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Egetis Therapeutics raises approximately SEK 180 million through an oversubscribed rights issue

Stockholm, Sweden, May 19, 2022. Egetis Therapeutics AB (publ) (“Egetis” or the “Company”) (Nasdaq Stockholm: EGTX) announced on March 21, 2022, that the Board of Directors had resolved to carry out a guaranteed issue of shares with preferential rights for the Company’s existing shareholders (the “Rights Issue”). The Board of Directors’ resolution of the Rights Issue was approved by an extraordinary general meeting held on April 13, 2022. Today, Egetis hereby announces that the Rights Issue was oversubscribed. Through the Rights Issue, Egetis will receive proceeds amounting to approximately SEK 180 million before transaction costs.

The result of the Rights Issue shows that 44,755,164 new shares, corresponding to approximately 90.4 percent of the Rights Issue, have been subscribed for with subscription rights. Additionally, 5,840,471 shares were subscribed for without subscription rights of which 4,765,404 shares, corresponding to 9.6 percent of the Rights Issue, have been allotted to investors that have subscribed for shares without subscription rights. The Rights Issue is thus oversubscribed.

The purpose of the Rights Issue is to finance the preparations for regulatory submissions for market approval in EU and US, initiate the establishment of a commercial infrastructure in Europe and US for Emcitate® and pre-launch activities, as well as general corporate purposes, in addition to providing financial flexibility.

Egetis’ CEO Nicklas Westerholm comments: *“We are pleased with the great interest to participate in our Rights Issue and are thankful for the continued support of our current shareholders, in particular specialist life science investor Flerie Invest AB for their commitment to increase their shareholding. Additionally, we further strengthen the specialist investor base in Egetis through Linc AB and welcome them as a new shareholder. The new funds contributed through the Rights Issue will be vital to continue the step-wise establishment of a commercial infrastructure in the EU and the US for Emcitate® as well as pre-launch activities.”*

As confirmation of allotment of shares subscribed for without subscription rights, a contract note will be sent on or about May 20, 2022. Subscribed and allotted shares must be paid in cash in accordance with the instructions in the contract note. The allotment of shares subscribed for without preferential rights has been carried out in accordance with the allotment principles set out in the prospectus published by Egetis on April 28, 2022. Nominee-registered shareholders will receive notice of allotment and payment in accordance with the procedures of the nominee.

Following the Rights Issue, Egetis’ share capital will increase by approximately SEK 2,606,347, from approximately SEK 8,687,822 to approximately SEK 11,294,169 and the number of shares in Egetis will increase by 49,520,568, from 165,068,560 shares to 214,589,128 shares.

The shares subscribed for with subscription rights are expected to be registered with the Swedish Companies Registration Office (the “SCRO”) on or about May 31, 2022 and are expected to begin trading on Nasdaq Stockholm on June 3, 2022.

The shares subscribed for without subscription rights are expected to be registered with the SCRO on or about May 31, 2022 and are expected to begin trading on Nasdaq Stockholm on June 3, 2022.

Advisors

Pareto Securities AB acts as Sole Manager and Bookrunner in connection with the Rights Issue. Advokatfirman Lindahl KB acts as legal advisor to Egetis, and White & Case Advokat AB acts as legal advisor to the Pareto Securities AB in connection with the Rights Issue.

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About Egetis Therapeutics AB

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 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. The recruitment target of 16 patients was reached in the beginning of April 2022. 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 an application for ODD was submitted in the EU in the first quarter of 2021. There is an ongoing dialogue with EMA on the appropriate scope of the indication for an ODD in the EU.

Egetis Therapeutics (STO: EGTX) is listed on the Nasdaq Stockholm main market. For more information, see www.egetis.com

Important information

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This press release does not identify or suggest, or purport to identify or suggest, the risks (direct or indirect) that may be associated with an investment in the new shares. Any investment decision in connection with the rights issue must be made on the basis of all publicly available information relating to the Company and the Company’s shares including the information to be contained in the prospectus. Such information has not been independently verified by Pareto Securities AB (“**Manager**”). The information contained in this press release is for background purposes only and does not purport to be full or complete. No reliance may be placed for any purpose on the information contained in this press release or its accuracy or completeness. The Manager is acting for the Company in connection with the transaction and no one else and will not be responsible to anyone other than the Company for providing the protections afforded to its clients nor for giving advice in relation to the transaction or any other matter referred to herein.

This press release does not constitute a recommendation concerning any investor’s option with respect to the rights issue. Each investor or prospective investor should conduct his, her or its own investigation, analysis and evaluation of the business and data described in this press release and publicly available information. The price and value of securities can go down as well as up. Past performance is not a guide to future performance.

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This announcement is not a prospectus for the purposes of Regulation (EU) 2017/1129 (the “**Prospectus Regulation**”) and has not been approved by any regulatory authority in any jurisdiction. The Company has not authorized any offer to the public of shares or rights in any member state of the EEA other than in Sweden.

In the United Kingdom, this press release and any other materials in relation to the securities described herein is only being distributed to, and is only directed at, and any investment or investment activity to which this document relates is available only to, and will be engaged in only with, “**qualified investors**” (within the meaning of the United Kingdom version of the EU Prospectus Regulation (2017/1129/ EU) which is part of United Kingdom law by virtue of the European Union (Withdrawal) Act 2018) who are (i) persons having professional experience in matters relating to investments who fall within the definition of “investment professionals” in Article 19(5) of the Financial Services and Markets Act 2000 (Financial Promotion) Order 2005 (the “**Order**”); or (ii) high net worth entities falling within Article 49(2)(a) to (d) of the Order (all such persons together being referred to as “**relevant persons**”). In the United Kingdom, any investment or investment activity to which this communication relates is available only to, and will be engaged in only with, relevant persons. Persons who are not relevant persons should not take any action on the basis of this press release and should not act or rely on it.

Forward-looking statements

This press release contains forward-looking statements that reflect the Company's intentions, beliefs, or current expectations about and targets for the Company's future results of operations, financial condition, liquidity, performance, prospects, anticipated growth, strategies and opportunities and the markets in which the Company operates. Forward-looking statements are statements that are not historical facts and may be identified by words such as "believe", "expect", "anticipate", "intend", "may", "plan", "estimate", "will", "should", "could", "aim" or "might", or, in each case, their negative, or similar expressions. The forward-looking statements in this press release are based upon various assumptions, many of which are based, in turn, upon further assumptions. Although the Company believes that the expectations reflected in these forward-looking statements are reasonable, it can give no assurances that they will materialize or prove to be correct. Because these statements are based on assumptions or estimates and are subject to risks and uncertainties, the actual results or outcome could differ materially from those set out in the forward-looking statements as a result of many factors. Such risks, uncertainties, contingencies and other important factors could cause actual events to differ materially from the expectations expressed or implied in this release by such forward-looking statements. The Company does not guarantee that the assumptions underlying the forward-looking statements in this press release are free from errors and readers of this press release should not place undue reliance on the forward-looking statements in this press release. The information, opinions and forward-looking statements that are expressly or implicitly contained herein speak only as of its date and are subject to change without notice. Neither the Company nor anyone else undertake to review, update, confirm or to release publicly any revisions to any forward-looking statements to reflect events that occur or circumstances that arise in relation to the content of this press release, unless it is not required by law or Nasdaq Stockholm's rule book for issuers.

This information is information that Egetis Therapeutics is obliged to make public pursuant to the EU Market Abuse Regulation. The information was submitted for publication, through the agency of the contact persons set out above, at 2022-05-19 08:00 CEST.

Attachments

[Egetis Therapeutics raises approximately SEK 180 million through an oversubscribed rights issue](#)