

VICORE PHARMA

Unlocking the potential of a new class of drugs

AGM 11 May 2023

Forward looking statement



This presentation may contain certain forward-looking statements and forecasts based on uncertainty, since they relate to events and depend on circumstances that will occur in the future and which, by their nature, will have an impact on Vicore Pharma's business, financial condition and results of operations. The terms "anticipates", "assumes", "believes", "can", "could", "estimates", "expects", "forecasts", "intends", "may", "might", "plans", "should", "projects", "will", "would" or, in each case, their negative, or other variations or comparable terminology are used to identify forward-looking statement.

There are a number of factors that could cause actual results and developments to differ materially from those expressed or implied in a forward-looking statement or affect the extent to which a particular projection is realized. Factors that could cause these differences include, but are not limited to, implementation of Vicore Pharma's strategy and its ability to further grow, risks associated with the development and/or approval of Vicore Pharma's products candidates, ongoing clinical trials and expected trial results, the ability to commercialize C21, technology changes and new products in Vicore Pharma's potential market and industry, the ability to develop new products and enhance existing products, the impact of competition, changes in general economy and industry conditions and legislative, regulatory and political factors.

No assurance can be given that such expectations will prove to have been correct. Vicore Pharma disclaims any obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise.



Vicore 2022 – Highlights



Q2 22

 First subject dosed in the phase 1 trial with ATRAG C106



Q4 22

- AlmeeTM pilot study showed a ~50% anxiety reduction. COMPANION trial initiated
- C103 selected as third ATRAG drug candidate
- Second Interim analysis of AIR trial reinforcing previous data, further strengthening benefit-risk profile
- Share issue raising 200 MSEK



Q1 22

Interim analysis of phase 2 AIR trial with C21 in IPF shows disease stabilization and increased lung function with excellent safety profile



Q3 22

- Study in healthy volunteers shows that C21 increases local blood flow.
- COVID-19 phase 3 trial (ATTRACT-3) reported and further clinical development discontinued.



Q1 23

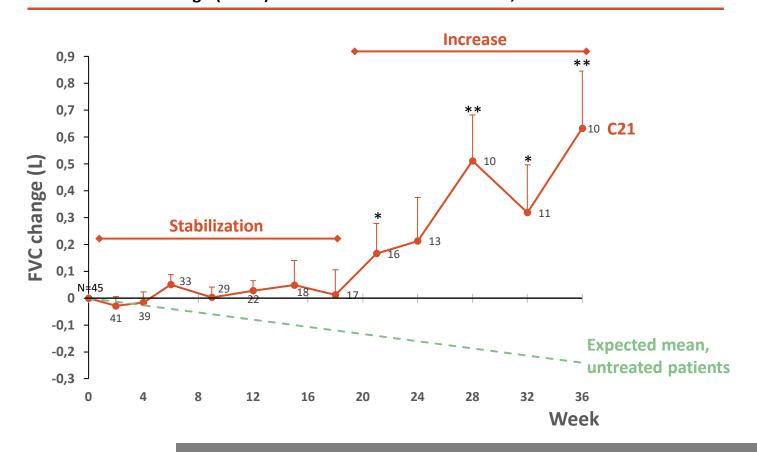
- **Innovation Passport** Designation in IPF by UK **MHRA**
- PoC study on endothelial dysfunction initiated



AIR interim analysis shows disease stabilization with later increase in FVC

Interim analysis October 2022

Mean change (±SEM) from baseline in FVC over time, observed values



Stabilizes disease over 36 weeks

Increased FVC in high responders

Safe and well tolerated – no GI signals

Encouraging patient testimonies

A placebo-controlled dose confirmatory trial (2b) will be the next step



Almee™ – Digital Therapy for Anxiety in Pulmonary Fibrosis

- alm
- > Treat symptoms of anxiety and improve quality of life in adults with pulmonary fibrosis
- Reduce costs for overburdened hospital systems (nurse/psychologist resources, hospitalizations, ER visits)

250 000 Pulmonary Fibrosis patients in the US

- Huge unmet need: 63% of patients with treatable levels of anxiety¹
- Current pharmacological treatments do not improve patients' quality of life
- Health care resource utilization two-fold versus controls

Almee™ – CBT-based digital therapy

- CBT has strong evidence base in anxiety
- Pilot study showed reduction of GAD-score by 49% after 4 weeks treatment

COMPANION – decentralized study

- US based RCT planned for 250 patients
- FPI December 2022
- Treatment period 9 weeks





Vicore 2023 – what is going on



- AIR trial
 - New update at ATS, May 21
- DDI trial
 - Completed during the year
- Endothelial dysfunction in diabetics
 - Completed during the year
- Companion Almee
 - Completed during the year
- C106 first in man trial
 - Completed during the year
- Preparations for phase 2b ANDAS
 - Start of study next year



Strong leadership team with extensive industry experience



CARL-JOHAN DALSGAARD, MD, PhD **CHIEF EXECUTIVE OFFICER**

Ex AstraZeneca R&D: Head of Therapy Area Pain Control, 10 years senior management. HealthCap: 19 years Venture Partner.







HANS JEPPSSON, PhD **CHIEF FINANCIAL OFFICER**

Cross-disciplinary background in finance and medicine. Ex Danske Bank: Equity analyst.









MIKAEL NYGÅRD, PhD VP BUSINESS DEVELOPMENT

Experienced healthcare Business Development executive, has led M&A and Corporate Development functions.







ELIN ROSENDAHL, MSc Pharm VP CLINICAL DEVELOPMENT

More than 20 years of global biopharmaceutical development at Pharmacia and SOBI. Solid experience of managing all clinical phases.







JOHANNA GRÄNS, PhD PROGRAM DIRECTOR, EARLY DEVELOPMENT

Extensive experience in preclinical R&D. Project management and regulatory affairs. Research experience in drug metabolism.



UNIVERSITY OF GOTHENBURG



ÅSA MAGNUSSON CHIEF COMMERCIAL OFFICER

More than 20 years of experience as a commercial executive in the pharmaceutical industry with focus on securing market access and launching rare disease medicines.





ROHIT BATTA, MBBS, MRCGP, MFPM **CHIEF MEDICAL OFFICER**

MD with extensive industry experience in Rare Diseases. Ex GSK: Led the global medical and clinical development of the world's first paediatric gene therapy.

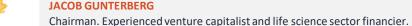


JOHAN RAUD, MD, PhD CHIEF SCIENTIFIC OFFICER

Ex AstraZeneca: Director Inflammation research.

25 years of experience in drug development. AstraZeneca 25







NINA CARLÈN CHIEF ADMINISTRATIVE OFFICER

More than 20 years of marketing and communications experience. Responsible for HR and company administration.



JESSICA SHULL. PhD **HEAD OF DIGITAL THERAPEUTICS**

More than 20 years of experience in the development and adoption of digital healthcare technologies.







STINE FURBO **HEAD OF CMC**

More than 20 years of experience with pharmaceutical drug product development & product supply from early development to launch.



HEIDI HUNTER

Biopharma, SOBI and Pharvaris)

HANS SCHIKAN

President Cardinal Health Specialty Solutions.

25 years in senior pharmaceutical development and commercialization positions.

25 years management experience in global pharmaceuticals (e.g. CEO of

Prosensa). Extensive board work in listed life science companies (e.g. Hansa

Board of Directors

SARA MALCUS

10 years experience in operational management and board work at AstraZeneca and GU Ventures.

MAARTEN KRAAN

Extensive experience in biomedicine, managerial roles at AstraZeneca.



CAROLINE SPEARPOINT, PhD THERAPY AREA LEAD RARE LUNG DISEASES

20 years industry experience from pharmaceutical, biotech and consulting, managing global cross-functional projects.







Summary



PIONEERING A NEW CLASS OF DRUGS

UNPRECEDENTED DATA IN IPF

TARGETING ALVEOLAR REPAIR AND ENDOTHELIAL DYSFUNCTION

STRONG CLINICAL PIPELINE

