

#### **PILA PHARMA AB**

Norra Vallgatan 72 211 22 Malmö Sweden

pilapharma.com

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# CEO Gustav H. Gram increases holding in Pila Pharma AB

Pila Pharma AB (publ) (FN STO: PILA) announces that newly appointed CEO Gustav H. Gram has purchased additional shares.

The purchases occurred in two tranches:

Friday May 10th, where 13.756 shares in the company were acquired. The shares were purchased at an average price of SEK 2,4615.

Monday May 13th, where 8.453 shares in the company were acquired. The shares were purchased at an average price of SEK 2,5976.

Mr. Gram's direct holdings in Pila Pharma AB now amounts to 45.441 shares.

## CEO Gustav H. Gram comments:

"We feel we have a great clinical-stage drug candidate for type 2 diabetes with further possible benefits on cardiovascular disease and potentially obesity. The market has developed immensely around this therapeutic area in recent years and we have a candidate that could potentially disrupt the way cardiometabolic conditions are treated. At a time with globally exploding numbers in patients with lifestyle disease it has never been more important to drive innovation, and the team and myself are adamant to show that we can succeed with bringing forward the first TRPV1-antagonist to proof-of-concept. On this basis I feel it's important, both as investor and as CEO, to increase my commitment and keep re-investing in the company."

To stay up to date on news, events & where to meet us, please see our website: <a href="https://www.pilapharma.com/">https://www.pilapharma.com/</a>

For more information:

Gustav H. Gram, CEO ghg@pilapharma.com

Pila Pharma's share ticker PILA is subject to trade on Nasdaq First North Growth Market, Sweden with Aqurat Fondkommission AB as Certified Adviser. Contact: M: <a href="mailto:ca@aqurat.se">ca@aqurat.se</a> - T: +46 (0)8 684 05 800



## About PILA PHARMA AB (Publ)

Pila Pharma is a Swedish biotech company based in Malmö, Sweden. The aim of the company is to develop TRPV1 antagonists as a novel treatment of type 2 diabetes and potentially of other diseases with an inflammatory background.

The Company owns a TRPV1 asset with data and chemical entities including the development candidate XEN-D0501. Further, the Company owns use-patents covering the use of TRPV1-antagonists as treatment of obesity and diabetes and intends to submit further patents regarding the synthesis, formulation, or use of XEN-D0501 or back-up compounds. In July 2022, the Company was awarded orphan drug designation ("Orphan drug designation") for XEN-D0501 as a treatment for erythromelalgia.

Pila Pharma currently focuses on 3 projects within Type-2 Diabetes, Erythromelalgia, and Abdominal Aorta Aneurism.

# About XEN-D0501 and TRPV1 antagonists

XEN-D0501 is a selective, synthetic potent small molecule TRPV1 antagonist that was in-licensed in 2016. TRPV1 antagonists that down-regulate neurogenic inflammation, has demonstrated applications across pain and inflammatory diseases and potentially plays a role in diabetes and potentially other metabolic disorders like obesity. Prior to in-licensing, XEN-D0501 had been found to have a good safety profile in other (non-diabetic) patient groups. Pila Pharma has to date completed two phase 2a clinical trials (PP-CT01 and PP-CT02), that both demonstrated that XEN-D0501 is well tolerated by type 2 diabetic patients. Further, PP-CT02, demonstrated that XEN-D0501 (administered as 4 mg bi-daily for 28 days) - with statistical significance versus placebo - enhanced the endogenous insulin response to oral glucose. Furthermore, ANP, a heart failure biomarker, was highly statistically significantly reduced. During 2023 we could report a very good tolerability of XEN-D0501 following 13 weeks administration of very high doses in 2 animal species, and XEN-D0501 can thus progress into longer clinical trials.

Recently, finances to sponsor a slim phase 2a dose-escalation study was secured and the study is being prepared with the objective of identifying the maximal tolerable dose of XEN-D0501 in overweight or obese people with type 2 diabetes. In addition to this, it's aimed to further identify (trends for) a reduction of HbA<sub>1c</sub>, body weight and ANP, a relevant marker of cardiovascular disease.

## **About Diabetes and Obesity**

Diabetes is a globally spanning pandemic with a staggering estimated prevalence of more than 537 million people living with diabetes corresponding to approximately 8-10% of the global adult population. Among these, its estimated that more than approximately 90 % of all diabetics suffer from type-2 diabetes, whilst approximately



less than 10% suffers from type-1 diabetes.

Despite recent therapeutic advances, large and growing unmet needs exist both from efficacy, safety, and accessibility standpoints.

Obesity is most often preceding the development of type 2 diabetes and a serious risk-factor for not only developing type 2 diabetes but also co-morbidities resulting in "whole body dysfunction" and subsequent development of several diseases. The accumulated effect is a year-long reduction in quality of life for obese people with or without diabetes. Obesity leads to an increased risk of developing cardiovascular disease that eventually results in premature death and shortening of life duration. Recent advances by "Big Pharma" in the development of effective anti-obesity drugs, has proven that pharmacological weight management is possible and leads to obvious quality-of-life and longevity benefits for people living with obesity. Even long-term, public health costs are expected to be reduced if the clinically negative effects of the obesity pandemic are limited. This has sparked a general interest in future potential oral treatments that can meet the accessibility criteria needed to stimulate growing demand and several acquisitions have been done in the obesity segment recently.

#### About Erythromelalgia

Erythromelalgia is a rare disease where neurogenic inflammation plays a role in the development of symptoms. The disease can cause near-constant or episodic pain (ranging from mild tingling to severe burning sensations), and redness to extremities. It most commonly affects the feet but may also occur in the hands, face, or other parts of the body with both nerves and blood vessels involved. Symptoms are frequently managed through avoidance of pain triggers. The disorder can be extremely debilitating, with a significant negative impact on quality of life and with potential to impact mortality rates among young people and the suicide rates among adults. Pila Pharma aims to conduct a small proof of concept study in persons with erythromelalgia to demonstrate an effect of XEN-D0501 on reducing perceived pain during "flare ups".

## About Abdominal Aorta Aneurism

Abdominal Aorta Aneurism is a cardiovascular disease with 'balooning' of the lower part of the main artery of the body, aorta. The cause is unknown, but risk factors are atherosclerosis, high blood pressure, cardiovascular inflammation and infection as well as trauma. It affects millions of people globally and accounts for the death of 1% of men over the age of 65. It develops gradually over several years up to a dilatation of more than 3mm in diameter when surgery to insert a stent to prevent rupture is then the only treatment option, which is both expensive and with possibility for complications. Currently no preventive treatment is available. In November 2023 a research collaboration was entered on investigating the effect of XEN-D0501 on Abdominal Aorta Aneurism growth in mice.