It is not a question of if, but when...





...the next Influenza-induced epidemic will kill 650.000 people and strike 1 billion patients



1



Investor/partner presentation 2023



BUILDING ON TWO DECADES OF SCIENCE

Headquarters

Tromsø, Norway

Established

2017

Key indication

Influenza in high-risk immunocompromised patients

Product offering

Gel and Nasal spray



Prominent research environment

Close ties to the Tromsø research environment, a prominent Norwegian research environment



Spin-off

Pharma Holdings is a spin-off of Lytix Biopharma, a company listed on Euronext Growth



Topical treatment

A mucoadhesive nasal spray formulation has successfully been developed.



Development stage

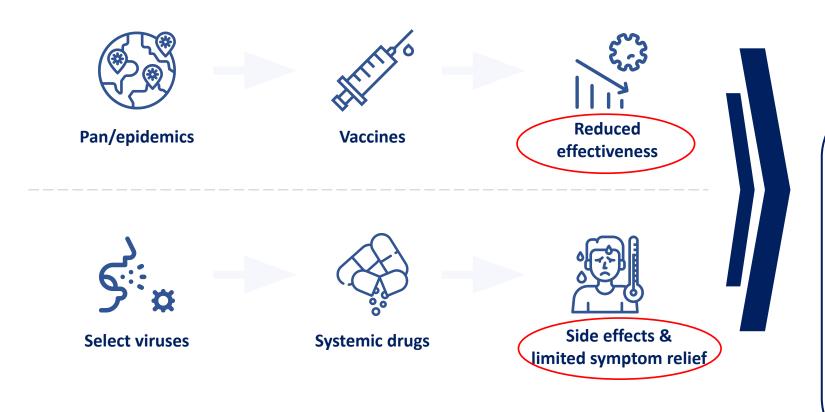
LTX-109 has completed initials trials showing a good safety profile when given topically.

Clinical trials in vulnerable high-risk patients with respiratory viral infections are planned



THE **UNMET NEED**: LTX-109 TO ADDRESS THE WEAKNESSES OF COMMON ANTIVIRAL APPROACHES AND TREATMENTS

Most prevalent treatment methods



Pharma Holdings' solution

Topical treatment

Selected benefits of topical therapeutics

- Direct virucidal effect as opposed to only viral replication inhibition
- Less prone to effect deterioration
- Able to address a variety of viruses
- Fewer side effects than systemic drugs

DEVELOPMENT DRIVEN BY EXPERIENCED TEAM

Christian Lütken CEO (FTE)

- Several years of experience from various fields within medicine as a medical doctor
- Previous experience as Head of Department in a private healthcare company
- · Experience from DNB's Global Healthcare Team as an industry expert financing global healthcare companies
- · Holds a MD from the University of Oslo

Management Team



Torsteinn Erlingsson CFO (FTE)

- More than 25 years of experience from Corporate Finance and Business development, where he worked on several biotech transactions
- Instrumental in securing soft-money and equity financing Holds a MSc from University of Tromsø







Johnny I. Ryvoll Vice President Projects (FTE)

- More than 25 years of experience from Corporate Finance, Project Management and Business Development
- Holds a BSc in Electrical Engineering from University of Utah and an MBA from University of Washington



Board



Bernt Endrerud Chairman

- Active innovator and investor
- Has established and led several successful businesses
- More than 30 years of experience as a business owner and developer



Lars Vorland Board Member

- Holds a MD and a PhD, where he specialized in molecular biology and medical microbiology
- Previous experience working for the Norwegian Institute of Public Health and Helse Nord RHF







Xavier Frapaise (MD) Chief Medical Officer

- 40 years experience from Pharma industry international experience from drug development, from early phase 1 to 3. both EU and US.
- Proven track record in securing funding for start up
- Experience in the space of infectious diseases, among them Covid-19.





Niklas Hammarstedt Chief Quality Officer (FTE)

- Several years' experience within Quality Assurance in diverse positions
- Chief Quality Officer in Atlas Antibodies
- Former Head of Production Documentation on Octapharma
- Holds a MSc in Chemistry and Pharmaceutical engineering from KTH Royal Institute of Technology

octapharma Tatlas antibodies





Edvard Christian Fuglset Dahl Chief Business Development Officer (FTE)

- Several years of Sales and Marketing experience within healthcare industry
- Business Development Specialist for Inven2
- Former Director of Business Development I Mentis Cura
- Director of Business Development in Media









Håvard Ebbestad **Board Member**

- More than 20 years of experience as a chief executive at various pharma companies (incl CEO of Norwegian subsidiary of Pfizer)
- Current position: CEO of Fürst laboratories



Øyvind K. Arnesen **Board Member**

- Holds a MD from University of Oslo and has more than 10 years of clinical practice
- Extensive experience from the pharmaceutical industry, including Medical Director at BMS, Boehringer Ingelheim and CEO of Ultimovacs



DEVELOPMENT SUPPORTED BY EXPERIENCED TEAM



Rene Bommer
Nasal Spray expert

- >30 years experience from academia and pharma industry
- Ph.D in chemistry
- Published several articles in medical device and packaging journals
- Has regulary been invited as speaker at international drug delivery conferences



Prof. David Smith (MD)
Scientific Advisor (US)

- Head of Division of Infectious diseases and public health at UC San Diego
- Awarded 37 mill USD in federal funding as PI
- Advisor to FDA in Covid-19 trials
- Speaker at Nobel Price symposium 2022 (Covid-19 therapeutic interventions)

Advisory Team (Contracted)



Cecil NickRegulatory Strategic Advisor

- Vice President at Parexel Consulting
- 40 years experience from clinical development and regulatory affairs.
- Particular experience in development of anti-infective medicines including having supported many Covid-19 therapeutic interventions.



Dr Med Lars Heggelund *Scientific Advisor (Nordics)*

- Specialist in Infectious Diseases in 2018
- Head of Research at Drammen Hospital
- Professor at University of Bergen
- 51 scientific publications



Ass Prof. Martin Hoenigl (MD)
Scientific Advisor (EU)

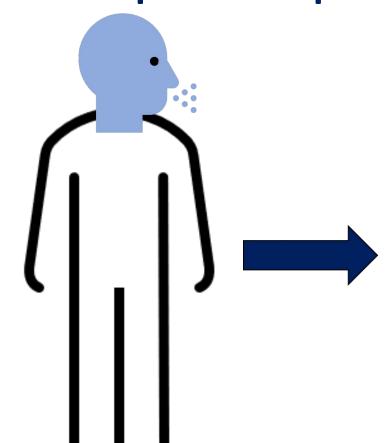
- Works at Division of Infectious disease at medical university of Graz, Austria.
- Awarded researcher of the year 2011
- Author on 250 publications in the field of infectious diseases.
- Particular experience in conducting research on clinical mycology, virology and respiratory viruses.



NORDIC MENTOR NETWORK for ENTREPRENEURSHIP PHARMA HOLDINGS is a member of Novo Nordisk mentor program

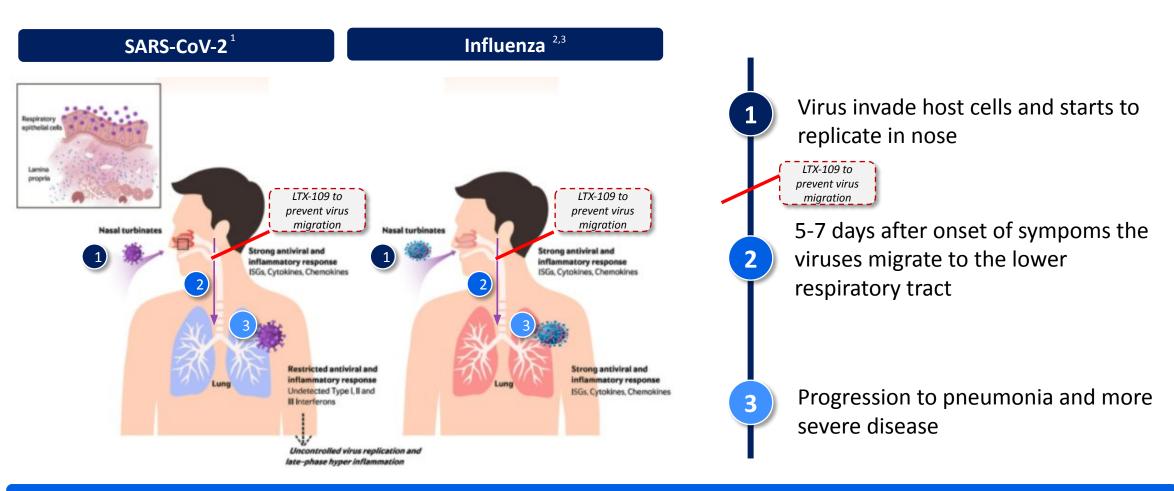
HIGH RISK PATIENTS AND UNMET NEED Immunocompromized patients

- High mortality
- Little/no effect of vaccines
- ✓ Longer hospital stays
- Prolonged viral shedding
- Prolonged course of disease
- Shortcomings of current SoC



LTX-109, as a direct virucidal, will demonstrate reduction in mortality rates, length of hospital stay and symptom duration in non-hospitalized early stage immunocompromized influenza patients, as an add-on to standard of care

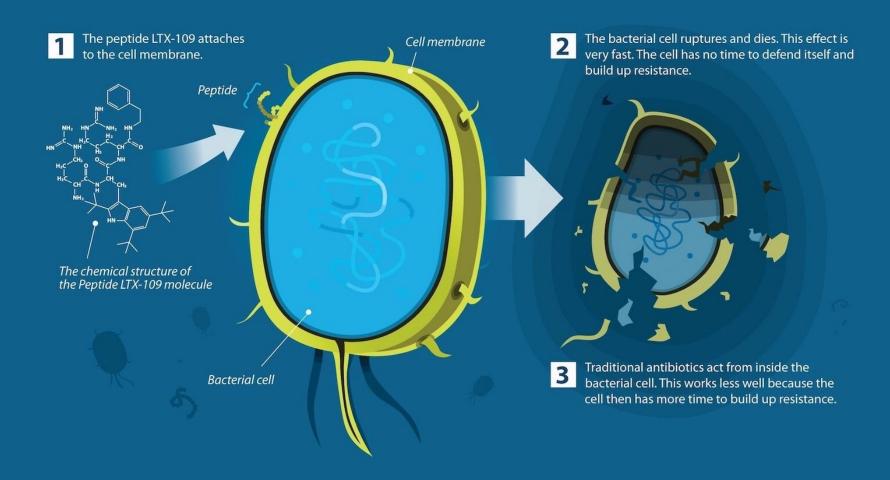
THE NOSE IS THE INTIAL AND MAIN SITE FOR VIRAL REPLICATION



High Influenza viral load titers (nasal and pharyngeal swabs) seem to correlate with worse outcome and increased risk of mortality⁴

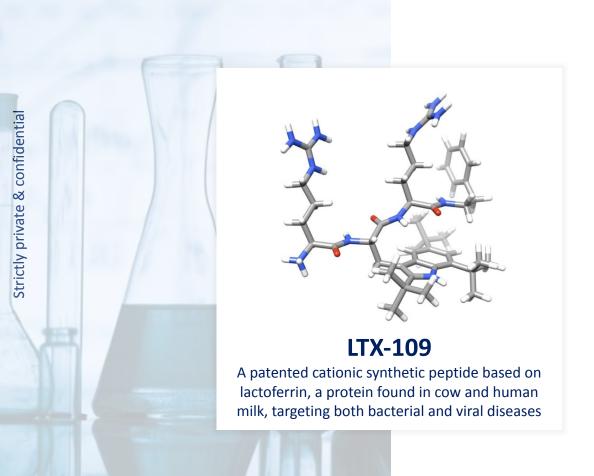


LTX-109 IN ACTION





MAIN FEATURES OF LTX-109





Rapid membrane lysing mechanism of action



Low propensity for resistance development



Broad spectrum of antiviral and antibacterial activity



Safe and well tolerated



PROMISING IN VITRO RESULTS AGAINST RESPIRATORY VIRUSES

Enveloped viruses:

Virus	1% LTX-109 tested against RSV, IVA and SARS-CoV-2 (1 hour incubation)	
	Log decrease	Percentage
RSV	3.25	99.949
Influenza A	3.42	99.969
SARS-CoV-2	4.33	99.995

Virus	3% LTX-109 tested against RSV, IVA and SARS-CoV-2 (60 sec incubation)	
	Log decrease	Percentage
RSV	2.92	99.88
Influenza A	1.42	96.19
SARS-CoV-2	2.27	99.50



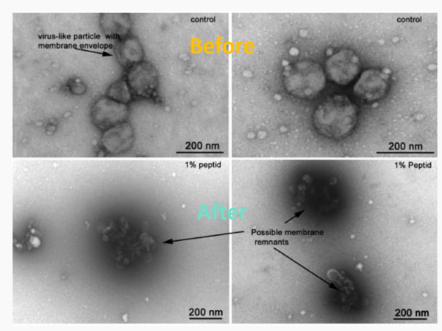


Figure 1: The image shows Lenti-virus-like particles with intact envelope (control) and dissolved envelope (1% LTX-109)

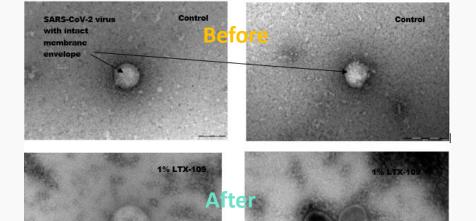


Figure 2: The loss of morphological integrity of the virus post-treatment with LTX-109, note the capsid of the virion without envelop

DEVELOPMENT STAGE AND MAJOR MILESTONES

Key milestones going forward Indication / activity Q3'21 Q4'21 2022 2023 2024 2025 2026 In vitro efficacy established Phase Ib Phase Influenza in High-Risk* Phase III In In vivo to be conducted in H1 Dose IIb(PoC) Registration patients **Nasal Spray** vivo 2023 finding IMP GMP ready **US** approval Phase III Proof of Cond Dose finding In vivo efficacy Registration **EMA** approval Efficacy against all common Phase Nasal decolonization pathogens of S. Aureus Gel Dose exploration **Proof of Concept** Additional EUR 1.5m in bridge funding if **Additional funding** Raised a total of EUR 2.8m since spin-off, funded through EUR 7m needed, funding in place to read phase Ib is (TBD) for pivotal Q3'21 through Q1'24* out of Phase IIa successful **Phase III studies** (~EUR 30m)

CMC

*Haematopoietic stem cell transplan (potentially also to include hematologic malignancy patients) Multiple batches of API produced

New manufacturing contracts in place

✓ Low COGs



Nasal spray
API scale-up



Nasal spray
DP scale-up

MARKET EXTENSION POTENTIAL

Primary market Influenza high risk US/EU:

Infection rate: 1:8 (~5m)

Immunocompromised patients: ~40m

Tentative pricing: ~200 USD/treatment***



Primary market revenues:

Market size: 1B USD

COGS per treatment*: 10EUR

Pharma Holdings' revenue per treatment: 40EUR

Potential revenues: 200M USD



MARKET EXTENSION:

- Other high-risk patients (elderly, COPD, heart failure etc)
 - Other relevant respiratory viruses: SARS-CoV-2, RSV etc



Market extension revenues:

Market size: XX BUSD

COGS per treatment*: 10EUR

Pharma Holdings' revenue per treatment: 40EUR

Potential revenues: Multibillion USD

GLOBAL SALES TAMIFLU+XOFLUZA 2022: 400M USD, EXPECTED GLOBAL SALES 2027: 650M USD**



^{***} Tentaive price etimate report by external company Pricia (Denmark)

^{**} Source global data

EXECUTIVE SUMMARY

Our need towards success:

- ☐ Funding of EUR 7m to secure runway through Phase Ib
- ☐ Funding of EUR 30-35m to secure runway through Phase IIb PoC
- ☐ Commercial partners to accelerate clinical development and secure market access



Market and medical unmet need

Urgent need to develop new antivirals – address a huge unmet medical need



Evidence

Preclinical studies have provided a strong signal of antiviral activity. Nasal spray adminitration limits toxiticity, side effects and drug – drug interactions



Influenza as proof-of-concept

Study to be initiated H1 2023 in ferrets will provide info required by regulators and IRBs, and will guide dose-regimen selection for the first human studies to be initiated H2 2023



Development towards market approval

The clinical development will be based on early signs of efficacy in FIH Phase Ib (H2 2023), then seamless transition through adaptive study design in the planned for Phase IIb/III





THANK YOU FOR YOUR ATTENTION

CHIEF BUSINESS DEVELOPMENT OFFICER
Edvard Christian Fuglset Dahl
Phone: +47 480 47 041

Email: edvard.dahl@pharmaholdings.no

