

Product supply

CMD24
CAPITAL MARKETS DAY

7 MARCH



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Forward-looking statements

Novo Nordisk's reports filed with or furnished to the US Securities and Exchange Commission (SEC), including the statutory Annual Report 2023 and Form 20-F, which both were filed with the SEC in January 2024 in continuation of the publication of the Annual Report 2023, this presentation, and written information released, or oral statements made, to the public in the future by or on behalf of Novo Nordisk, may contain forward-looking statements. Words such as 'believe', 'expect', 'may', 'will', 'plan', 'strategy', 'prospect', 'foresee', 'estimate', 'project', 'anticipate', 'can', 'intend', 'target' and other words and terms of similar meaning in connection with any discussion of future operating or financial performance identify forward-looking statements. Examples of such forward-looking statements include, but are not limited to:

- Statements of targets, plans, objectives or goals for future operations, including those related to Novo Nordisk's products, product research, product development, product introductions and product approvals as well as cooperation in relation thereto,
- Statements containing projections of or targets for revenues, costs, income (or loss), earnings per share, capital expenditures, dividends, capital structure, net financials and other financial measures,
- Statements regarding future economic performance, future actions and outcome of contingencies such as legal proceedings, and
- Statements regarding the assumptions underlying or relating to such statements.

These statements are based on current plans, estimates and projections. By their very nature, forward-looking statements involve inherent risks and uncertainties, both general and specific. Novo Nordisk cautions that a number of important factors, including those described in this presentation, could cause actual results to differ materially from those contemplated in any forward-looking statements.

Factors that may affect future results include, but are not limited to, global as well as local political and economic conditions, such as interest rate and currency exchange rate fluctuations, delay or failure of projects related to research and/or development, unplanned loss of patents, interruptions of supplies and production, including as a result of interruptions or delays affecting supply chains on which Novo Nordisk relies, shortages of supplies, including energy supplies, product recalls, unexpected contract breaches or terminations, government- mandated or market-driven price decreases for Novo Nordisk's products, introduction of competing products, reliance on information technology including the risk of cybersecurity breaches, Novo Nordisk's ability to successfully market current and new products, exposure to product liability and legal proceedings and investigations, changes in governmental laws and related interpretation thereof, including on reimbursement, intellectual property protection and regulatory controls on testing, approval, manufacturing and marketing, perceived or actual failure to adhere to ethical marketing practices, investments in and divestitures of domestic and foreign companies, unexpected growth in costs and expenses, strikes and other labour market disputes, failure to recruit and retain the right employees, failure to maintain a culture of compliance, epidemics, pandemics or other public health crises, the effects of domestic or international crises, civil unrest, war or other conflict and factors related to the foregoing matters and other factors not specifically identified herein.

For an overview of some, but not all, of the risks that could adversely affect Novo Nordisk's results or the accuracy of forward-looking statements in the Annual Report 2023, reference is made to the overview of risk factors in 'Risk Management' of the Annual Report 2023.

Unless required by law, Novo Nordisk has no duty and undertakes no obligation to update or revise any forward-looking statement after the distribution of the Annual Report 2023, whether as a result of new information, future events, or otherwise.

Important drug information

Victoza® and Ozempic® are approved for the management of type 2 diabetes only
Saxenda® and Wegovy® are approved for the treatment of obesity only

Strategic aspirations 2025



Purpose and sustainability (ESG)

- Progress towards zero environmental impact
- Being respected for adding value to society
- Being recognised as a sustainable employer



Innovation and therapeutic focus

- Further raise the innovation-bar for diabetes treatment
- Develop a leading portfolio of superior treatment solutions for obesity
- Strengthen and progress the Rare disease pipeline
- Establish presence in Cardiovascular & emerging therapy areas



Commercial execution

- Strengthen Diabetes leadership - aim at global value market share of more than 1/3
- More than 25 billion DKK in Obesity sales by 2025
- Secure a sustained growth outlook for Rare disease



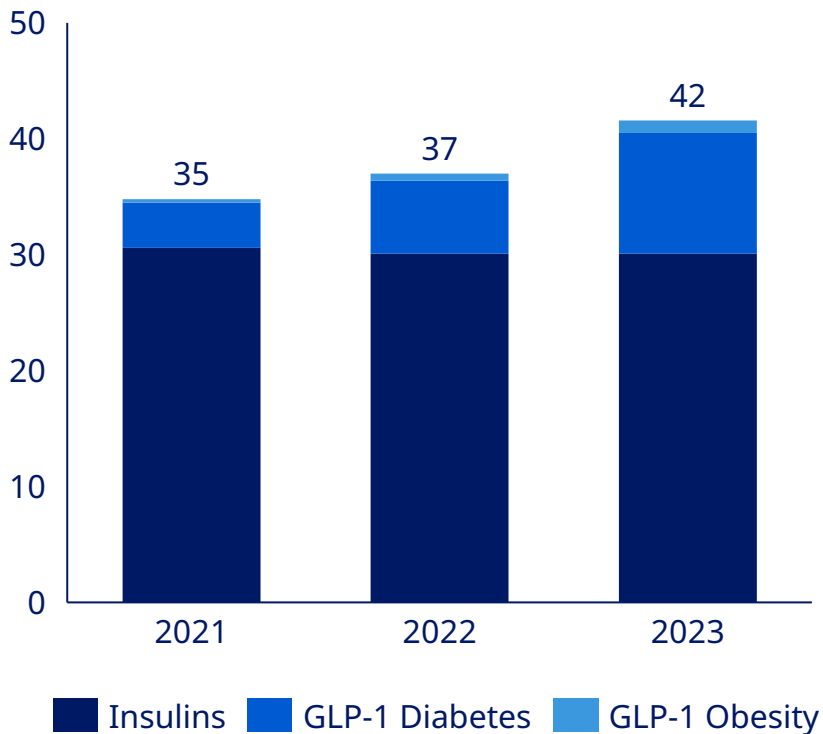
Financials

- **Deliver solid sales and operating profit growth**
- **Drive operational efficiencies across the value chain to enable investments in future growth assets**
- **Deliver free cash flow to enable attractive capital allocation to shareholders**





Product supply has continued step-up in investments and employees to support growth

Patient reach has accelerated since 2021

Million patients on NN products



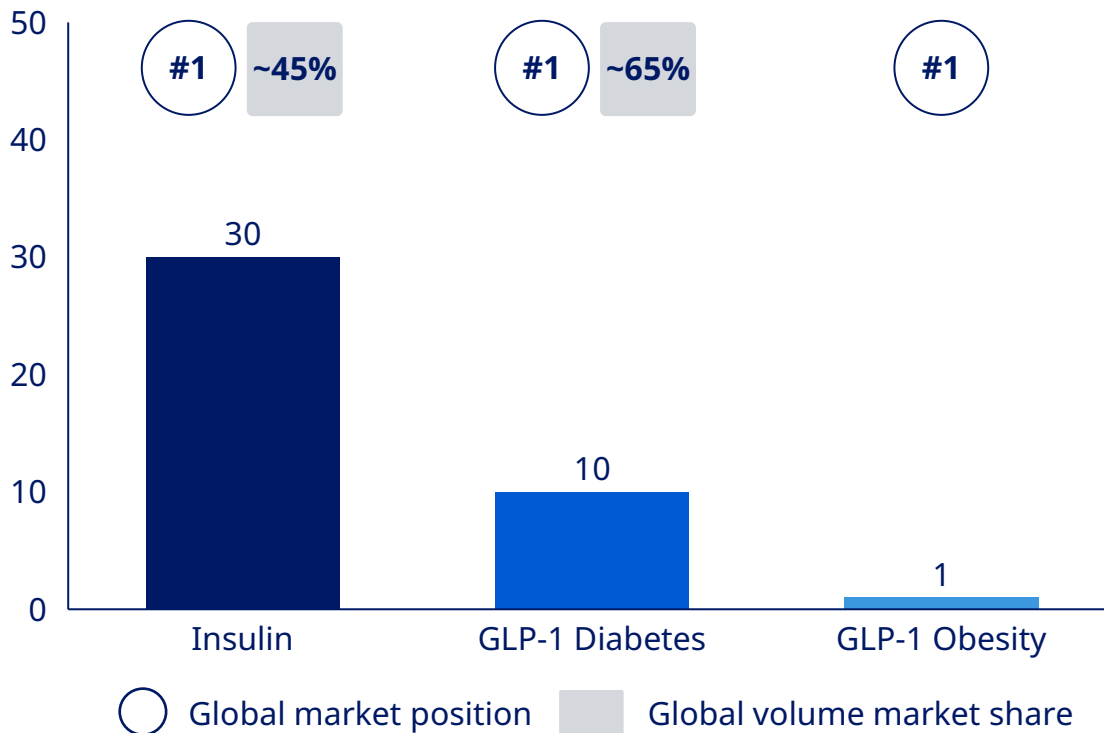
Product supply has expanded to enable the current growth

	2021	2023
 Number of employees	~16,000 employees	~25,000 employees
 CAPEX investment level CAPEX to sales ratio	6 bDKK 4%	26 bDKK 11%
 Ozempic® devices	Index 100	Index ~300
 Semaglutide API	Index 100	Index ~400

Manufacturing scale and expertise within biologics is a competitive advantage for Novo Nordisk

The world's largest manufacturer of insulin and GLP-1

Million patients on NN products in 2023



Novo Nordisk competitive advantages in manufacturing



Decades of experience with high volume production of core yeast and mammalian API platforms

API scalability and yield optimisation driven by continuous production technology

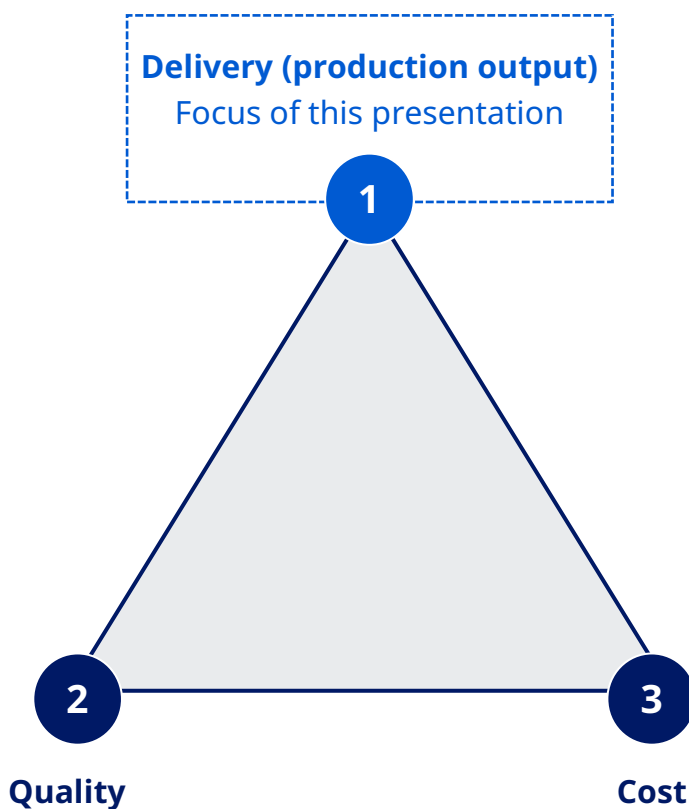


High volume installed capacity for biologics

In-house expertise in the development and manufacturing of devices

Manufacturing strategy focuses on quality, cost and delivery

Manufacturing focus areas



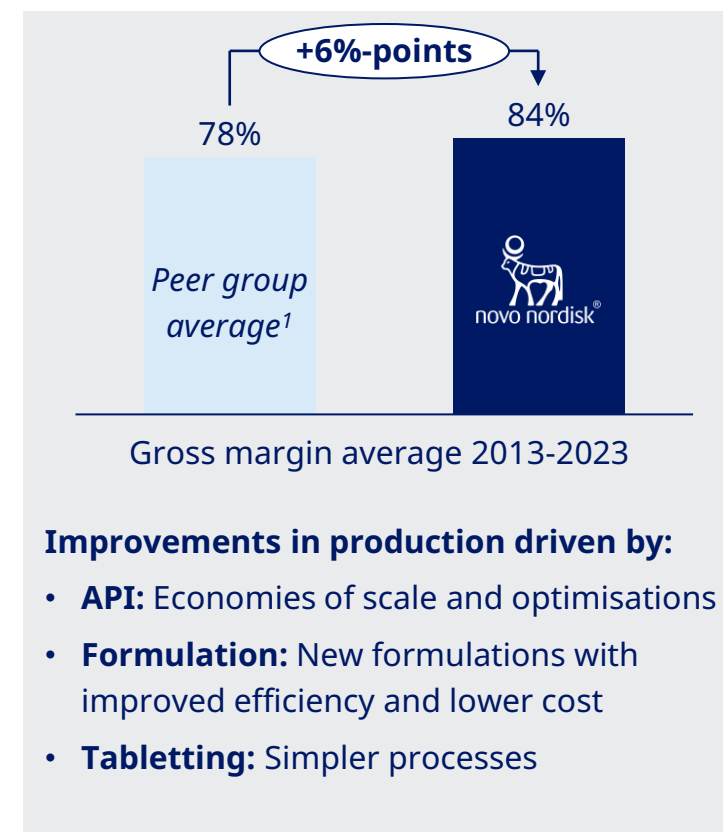
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Maintain highest quality



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Drive constant improvements



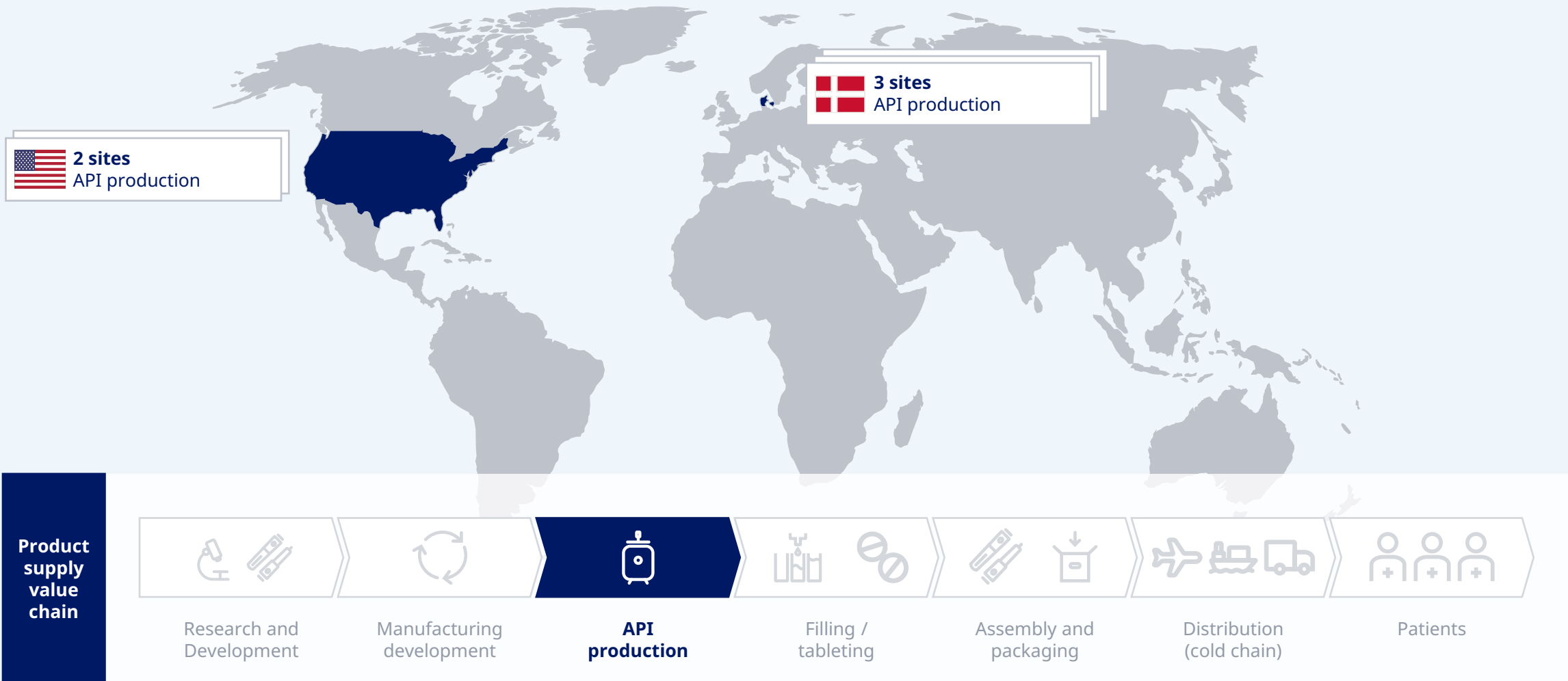
¹Peers include AbbVie, Amgen, AstraZeneca, Biogen, Boehringer Ingelheim, BMS, Eli Lilly, Gilead, GlaxoSmithKline, Johnson & Johnson, Merck, Novartis, Pfizer, Roche, Sanofi. NAI average also incl. BioGenetics, Celgene, Lundbeck, Shire HGT, Teva, UCB

²No action indicated (NAI) ratio calculated as number of times a company got no actions from the FDA during an inspection divided by the total number of inspections during 2013-2023

API: Active pharmaceutical ingredient; FDA: US Food and Drug Administration; NAI: No action indicated

Sources: FDA data dashboard; Evaluate Pharma


The strategically important active pharmaceutical ingredient sites in Novo Nordisk are based in Denmark and the US



API platforms in Novo Nordisk are mostly yeast based and the key to expand capacity is investments and optimisations



API production platforms

Scale	Platform	Therapy areas
Largest  Smallest	Yeast	<ul style="list-style-type: none"> Diabetes Obesity
	Mammalian cells	<ul style="list-style-type: none"> Rare disease Cardiovascular and emerging therapy areas
	E. Coli	<ul style="list-style-type: none"> Rare disease
	Organic synthesis	<ul style="list-style-type: none"> Cardiovascular and emerging therapy areas

Strategy for API expansion



CAPEX investments to expand in-house capacity across strategic sites in DK and US

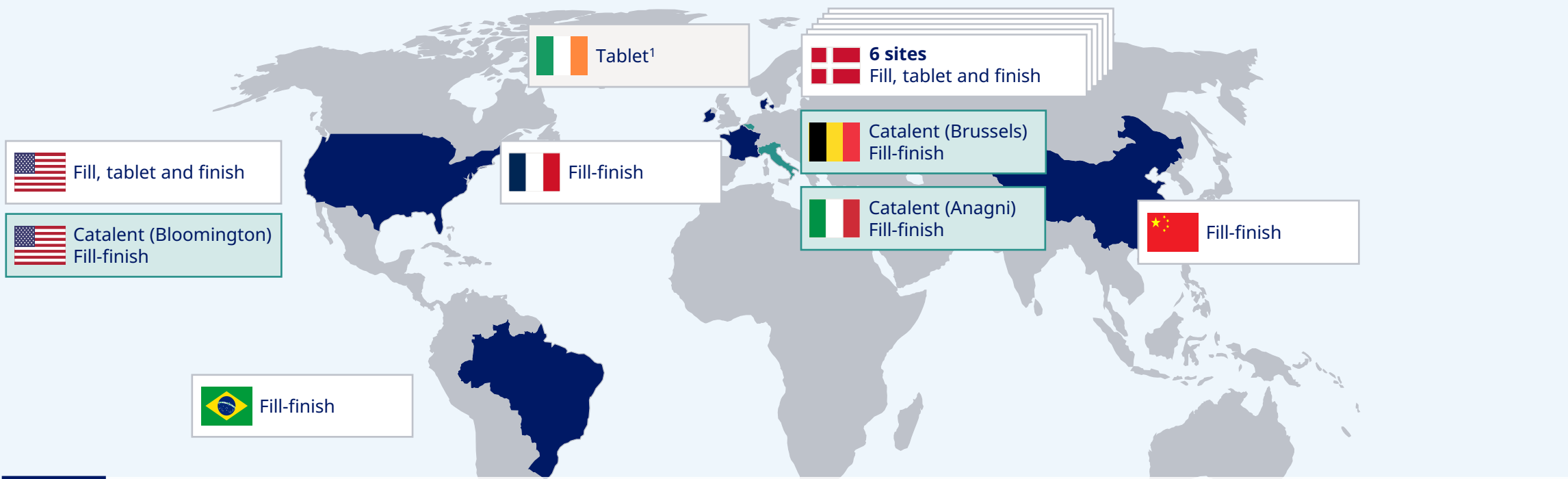


Continuously optimise yield with technology upgrades and simpler processes



Build for flexibility to cater for oral and injectable portfolio depending on demand

The global fill-finish footprint is expected to expand from 11 to 14 sites with the acquisition of the three Catalent sites



¹The Alkermes transaction (Dec 2023): Expected to close in mid-2024
 API: Active pharmaceutical ingredient
 Note: There are local production facilities in Algeria, Iran, Japan, and Russia

New sites pending closing of the Catalent transaction

Catalent fill-finish sites are expected to start adding additional capacity from 2026

The three Catalent fill-finish sites



Bloomington site (Indiana, US)



Brussels site (Belgium)



Anagni site (Italy)



After closing, Novo Nordisk will honour all customer obligations at the three Catalent sites that Novo Nordisk is acquiring

The acquisition will help expand capacity faster

- Will help reach more patients with current and future treatments
- Enables faster expansion of manufacturing capacity at scale, while providing future optionality and flexibility
- The three sites are fully operational and employ >3,000 people
- The acquisition is expected to gradually increase Novo Nordisk's fill-finish capacity from 2026 and onwards

The acquisition is expected to be completed towards the end of 2024 upon satisfaction of various customary closing conditions

Novo Nordisk has several device platforms and plans to expand capacity both internally and externally



Device platforms (finished product)

Durable¹

Vials

FlexTouch® multi dose²

Single dose

Tablets



Indicative device use per year (using GLP-1 as example)

1

~12

13

52

365

Strategy for fill-finish expansion



Step-up CAPEX investments to expand in-house capacity across strategic sites



CMO partnerships to strengthen network flexibility and speed to establish new capacity



Transition to less frequent dosing

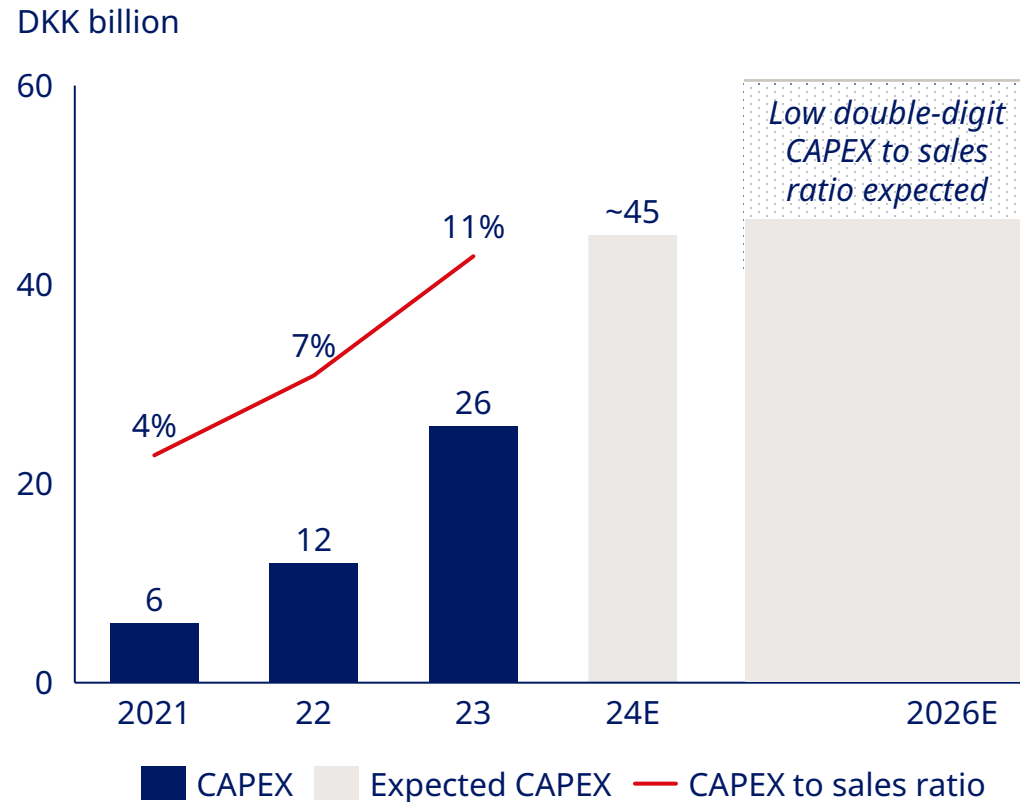


Building for flexibility to cater for oral and injectable portfolio depending on demand

¹The device shown here is the Novopen®6 device currently used for insulin products; ² This includes the older generation multi-dose FlexPen® device currently used for insulin and liraglutide
 CAPEX: Capital Expenditure; CMO: Contract Manufacturing Organisation

Significant step-up in CAPEX investments across the full value chain to enable growth for current and future products

CAPEX investments



Several large investments announced since 2021

Announced	Site	Scope	Investment
2021 December	Kalundborg Denmark	Mainly API	17 bDKK
2022 November	Bagsværd Denmark	Clinical API	5 bDKK
2023 June	Hillerød Denmark	API for CETA	16 bDKK
2023 November	Kalundborg Denmark	Mainly API	42 bDKK
2023 November	Chartres France	Fill-Finish	16 bDKK
2023 December	Athlone Ireland	Oral portfolio	1 bDKK

Typical construction timelines: API: 5+ years | Fill-finish: 3+ year

API: Active pharmaceutical ingredient; CAPEX: Capital expenditures; CETA: Cardiovascular and emerging therapy areas
 Note: Investment figures have been rounded

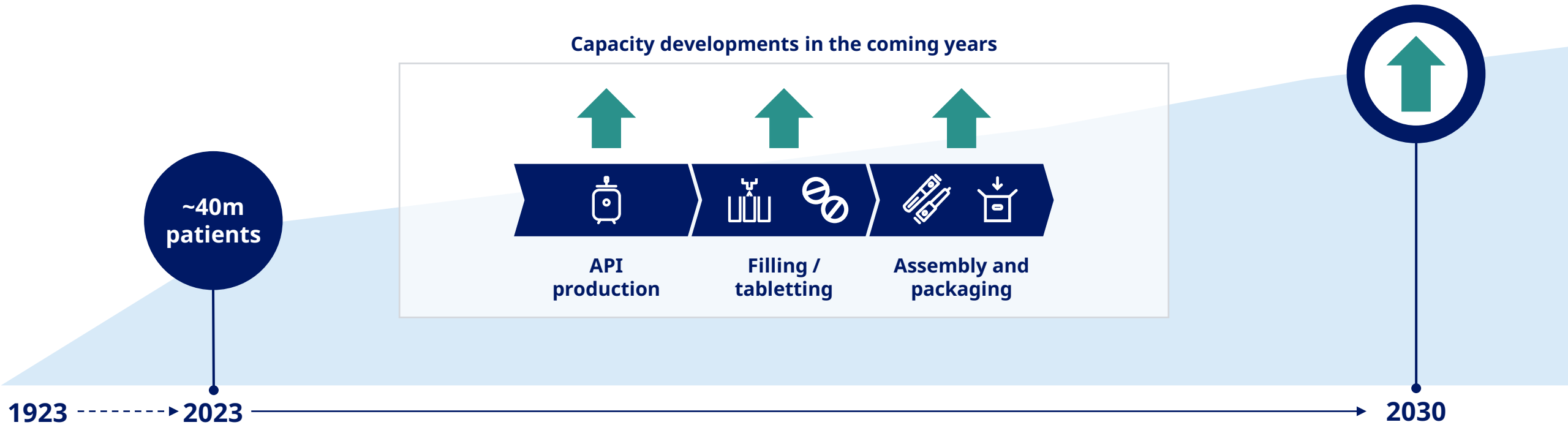
Plans are in place for scaling portfolio of GLP-1 treatments



¹External API contracts in place for cagrilintide
 API: Active Pharmaceutical Ingredient; sc: Subcutaneous; Sema: Semaglutide; SDD: Single-dose device

Investments across the full manufacturing value chain to significantly increase patient reach towards 2030

ILLUSTRATIVE



Closing remarks

Manufacturing scale and expertise with biologics is a competitive advantage for Novo Nordisk

Continued CAPEX investments and scaling across the full value chain for current and future products

Plans in place expected to significantly increase patient reach towards 2030

