HERANTIS

Toward clinical breakthroughs based on leading science

Danske Bank Small & Mid Cap Day

LSX Nordic

Pareto Securities 11th Healthcare Conference

1-3 September 2020

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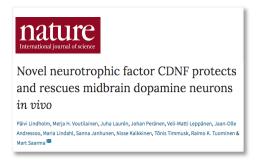
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High Impact Science In High Impact Diseases

- Highly differentiated assets with novel and distinct scientific approaches
- Targeting unique therapeutic opportunities in despairing diseases
- Potentially disease-modifying and curative in their current indications, and beyond
 - CDNF (Cerebral Dopamine Neurotrophic Factor) biologic for Parkinson disease
 - Lymfactin® VEGF-C gene therapy for secondary Lymphedema
- Both assets safely into humans, in clinical proof of concept studies, funded to data readouts

World class research published in Nature and Science









CDNF For Parkinsons Disease (PD)

- PD an incurable brain disease caused by degeneration of dopaminergic neurons in midbrain
- CDNF a natural protein acts in several ways to potentially modify the diseases process
- Massive medical need with 7-10 million patients worldwide, societal costs \$25bn US alone

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- Highly promising preclinical & early clinical data confirms safety, suggests biological activity and potential slowing of disease progression

Phase I-II completed successfully Aug 2020



Lymfactin® For Lymphedema (LE)

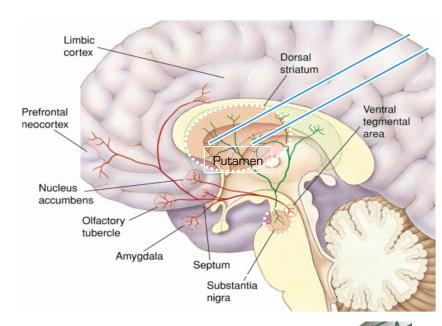
- LE is severe chronic swelling and accumulation of lymph due to damaged lymph drainage
- Lymfactin[®] is a local gene therapy, which stimulates natural growth of lymphatic vessels
- Huge medical need with over 140 million patients worldwide
 - Lymfactin® is targeted as the first curative treatment of secondary BCAL Lymphedema by reconstituting the lymphatic system
- Phase I completed, Phase II ongoing with read-out expected Q1/2021

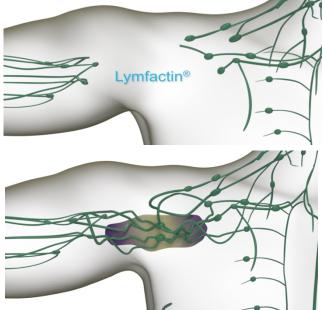


How Our Drugs Work

 CDNF biologic therapy is dosed directly into the brain where it promotes functional recovery of dopaminergic neurons in the putamen, and protects the dopaminergic cell bodies in the substantia nigra

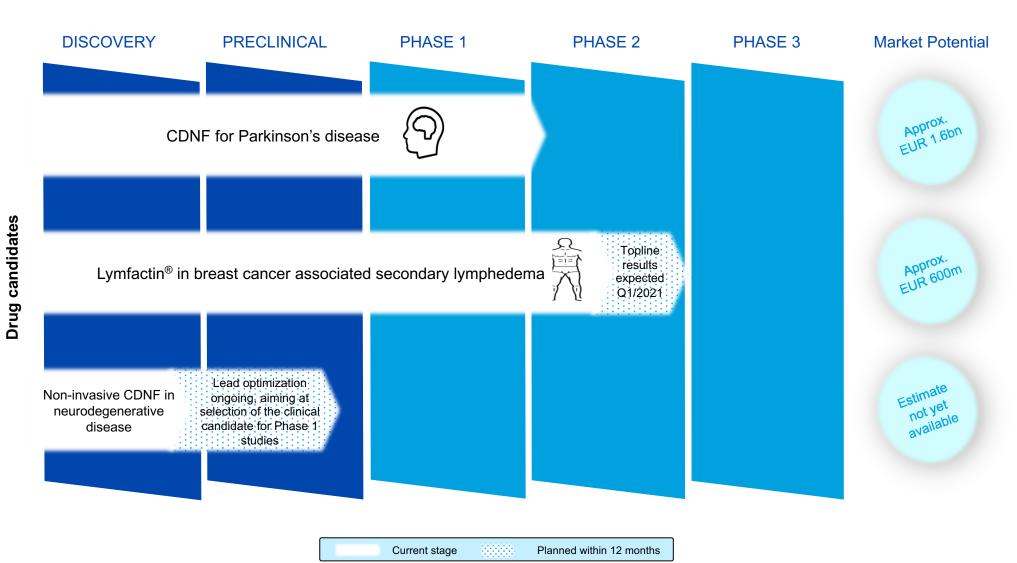
 Lymfactin® VEGF-C gene therapy is administered locally into the armpit together with lymph node transfer surgery as a single-dose treatment where it promotes reconstitution of a functional lymphatic system







Rich Pipeline Progressing Towards Significant Markets



CDNF Strong Safety Data And Promising Efficacy Potential

- Successful FIH Study Just Completed August 2020
- Primary objective to establish safety

Data confirms CDNF safe and well tolerated

Secondary objective to explore efficacy indicators

Data varied but suggests potential slowing of disease progression in some patients, as well potential signals of biological activity in some patients (PET DAT)

Promising data in FIH study where safety is focus, patients have advanced disease, most dopaminergic neurons are lost, CDNF doses are low, and observed improvements in efficacy scores not powered for statistical evaluation

Study results to be presented via company webinar on 15th September – please consult www.herantis.com

"In addition to the study's favourable safety profile, it is encouraging to observe an improvement in the UPDRS motor score as well as an indication of a biological response compared to baseline, which may be clinically relevant. In chronic diseases, such as Parkinson's, where patients continue to decline over time despite receiving treatment, the differences observed in this study present a potential opportunity for the development of meaningful therapeutic effects."

- Prof Per Svenningsson, Karolinska University Hospital (Study PI)



CDNF Results Compelling Platform For Next Gen xCDNF

- Common challenge in CNS diseases is drug delivery to the brain
 - CDNF currently administered via surgically implanted device
- xCDNF, i.e. smaller selected fragments of CDNF
 - retain the biological activity of CDNF (comparable efficacy in PD models; same target!)
 - o penetrate the BBB → administered subcutaneously no need for medical device
 - highly potent molecules

 xCDNF optimisation at final stages of compound selection to take forward for development

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Lymfactin Established Safe, Phase II Underway

Phase I study completed 2019 (follow-up continues until 2023)

Data confirms Lymfactin® safe and well tolerated by patients

 Promising improvements observed in signs and symptoms of LE after 12-month follow-up (uncontrolled data)

- Randomized Phase II study ongoing
 - Results expected Q1 2021



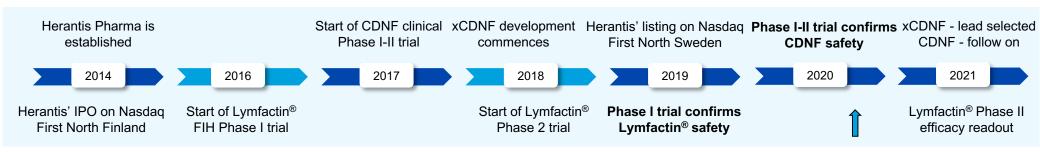
"Breast cancer-associated lymphedema is a highly debilitating disease that severely impacts quality-of-life of patients that have already experienced a significant impact on their day-to-day health. Addressing this unmet medical need is an important endeavor to which we are excited to contribute. Lymfactin® has very interesting product profile addressing underlying root cause by restoring functionality of lymphatic vasculature, which will be critical to evaluate going forward."

- Associate Professor Anne Saarikko of Hospital of Helsinki (Study PI)



Herantis Proposition Summary

- Groundbreaking science at leading edge of biological and gene therapy innovation
- Potential to impact significant diseases that have shown limited progress in decades
- Confirmed successful translation from discovery into humans
- Highly skilled team who know what is needed, and how to do it
- Clear path to potential partnerability and commercialization





Thank You

