owned Commercial Products** 

24

Approved ANDA's

51

**Pipeline Products** 

28

**Pipeline CGT Opportunity Products** 

21

CDMO/CMO Commercial Products

69

CDMO/CMO Pipeline Products

**Emerging Markets** 

237

**Approved Products** 

537

**Products Under Registration** 

**40**+

**Countries Present** 

# Key Financial Highlights – Q3 & 9M FY25\*

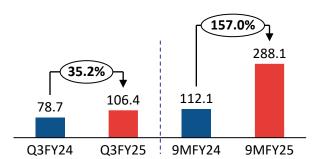


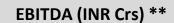
Income up by 157%
Y-o-Y for 9MFY25

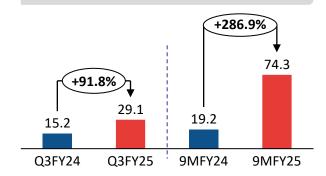
**EBIDTA** up by **287%**Y-o-Y for 9MFY25

PAT after MI up by 162%
Y-o-Y for 9MFY25

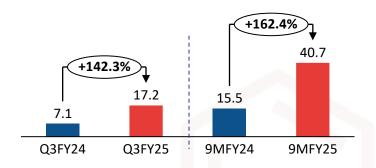
### Income (INR Crs)



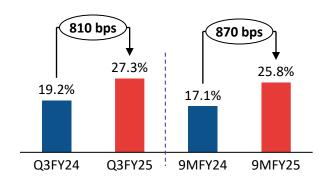




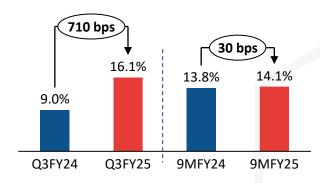
### PAT after MI (INR Crs)



### EBITDA Margin (%) \*\*



### PAT Margin after MI (%)



<sup>\*</sup>Q3FY24 includes financials of Ratnatris Pharmaceuticals for only 17 days on account of the acquisition of a majority stake on 14th December 2023

<sup>\*\*</sup> EBITDA includes Other Operating Income but excludes non-operating income.

# Q3 & 9M FY25 Profit & Loss Account \*



Profit and Loss (INR Crs)	Q3 FY25	Q3 FY24	YoY	9M FY25	9M FY24	YoY
Revenue	103.0	78.7	30.9%	284.0	112.1	153.4%
Other Operating Income	3.4	0.0		4.1	0.0	
Total Income	106.4	78.7	35.2%	288.1	112.1	157.0%
Total COGS	40.7	48.3		123.3	60.7	
Gross Profit	65.7	30.4	116.2%	164.9	51.3	221.1%
Gross Margin (%)	61.7%	38.6%	2,310 bps	57.2%	45.8%	1,140 bps
Employee Cost	19.0	8.2		45.7	17.2	
Other Expenses	17.6	7.1		44.8	14.9	
EBITDA	29.1	15.1	91.8%	74.3	19.2	287.0%
EBITDA Margin (%)	27.3%	19.2%	810 bps	25.8%	17.1%	870 bps
Depreciation	4.2	3.2		11.6	6.3	
Other Income	1.8	0.8		3.4	0.9	
EBIT	26.6	12.7	109.7%	66.1	13.8	380.8%
Finance Cost	6.2	2.4		16.2	6.4	
Profit before Tax	20.5	10.3	99.7%	49.9	7.3	580.2%
Tax + Minority Interest	3.3	3.2		9.2	-8.2	
Profit after Tax and Minority Interest	17.2	7.1	142.2%	40.7	15.5	162.4%
Profit after Tax and Minority Interest Margin (%)	16.1%	9.0%	710 bps	14.1%	13.8%	30 bps
EPS (INR)	3.5	4.0		12.1	7.5	

# 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 December 31, 2024	Amount Unutilized as on December 31, 2024	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		
2	Re-payment/pre-payment, in full or in part, of certain borrowings availed by our Company	73.5	-	73.5	Out of the unutilized amount Rs 200 crores parked as Fixed Deposit	
3	Investment in our Subsidiary, namely, Havix, for repayment/pre-payment in full or in part, of certain borrowings availed by such Subsidiaries	20.2	-	20.2		
4	Funding the working capital requirements of our Company	43.3	-	43.3	with HDFC Bank, Rs 250 crores parked as Fixed Deposit with ICICI Bank and balance held in HDFC bank account	
5	Investment in our Subsidiaries, namely, SPI to fund their working capital requirements	28.8	-	28.8		
6	Investment in our Subsidiaries, namely Ratnatris to fund their working capital requirements	30.7	-	30.7		
7	Funding inorganic growth through acquisition and other strategic initiatives and general corporate purposes	154.4	-	154.4		
8	Offer expenses	42.2	10.0	32.2		
	Total	500.0	10.0	490.0		

About Senores
Pharmaceuticals
Limited

# Introduction to Senores Pharma



### **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**

1USFDA & DEA approved formulation facility in the US

### **Partnerships**

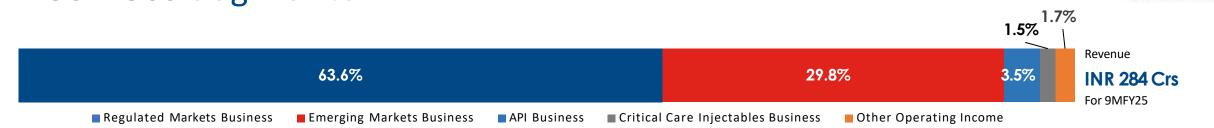
Partnerships with leading pharmaceutical companies across Regulated as well as Emerging Markets

### 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

**24** Approved ANDAs

**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



Approval for manufacturing facility from regulatory bodies of 10 countries

Product registration for **267 Products** 

Product applications filed for **537 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 **35** employees

Launched 55 Products

# **Key Events & Milestones**





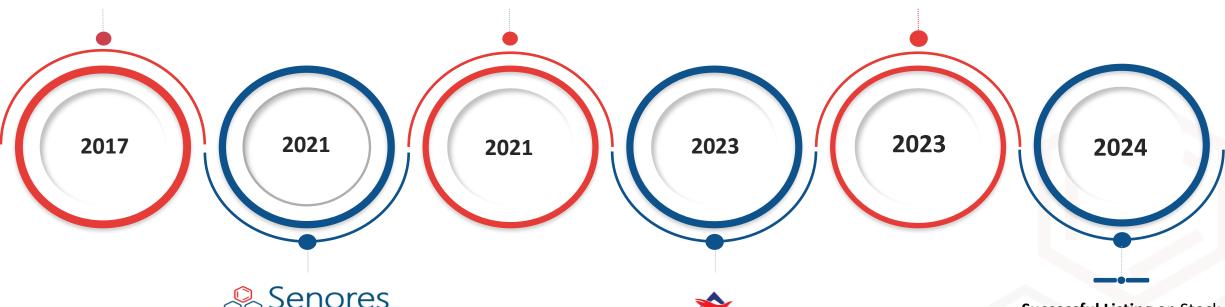
Senores Pharmaceuticals Inc.

Ratnatris
Pharmaceuticals
Your Health, Our Priority!

Incorporated as "Senores Pharmaceuticals Pvt. Ltd." in India.

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

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



Entered into a new segment of API with acquisition of Ratnagene Lifescience Pvt. Ltd.<sup>1</sup>

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

Successful Listing on Stock
Exchanges in India on BSE & NSE



# Key Strengths of the Company



1 Distinct specialty and complex product portfolio built in a short span for Regulated Markets



2 Long-term **marketing arrangements** with pharmaceutical companies in the Regulated Markets



3 Presence in the **Emerging Markets** with portfolio, including **specialty** and **complex products** 



4 Strong R&D capabilities driving differentiated portfolio of products



5 US FDA approved facility in the US catering to the Regulated & Semi-Regulated Markets and WHO-GMP compliant facility in India catering to Emerging Markets



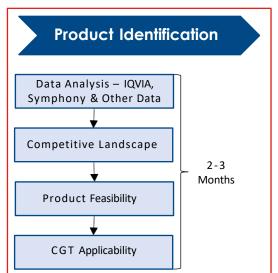
6 Professional & dedicated management teams for the diverse business verticals

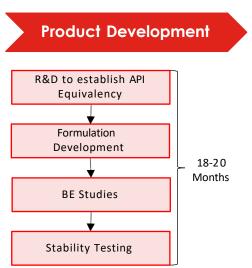
# Distinct Specialty & Complex Product Portfolio Built in a



# Short Span for Regulated Markets

Demonstrated Capability to propel Products from Conception to Commercialization





# Partner Finalization Concurrent Process Dossier Compilation 1-2 Months Filing Immediate Post Compilation Approval 10-15 months post filing Launch Immediately Post Approval

### **ANDA Products**



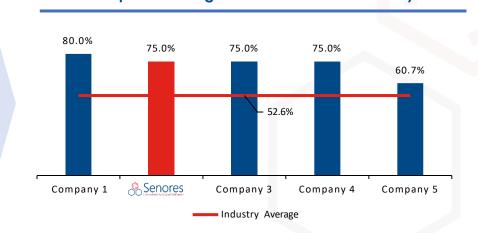
### 2nd Highest in terms of proportion of CGT Exclusivity amongst Peers

- 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 30<sup>th</sup> 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
- **≥ 24** Approved **ANDAs**

### 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

345



**Specialty and Complex Products** 

### **Special Focus**

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



804 Total **Products** 

267 Product Registrations

537 Product **Applications Filed** 



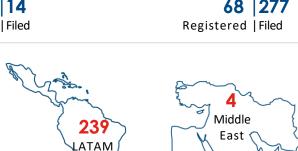
132

Africa

25 | 107

Registered | Filed

South East 7 \_ Asia 70 | 14 68 | 277 Registered | Filed







**Ferric** Carboxymaltose



**Tofacitinib** 



Sacubitril + Valsartan



**Apixaban** 



Eltrombopag **Olamine** 



**Diverse Business Models** 

P2P Model

**CDMO Model** 

**Distributor Model** 

Own Brands<sup>2</sup>

# Top Emerging Markets<sup>1</sup>

**Philippines** 

Peru











# Partners to Global Pharmaceutical Companies in Regulated & Emerging Markets



























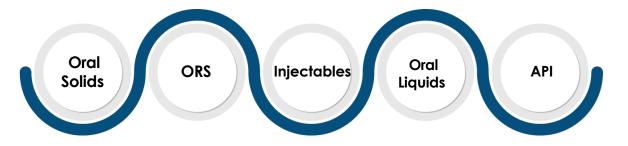




# Strong R&D Capabilities Driving Differentiated Product



## 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

- **54** 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**

Manufacturing Lines

185,300 Sq. Ft. Total Area

### **OSD**

### 1.2 Bn units

**Annual Installed** Capacity<sup>1</sup>



### 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

**Dosage Forms** 

### **General Oral** Dosage

1397.96 Mn

**Annual Installed** Capacity<sup>1</sup>

### **Injectables**

49.92 Mn

**Annual Installed** Capacity<sup>1</sup>

### **Beta Lactum** Orals

511.68 Mn

**Annual Installed** Capacity1





FDA

**PHILIPPINES** 

**NIGERIA NAFDAC** 

MOH



**PERU** 

**DIGEMID** 





**ZAMBIA** МОН

**LIBERIA** \* МОН

# **Corporate Structure**





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





Havix Group Inc
Caters to Regulated Market

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

66.58%<sup>1</sup> Subsidiary



Ratnatris Pharmaceuticals Pvt. Ltd.
Caters to Emerging Market

Development, Manufacturing and Marketing of Generic Pharmaceutical Formulations

69% Subsidiary

Senores Pharmaceuticals Inc.

Marketing cum ANDA

Holding Company

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

100% Subsidiary

# Professional and Dedicated Management Team





Swapnil Jatinbhai Shah
Promoter & Managing Director

- 16+ 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

- 39+ 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

- 24+ 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



# **Key Strategies**





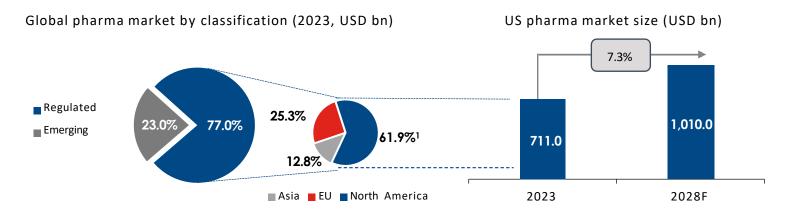
# Enhance Market Presence of Marketed Products in North

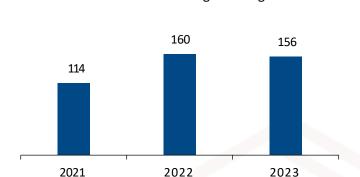


# America and Other Regulated Markets

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





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





Source: F&S Report

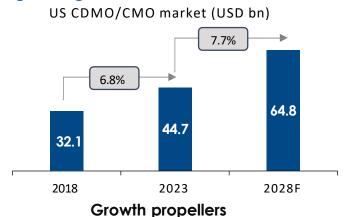
25 Note: 1: Pharma market value for North America is USD 778.8 bn, Europe is USD 318.5 bn & Asia is USD 161.3 bn

# Entering into Strategic Alliance for CMO/CDMO in Regulated



# Markets

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



# Growing drug complexity Rapid technological turnaround Upcoming loss of exclusivity for drugs Business model shift from Capex to Opex Expected Increase in average outsourcing penetration US accounts for 40-45% of CDMO markets

### Strategic alliance for CMO/CDMO in Regulated Markets

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



### Offering benefits such as

Focus on core competencies

Cost Advantages

Early to Market advantage

Flexibility & Scalability

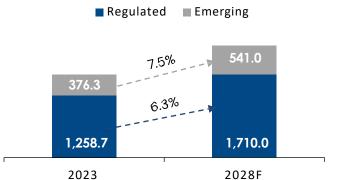
Access to global talent & advanced technologies

# **Expanding into New Regulated & Emerging Markets**

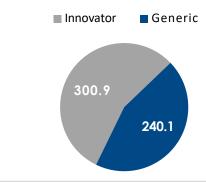


### Regulated & Emerging markets witnessing strong growth

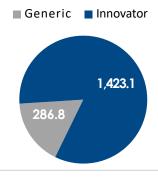
Global pharma market by classification (USD bn)



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

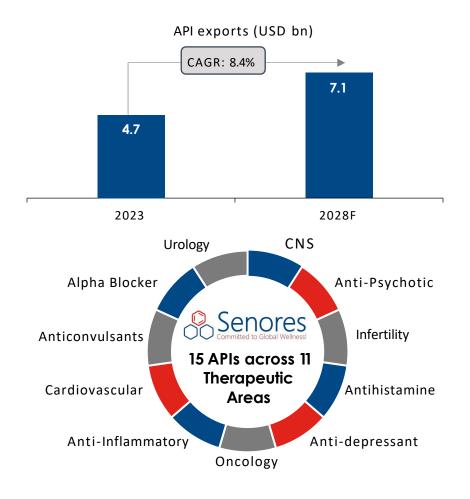


Multiple marketing and distribution models to enhance presence in Emerging market countries

# **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







Flexibility & Innovation





Diversified revenue stream & business resiliance

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



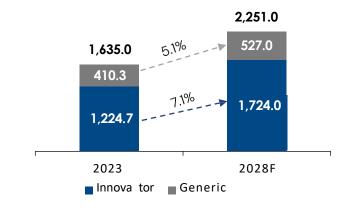
## Global Pharma Market



30

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

Global pharma market by product type (USD bn)

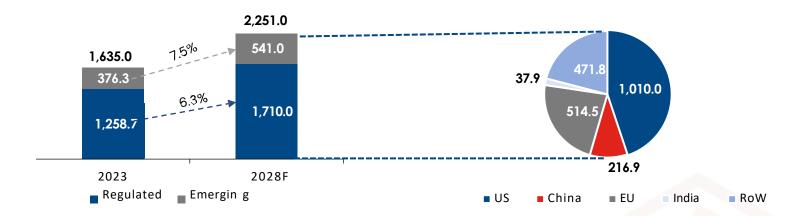


# ...regulated market will continue to dominate the global pharma market

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

 $^{\sim}$  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

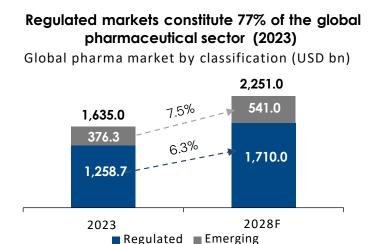
### Demand from developing nations

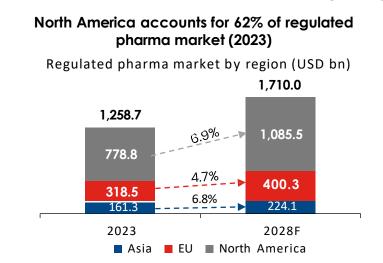
Rising instances of chronic conditions and persistent burden of infectious diseases drive dual demand

# Regulated Pharma Market



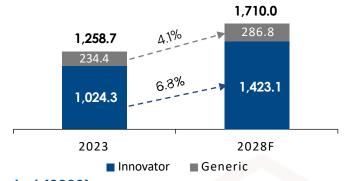
Regulated markets to dominate the global pharmaceutical sector, driven by their access to a growing innovative drug market & a thriving generics market





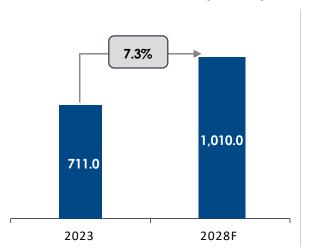
# New generics and patent expirations to drive drug market growth

Regulated pharma market by product type (USD bn)

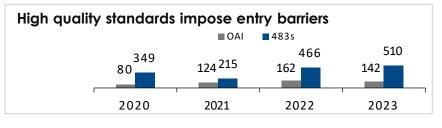


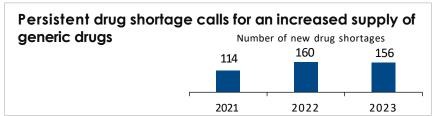
US accounts for 43% of the global pharma market, 56% of the regulated market and 91% of North American market (2023)

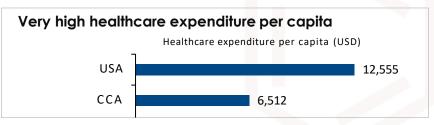
US Pharma Market size (USD bn)

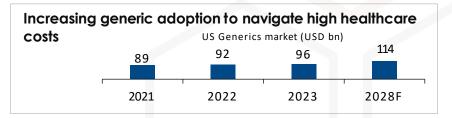


### Characteristics of US healthcare market









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

Source: F&S Report

CCA: Comparable Country Average, OAI: Official Action Indicated

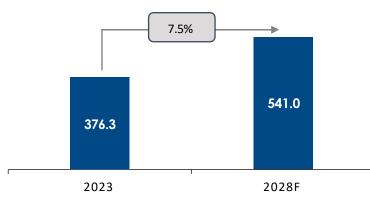
# **Emerging Pharma Market**



### **Emerging pharma market**

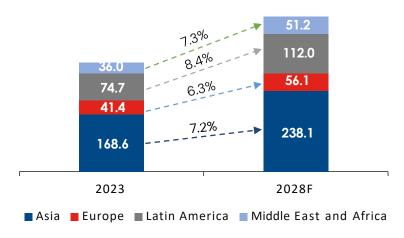
### Emerging markets to outpace developed ones...

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)



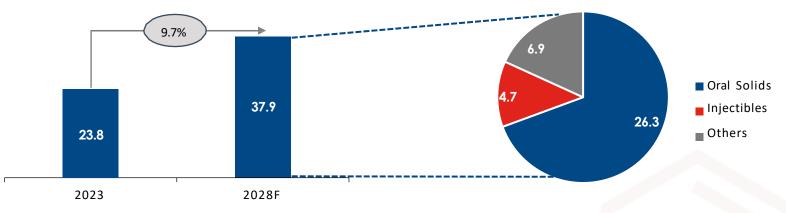
### Overview of India pharma market

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

Indian pharma market size (USD bn)

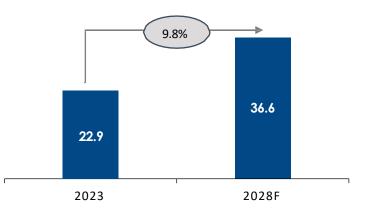
### Oral solids will dominate IPM

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



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

Indian generic pharma market size (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

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