

Egetis Therapeutics to Participate in Upcoming November Investor Conferences

Stockholm, Sweden, November 6, 2025. Egetis Therapeutics AB (publ) ("Egetis" or the "Company") (NASDAQ Stockholm: EGTIX), today announced that management will participate in upcoming investor conferences in November 2025.

Stifel 2025 Healthcare Conference

Date: November 12, 2025

Location: New York, NY, USA

Time of Presentation: 2:00 - 2:30 PM EST (8:00 - 8:30 PM CET)

Format: Presentation by Nicklas Westerholm (CEO), 1x1 Meetings

The presentation will be available via webcast [here](#)

LifeSci Capital and Sofinnova Partners Growth & Innovation Summit

Date: November 17, 2025

Location: London, UK

Format: 1x1 Meetings

DNB Carnegie Nordic Healthcare Conference

Date: November 25, 2025

Location: Oslo, Norway

Time of Presentation: 3:35 - 3:55 PM CET

Format: Presentation by Yilmaz Mahshid (CFO), 1x1 Meetings

Aktiespararnas Stora Aktiedagar

Date: November 26, 2025

Location: Stockholm, Sweden

Time of Presentation: 5:40 - 6:05 PM CET

Format: Presentation by Karl Hård (VP Investor Relations)

The presentation (in Swedish) will be available via webcast [here](#)

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 February 2025 the European Commission approved Emcitate® as the first and only treatment for MCT8 deficiency in EU. Egetis initiated the launch of Emcitate® in Germany on May 1, 2025.

The Company has agreed with the FDA to submit a rolling NDA for Emcitate® (tiratricol), commencing in December 2025 targeting a complete NDA submission in early 2026 and completion of FDA's review process in the third quarter of 2026.

Based on feedback from the FDA, the NDA for Emcitate® (tiratricol) for treatment of MCT8 deficiency will be based on currently available clinical data from Triac Trial I, Triac Trial II, ReTRIACt, EMC Cohort Study, EMC Survival Study and the US Expanded Access Program.

Tiratricol holds Orphan Drug Designation (ODD) for MCT8 deficiency and resistance to thyroid hormone beta (RTH-beta) in the US and the EU. MCT8 deficiency and RTH-beta are two distinct indications, with no overlap in patient populations. Tiratricol has been granted Breakthrough Therapy Designation and Rare Pediatric Disease Designation (RPDD) by the FDA, which gives Egetis the opportunity to receive a Priority Review Voucher (PRV) in the US, after approval.

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 development program for Aladote® has been parked 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 is listed on the Nasdaq Stockholm main market (Nasdaq Stockholm: EGTX).

For more information, see www.egetis.com

Attachments

[Egetis Therapeutics to Participate in Upcoming November Investor Conferences](#)