PRESS RELEASE
01 December 2023 07:00:00 CET

Save the Date: Egetis to host Investor Day on December 19, 2023

Stockholm, Sweden, December 1, 2023. Egetis Therapeutics AB (publ) (Nasdaq Stockholm: EGTX) today announced that the Company will host an Investor Day on Tuesday December 19, 2023, in Stockholm, Sweden, for investors, analysts and media.

Time: Tuesday, December 19, 2023, at 3:00 pm - 6:00 pm CET (9:00 am – 12:00 pm ET)
Venue: Redeye, Mäster Samuelsgatan 42, Stockholm, Sweden

The event will also be accessible through a live webcast.

During the event, the Company will provide an update and review of its strategy and pipeline. The presentations will focus on the unmet medical need, development plans, pre-launch activities and commercialization plans for the investigational drug *Emcitate* (tiratricol), being developed as a possible treatment for MCT8 deficiency, as well as additional activities to create and enhance long-term shareholder value. Presentations will be made by members of Egetis' management team and invited key opinion leaders.

A complete agenda and a link to the live webcast will be shared in advance of the event.
Presentations will be held in English.

If you would like to attend the event in person in Stockholm, please register by email to info@egetis.com

For further information, please contact:

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About Egetis Therapeutics

Egetis Therapeutics is an innovative and integrated pharmaceutical company, focusing on projects in late-stage development for commercialization for treatments of serious diseases with significant unmet medical needs in the orphan drug segment.
The Company's lead drug candidate *Emcitate* is under development for the treatment of patients with monocarboxylate transporter 8 (MCT8) deficiency, a highly debilitating rare disease with no available treatment. In previous studies (Triac Trial I and a long-term real-life study) *Emcitate* has shown highly significant and clinically relevant results on serum thyroid hormone T3 levels and secondary clinical endpoints. Egetis submitted a marketing authorisation application (MAA) for *Emcitate* to the European Medicines Agency (EMA) in October 2023.
After a dialogue with the FDA, Egetis is conducting a small randomized, placebo-controlled pivotal study in 16 patients to verify the results on T3 levels seen in previous clinical trials and publications. Egetis intends to submit a new drug application (NDA) in the US for *Emcitate* in mid 2024 under the Fast-Track Designation granted by FDA.
Emcitate holds Orphan Drug Designation (ODD) for MCT8 deficiency and resistance to thyroid hormone type beta (RTH-beta) in the US and the EU. MCT8 deficiency and RTH-beta are two distinct indications, with no overlap in patient populations. *Emcitate* has been granted Rare Pediatric Disease Designation (RPDD) which gives Egetis the opportunity to receive a Priority Review Voucher (PRV) in the US, after approval. This voucher can be transferred or sold to another sponsor.
The drug candidate *Aladote* is a first in class drug candidate developed to reduce the risk of acute liver injury associated with paracetamol (acetaminophen) overdose. A proof of principle study has been successfully completed and the design of the upcoming pivotal Phase IIb/III study with the purpose of applying for market approval in the US and Europe for *Aladote* has been finalized after completed interactions with FDA, EMA and MHRA and study start is planned after *Emcitate* submissions have been completed. *Aladote* has been granted ODD in the US and in the EU.
Egetis Therapeutics (STO: EGTX) is listed on the Nasdaq Stockholm main market. For more information, see www.egetis.com

Attachments

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