

42nd Annual J.P. Morgan Healthcare Conference

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January 2024

Disclaimer and safe harbor

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The implementation of the strategy has to be submitted to the relevant staff representation authorities in each country concerned, in compliance with the specific procedures, terms and conditions set forth by each national legislation.

In those countries in which public or private-health cover is provided, Ipsen is dependent on prices set for medicines, pricing and reimbursement-regime reforms and is vulnerable to the potential withdrawal of certain medicines from the list of reimbursable medicines by governments, and the relevant regulatory authorities in its locations.

Ipsen operates in certain geographical regions whose governmental finances, local currencies or inflation rates could erode the local competitiveness of Ipsen's medicines relative to competitors operating in local currency, and/or could be detrimental to Ipsen's margins in those regions where Ipsen's sales are billed in local currencies.

In a number of countries, Ipsen markets its medicines via distributors or agents; some of these partners' financial strengths could be impacted by changing economic or market conditions, potentially subjecting Ipsen to difficulties in recovering its receivables. Furthermore, in certain countries whose financial equilibrium is threatened by changing economic or market conditions, and where Ipsen sells its medicines directly to hospitals, Ipsen could be forced to lengthen its payment terms or could experience difficulties in recovering its receivables in full.

Ipsen also faces various risks and uncertainties inherent to its activities identified under the caption 'Risk Factors' in the Company's Universal Registration Document.

All of the above risks could affect Ipsen's future ability to achieve its financial targets, which were set assuming reasonable macroeconomic conditions based on the information available today.

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Achievements since 2020

A pure-play biopharmaceutical company

OUR VISION

To be a leading global mid-sized biopharmaceutical company with a focus on transformative medicines in **Oncology**, Rare Disease & Neuroscience

SIGNIFICANT PROGRESS

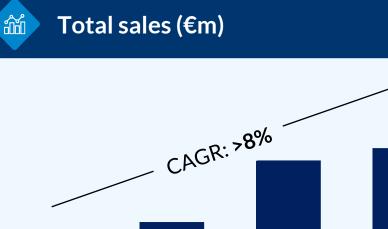
- **Double-digit performances by** growth platforms
- **Divestment** of Consumer HealthCare
- **External innovation**: >20 new programs & three late-stage transactions

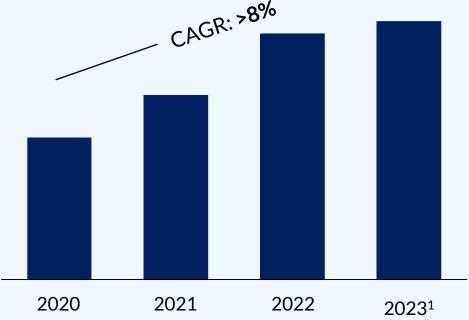




Epizyme^{*}







CAGR: compound annual growth rate.¹ Based on FY 2023 guidance. Total sales here exclude new medicines (Tazverik & Bylvay) & are adjusted for divestment of Consumer HealthCare & are at constant exchange rates.

Our growth journey

Next phase of transformation built on solid foundations

2020-2023

Setting foundations

New strategy

Focus on **Specialty Care**

2024-2027

Dynamic growth

Several launches

>> Further pipeline expansion

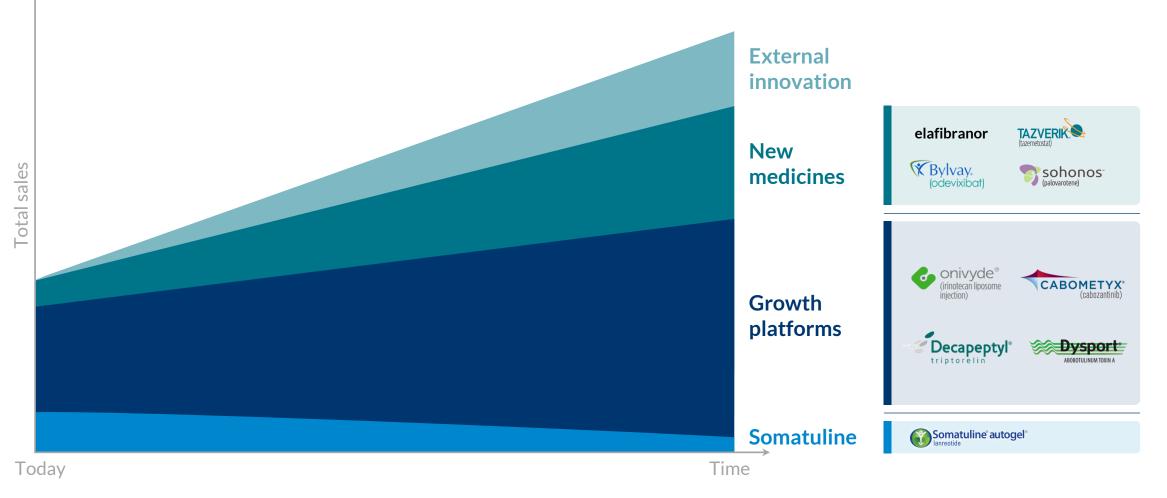
2028+

Lasting momentum

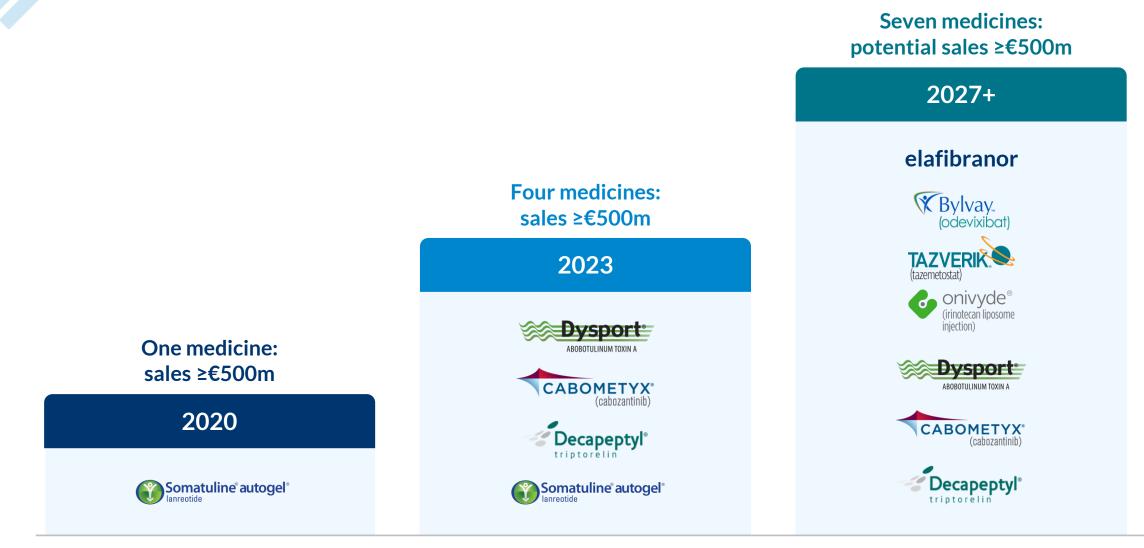
- Balanced & diversified portfolio across three therapy areas
- Sustained growth, supported by pipeline & external innovation

A strong platform for growth

Growth platforms & new medicines continue to drive momentum

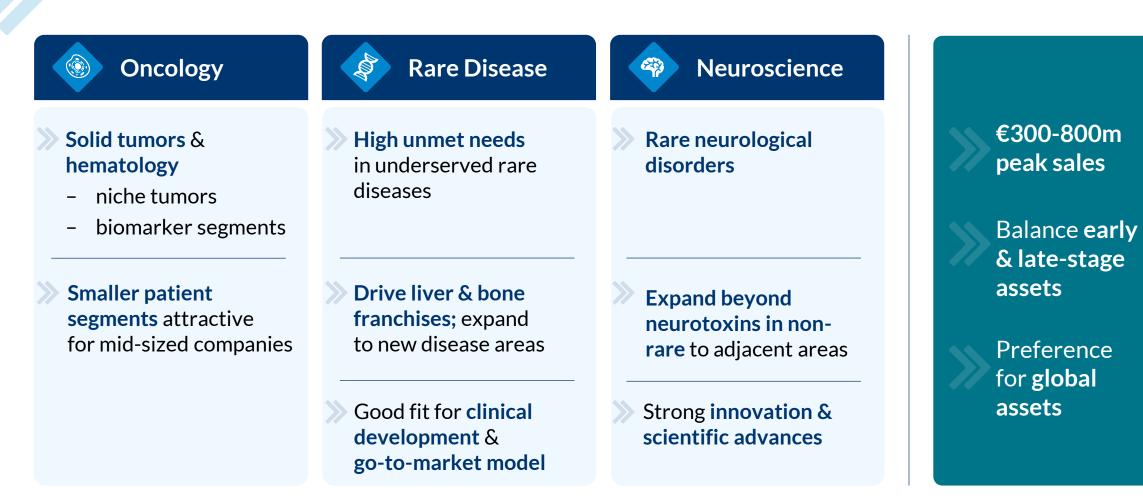


Increasingly diversified portfolio



2023 sales based on latest available consensus forecasts.

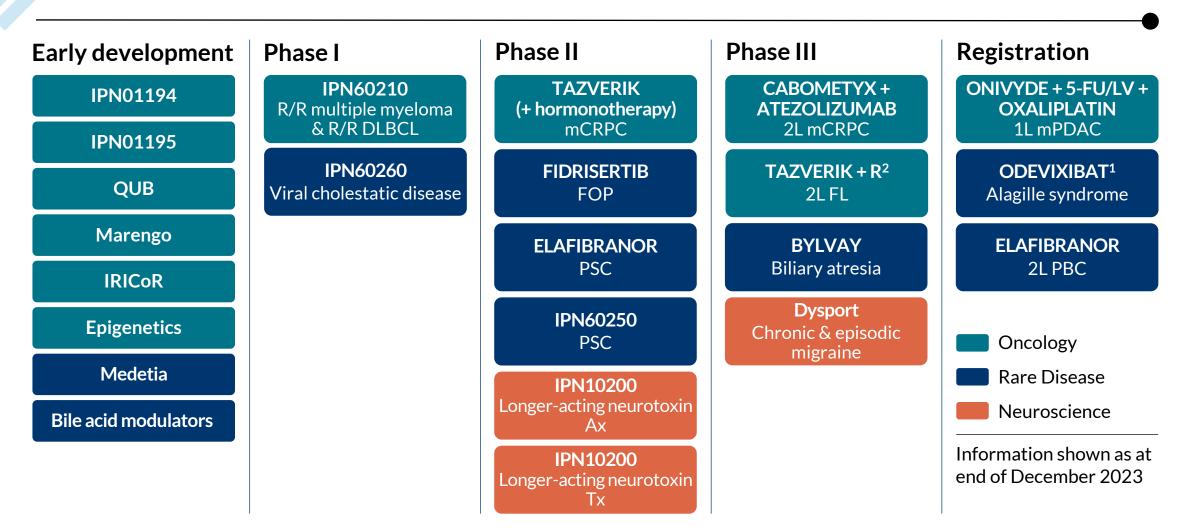
Clear strategy to continue external innovation



Cumulative firepower of up to €5bn¹ by 2027

Fuelling a high-value, sustainable pipeline

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R/R: relapsed/refractory; DLBCL: diffuse large B-cell lymphoma; mCRPC: metastatic castration-resistant prostate cancer;

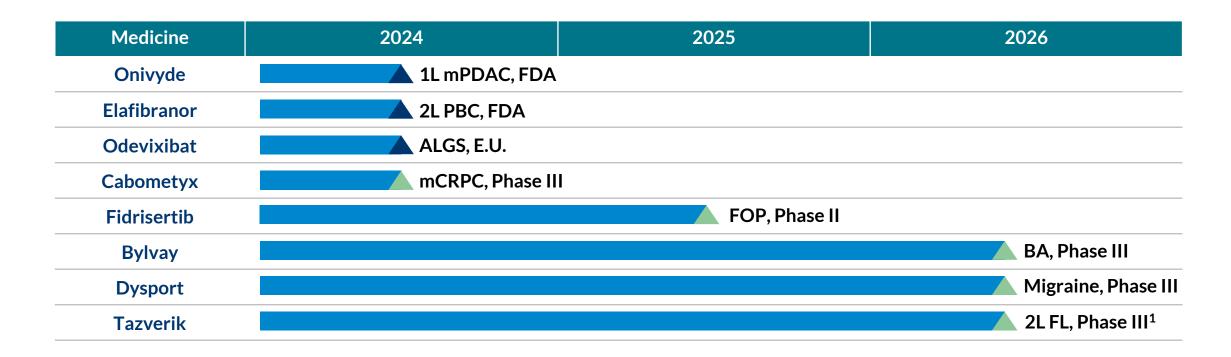
FOP: fibrodysplasia ossificans progressiva; PSC: primary sclerosing cholangitis; Ax: aesthetics; Tx: therapeutics; R²: lenalidomide + rituximab; 2L: second line;

FL: follicular lymphoma; 1L: first line; mPDAC: metastatic pancreatic ductal adenocarcinoma; PBC: primary biliary cholangitis. ¹ E.U.

Near to mid-term outlook



Key milestones



1L: first line; mPDAC: metastatic pancreatic ductal adenocarcinoma; FDA: U.S. Food & Drug Administration; 2L: second line;

PBC: primary biliary cholangitis; **ALGS**: Alagille syndrome; **mCRPC**: metastatic castration-resistant prostate cancer;

FOP: fibrodysplasia ossificans progressiva; **BA**: biliary atresia; **FL:** follicular lymphoma. ¹ Early data readout anticipated, pending regulatory agreement. Disclaimer: trials are event-driven & timings can change.

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Launching four new medicines or new indications in near term

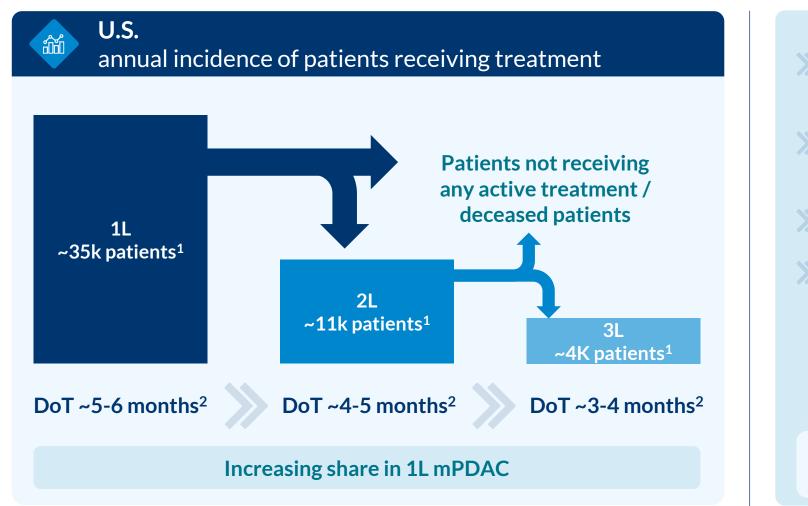
Building Rare Disease franchise & strengthening Oncology

Medicine	Indication	Market	Expected regulatory-decision date
cirinotecan liposome injection)	1L mPDAC	U.S. only	FDA: 13 February 2024
(odevixibat)	ALGS	Global ¹	U.S. launch underway EMA: 2024
elafibranor	2L PBC	Global ²	FDA: 10 June 2024 EMA: H2 2024
sohonos ⁻ (palovarotene)	FOP	U.S. & selected RoW	U.S. launch underway

1L: first line; mPDAC: metastatic pancreatic ductal adenocarcinoma; ALGS: Alagille syndrome; 2L: second line; PBC: primary biliary cholangitis; FOP: fibrodysplasia ossificans progressiva; RoW: Rest of World; FDA: U.S. Food & Drug Administration; EMA: European Medicines Agency. ¹ Excludes Japan. ² Excludes China, Taiwan, Hong Kong & Macau.

Onivyde: significant potential in 1L mPDAC





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High unmet medical need & low one-year survival rate

Phase III NAPOLI-3 trial: positive results

Immediate launch planned

Potential to become new SoC in 1L mPDAC by gaining market share in all segments

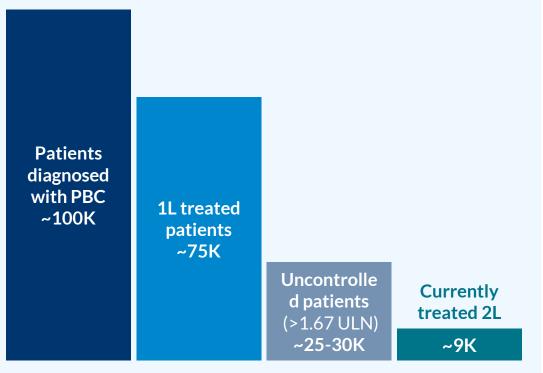
Peak sales expected to exceed €500m³

1L: first line; mPDAC: pancreatic ductal adenocarcinoma; 2L: second line; 3L: third line; DoT: duration of treatment; gem: gemcitabine; PDUFA: Prescription Drug User Fee Act. Sources: ¹ IQVIA Market Sizing report Aug 2022 to Jul 2023; ² Kantar, CancerMPact, Pancreatic Cancer, Treatment Architecture, September 2023. ³ Assumes approval in potential 1L mPDAC indication.

Elafibranor: significant potential

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U.S. example: 2L PBC patient flow: number of U.S. patients



2L-eligible patients

Underdeveloped global 2L PBC market

Significant unmet medical need

- Dissatisfaction with current treatment options
- Uncontrolled disease
- Limited share (20-40%) of eligible patients receiving 2L treatment today
 - Patient eligibility not well defined by HCPs
- New entrants to expand market by accelerating number of patients under 2L treatment
- Global 2L PBC market estimated at ~€1.5bn (2030)
- Customer overlap in rare liver disease with Bylvay

Peak sales expected to exceed €500m¹

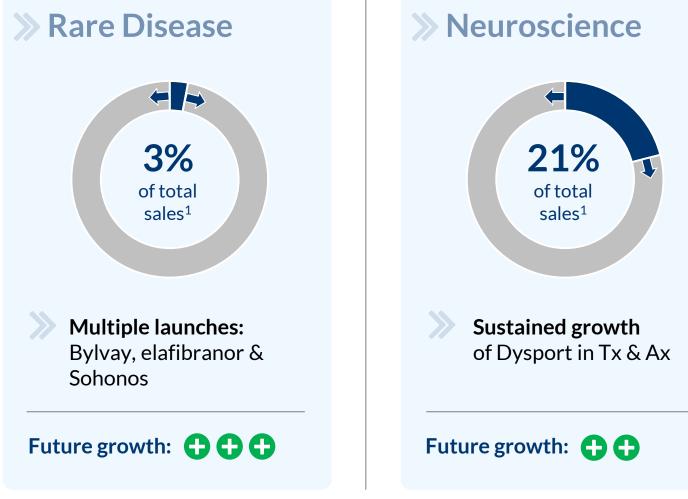
2L: second line; PBC: primary biliary cholangitis; 1L: first line; ULN: upper limit normal; HCPs: healthcare professionals. Source: Lu et al., 2018; Webb et al., 2021; Dahlqvist et al, 2017; Sebode et al, 2020; Pla et al, 2007; Marzioni et al, 2019. ¹ Based only on the potential PBC indication.

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More balanced split of sales by three therapy areas



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1L: first line; mPDAC: metastatic pancreatic ductal adenocarcinoma; Tx: therapeutics; Ax: aesthetics. ¹ Based on September year-to-date 2023 total sales.

Global leader with growth across all regions



Future growth:

Future growth: 🔁

Asia-Pacific & Latin America

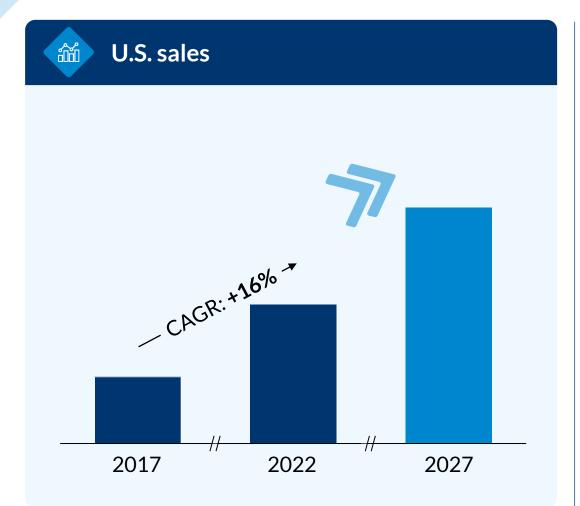
Future growth: **C**

¹Based on September year-to-date 2023 total sales.

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Europe is defined in this presentation as the E.U., the U.K., Iceland, Liechtenstein, Norway and Switzerland.

Strong U.S. growth driven by multiple potential launches





Significant opportunities

Growing footprint in Oncology

- Onivyde 1L mPDAC
- Tazverik

Becoming established in Rare Disease

- Building franchise in rare liver: elafibranor in 2L PBC, Bylvay in PFIC & ALGS
- Launching first treatment in FOP

Growing Dysport Tx in Neuroscience

Driving operating leverage

1L: first line; mPDAC: metastatic pancreatic ductal adenocarcinoma; 2L: second line; PBC: primary biliary cholangitis; PFIC: progressive familial intrahepatic cholestasis; ALGS: Alagille syndrome; FOP: fibrodysplasia ossificans progressiva; Tx: therapeutics.

Prior performance at actual rates.

2027 mid-term outlook

Excluding potential additional late-stage¹ external-innovation opportunities





Growth platforms

Somatuline erosion





Improved SG&A expenses-to-sales ratio

Sustained R&D expenses-to-sales ratio

Conclusion



Sustainability roadmap

Generation lpsen: for positive change

Net zero by 2045

Maintaining progress across Environment, Patients, People & Governance



Excellence in execution

Commercial & medical execution underpinning attractive opportunities

- Launching four new medicines or new indications in near term
- Increasingly balanced business



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Questions

January 2024

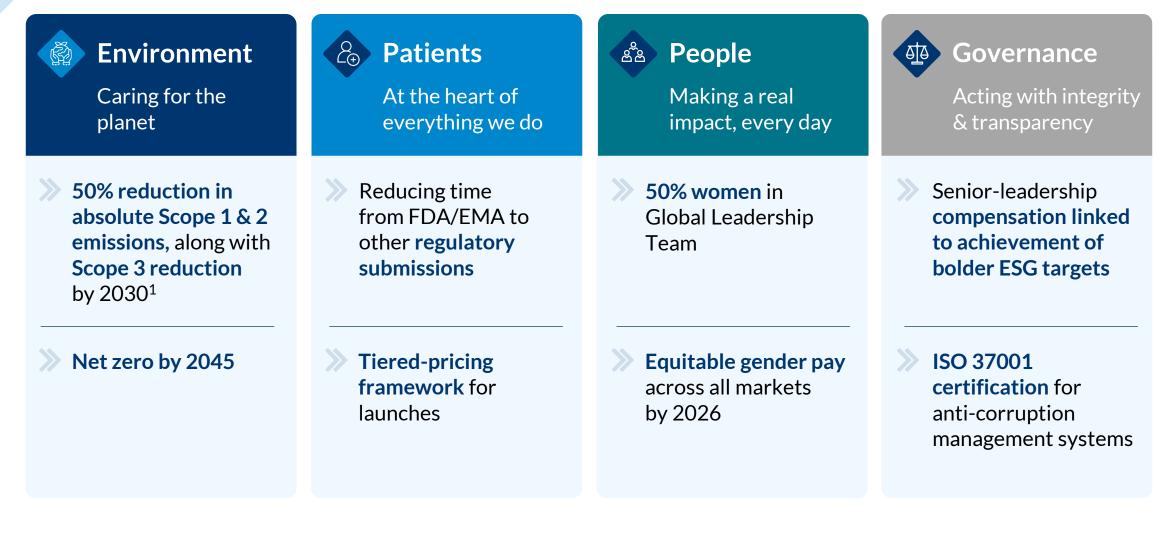
Diana Living with post-stroke spasticity Sintra, Portugal



Appendix

SIPS Innovation -

Generation Ipsen: for positive change



FDA: U.S. Food & Drug Administration; **EMA**: European Medicines Agency. ¹ Vs. baseline year, 2019.

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Thank you



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