

Q4 Report Webcast

February 17, 2022

Disclaimer

IMPORTANT: You must read the following before continuing. The following applies to this document, the oral presentation of the information in this document by Oncopeptides AB (the “Company”) or any person on behalf of the Company, and any question-and-answer session that follows the oral presentation (collectively, the “Information”).

Oncopeptides is a biotech company focused on research and development of therapies for difficult-to-treat hematological diseases. The company uses its proprietary peptide-drug conjugate (PDC) platform to develop compounds that rapidly and selectively deliver cytotoxic agents into cancer cells. The first drug coming from the PDC platform, Pepaxto® (INN melphalan flufenamide), also called melflufen was granted accelerated approval in the U.S., on February 26, 2021 in combination with dexamethasone, for treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy and whose disease is refractory to at least one proteasome inhibitor, one immunomodulatory agent, and one CD38-directed monoclonal antibody. Oncopeptides voluntarily withdrew the drug from the U.S. market on October 22, 2021, due to worse overall survival data in the phase 3 OCEAN study. The study was a post-approval requirement under the accelerated approval program. Oncopeptides is developing several new compounds based on the PDC platform. Melflufen is not approved by any other registration authorities.

The Information contains forward-looking statements. All statements other than statements of historical fact included in the Information are forward-looking statements. Forward-looking statements give the Company’s current expectations and projections relating to its financial condition, results of operations, plans, objectives, future performance and business. These statements may include, without limitation, any statements preceded by, followed by or including words such as “target,” “believe,” “expect,” “aim,” “intend,” “may,” “anticipate,” “estimate,” “plan,” “project,” “will,” “can have,” “likely,” “should,” “would,” “could” and other words and terms of similar meaning or the negative thereof. Such forward-looking statements involve known and unknown risks, uncertainties and other important factors beyond the Company’s control that could cause the Company’s actual results, performance or achievements to be materially different from the expected results, performance or achievements expressed or implied by such forward-looking statements. Such forward-looking statements are based on numerous assumptions regarding the Company’s present and future business strategies and the environment in which it will operate in the future.

No representation, warranty or undertaking, express or implied, is made as to, and no reliance should be placed on, the fairness, accuracy, completeness or correctness of the Information or the opinions contained therein. The Information has not been independently verified and will not be updated. The Information, including but not limited to forward-looking statements, applies only as of the date of this document and is not intended to give any assurances as to future results. The Company expressly disclaims any obligation or undertaking to disseminate any updates or revisions to the Information, including any financial data or forward-looking statements, and will not publicly release any revisions it may make to the Information that may result from any change in the Company’s expectations, any change in events, conditions or circumstances on which these forward-looking statements are based, or other events or circumstances arising after the date of this document. Market data used in the Information not attributed to a specific source are estimates of the Company and have not been independently verified.

Participants



Jakob Lindberg
Chief Executive Officer



Klaas Bakker
Chief Medical Officer



Annika Muskantor
Chief Financial Officer



Status update

Jakob Lindberg, CEO

Highlights

- Q4 2021
 - Pepaxto voluntarily withdrawn from the US market
 - Refocus on R&D, close commercial operations, and scale down Sweden based organization
 - Focused clinical program to support ongoing EMA-review
 - Compassionate use program in the US established
 - Melflufen data presented at ASH meeting
 - Cash position of SEK 362 M by year-end 2021
- After reporting period
 - OCEAN data published in the Lancet Haematology
 - Further analyses of survival data led to a recission of the voluntary withdrawal letter of Pepaxto in the US



Near-term objectives Q4-2021



Near-term objectives – current status



Secure cash runway

→ Target achieved

- Closed commercial organizations in US and EU
- Significantly downsized the Sweden based HQ organization
- Decreased operational burn rate
- Reduced clinical trial activity



Regulatory process

Klaas Bakker, CMO

Rescission of voluntary withdrawal letter

- No intention to market Pepaxto in the US at this time
- Reach mutual understanding with the FDA on interpretation of the OCEAN study
- Dialogue with the FDA continues
- Oncopeptides will continuously work with the FDA to make melflufen available for those patients in the US currently being treated with the drug

EMA filing: CHMP opinion for melflufen

- Ongoing application process
 - Further analyses of survival data shared with EMA
 - 180-day questions end of March
- Unmet need: 70 patients included in Early Access Program in Europe
- Based on EMA process, market access preparations in Germany ongoing



Financial highlights

Annika Muskantor, CFO

Financial highlights October – December 2021



- Operating loss decreased to SEK 389.8 M (loss: 511.6)
 - Closing US- and European organizations and downsizing all global units
 - R&D decreased primarily due to less cost in the OCEAN project
 - OCEAN SEK 31 M (67)
 - Number of co-workers decreased from 321 to 162 (to be reduced further during Q1 2022 as notice periods end)
- Cash flow from operating activities neg. SEK 446.5 M (neg. 357.2)

Financial highlights January – December 2021



- Operating loss decreased to SEK 1,420.9 M (loss: 1,591.3) for Jan-Dec
 - R&D decreased primarily due to less cost in OCEAN and HORIZON studies
 - OCEAN SEK 140 M (314)
 - Build-up of commercial- and medical affairs in the US and EU through September - and closing thereof in Q4 – drove increase in M&S
 - Number of co-workers decreased to 162 (280) as of Dec 31₂₀₂₁ (To be reduced further during Q1 -22 as notice periods end)
- Cash flow from operating activities neg. SEK 1,516.4 M (neg. 1,296.5)
- Cash position was SEK 362.2 M (840.3) as of Dec 31, 2021
 - Directed share issue of net SEK 1,039 M in April 2021



Q&A



bringing hope through science