PRESS RELEASE
24 February 2026 11:00:00 CET

Egetis Therapeutics to Present at Upcoming Investor Events

Stockholm, Sweden, February 24, 2026. Egetis Therapeutics AB (publ) ("Egetis" or the "Company") (NASDAQ Stockholm: EGTX), today announced that Nicklas Westerholm (CEO) and Yilmaz Mahshid (CFO) will present at upcoming investor events in February and March.

Cantor Rare Disease Month
Date: February 26, 2026
Location: Virtual (register here)
Time of Presentation: 10:00-11:00 am EST (4:00-5:00 pm CET)
Format: Fireside chat with Nicklas Westerholm (CEO)

Leerink Global Healthcare Conference
Date: March 11, 2026
Location: Miami, FL, USA
Time of Presentation: 1:00-1:30 pm EDT (6:00-6:30 pm CET)
Format: Presentation by Nicklas Westerholm (CEO), 1x1 meetings
The presentation will be available via webcast here

DNB Carnegie Nordic Healthcare Conference
Date: March 12, 2026
Location: Stockholm, Sweden
Time of Presentation: 9:50-10:20 am CET
Format: Fireside chat with Yilmaz Mahshid (CFO), 1x1 meetings
No webcast

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 developed 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. Emcitate® (tiratricol) is not approved in the USA.

The Company completed a rolling New Drug Application (NDA) for Emcitate® (tiratricol) in the USA on January 29, 2026. The FDA is expected to confirm within 60 days that the NDA submission is complete. As a designated Fast Track and Breakthrough Therapy, Egetis has requested Priority Review, and if granted, the FDA review should be completed within six months following the 60-day filing review period.

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. 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

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