

Xbrane Biopharma AB Presentation Interim Report January – June 2022



*“Production for the launch of
Xlucane™ in Europe Q1 2023, is
being initiated*”*

Martin Åmark, CEO

Solna

July 22nd, 2022

***Post expiry of originator patents**

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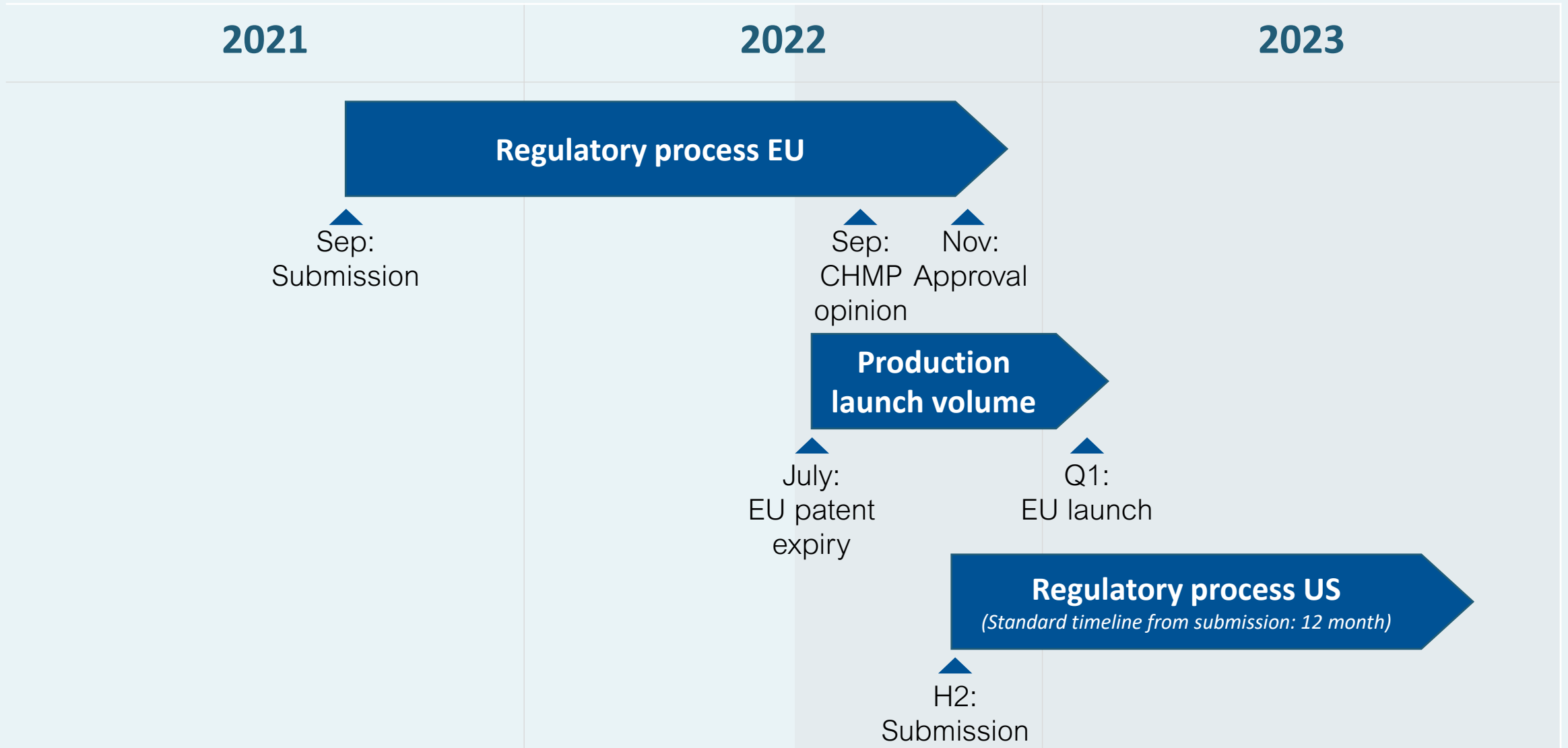
Xbrane has a portfolio addressing €53 billion of peak reference product sales

Candidate	Reference Product	Indication	Patent Expiry	Peak sales estimate*	Development Phase	Next milestone	Commercialization Partner
Xlucane™		<ul style="list-style-type: none"> Age-related macular degeneration Diabetic macular edema, Diabetic related retinopathy 	2020/22 (US/EU)	€3.2b	Registration	H2 2022 : EU approval H2 2022: Submission of Biologics License Application in US Q1 2023: Product launch in Europe	 
BIIB801		<ul style="list-style-type: none"> Rheumatoid arthritis Psoriasis Crohn's disease 	2024/25 (US/EU)	€2b	Process Scale-up	2023 : Initiation of clinical trial	
Xdivane™		<ul style="list-style-type: none"> Multiple oncology indications including Lung, liver, head & neck, kidney, colorectal cancer and melanoma Multiple Myeloma 	2028/30 (US/EU)	€13b	Process development	2022 : Demonstrate analytical comparability	Oncology portfolio with potential to out-license in one deal
Xtrudane™			2029/31 (US/EU)	€26b	Cell-line development	2023 : Demonstrate analytical comparability	
Xdarzane™			2029/31 (US/EU)	€9b	Cell-line development		

*Evaluate Pharma

Ambition to initiate at least one new biosimilar development program per year

Xlucane™ updated timeline to launch

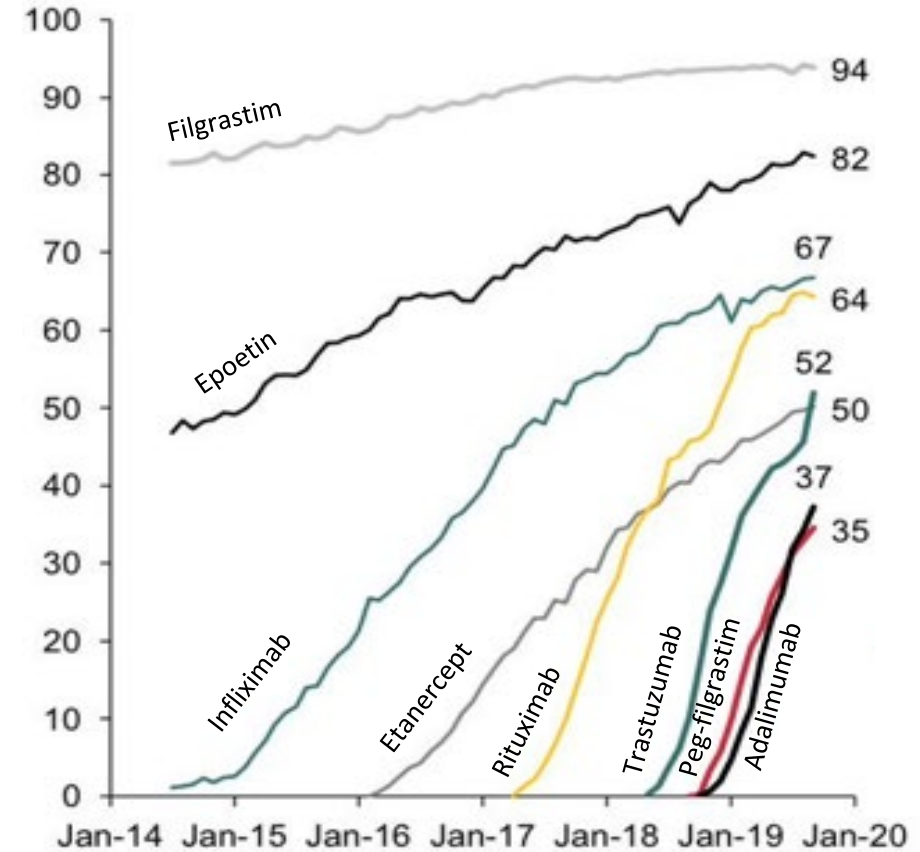


Xlucane™ – prospects of launch in Europe

Xlucane™ is addressing a €5.1 billion market opportunity in Europe

- Retinal VEGF inhibitor market in Europe generated sales of € 5.1 billion in 2021 – Lucentis® sales of approx. € 2 billion
- Xlucane™, one of three expected Lucentis® biosimilars to be launched
- STADA, with vast experience in biosimilars, will sell and market Xlucane™ across Europe
- Xbrane entitled to 50% of profits generated

+35% volume market share achieved at month 12 in recent biosimilar launches in Europe



Ambition to be the most attractive employer within our field

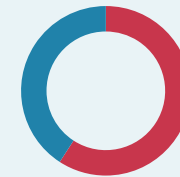
Certified as a Great Place to Work®

No of employees +30% YoY



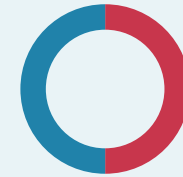
*) Parent Company only, previous quarters adjusted

Equality



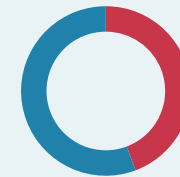
■ Women ■ Men

Management team



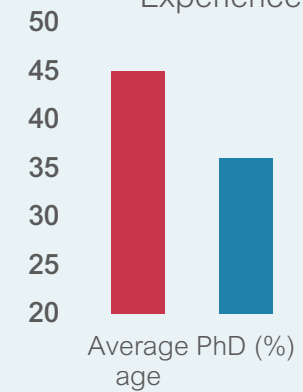
■ Women ■ Men

International



■ Foreign ■ Sweden

Experience



Upcoming Capital Market Events 2022



Xbrane Capital Market Day

August 2022
(invitations within short)



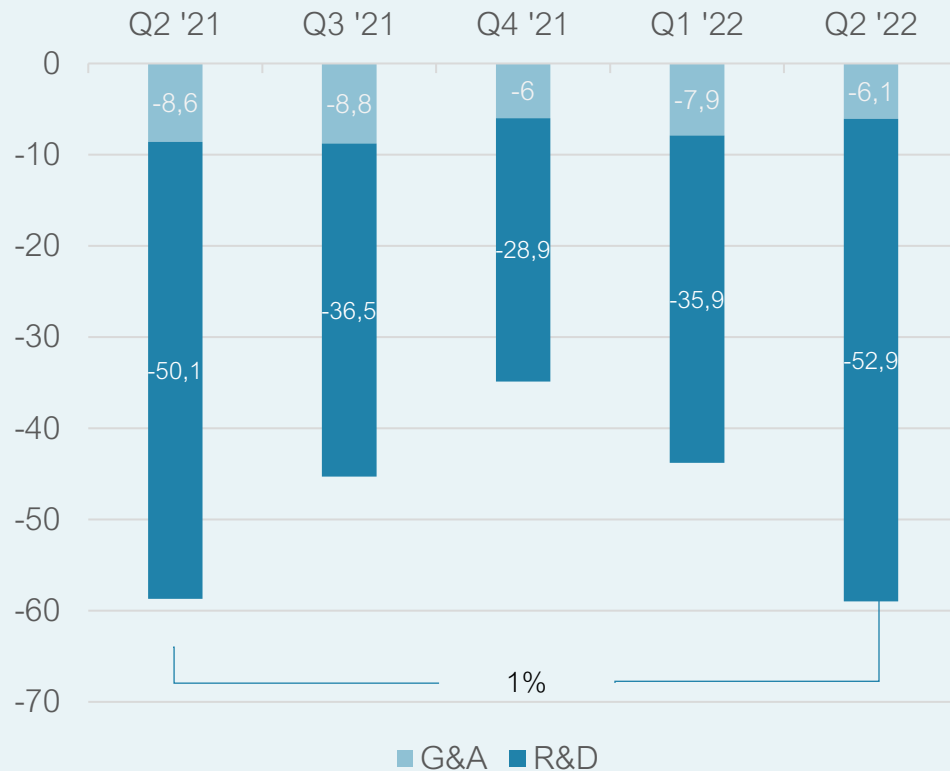
13th Annual Healthcare
Conference

September 2022



Interim Report January – June 2022, Financials

Company Expenses (G&A and R&D)

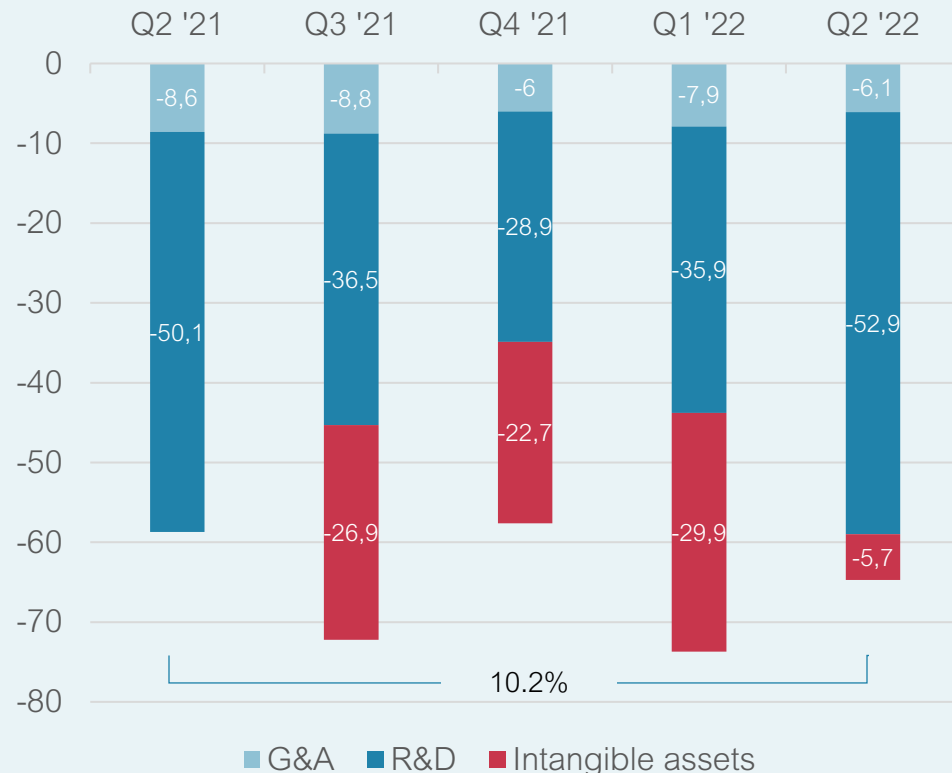


Total Company Expenses in the quarter was in line with last year even if the R&D costs increased somewhat and the overall G&A costs showed a decline.

The expensed R&D costs for the quarter amounted to net of SEK -52.9m (-50.1), representing 91 percent (86%) of total Operating Costs.

The R&D costs expensed in the P&L includes regulatory work and establishment of Supply Chain for Xlucane™, in addition with an expanding portfolio. More specifically, start-up costs for BII801 as well as exploratory work for the new Oncology portfolio.

Company Expenses (G&A and R&D)



Comparing “like-for-like” year-on-year, the total Operating Costs have increased with 10,2%

The total R&D costs (incl. Intangible Asset of SEK 5.7m) amount to SEK 58.6m, 90.6% of the total costs in the second quarter of 2022

Noteworthy is the reclassification of the development costs for Xlucane™ in accordance with IAS 38 that started July 1st, 2021.

During the second quarter, a total of SEK 5.7m was capitalized on the Balance Sheet, which is much lower compared to previous quarters, as expected as the development phase is coming to an end.

A total of SEK 86.2m is now capitalized on the Balance Sheet referring to Xlucane™.

Cash and Cash Equivalents



Shareholders' Equity



At the end of the second quarter 2022, Cash amounts to 250.1 MSEK.



Q&A