



Interim report January – September 2021

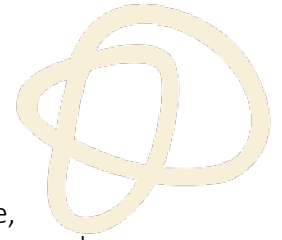
November 9th, 2021 at 3:00 p.m. CET.

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Anna Ljung, CEO



Disclaimer



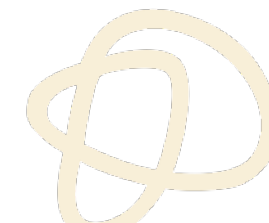
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2021 EU submission for potential new global market leader



MOB-015 has demonstrated world-leading ability to kill nail fungus

Partnerships in place – potential milestones of USD 120m

On track for launch – capturing full value potential

- 76%¹ of patients became fungus free, in two phase 3-studies including 800+ patients
- Additional de-risked US phase 3 study based on completed phase 3 studies to enable US approval and strengthen claims globally
- Targeting category leadership with USD 250-500m potential global product sales



EU



TAISHO PHARMACEUTICAL

Japan



Republic of Korea



Canada

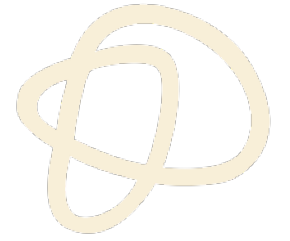


Scandinavia

- EU submission 2021
Product launch 2023
- Proven commercial track record from Kerasal Nail® – built SEK 440 million franchise with 30% market share in the US
- Commercialization process to be repeated for MOB-015

1) Other topical treatments demonstrating 30-54%.

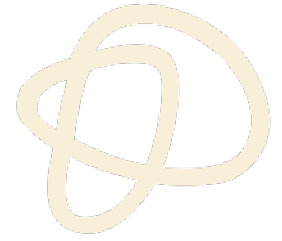
Significant events during Q3 2021



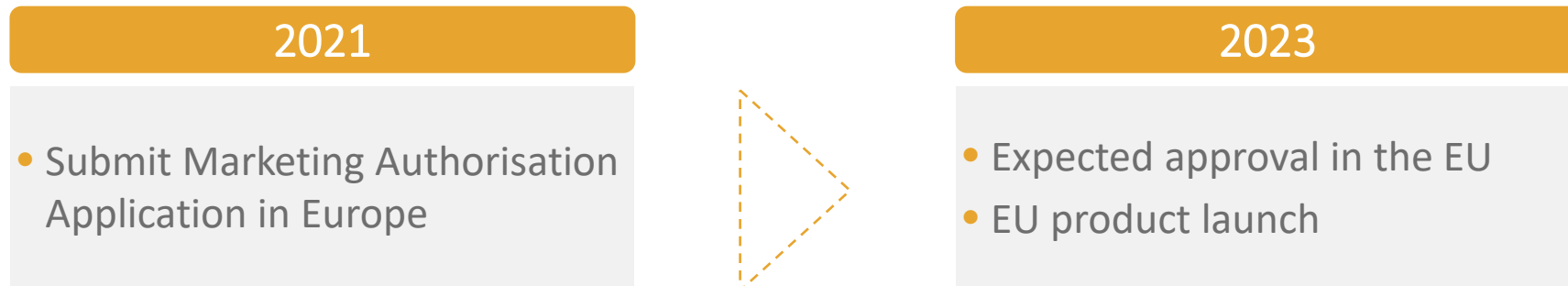
- Received approval for the pediatric plan from EMA
- Partnership with Allderma ahead of Scandinavian launch
- Preparations ongoing for the next Ph3-study in U.S.
- Agneta Larhed, VP Pharmaceutical Innovation & Development, joined the management team in September
- The Swedish Tax Agency has communicated acquisition cost of shares in Moberg Pharma AB after the distribution of shares in OncoZenge



On track to file for EU approval and product launch



- EMA's Paediatric Committee approval in September paves way for EU submission
 - Supplementary pediatric study during and after approval process for MOB-015
 - Enabling Moberg Pharma to pursue a full marketing authorization application
 - Opportunity to get data exclusivity for up to 10 years after market approval



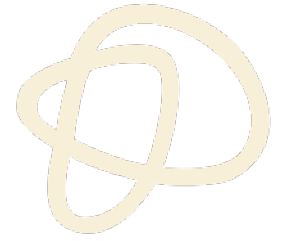
- Progressing US development plan in parallel

Partnership with Allderma

- Partnership with Allderma for Sweden, Norway and Denmark
 - Allderma is responsible for marketing, distribution and sales
 - Moberg Pharma is responsible for the manufacturing and product delivery
- Complements the existing licensing agreement for MOB-015 in Europe
 - Our European partner retains the right at a later date to assume the license in these markets.
- Allderma is managed by the team responsible for the successful Nordic launch of Nalox[®], our first-generation nail fungus product
- Great benefit in being directly involved in the launch of MOB-015 in our home market prior to additional launches with our partners

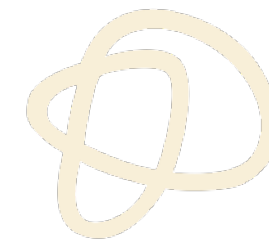


Preparations ongoing for the next Ph3-study in U.S.



- Purpose of the new study:
 - Enable market approval in the U.S.
 - Strengthen the product's clinical data and marketing claims globally
- The risk in the new study is significantly reduced through the experience gained from the previous studies
 - Cooperation with the same CRO and lead investigator as in the previous North American study
- Goal to submit documentation on the new study to the FDA and ethical committee in Q1 2022

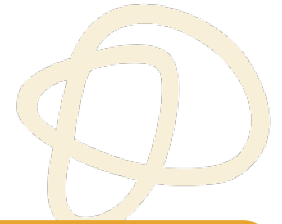
Key Financials



Last five quarters

(SEK thousand)	Jul-sep 2021	Apr-Jun 2021	Jan-Mar 2021	Oct-Dec 2020	Jul-Sep 2020	
Continuing operations						
Net revenue	-	-	-	-	-	
Gross profit	-	-	-	-	-	
Selling expenses	-	-	-	-4	-22	Expenses in line with previous periods
Business development and administrative expenses	-4,435	-3,702	-5,688	-5,405	-4,138	
Research and development costs	-600	-936	-1,207	-593	-1,143	
Other operating items	397	526	827	33	113	
Operating profit (EBIT)	-4,653	-4,119	-6,068	-5,969	-5,190	
Total profit for the period	-3,910	-3,324	18,639	-6,720	-4,666	Gain from BUPI spin off in Q1 2021
Cash and cash equivalents	111,407	124,195	133,611	29,285	30,006	Strong cash holdings from rights issue issued in Q1 2021
Investments in MOB-015	9,700	10,294	4,680	2,289	8,584	
Total Assets	453,512	456,488	463,209	479,704	364,060	

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