Nanoform Capital Markets Day

June 4th, 2021

14.00-16.00 Helsinki time



Disclaimer

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Agenda – all presentations in real time!

- 1. Prof. Edward Hæggström CEO
- 2. Miguel Calado Chairman of the Board

3. Live Tour:

Dr. Niklas Sandler Chief Technology Officer Tuomas Puranen Process Engineer Director Antonio Da Silva Quality Control Director Dr. Eric Kissi Quality Control Scientist Dr. Satu Lakio Director Pharmaceutical Development Dr. David Rowe Head of Manufacturing Peter Blackburn Manufacturing Operations Manager Rossella Dal Maso Production Team Leader Marco Minerva Operational Excellence Specialist

- 4. CCO Christian Jones Commercial & Team
- 5. Mike Rea Strategic Innovation Advisor
- 6. Dr. Maria Lume Biologics
- 7. Dr. Elisabetta Micelotta STARMAP®
- 8. CFO Albert Hæggström Finance
- 9. Q&A
- 10. CEO Prof. Edward Hæggström Closing remarks



Prof. Edward Hæggström CEO





Nanoform 12-month rapid progress





Nanoform Clients & Collaborations



Nanoform Global Commercial Team



Nanoform new mid-term business targets 2025





API = Active Pharmaceutical Ingredient GMP = Good Manufacturing Practice

Miguel Calado Chairman of the board





Live Tour

1. Research & Development

CESS® Technology

2. Quality Control

Solid State Characterization

- 3. Pharmaceutical Development
- 4. GMP Manufacturing
 - GMP Nanoform Line and Personnel airlock

Materials Storage, Sampling, Dispensing

Tiered Accountability Process

Dr. Niklas Sandler Chief Technology Officer **Tuomas Puranen Process Engineer** Director Antonio Da Silva Quality Control Director Dr. Eric Kissi Quality Control Scientist Dr. Satu Lakio Director Pharmaceutical Development Dr. David Rowe Head of Manufacturing Peter Blackburn Manufacturing Operations Manager Rossella Dal Maso Production Team Leader Marco Minerva Operational Excellence Specialist



Christian Jones Chief Commercial Officer

Commercial







> Experienced global sales team driving momentum and the shift in company focus from technology development to commercialization



Nanoform unlocks possibilities for patient-centric innovation

Adherence to longterm therapy for chronic illnesses in developed countries averages 50%

In developing countries, attrition rates are









Setting the North Star

Nanoforming empowers the adoption of "North Star TPPs" that meet patient demands from the start of development. True patient-centric innovation need not be the provision of the fast-follower...

...Greater choice

Decreased NCE/NBE attrition to accelerate development of next generation therapeutics

...More empowerment

Novel administration options that align with patient preferences to put them in control of their medicine journey

...Improved compliance

Potentially reduced AEs, improved toxicity and modulated release profiles to improve outcomes





2012 2026 61% 45%



small molecules launched

New launch "balance" increasingly weighted towards NCEs and away from re-formulations, but the reformulation market represents a large volume opportunity



biologics launched

Total biologics Biobetter

Although currently a small amount, we expect to see more "Biobetters" reach the market through the next decade as blockbuster biologics approach LoE and biosimilar players seek to differentiate



We know where opportunity is and how our technology can support it...

Oral

Inhaled

Transnasal

NxEs n=554 6% 46% Preference for innovators to pursue "tried and tested" oral and IV routes 38%



Reformulations n=42

More heterogeneous approaches adopted

Strategic approach

Increase PoS in customer-preferred routes while fostering confidence to set a patient-centric North Star TPP from the start

Encourage new, and de-risk ongoing patient-centric re-formulation programmes



nanoform.com @nanoformf Source: PharmaProjects accessed May 2021. All molecules entering preclinical or Phase I development since Jan 2019. "Non-disclosed" routes of admin excluded

YTD Opportunities by Clinical Phase n=57

NF opportunities **closely** mirror the market in terms of development phase with most emphasis on tackling solubility/bioavailability issues in **preclinical and phase 1** and adding value to marketed products in lifecycle extension and drug reformulation



small is powerful*

Small is a platform enabler across drug delivery





Mike Rea

Nanoform

- Strategic Innovation Advisor

CEO, IDEA Pharma

An overview of the opportunities in biologics





Nanoforming a potential game-changer in Biologics too





Dr. Maria Lume Biologics Team Leader

Processing of biologics at Nanoform





The BIO team

- Production of bio-nanoparticles
- Formulation of bio-nanoparticles
- Characterization and qualitative analysis of the bio-nanoparticles





Nanoforming biologics

Gentle, bottom-up process

So far used for processing peptides and proteins ranging from 6 to 140 kDa in size (range expanded from 6 to 66 kDa during Q2/2021)

Nanoparticles as small as 50 nm are produced



Nanoparticles of biologics





Tuning the particle size of HSA





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Assessment of insulin activity in cellular assays

The activity of bulk unprocessed insulin and nanoformed insulin was compared *in vitro*, using HepG2 cell-line



HepG2 cells on a dish

Upon activation, protein's phosphorylation levels change Phosphorylated protein = active signal-transmitting protein



pAKT was used as a read-out for the insulin activity assay



Processed insulin 1-month stability assessment

Fresh samples

1-month old processed samples



Both unprocessed and processed insulin activate IR and the downstream pathways

Nanoformed insulin is active after 1 month at + 4°C



LDH – Lactate dehydrogenase

Lactic dehydrogenase

Comprises of 4 subunits, molecular weight 140kDa

LDH catalyses the interconversion of pyruvate and lactate.

Highly sensitive to shear and temperature

Used to evaluate the presence of damage and toxicity of tissue and cells.

LDH is elevated in certain pathological conditions such as cancer







Assessing LDH size and retained activity





First results on LDH activity assay



Lyophilization min/max adopted from: Strøm Larsen, 2019, Pharm Dev Technol Tuderman, 2018, Int J Pharm Fang, 2018, J Pharm Sci Ogawa, 2016, J Oleo Sci Al-Hussein, 2015, Pharm Dev Technol Al-Hussein. 2013. J Pharm Sci Kawai, 2007, Pharm Res Iwai, 2007, J Pharm Sci

The activity assay depicts the conversion rate from lactate to pyruvate (Concentration of NADH)



Future prospects

smallis powerful®

Building of additional lines to support commercial projects.

Broadening the product portfolio capabilities e.g., by processing antibodies etc.

Supporting expansion towards GMP for biologicals.



Nanoform the stars that will shine the brightest with...



Dr. Elisabetta Micelotta Science & Technology Team Leader





- Molecules predicted to be best amenable to CESS[®]-powered nanoforming.
- Molecules that exhibit optimal production characteristics.

- **Revisiting drug candidates** unnecessarily discarded by AIs trained on old particle engineering techniques.
- Rapidly **picking winners** among **new drug** candidates.











Inputs and outputs





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Solubility Prediction in sc-CO₂

Very good match between predictions & measurements

Solid lines \rightarrow **Solubility predictions.**

Filled circles → Solubility measurements.

Shaded areas → Uncertainty in predictions.

Different colors \rightarrow Different temperatures of CO₂

tarmap^{2.0}

Illuminating the future of Pharma

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A continuous growth

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Albert Hæggström Chief Financial Officer

Nanoform is here to fill the gap

Global drug R&D pipeline size and growth

Global drug development by phase, 2021

*Launched and in development (7%), Pre-registered (1%), Registered (1%) and Suspended & N/A (1%)

Global clinical drug development phase trends, 2007-2021

Global number of companies with active pipelines

Share of pipeline contributed by top 10 companies, top 25 companies and companies with just 1 or 2 two drugs, 2011-2021

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Distribution of R&D companies by HQ country/region, 2021

Global drug development by category, 2021

Biological versus non-biological drugs as a % of pipeline

R&D pipeline by therapy group, 2020 and 2021

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Nanoform new mid-term business targets 2025

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API = Active Pharmaceutical Ingredient GMP = Good Manufacturing Practice

Q&A

Questions through:

1) Dial-in:

FIN: +358981710520 SE: +46850558358 UK: +443333009269 US: +1 6319131422

PIN: 12138908#

- 2) Type in the question in the question box below
- 3) E-mail to IR@nanoform.com

Prof. Edward Hæggström CEO

Closing remarks

Thank you!

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