





Forward-looking statements

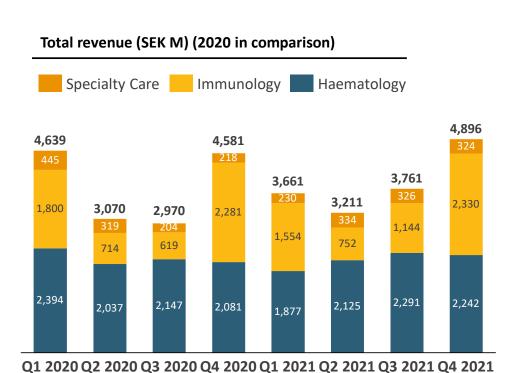
In order to utilise the 'Safe Harbor' provisions of the United States Private Securities Litigation Reform Act of 1995, Swedish Orphan Biovitrum AB (publ) (Sobi®) is providing the following cautionary statement: This presentation contains forward-looking statements with respect to the financial condition,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







2021: revenue and EBITA delivered in line with outlook



	Q4	Q4		FY	FY	
Amounts in SEK M	2021	2020	Change	2021	2020	Change
Total revenue	4,896	4,581	7%	15,529	15,261	2%
Gross profit	3,880	3,718	4%	12,045	12,036	0%
Gross margin ¹	79%	81%		78%	79%	
EBITA adjusted ^{1,2}	2,002	2,177	-8%	5,575	6,301	-12%`
EBITA margin adjusted ^{1,2}	41%	48%		36%	41%	
Profit for the period	1,241	1,502	-17%	2,679	3,245	-17%
Earnings per share, SEK adjusted ^{1,2,3}	4.21	3.74	12%	9.08	9.66	-6%
Operating cashflow	2,121	716	196%	5,470	4,926	11%
Net debt (+)/net cash (-)	9,500	13,748		9,500	13,748	

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

^{2.} EBITA 2020 excluding non-recurring items; other operating income related to the reversal of the CVR liability of SEK 399 M.

^{3.} EPS full-year 2020 excluding the reversal of the CVR liability of SEK 399 M.



2021: solid growth with increased diversification across both the disease areas and global regions

	Q4 '21	change	ratio	FY '21	change	ratio
	SEK M	%	%	SEK M	%	%
Haematology	2,242	8	46	8,536	3	55
– Haemophilia	1,936	3	40	7,419	-5	48
Immunology	2,330	6	47	5,780	15	37
Speciality Care	324	46	7	1,213	8	8
Total	4,896	9	100	15,529	7	100

	Q4 '21	change	ratio	FY '21	change	ratio
	SEK M	%	%	SEK M	%	%
Europe	1,806	6	37	7,011	-5	45
North America	2,339	6	48	6,120	21	40
Rest of world	434	59	9	1,147	40	7
Other ¹	317	-1	6	1,251	3	8
Total	4,896	9	100	15,529	7	100

Royalty revenue

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2022 Q1: a good start

- Revenue up 24%, incl. phasing of Doptelet® sales to China
- Launch medicines¹ up 126%
- EBITA margin 26%, incl. provision for expected credit losses in Russia and restructuring (contract manufacturing closure, site simplification, efficiency programmes)
- EBITA margin adjusted 40%
- Efficiency programmes will focus resources into core areas, simplify the organisation and adjust the cost base to enable Sobi to continue sustainable growth and margin improvement over time
- Pipeline progressed with first efanesoctocog alfa phase 3 data
- Significant pipeline news flow over 2022 and 2023 timeframe
- 2022 outlook underpinned

Strategy on track:

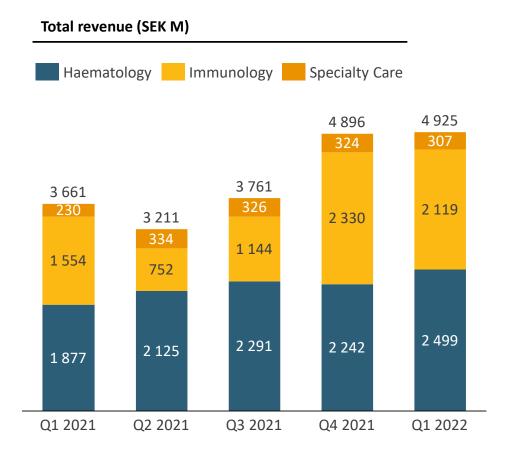
Good start in 2022 with delivery on strategic agenda and the 2022 outlook is underpinned



^{1.} Launch medicines include Doptelet, Aspaveli®/Empaveli™ and Gamifant®. Change at constant exchange rates.



2022 Q1: revenue, EBITA margin adjusted underpin outlook



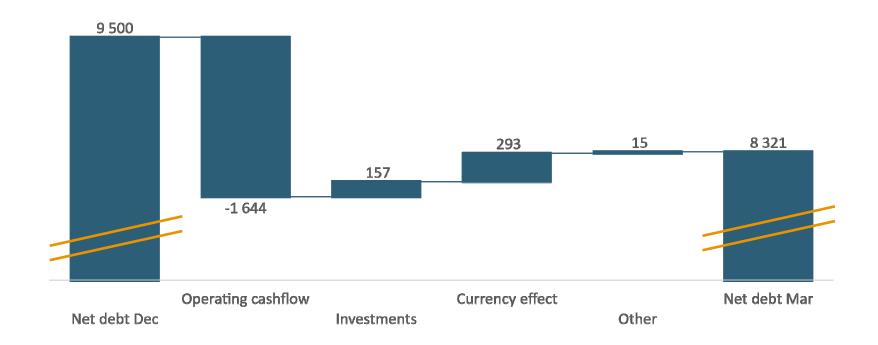
	Q1 2022	Q1 2021	Change	Full-year 2021
Total revenue	4,925	3,661	35%	15,529
Gross profit	3,409	2,935	16%	12,045
Gross margin ¹	69%	80%		78%
EBITA ¹	1,290	1,484	-13%	5,575
EBITA adjusted ^{1,2}	1,951	1,484	31%	5,575
EBITA margin ¹	26%	41%		36%
EBITA margin adjusted ^{1,2}	40%	41%		36%
Profit	543	696	-22%	2,679
Earnings per share (EPS), before dilution, SEK ¹	1.84	2.36	-22%	9.08
EPS, before dilution, SEK adjusted	3.67	2.36	55%	9.08
Operating cashflow	1,644	1,699	-3%	
Net debt (+)/net cash (-)	8,321	12,674		9,500

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

^{2.} Items affecting comparability in Q1 2022, see page 3 in report for further information.



2022 Q1: net debt - continued strong cash generation



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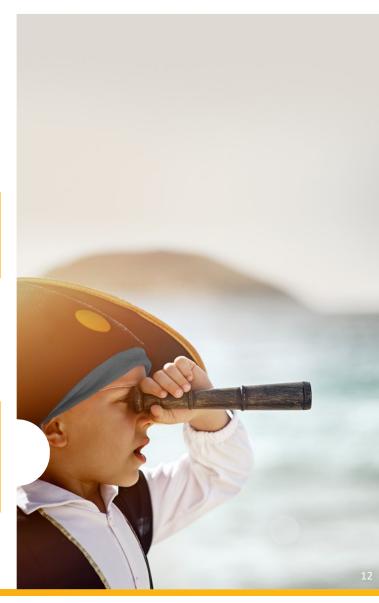
2022 outlook

Revenue

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

EBITA margin

Anticipated to be at a low 30s percentage of revenue (now based on EBITA margin adjusted²)





Pipeline: future growth

Phase 1		Phase 2	Phase 3	Registration
		Aspaveli/Empaveli (pegcetacoplan)	BIVV001 (efanesoctocog alfa)	Doptelet (avatrombopag)
		Transplant-associated	Haemophilia A	Chronic liver disease
		thrombotic microangiopathy after allogenic haematopoietic stem cell	Gamifant (emapalumab)	Aspaveli/Empaveli (pegcetacoplan)
	transplantation Aspaveli/Empaveli (pegcetacoplan)	Macrophage activation syndrome in rheumatological diseases	Paroxysmal nocturnal haemoglobinuria	
		Amyotrophic lateral sclerosis	SEL-212	Kineret (anakinra)
		Chronic refractory gout	COVID-19 patients at risk of developing severe respiratory	
		MEDI8897 (nirsevimab)	failure	
			Respiratory syncytial virus prevention	



Pipeline: continued news flow

Anticipated major upcoming pipeline news flow

H1 2022

Aspaveli/Empaveli – IC-MPGN and C3G¹: phase 3 study first patient

dosed (by Apellis)

Aspaveli/Empaveli – CAD²: phase 3 study first patient dosed

SEL-212 – CRG³: DISSOLVE II phase 3 study enrolment completion

H2 2022

Efanesoctocog alfa – haemophilia A: regulatory submission (US) (by

Sanofi in mid-2022)

Nirsevimab – RSV⁴ prevention: regulatory submission (US) (by AstraZeneca/Sanofi) (financial participation by Sobi)

Kineret® – COVID-19: regulatory decision, emergency use (US)

Gamifant – MAS⁵ in rheumatological diseases: EMERALD phase 3 study data readout

SEL-212 – CRG: phase 3 studies data readout

2023

Efanesoctocog alfa – haemophilia A (paediatric): XTEND-Kids phase 3 study data readout

Efanesoctocog alfa – haemophilia A: regulatory submission (EU)

Doptelet – CLD⁶: regulatory decision (JP)

Aspaveli/Empaveli – ALS⁷: MERIDIAN phase 2 study data readout (by Apellis in mid-2023)

Gamifant – MAS in rheumatological diseases: regulatory submission (US)

SEL-212 – CRG: regulatory submission (US)



^{1.} Immune complex-mediated membranoproliferative glomerulonephritis and C3 glomerulopathy 2. Cold agglutinin disease 3. Chronic refractory gout

^{4.} Respiratory syncytial virus 5. Macrophage activation syndrome 6. Chronic liver disease 7 Amyotrophic lateral sclerosis. Status as of 27 April 2022.



Sustainability: we have established a strong strategy, and our progress is recognised by external stakeholders









Our R&D is ethical and focused on medical need

We expand access to treatment

We are patient centric & engage with our communities

We contribute to knowledge to enhance the practice of medicine

We focus on patient safety













We help our people develop and keep them safe and healthy

We have zero tolerance for corruption

We source responsibly

We reduce our environmental footprint





13 point improvement in EcoVadis score



Commitment to the 2030 Agenda, the UN Sustainable Development Goals and the Paris Agreement



Sustainability: examples of our progress in 2021

