

Interim Report January – March 2024

May 7th, 2024



Disclaimer

The purpose of this presentation (the "**Presentation**") is to provide an overview of Moberg Pharma AB (publ) (the "**Company**"). For the purposes of this notice, "Presentation" means this document, its contents or any part of it, any oral presentation, any question or answer session and any written or oral material discussed or distributed during the Presentation meeting.

This Presentation is not a prospectus or similar offer document. This Presentation does not purport to contain comprehensive or complete information about the Company and is qualified in its entirety by the business, financial and other information the Company is required to publish in accordance with the rules, regulations and practices applicable to companies listed on Nasdaq Stockholm (the "Exchange Information"). Any decision to invest in any securities of the Company should only be made on the basis of a thorough examination of the Exchange Information and an independent investigation of the Company itself and not on the basis of this Presentation. Neither this Presentation nor any of the Exchange Information has been independently verified by any other person unless expressly stated therein. No representation or warranty, express or implied, is made as to, and no reliance should be placed on, the fairness, accuracy or completeness of the information or opinions contained in this Presentation.

Except where otherwise indicated in this Presentation, the information provided herein is based on matters as they exist at the date of preparation of this Presentation and not as of any future date. All information presented or contained and any opinions expressed in this Presentation are subject to change without notice. None of the Company or any of its directors, officers, employees, agents, affiliates or advisers is under any obligation to update, complete, revise or keep current the information contained in this Presentation to which it relates or to provide the recipient of with access to any additional information that may arise in connection with it.

This Presentation contains "forward-looking" statements. These forward-looking statements can be identified by the fact that they do not relate only to historical or current facts. In particular, forward-looking statements include all statements that express forecasts, expectations, plans, outlook and projections with respect to future matters, including trends in results of operations, margins, growth rates, overall market trends, the impact of interest or exchange rates, the availability or cost of financing, anticipated cost savings or synergies, the completion of strategic transactions and restructuring programmes, anticipated tax rates, expected cash payments, and general economic conditions. By their nature, forward-looking statements involve risk and uncertainty because they relate to events and depend on circumstances that will occur in the future and they are subject to change at any time. There are a number of factors that could cause actual results and developments to differ materially from those expressed or implied by these forward-looking statements, including risks associated with the inherent uncertainty of pharmaceutical research and product development, manufacturing and commercialization, the impact of competitive products, patents, legal challenges, government regulation and approval, the Company's ability to secure new products for commercialization and/or development and other risks and uncertainties detailed from time to time in the Company's interim or annual reports, prospectuses or press releases and other factors that are outside the Company's control. Any forward-looking statements made by or on behalf of the Company speak only as of the date they are made. The Company does not undertake to update forward-looking statements to reflect any changes in the Company's expectations with regard thereto or any changes in events, conditions or circumstances on which any such statement is based.

Potential new global market leader in Onychomycosis



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

Partnerships in place – potential milestones of USD 70m

Launch ongoing under brand name
Terclara®

- 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



Canada



Republic of Korea



Israel



Scandinavia

- National approvals in 13 EU
 countries 7 OTC, 6 Rx
- 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

3

Significant events during 2024 to date





MOB-015 is launched in Sweden by our partner Allderma under the brand name Terclara® and started to be available on Swedish pharmacy shelves in February. In parallel, work was ongoing in February and March to inform physicians and pharmacists about the unique benefits of Terclara®. Now, the focus is shifting to end consumers and the pharmacy chains are increasing their orders after consumer marketing began around the end of March.

Following the positive outcome of the Decentralized Procedure in June 2023, national approvals have been received in all 13 EU countries.

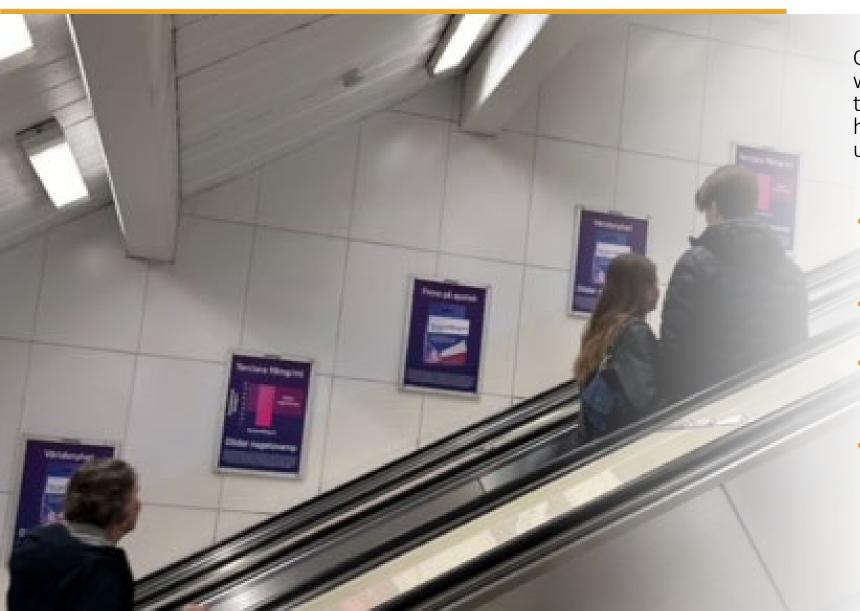
Progress in efforts to secure long term supply of terbinafine, **application** submitted, approval is expected before year-end.

In the North American study, half of the patients have now completed their treatment; 384 patients have been randomized at 33 study centers in the U.S. and Canada. Topline results are expected in January 2025.

Collaboration with Back Bay Life Science Advisors, which has conducted indepth interviews with U.S. payer representatives and is **organizing our process to find the best partner** for targeting U.S. dermatologists.

4

Terclara[®] launch ongoing- pharmacies report major demand for the new medication



Our partner Allderma AB, a company which specialises in the sale of overthe-counter (OTC) pharmaceuticals – has now launched sales of MOB-015 under the Terclara® brand.

- A majority of Swedish pharmacies have the product available on the shelf.
- TV marketing started as planned on April 1
- The pharmacy chains are increasing their orders after consumer marketing began
- Sales in the quarter mainly reflect the initial pharmacy orders. It is not until consumer marketing begins that demand from patients will affect the sales figures.

MOB-015 approval in EU >50% of countries approved as OTC already from start

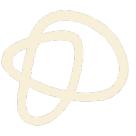
9

National approval in 13 European countries

- The approval in the European Union represents the first marketing authorization for Moberg Pharma's new onychomycosis treatment worldwide.
- Approval supported by two Phase 3 trials where MOB-015 demonstrated superior levels
 of mycological cure (76% vs 28% to 42% for comparators), and a significantly better
 complete cure rate compared to vehicle, without any serious adverse reactions.
- MOB-015 a topical formulation of terbinafine, enabling effective concentrations of terbinafine to the nail and nail bed while avoiding the risk of systemic exposure seen with oral terbinafine use.
- National approvals received:
 - National approvals for prescription sales (Rx) in Czech Republic, Denmark, Finland, France, Ireland and Spain
 - National approvals for over-the-counter sales (OTC) in Austria, Belgium, Hungary, Italy Netherlands, Norway and Sweden



Commercialization rollout of MOB-015



- Two-step process, driven by:
- 1. Moberg Pharma believe the results in the ongoing North American study is likely to strengthen the product claims further, including a **shorter dosing regimen** with the potential to deliver **superior complete cure** rates
- 2. Need to secure **sufficient API** for a pan-European launch
- 3. An early Scandinavian launch enables us to **gain valuable insights** into consumer behavior, collecting patient feedback and provide user data to support direct to OTC/OTC-switches in more countries

Additional phase 3 study in North America ongoing



The North American Phase 3 study is progressing according to plan

- Similar design as the already completed North American study
 - Multi-center, double-blind, randomized, vehicle-controlled study
 - Includes 384 patients in North America, 50% completed as of report date
 - 33 clinics in the U.S. and Canada are treating patients
 - Topline data expected January 2025
- Purpose of the new study:
 - Enable market approval in the U.S.
 - Strengthen the product's clinical data and marketing claims globally
- The new study builds on the experience gained from the previous studies
 - Cooperation with the same CRO and lead investigator as in the previous North American study

US commercialization ongoing



- US Strategy utilize the full potential and the knowledge from our first-generation product Kerasal Nail®
 - Build our own presence in the US market for podiatrists
 - Collaborate with a company with an established sales force for dermatologists
- Collaboration with Back Bay Life Science Advisors initiated:
 - Payer interviews reaffirms US market potential and high payment per treatment cycle for onychomycosis patients
 - Back Bay will organize our process for finding the best collaboration partner towards American dermatologists

Key Financials



Last five quarters

(SEK thousand)	Jan-Mar	Oct-Dec	Jul-sep	Apr-Jun	Jan-Mar	
	2024	2023	2023	2023	2023	
Net revenue	820	-	-	-	-	Launch o
Cost of goods sold	-328	-	-	-	-	
Gross profit	492	-	-	-	-	February
						COGS re
Selling expenses	-1,108	-1,167	-912	-817	-361	
Business development and administrative expenses	-6,983	-6,288	-5,509	-4,392	-5,414	Costs on
Research and development costs	-921	-1,037	-693	-1,109	-818	Costs co
Other operating items	624	257	-147	1,024	-80	which in
Operating profit (EBIT)	-7,896	-8,235	-7,261	-5,294	-6,673	to the la
Total profit for the period	-6,497	-6,445	-5,766	-3,852	-5,030	Less R&I
Cash and cash equivalents	38,631	60,555	101,504	51,951	84,540	MOB-01
Investments in MOB-015	17,822	33,215	33,642	22,761	34,498	phase 3 finalized
Total Assets	632,029	634,732	644,179	549,719	551,296	

Launch of Terclara in Sweden in February. Initial revenue and COGS reported*

Costs consistent with Q4 2023 which includes items attributed to the launch

Less R&D investments in MOB-015 in the ongoing US phase 3 study as recruitment finalized Q4 2023

^{*}Sales in the quarter mainly reflect the initial pharmacy orders. It is not until consumer marketing begins that demand from patients will affect the sales figures.

Potential new global market leader in Onychomycosis



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

Partnerships in place – potential milestones of USD 70m

Launch ongoing under brand name
Terclara®

- 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







Canada



Republic of Korea



Israel



Scandinavia

- National approvals in 13 EU
 countries 7 OTC, 6 Rx
- 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%.





Moberg Pharma AB (Publ) Gustavslundsvägen 42, 5 tr. 167 51 Bromma

mobergpharma.se