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## Today's presenters and agenda



Henrik Stenqvist
Chief Financial Officer



Cecilia Nord
Global Head of Sustainability



Erik Krohn
Head of Treasury and Tax



**Overview of Sobi** 



Financials and operational highlights



**Sustainability at Sobi** 



Financing and contemplated transaction



**Appendix** 







### Sobi at a glance



Biopharmaceutical company with a focus on rare diseases

Business areas:



- Haematology
- Immunology
- Specialty Care



Present around 30 countries, delivering treatments to patients in many more



7 assets across 9 projects in development



Head office in Stockholm, Sweden



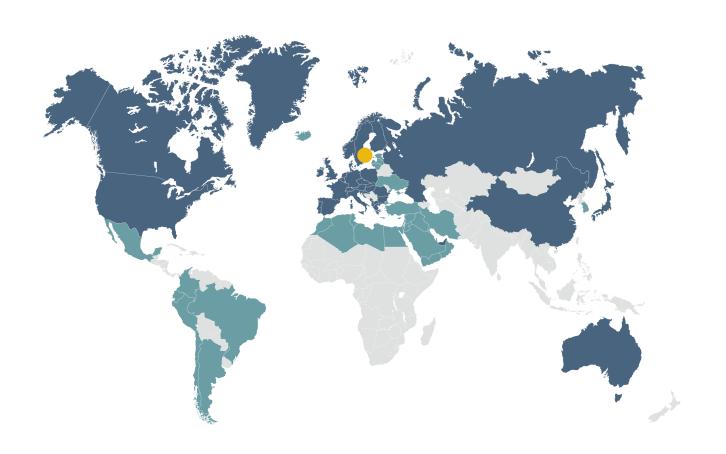
~1,800 employees



Strong main shareholder



SEK 22.1B 2023 revenue





# Sobi's mission is to transform the lives of people with rare and debilitating diseases

#### **Sobi strategy**



**Lead in Haematology** 



Capture the value of the pipeline



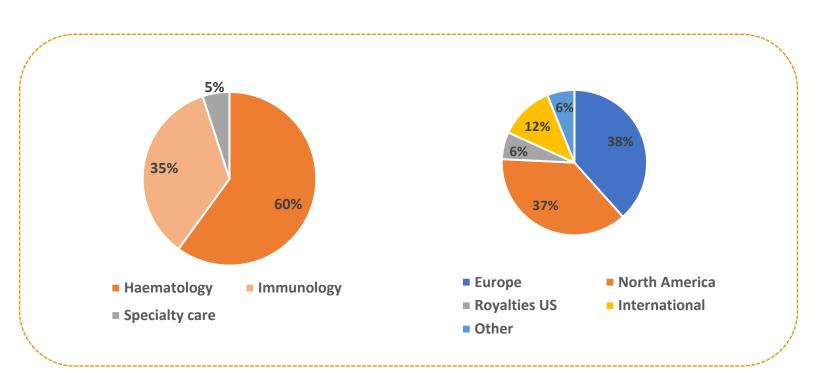
**Grow Immunology** and **Specialty Care** 



Go Global

2023: Revenue: SEK 22,123 M (+12% at CER)

**EBITA** margin adjusted 34%





### What are rare diseases?



The European Commission on Public Health defines rare diseases as lifethreatening or chronically debilitating diseases which affect fewer than 1 in 2,000 people.<sup>1</sup>



The Orphan Drug Act defines a rare disease as a disease or condition that affects less than 200,000 people in the United States. <sup>2</sup>

### Rare disease in numbers<sup>3-5</sup>

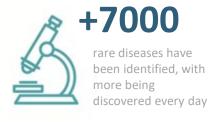
Rare diseases affect ~300m people worldwide

80% of rare diseases have a genetic basis, and are often chronic and affecting patients from birth

Only ~5% of rare diseases currently have an approved treatment



>80% of rare diseases are caused by faulty genes underscoring the need for effective treatment rather than preventative measures



#### Source:

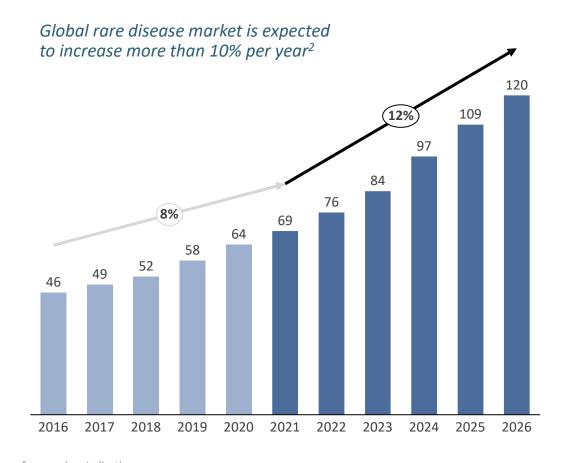
- 1. https://ec.europa.eu/health/ph information/documents/ev20040705 rd05 en.pdf
- 2. https://www.fda.gov/patients/rare-diseases-fda
- 3. National Organizations for Rare Disorders (NORD) https://rarediseases.org/understanding-rare-disease/rare-disease-facts-and-statistics/
- 4. National Human Genome Research Institute https://www.genome.gov/dna-day/15-ways/rare-genetic-diseases
- 5. Global Genes https://globalgenes.org/learn/rare-disease-facts



### Why rare diseases?

- High medical unmet need: 30% of children born with a rare disease die before their 5<sup>th</sup> birthday
- Government and Regulatory support: incentives such as fast regulatory approval with orphan designation
- Robust payer coverage: payers recognize the impact of the drugs
- Motivated patient populations: strong patient organizations support access and treatment
- Small number of KOLs<sup>1</sup> and prescribers: requires specialized but much smaller medical and sales teams than more traditional pharmaceuticals

#### Rare disease market, USD Bn



Source: Evaluate Pharma

<sup>1.</sup> Key Opinion Leaders

<sup>2.</sup> Excludes orphan oncology products and medicines with orphan indications where <50% of lifetime sales are expected to come from orphan indications



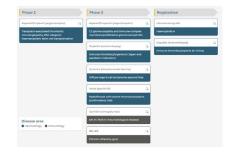
Business model - Source, develop and commercialise in the rare disease space

2018

Acquisitions **ADC Therapeutics** Sanofi Source **Novimmune** Apellis CTI Pharma efanesoctocog alfa Zynlonta Gamifant Doptelet Licensing Vonio Apellis AstraZeneca Aspaveli/Empaveli Synagis and nirsevimab (royalty) Selecta SEL-212

Develop

7 medicines in 9 projects in development (phase 2 through registration)



2022

2023

Commercialise

2023: Revenue: SEK 22,123 M (+12% at CER)



2019

2020

Europe +6% North America + 6% International +7% Other -6%



# Strong record of successful achievements in delivering medicines to rare disease patients around the world

#### Selected examples from the Sobi portfolio gamifant® S ASPAVELI\* **VONJO** Doptelet. (avatrombopag) tablets pegcetacoplan Apellis (Apellis CTI Pharma (CTf) Dova ... Novimmune ITP - FDA approval 2018 PNH - EU approval 2021 MF – FDA approval 2022 pHLH - FDA approval 2018 **EU approval 2021 International expansion** CTI integrated, launch China approval 2022 **International expansion** ongoing ongoing sHLH/MAS Phase 3 data **Future development options Nephrology indications**ongoing expected 2024 Paediatric data 2024 Phase 3 data in 2024 in planning

Commercialise

Source

Develop

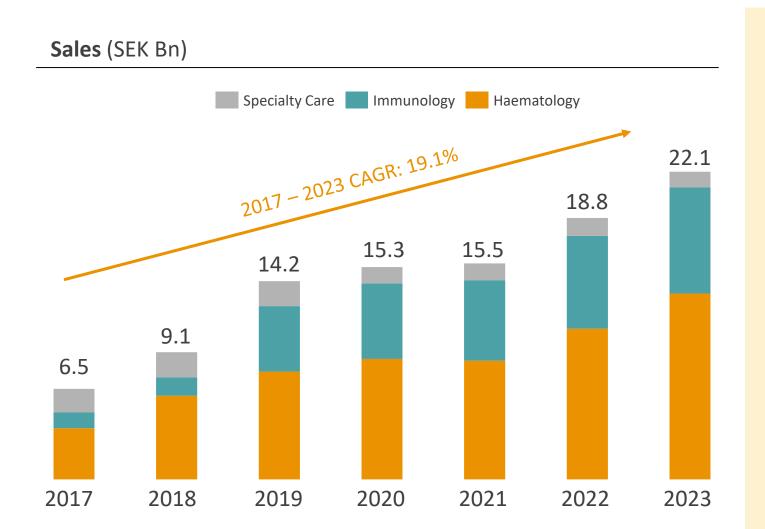
SEK 1,645 M +77% SEK 2,997 M +13% +62% (ex-China)

SEK 594 M +>200% SEK 706 M First year of launch, Jun-Dec

Revenue 2023, % growth, year over year at CER



### Strong track record of operational performance



#### Haematology – 2023

Accounted for 60% of total revenue 2023

In June, the acquisition of CTI was completed and Vonjo was added to the portfolio.

Strong sales growth for Elocta, Alprolix and Doptelet

#### Immunology – 2023

Sales of Gamifant increased by 77 per cent at CER (2023)

Kineret was approved for FMF<sup>1</sup> and CAPS<sup>2</sup> in China during the year and launched in Saudi Arabia.

Stable revenue increase from Immunology the past years

#### **Specialty Care – 2023**

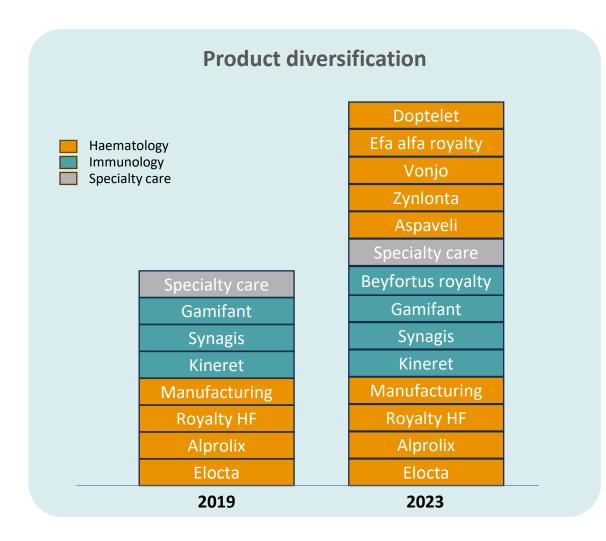
As anticipated, sales in Specialty Care decreased by 17 per cent at CER to SEK 1,119 M during the year

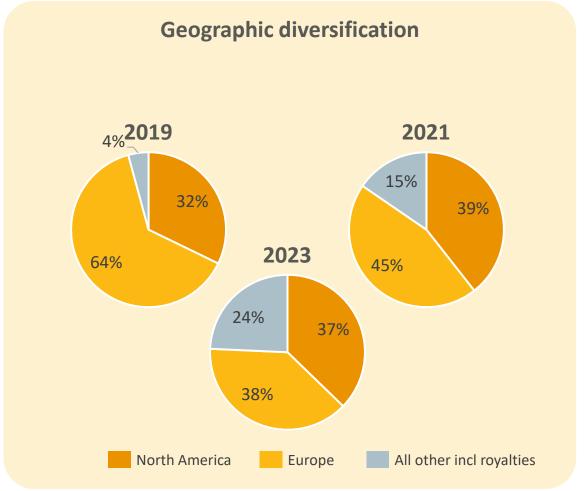
<sup>1.</sup> Familial Mediterranean Fever

<sup>2.</sup> Coding Atlas for Pharmaceutical Substances



### Increased diversification in products and the global regions









### Strategic portfolio delivering top-line growth in all regions



### Sobi strategy

#### Q1 2024 High double-digit performance at CER

**Revenue** Q1 - SEK 6,256 M, +20%

**EBITA** margin adjusted 37%

Operating cash flow SEK 2,256 M



- Vonjo<sup>®</sup> (SEK 320 M)
- Beyfortus<sup>™</sup> royalties SEK 318 M
- Altuviiio™ royalties SEK 108 M

- Continued strong growth
  - Doptelet® (SEK 756 M, +59%)
  - Gamifant® (SEK 438 M, +100%)
  - Aspaveli/Empaveli® (SEK 240 M, +155%)

#### Key milestones for late-stage pipeline

- Doptelet: positive phase 3 paediatric data
- Aspaveli / Empaveli: 1-L positive CHMP opinion in PNH

#### SEL-212: FDA fast-track designation

Kineret: Approved for Still's disease in China

#### 2024 outlook - Unchanged

**Revenue:** anticipated to grow by a high single-digit percentage at CER

Adjusted EBITA margin: anticipated to be in the mid-30s per cent of revenue



Lead in Haematology



**Capture the value** of the pipeline



**Grow Immunology** and Specialty Care



Go Global



## Significant events ahead in 2024

Anticipated major pipeline news flow

EU decision

SEL-212 & Gamifant
US submissions

Doptelet
US & EU (paediatric)
CN, JP (ITP)

2024 H1

2024 H2

**Efanesoctocog alfa** – Haemophilia A:

Regulatory decision in EU



**Gamifant** – sHLH / MAS in rheumatological diseases:

 Regulatory submission in the US (Still's disease cohort)



**Doptelet** – ITP:

Regulatory decision in China



**Doptelet** – ITP:

- Regulatory submission in Japan
- Paediatric submission in US & EU



**SEL-212** – Chronic Refractory Gout:

Regulatory submission in the US



Aspaveli/Empaveli – C3G & IC-MPGN:

VALIANT phase 3 study data readout





ITP: immune thrombocytopenia. C3G and IC-MPGN: Complement 3 glomerulopathy and immune-complex membranoproliferative glomerulonephritis. sHLH / MAS: secondary hemophagocytic lymphohistiocytosis / macrophage activation syndrome in patients with underlying rheumatological diseases, specifically Still's disease and systemic lupus erythematosus







Sobi's sustainability strategy supports our strategic

business priorities



Lead in Haematology

Grow Immunology

Go global

Capture the value of our pipeline

Maintain commitment to patients

Always act responsibly

Commitment to the UN Global Compact. Contribution to the 2030 Agenda with the UN Sustainable Development Goals and the Paris Agreement.

### Maintain commitment to patients

### **Access to treatment**

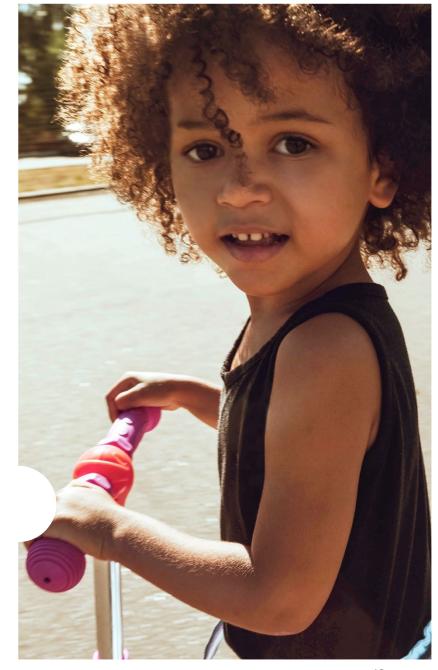
**36,000+** patients with rare or debilitating diseases treated with Sobi medicines.

Sobi's medicines available in **ten** new markets.

### **Humanitarian aid**

**810 M IU factor donated** in ten-year commitment to World Federation of Hemophilia to donate **1 Billion IU** 2015-2025.

Over **22,000 patients** reported treated by 2023.





### Always act responsibly

#### Responsible sourcing process pillars

- 1 Alignment of principles
- 2 Risk assessment & qualification
- 3 Performance management

**95%** of contract manufacturers scored by EcoVadis with a 64 point average.

>80% of contract manufacturers currently with approved Science Based Targets or committed to setting them (share of spend).





# The progress on our sustainability strategy is recognised by external stakeholders

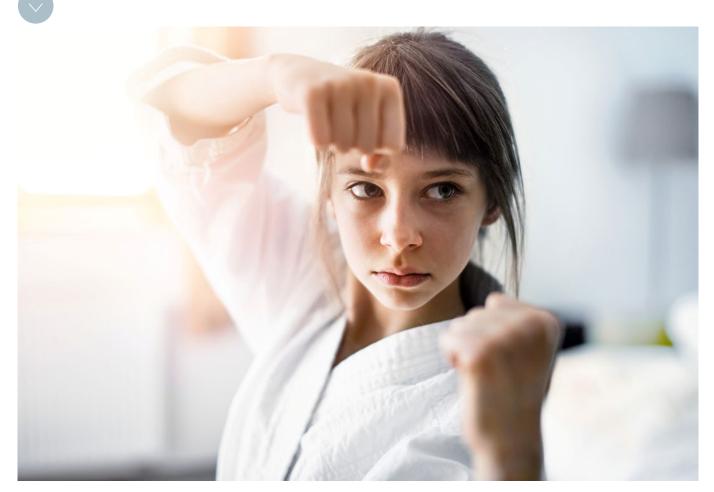
**One of ten** companies in DJSI Europe within Pharmaceuticals, Biotech and Life Sciences

Member of

### Dow Jones Sustainability Indices

Powered by the S&P Global CSA

#### Stable performance in analyst indices









### Financing objectives

Diversify funding sources

#### **Enter the domestic debt capital market**

- Access to new liquidity
- Multiple sources to reduce refinancing risk
- Reduce dependency of banks

Enable future BD\* activity

#### Support Sobi's growth journey

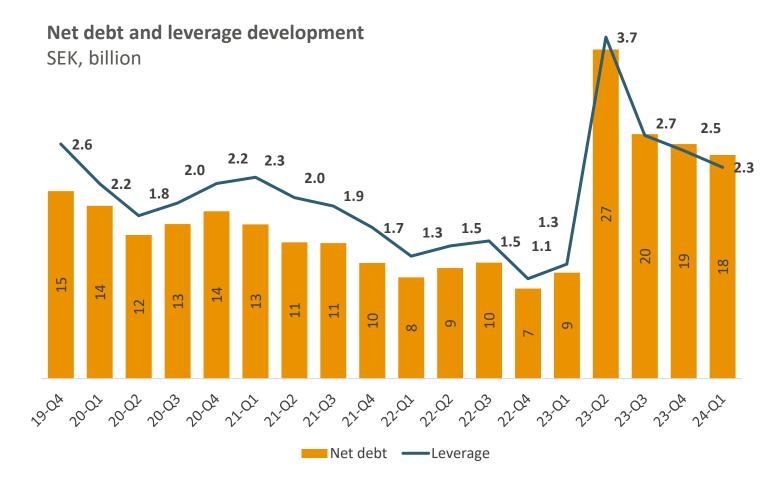
- Maintain liquidity for mid-size acquisitions and in-licensing
- Increase flexibility for take-out of major financing
- Maintain ability to lever up to 3-4x in M&A scenario

\* Business Development



### Historical leverage development

- Debt increases with BD transactions and declines by strong cash flow and growing EBITDA
- Several funding sources
- Equity used to partly finance previous M&A (Synagis and CTI), maintaining potential to further invest in the business

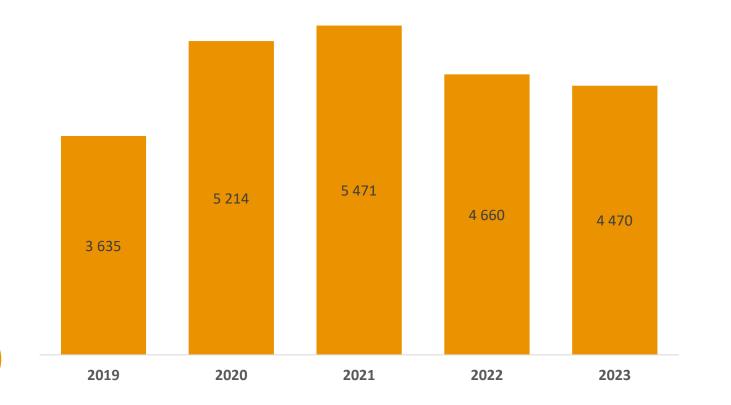


Leverage defined as Net debt / EBITDA adjusted



### Strong cash flow with deleveraging capability





Strong cash conversion

Highly cash generative business model

Capacity to delever

### Current maturity structure







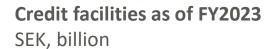
Balanced maturity profile with several extension options

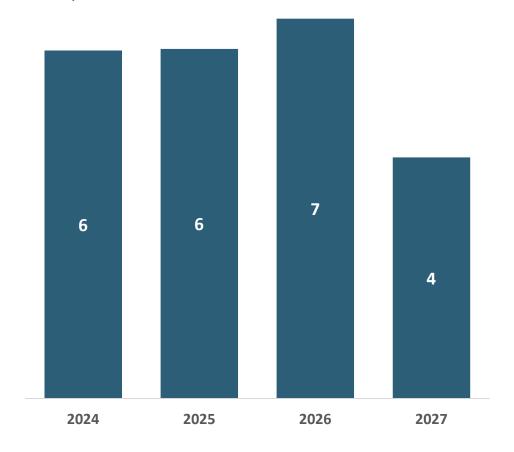


Maximum maturities within 12 months defined in policy



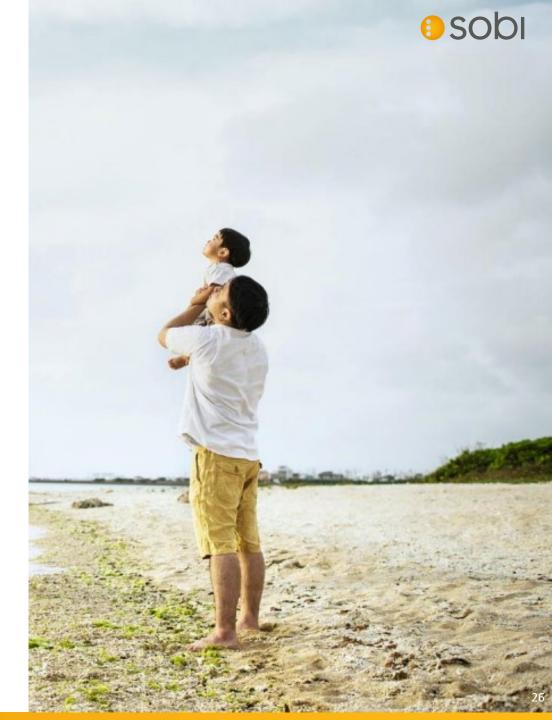
Financing historically driven by M&A with limited need to refinance





## Proposed transaction

Issuer	Swedish Orphan Biovitrum AB (publ) ("SOBI")
Use of proceeds	General corporate purposes
Tenors	3 & 5 years expected
Interest	FRN and/or FXD
Exp. Combined Size	TBD
Issuer call	1m (on 3y tranche) & 3m (on 5y tranche) Par Calls
Denominations	2m x 2m
Target market	Eligible counterparties, professional clients and retail clients
Documentation	Issuer's MTN Programme dated April 26, 2024
Law	Swedish Law
Joint Bookrunners	Nordea and SEB





## Summary and key credit highlights



Rare diseases are an attractive market and expected to grow faster than general pharma



Strong portfolio of medicines and solid pipeline driving future growth



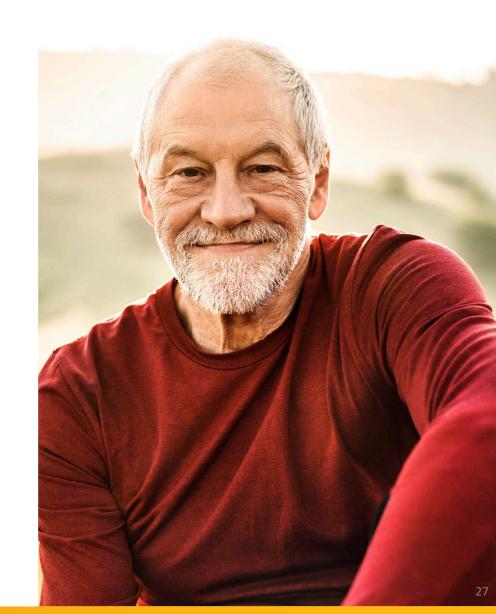
Globally diversified, present in around 30 countries, delivering treatments to patients in many more

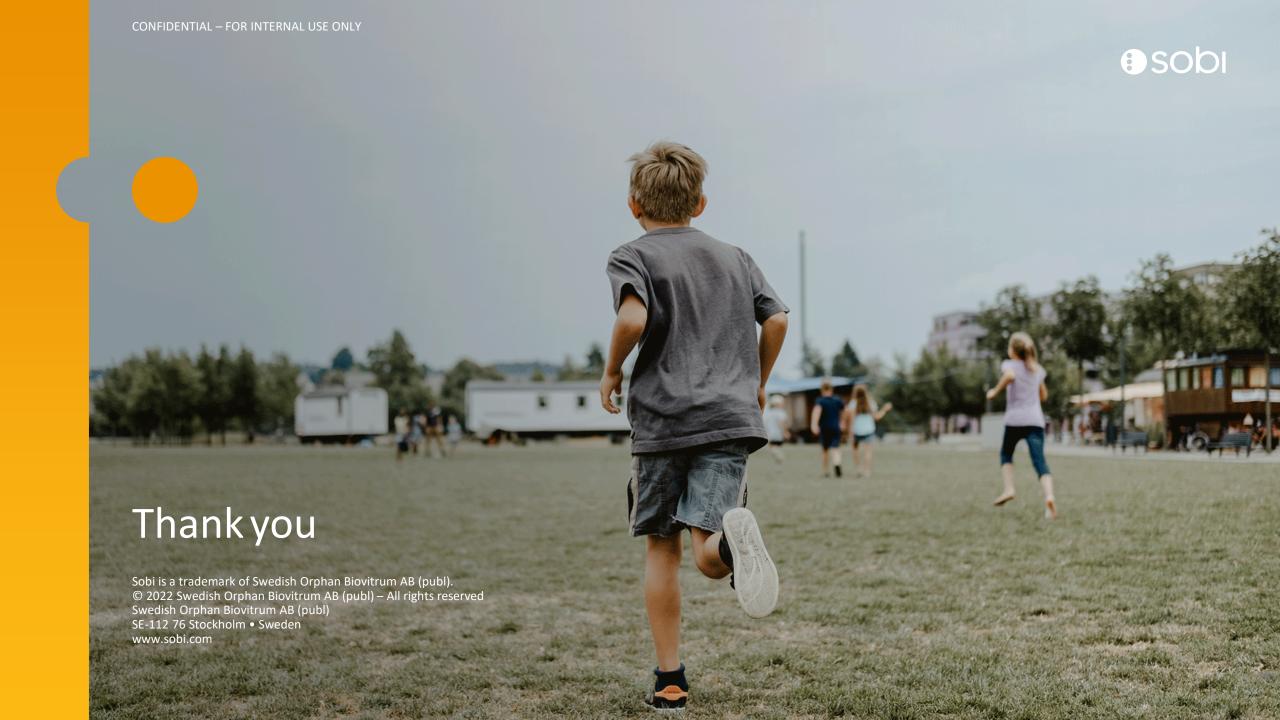


Highly cash generative business model supporting deleveraging



Strong main shareholder



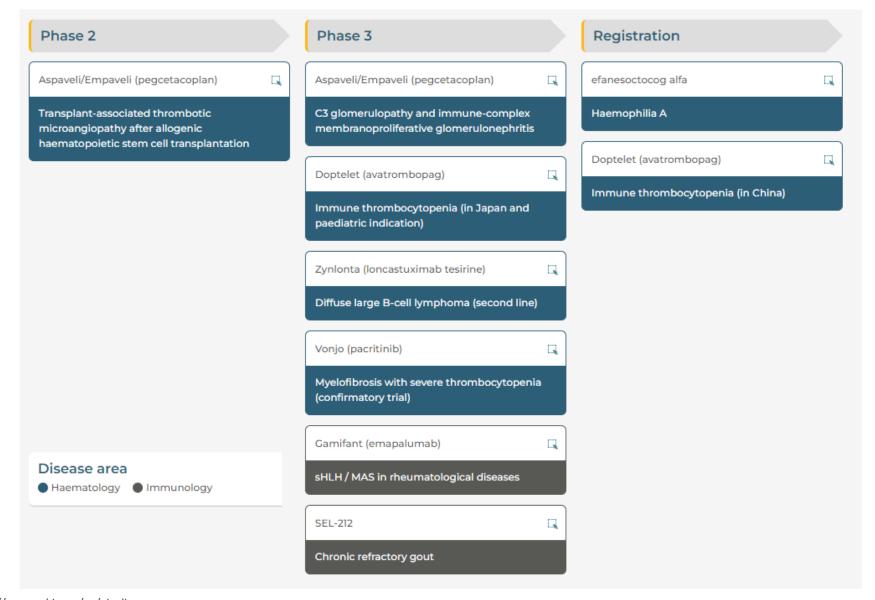








### **Current Development pipeline**



Source: https://www.sobi.com/en/pipeline.

Visionary founding partner

## Humanitarian Aid Program



Over 22,000 people reported treated since programme start



Over 9,500 acute bleeds treated in 2023



Over 1,000 surgeries in 2023



Over 810 million IU factor donated in total

