



Guido Oelkers, CEO Mats-Olof Wallin, CFO

22 February 2018

Q4/FY 2017 results presentation

Forward looking statements



In order to utilize the 'Safe Harbor' provisions of the United States Private Securities Litigation Reform Act of 1995, Swedish Orphan Biovitrum AB (publ) is providing the following cautionary statement. This presentation contains forward-looking statements with respect to the financial condition, results of operations and businesses of Swedish Orphan Biovitrum AB (publ). By their nature, forward-looking statements and forecasts involve risk and uncertainty because they relate to events and depend on circumstances that will occur in the future. There are a number of factors that could cause actual results and developments to differ materially from that expressed or implied by these forward-looking statements. These factors include, among other things, the loss or expiration of patents, marketing exclusivity or trade marks; exchange rate fluctuations; the risk that R&D will not yield new products that achieve commercial success; the impact of competition, price controls and price reductions; taxation risks; the risk of substantial product liability claims; the impact of any failure by third parties to supply materials or services; the risk of delay to new product launches; the difficulties of obtaining and maintaining governmental approvals for products; the risk of failure to observe ongoing regulatory oversight; the risk that new products do not perform as we expect; and the risk of environmental liabilities.

Highlights 2017



Haemophilia

- ✓ Outstanding sales growth
- ✓ rFIXFc-XTEN added to Bioverativ collaboration agreement
- ✓ Improved joint-health for haemophilia A patients after prophylactic treatment with Elocta ®
- ✓ New dosing regimen for Alprolix to dose 14 days or longer was approved
- ✓ FPI in 24 months real world study ASURE of Elocta
- ✓ FPI in the RelTIrate study to evaluate immune tolerance induction with Elocta
- ✓ Bioverativ initiated first in human phase 1/2 trial with next generation EHL product in haemophilia A, rFVIIIFc-VWF-XTEN (BIVV001)
- ✓ Elocta launched in 22 countries and Alprolix in 14 (Year end 2017)

Specialty Care

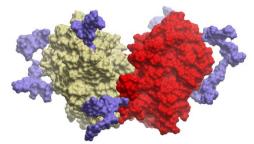
- ✓ Established in Q2
- ✓ Double digit growth for Kineret® and Orfadin®
- ✓ Solid uptake for Orfadin's new formulations oral suspension and 20mg
- ✓ New Orfadin formulations approved in Saudi Arabia and Canada
- ✓ Kineret approved in Canada for the treatment of NOMID
- ✓ The studies- anaGO and anaSTILLs to evaluate the safety and efficacy of anakinra (Kineret) as a treatment for acute gout and Still's disease were initiated
- ✓ SOBI003 for the potential treatment of MPSIIIA recieved orphan designation in the US

Events after the reporting period



- FDA accepted investigational new drug application and granted Fast Track status for SOBI003 for the treatment of MPS IIIA
- Ireland switching all people with haemophilia A & B treated with replacement clotting factors to Sobi's extended half-life therapies









Financial highlights Q4

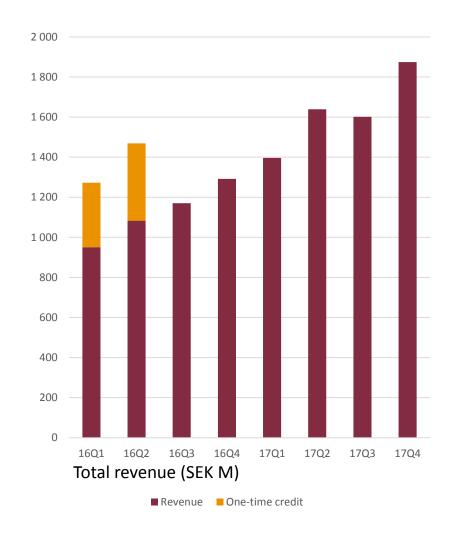


Revenue SEK 1,875 M (1,292) +45%

Gross margin 71% (67)

EBITA SEK 619 M (210) +195%

Cash flow from operations SEK 257 M (26)



Financial highlights FY

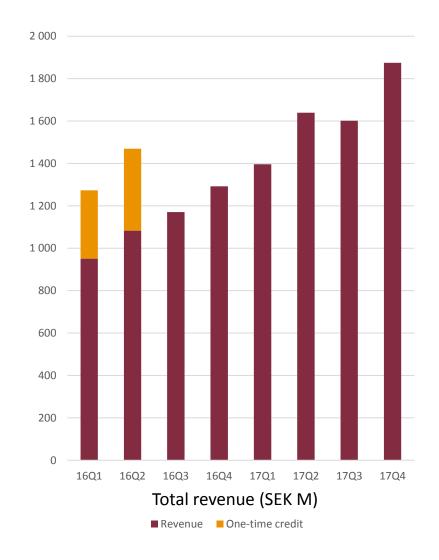


Revenue SEK 6,511 M (5,204) +25% (+ 45 % excl. one-time credits in 2016)

Gross margin 72% (70)

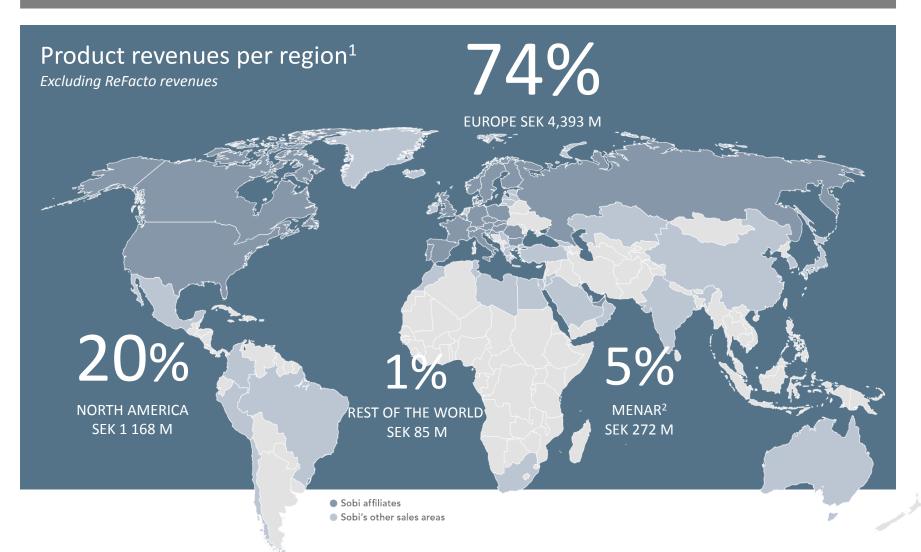
EBITA SEK 2,053 M (1,543) +33%

Cash flow from operations SEK 1,333 M (343)



Sales per region





¹Revenues from legal companies registered in each region, FY 2017

²Middle East, North Africa, Russia

Commercial results

Haemophilia – delivering stellar growth

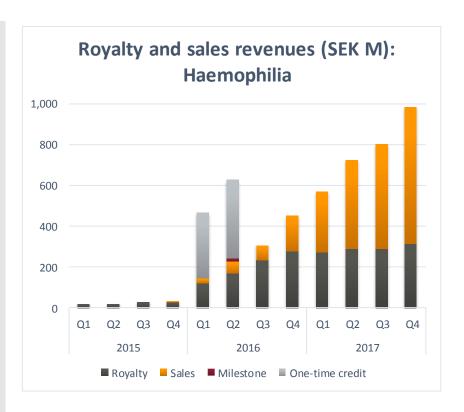


Q4 revenue SEK 985 M (451)

- SEK 671 M (174) in sales revenue
- SEK 314 M (277) in royalty revenue

FY revenue SEK 3,088 M (1,853)

- SEK 1,920 M (327) in sales revenue
- SEK 1,168 M (1,525) in royalty revenue
- Increase of SEK 1,943 M when one-time credits in 2016 are excluded



Elocta – increasingly the standard of care

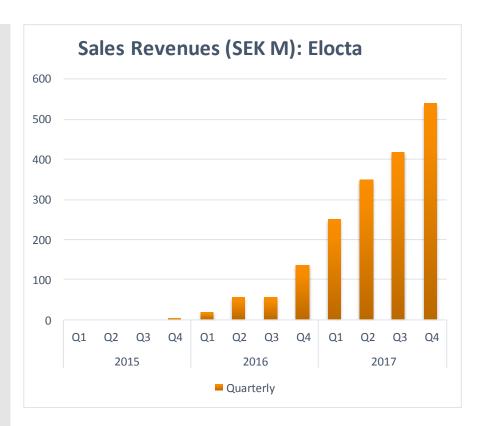


Q4 sales revenue of SEK 540 M (135)

- SEK 123 M (29%) growth compared to Q3 2017
- More than 50 % of the growth derived from France and Germany

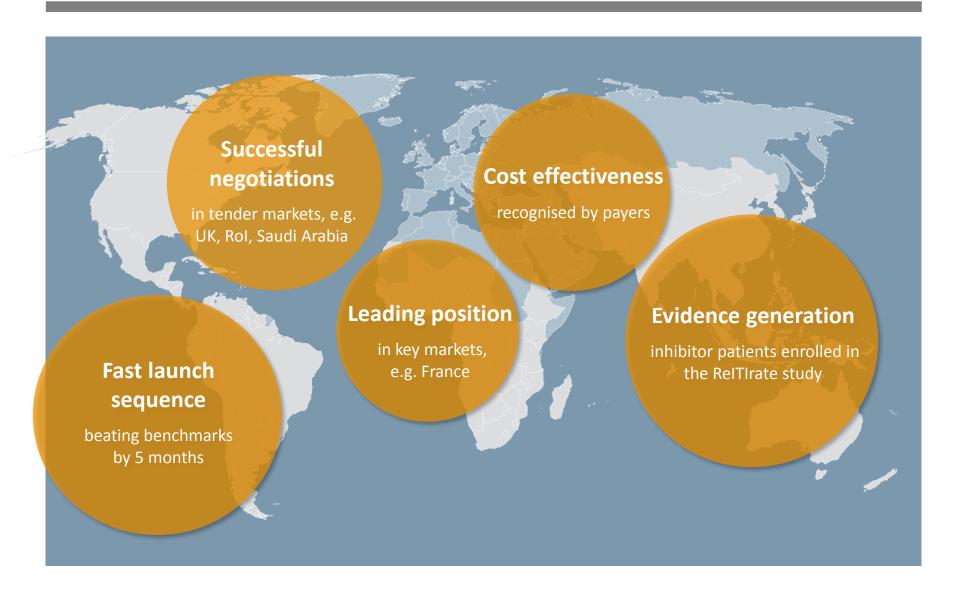
FY sales revenue SEK 1 557 M (267)

Launched in 22 countries



Landmark commercialisation of Elocta





Alprolix – uptake continues strongly



Q4 sales revenue of SEK 131 M (39)

- SEK 34 M (34%) growth compared to Q3 2017
- More than 50% of the growth derived from Germany, Ireland and the Netherlands

FY sales revenue SEK 363 M (60)

Launched in 14 countries



Haemophilia strategic imperatives – driving success



Real-world evidence and experience supporting safety and efficacy

Established real-world safety profile in thousands of patients



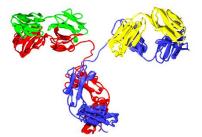
Provide individual choice for optimal therapeutic outcome

Elocta and Alprolix indicated for all ages, in prophylaxis, OD and surgery



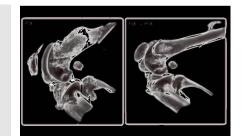
Provide further evidence on elimination of inhibitors in Haemophilia A

• Inhibitor eradication remains the first goal of care for inhibitor patients



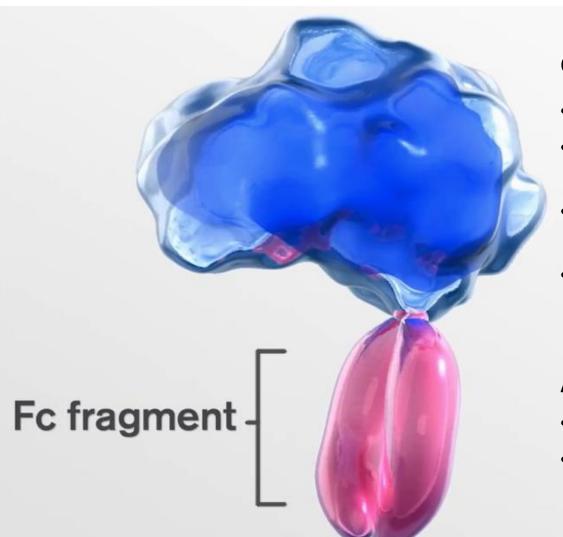
Driving the **Joint Health** agenda with evidence

 A-LONG and ASPIRE data show improvement in joint health scores for haemophilia A patients on prophylactic dosing with Elocta over a period of 4 years



Significant potential for development





Growth based on innovation

- Fc fusion technology
- Real world evidence supporting differentiating efficacy and safety
- Priced to enable wide access and adoption
- Extended half-life products allow reduced treatment burden

Alprolix

- 14 days interval
- Emerging evidence of accumulation in joints

Specialty Care – three core strengths



World-class commercialisation platform



Lifecycle management & indication expansion



Partner product portfolio with room to grow



Specialty Care – strong business platform for continued growth



Q4 revenue SEK 761 M (692) +10%

YTD revenue SEK 2,829 M (2,695) +5%

 Adjusted for discontinued products (mainly Cometrig) increase of 9%

Double-digit growth for Kineret and Orfadin



Orfadin – double digit growth with increased generic competition



Q4 revenue SEK 223 M (197) +13%

FY revenue SEK 862 M (770) +12%

Double digit growth in North America and EMENAR

- · Patient insights and market understanding
- Growth in North America mainly due to launch of 20 mg and oral suspension formulations
- Strong performance across all EMENAR markets, especially Russia, Middle East and North Africa.



Kineret – strong performance driven by patient demand



Q4 revenue SEK 307 M (266) +15%

YTD revenue SEK 1,142 M (1,001) +14%

Strong double digit value and volume growth in both North America and FMFNAR

 EMENAR sales were positively impacted by phasing of shipments from third quarter to the Middle East.



Kineret – a unique mechanism of action

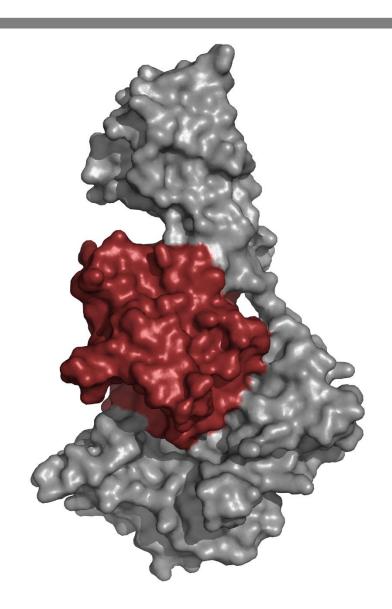


IL-1 pathway

Two new clinical trials in 2017 to explore indication expansion

- anaGO acute gout
- anaSTILLs Still's disease

Anakinra, in red, bound to the interleukin 1 receptor, in grey.



ReFacto – a solid revenue base

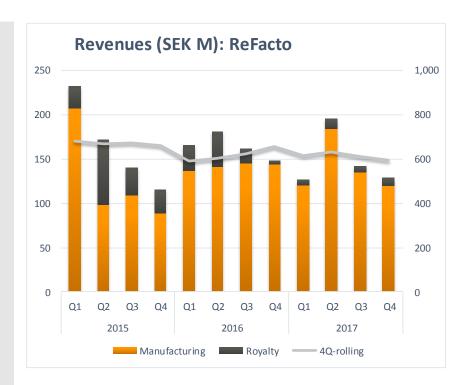


Q4 Revenue for manufacturing and royalty SEK 128 M (148)

• Phasing impacts delivery between the quarters

Q4 Manufacturing revenue SEK 120 M (145)

FY was SEK 594 M (656)

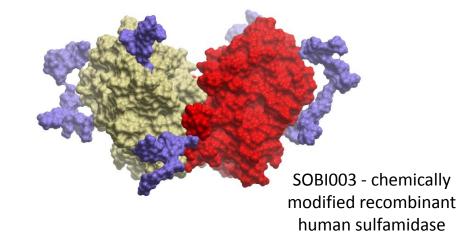


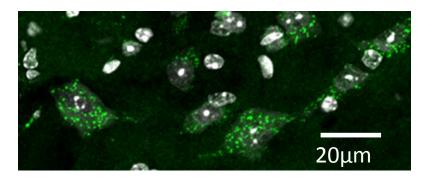
Pipeline

Pipeline



- FPI in phase 2 study anaGO to evaluate anakinra for the treatment of acute gout
- FPI in the anaSTILL's study evaluating anakinra for the treatment of Still's disease
- Fast Track status for SOBI003 for the treatment of MPS IIIA
- The majority of early stage clinical trials in the US





Distinct intracellular SOBI003 fluorescence (Green) indicates uptake into lysosomes in nerve cells in the CNS.

Financial results Mats-Olof Wallin

Profit & loss statement



Amounts in SEK M	Q4-17	Q4-16	Full year-17	Full year-16
Total revenues	1,875	1,292	6,511	5,204
Gross profit	1,337	860	4,657	3,651
Gross Margin	71%	67%	72%	70%
Sales and Administration	-477	-399	-1,644	-1,366
Research and development	-228	-258	-908	-778
Other operating revenues/expenses	-13	7	-52	36
EBITA	619	210	2,053	1,543
Amortisations and write-downs	-110	-110	-453	-410
EBIT	509	100	1,600	1,133
Financial income/expenses	-15	-12	-68	-85
Profit before tax	494	88	1,532	1,048
Income tax expense	-137	-15	-384	-246
Profit for the period	357	73	1,149	802

Balance sheet



Amounts in SEK M	Dec 2017	Sep 2017	Dec 2016
ASSETS			
Intangible	6,445	6,535	6,806
Tangible and other	301	277	257
Total non-current assets	6,746	6,812	7,063
Inventories	1,053	1,095	870
Accounts receivable	1,129	941	769
Other Receivable	496	469	487
Cash and equivalent	1,478	1,758	786
Total current assets	4,157	4,263	2,911
Total Asset	10,903	11,075	9,974
EQUITY AND LIABILITIES			
Equity	6,701	6,352	5,365
Long-term debt	5	503	502
Long-term liabilities	1,832	1,880	2,349
Short-term liabilities	2,365	2,341	1,758
Total liabilities	4,202	4,724	4,609
Total equity and liabilities	10,903	11,075	9,974

Summary Guido Oelkers

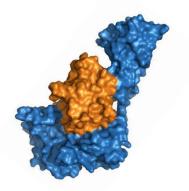
Outlook 2018



- Sobi expects total revenues for the full-year to be in the range of SEK 7,500 – 7,700 M
- Gross margin is expected to be at least 70 per cent
- Sobi expects EBITA for the full-year to be in the range of SEK 2,500 – 2,700 M









Our strategic position today



Accomplished successful inroads in multiple markets in Haemophilia



Drive commercial effectiveness agenda in Haemophilia



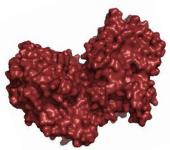
Establish a Specialty Care business and strengthen its focus



Establish late stage pipeline

Expansion in NA









Our strategic direction





Further internationalisation and commercialisation of Haemophilia



Build Specialty Care as a preferred partner



Strengthen position in the US and EMENAR



Build pipeline and self-sustained R&D

Vision

Global leader
in providing access
to innovative treatments
that make a significant
difference for individuals
with rare diseases

Q&A

