PRESS RELEASE
31 October 2024 16:15:00 CET

Egetis' Nomination Committee for the 2025 Annual General Meeting

Stockholm, Sweden, October 31, 2024. Egetis Therapeutics AB (publ) (NASDAQ Stockholm: EGTX) today announced the composition of the Nomination Committee for the 2025 Annual General Meeting (AGM) to be held on May 6, 2025.

The Nomination Committee, which has been appointed in accordance with the principles adopted by the Annual General Meeting held on May 6, 2024, comprises the following members:

James Brush, appointed by Frazier Life Sciences
Peter Lindell, appointed by Cidro Förvaltning AB
Peder Walberg, appointed by Cetoros AB

Mats Blom, Chairman of the Board of Directors, will co-opt to the Nomination Committee.

The Committee's assignment is to present proposals regarding Chairman and other members of the Board, as well as remuneration to the Board´s members, to the AGM. The Nomination Committee shall also submit proposals for appointment and remuneration of auditors. Further, the Committee shall submit proposals regarding the process to appoint the Nomination Committee to the AGM in 2026.

Shareholders who wish to submit proposals to the Nomination Committee can do so by email to info@egetis.com (please label emails 'Nomination Committee'). Proposals should be submitted no later than January 17, 2025.

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® (tiratricol) 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 randomized, placebo-controlled pivotal study in 16 evaluable patients to verify the results on T3 levels seen in previous clinical trials and publications. Egetis will update the market as soon as recruitment has been completed and at that point inform about the timing of availability of top-line results, and the expected timing of the subsequent NDA filing.

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® (calmangafodipir) 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. The design of a pivotal Phase IIb/III study (Albatross), with the purpose of applying for market approval in the US and Europe, has been finalized following interactions with the FDA, EMA and MHRA. The study start has been postponed until *Emcitate* marketing authorization submissions for MCT8 deficiency 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

Egetis' Nomination Committee for the 2025 Annual General Meeting