



Forward-looking statements

In order to utilise the 'Safe Harbor' provisions of the United States Private Securities Litigation Reform Act of 1995, Swedish Orphan Biovitrum AB (publ) (Sobi®) is providing the following cautionary statement: This presentation contains forward-looking statements with respect to the financial condition, results of operations and businesses of Sobi. By their nature, forwardlooking 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.



Overview and business	Guido Oelkers, Chief Executive Officer
Financials	Henrik Stenqvist, Chief Financial Officer
Pipeline	Tony Hoos, Head of RDMA ¹ , Chief Medical Officer
Summary and Q&A	All

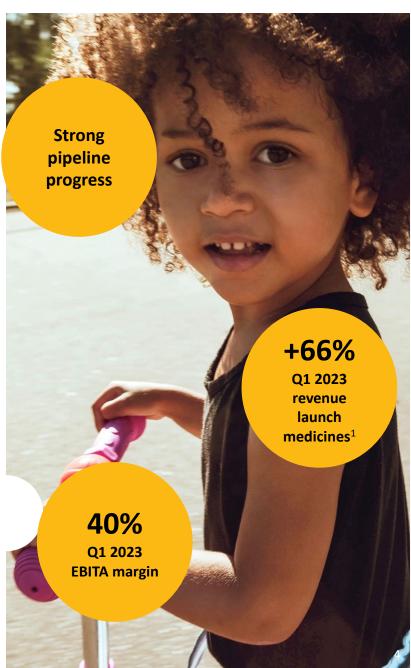
1. Research & Development and Medical Affairs.



Overview: Continued solid progress

- Revenue +6% and -2% at CER. Revenue increased by +8% excluding Kineret® COVID-19 related sales and Doptelet® sales in China
- Launch medicines¹ +66% in Q1, driven by Doptelet
- Haemophilia grew by +10%, Aspaveli® launch progressing well, SEK 95 M in Q1
- Immunology held back by COVID-19 comparison and softer Gamifant® sales
- Opex disciplined cost control
- EBITA margin 40% in Q1
- **Pipeline** progress:
 - efanesoctocog alfa US approval (Sanofi) and positive phase 3 study data readout
 - **SEL-212** positive phase 3 studies readout
 - nirsevemab simplified economics and creating strategic flexibility by royalty agreement
 - Empaveli PNH/Doptelet CLD approvals in Japan
 - Gamifant: complete enrollment in the cohort in study 14 relevant for US filing
- 2023 outlook unchanged

Continued performance: strong earnings and progress of our pipeline





Business: Growth at act. currency of 6%, at CER of -2%

Underlying growth of 8%, excluding the phasing effect of sales to our partner in China and missing COVID-19 business

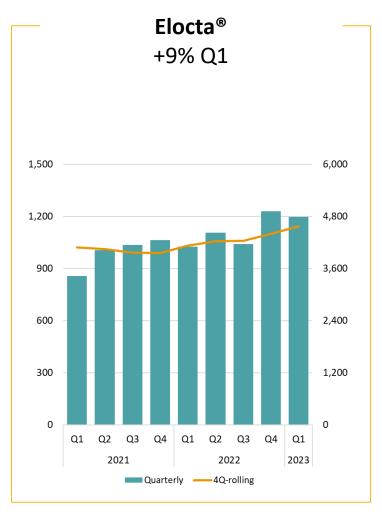
	Q1 '23	change	ratio		Q1 '23	change	ratio
	SEK M	%	%		SEK M	%	%
Haematology	2,815	5	55	Europe	2,142	12	42
– Haemophilia	2,053	7	40	North America	2,330	-2	43
Immunology	2,151	-9	40	Rest of world ¹	424	-42	8
Specialty Care	273	-17	5	Other ²	343	-7	6
Total	5,239	-2	100	Total	5,239	-2	100

^{1.} Rest of world grew 54 per cent, excluding Doptelet China and COVID-19 Kineret sales

hic area). 2. Royalty revenue



Haematology: haemophilia up in the quarter driven by increased patients and consumption



Haemophilia continued stability in Q1 2023

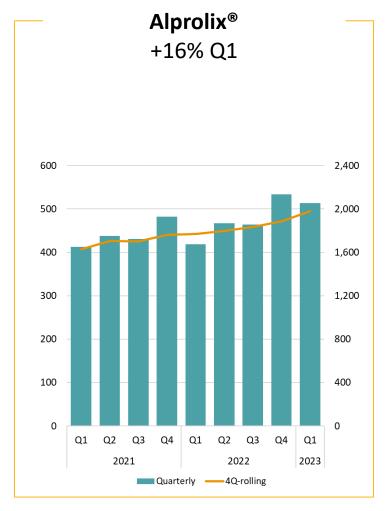
Elocta

 Growth in patients, consumption and geo expansion somewhat offset by price

Alprolix

Growth in patients and geo expansion offset by price



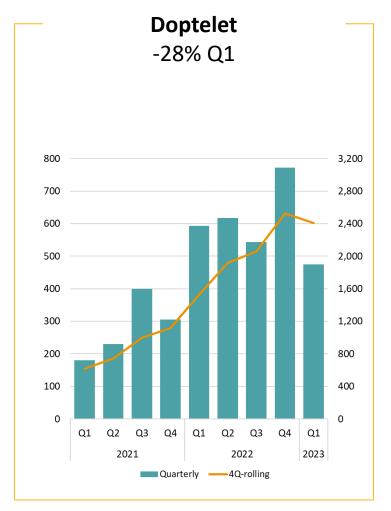


Sales in SEK million at actual exchange rates; change at constant exchange rates.

Sales in SEK million at actual exchange rates; change at constant exchange rates.



Haematology: Doptelet up 78% in Q1 in markets outside China



- US: new patients, new prescribers, higher market share and longer duration of treatment
- Europe: strong growth from Germany and recent reimbursements, e.g. Italy, Spain
- China: no sales in Q1 (SEK 352M)





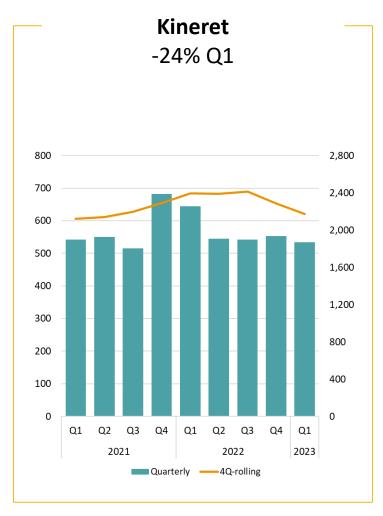
Sales in SEK million at actual exchange rates; change at constant exchange rates.







Immunology: Kineret COVID-19 impact, Gamifant patient growth



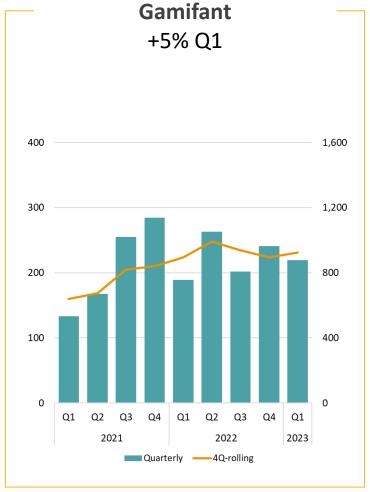
Kineret

 Sales performance impacted by Q1 2022 COVID-19 related sales. Rebasing to be completed by next quarter

Gamifant

 Increased adoption in the US offset by lower use in adults



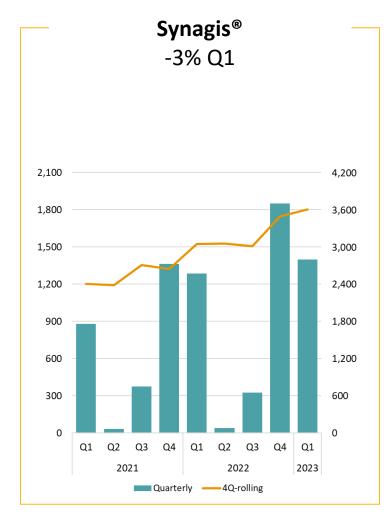


Sales in SEK million at actual exchange rates; change at constant exchange rates.

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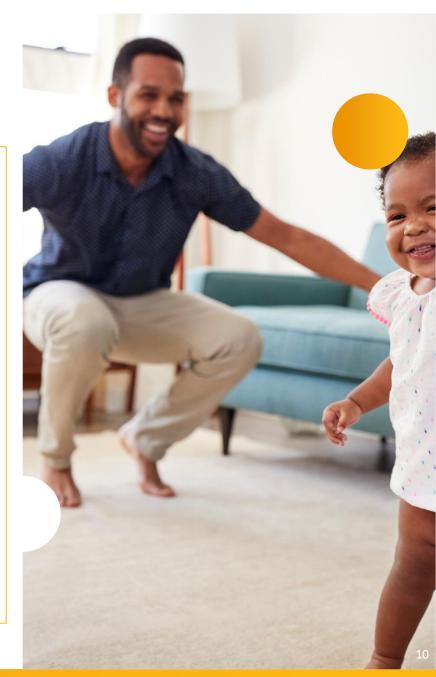


Immunology: Synagis quarter sales in line with plan



Sales in SEK million at actual exchange rates; change at constant exchange rates.

- SEK 1,398 M sales (1,286)
 increased by +9 per cent and
 decreased by -3 per cent at CER
- Sales reflect expected seasonal variability following strong demand earlier in the season
- Favourable price development
- Nirsevimab royalties¹ and AZ agreement termination¹ enabling simplification and increased strategic flexibility

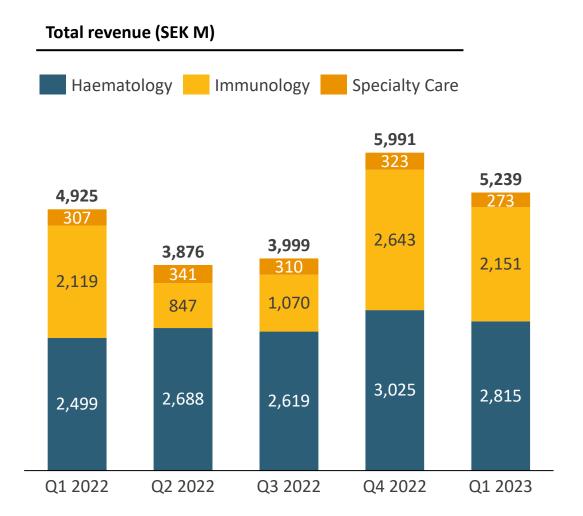




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Revenue and profit & loss



Amounts in SEK M	Q1 2023	Q1 2022	Change	Full-year 2022
Total revenue	5,239	4,925	6%	18,790
Gross profit	4,172	3,409	22%	14,014
Gross margin ^{1,2}	80%	69%		75%
EBITA ^{1,2}	2,121	1,290	64%	5,930
EBITA adjusted ^{1,2}	2,121	1,951	9%	6,605
EBITA margin ^{1,2}	40%	26%		32%
EBITA margin adjusted ^{1,2}	40%	40%		35%
Profit for the period	1,067	543	96%	2,638
Earnings per share, before dilution, SEK ^{1,2,}	3,60	1.84	96%	8.92
Earnings per share, before dilution, SEK adjusted ^{1,2}	3,60	3.67	-2%	10.77
Operating cashflow	1,983	1,644	21%	4,576
Net debt (+)/net cash (-)	8,708	8,321		7,406

^{1.} Alternative Performance Measures (APMs); see Q1 report for further information

^{2.} Items affecting comparability in 2022, see Q1 report for further information



Outlook 2023 – unchanged

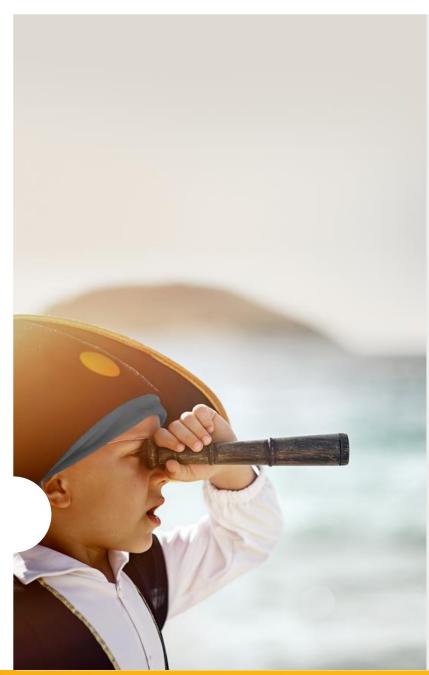
Outlook excludes Sobi's right to royalty on net sales of nirsevimab in the US

Revenue

Anticipated to grow by a low-to-mid-single-digit percentage at CER¹

EBITA margin adjusted²

Anticipated to be at a low 30s percentage of revenue





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Pipeline: continued milestone progress

Major pipeline milestones since the previous report

efanesoctocog alfa	haemophilia A	regulatory approval in the US (by Sanofi)
efanesoctocog alfa	haemophilia A (paediatric)	positive XTEND-Kids phase 3 study topline data readout
Kineret	CAPS	regulatory submission in China
SEL-212	CRG	positive phase 3 studies topline data readout
Doptelet	CLD	regulatory approval in Japan
Empaveli	PNH	regulatory approval in Japan



Pipeline news flow increasing

Sobi

Anticipated major upcoming pipeline news flow

H2 2023

2024

efanesoctocog alfa – haemophilia A: regulatory submission in the EU

Doptelet – ITP¹: regulatory decision in China

Kineret – FMF²: regulatory decision in China

Gamifant – MAS in rheumatological diseases: EMERALD phase 3 study data readout (Still's disease cohort)

Gamifant – MAS in rheumatological diseases: regulatory submission in the US (Still's disease cohort)

nirsevimab – RSV prevention: regulatory decision in the US (by AstraZeneca/Sanofi)

Doptelet – ITP: regulatory submission in Japan

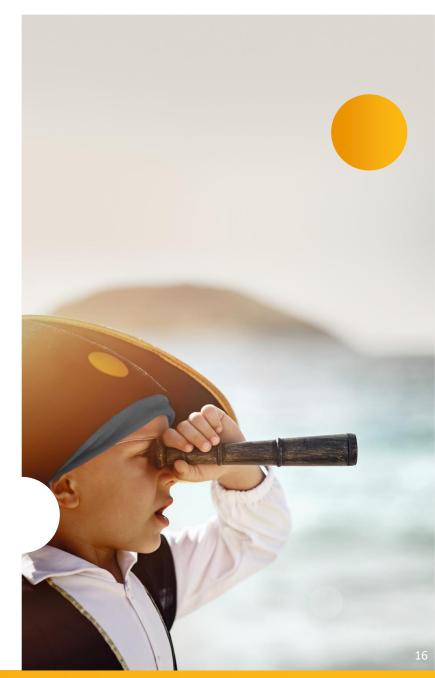
Aspaveli/Empaveli – IC-MPGN³ and C3G³: VALIANT phase 3 study data readout

Aspaveli/Empaveli – TA-TMA⁴: phase 2 study data readout

Kineret – Still's disease: regulatory decision in China

Kineret – CAPS: regulatory decision in China

SEL-212 – CRG: regulatory submission in the US (in first half 2024)





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Summary Q1 2023

- Continued solid progress
- Strong earnings
- Strength in major product areas, accelerating launches¹
- Several major pipeline milestones achieved
- Outlook 2023 unchanged²
 - Revenue to grow by a low-to-mid-single-digit percentage at CER³
 - EBITA margin adjusted⁴ to be at low 30s percentage of revenue



^{1.} Launch medicines include Doptelet (outside China), Aspaveli, Gamifant and Zynlonta 2. Outlook excludes Sobi's right to royalty on net sales of nirsevimab in the US. 3. Constant exchange rates 4. Excluding items affecting comparability







Appendix: Q1 2023 sustainability performance

Highlights in Q1 2023



- Milestones toward increased access
 - Positive results from phase 3 studies on SEL-212 used to treat patients suffering from chronic refractory gout.
 - FDA-approval of efanesoctogog alfa for treatment of haemophilia A and positive topline results in phase 3 paediatric study in children under 12.
- Raising awareness and supporting patients
 - Sharing knowledge at EAHAD* 16th Annual Congress.
 - Commemorating World Rare Disease Day through events in many Sobi markets together with patient organisations and key opinion leaders.



Maintain commitment to patients



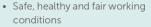
- Access to treatment
 - · Patient centricity and engagement
 - · Patient and product safety
 - · Ethical marketing and sales
 - Transparent and ethical R&D

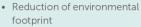


Always act responsibly



• An inclusive and diverse workplace that grows people

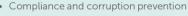














Commitment to the UN Global Compact. Contribution to the 2030 Agenda, the UN Sustainable Development Goals and the Paris Agreement

Member of

Dow Jones **Sustainability Indices**

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Highlights in Q1 2023



- Caring for employees
 - Choosing and celebrating the two winners of the annual Sobi Rare Strength Award, instituted to recognise company role models embodying Sobi values.
- Reduction of environmental footprint
 - Completing the first mapping of Sobi Scope 3 emissions.



8 SOOI rare strength