PRESS RELEASE
18 August 2022 08:00:00 CEST

Save the Date: Egetis to host a Capital Markets Day in Stockholm on October 13, 2022

Stockholm, Sweden, August 18, 2022. Egetis Therapeutics AB (publ) (Nasdaq Stockholm: EGTX) today announced that the Company will host a Capital Markets Day on Thursday, October 13, 2022, in Stockholm, Sweden, for investors, analysts and media.

Time: Thursday, October 13, 2022, at 1:00 pm - 5:00 pm CEST
Venue: Erik Penser Bank, Apelbergsgatan 27, 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, commercial opportunities, and additional activities to create and enhance long-term shareholder value. Presentations will be made by members of Egetis' management team and Board, as well as 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.

To attend in person in Stockholm, please sign up here
http://enews.penser.se/public/event/RegistrationForm/4342584A794541594A7240

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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 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 T3 levels and secondary clinical endpoints. As a result of fruitful regulatory interaction Egetis intends to submit a marketing authorisation application (MAA) for *Emcitate* to the European Medicines Agency (EMA) in the first half of 2023 based on existing clinical data.
In the US, after discussions with the FDA, Egetis will conduct a small randomized, placebo-controlled 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-2023 under the Fast-Track Designation granted by FDA.
Emcitate is currently being investigated in the fully recruited Triac Trial II, a Phase II/III study in very young MCT8 deficiency patients (<30 months of age) exploring potential disease modifying effects of early intervention from a neurocognitive and neurodevelopmental perspective. Results are expected in the first quarter of 2024 and are expected to be submitted post-approval to regulatory authorities shortly thereafter.
Emcitate holds Orphan Drug Designation (ODD) for MCT8 deficiency and resistance to thyroid hormone type beta (RTH- #) in the US and the EU. *Emcitate* has been granted Rare Pediatric Disease Designation (RPD) which gives Egetis the opportunity to receive a Priority Review Voucher (PRV) in the US, after approval.
The drug candidate *Aladote* is a first in class drug candidate developed to reduce the risk of acute liver injury associated with paracetamol (acetaminophen) poisoning. 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. *Aladote* has been granted ODD in the US and has received a positive opinion for ODD 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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