PRESS RELEASE

Acticor Biotech provides an update on the current receivership proceedings

Paris, France, November 06, 2024 – 8:30am CET - ACTICOR BIOTECH (FR0014005OJ5 - ALACT), a clinical stage biopharmaceutical company developing glenzocimab, an innovative drug for the treatment of cardiovascular emergencies, today provides an update on the current receivership proceedings.

To enable the Company to continue its search for new investors or partners to support its new development plan in STEMI (ST-Elevation Myocardial Infarction), the court-appointed administrator had granted mid-October 2024 an extension until Tuesday November 5, 2024, to the call for bids published on September 13, for a continuation or disposal plan.

To date, the court-appointed administrator has received no proposal from a buyer. **The Company is therefore pursuing its search for a solution to support the development of glenzocimab in STEMI, and the 6-month observation period opened on August 6, 2024, is ongoing.**

As a reminder, on 6 August 2024, the Paris Commercial Court had ordered the opening of receivership proceedings. The purpose of these proceedings is to enable the Company to assess all the options available to it to pursue its development, the development of its product, glenzocimab, and its search for financing and partners. As already mentioned, this procedure will also enable the Company to finance its operations until January 2025.

The Company will continue to keep the market informed of developments in the current receivership proceedings and calls on investors to follow closely future communications concerning its progress.

About ACTICOR BIOTECH

ACTICOR BIOTECH, a clinical-stage biopharmaceutical company founded in 2013 from the work of INSERM, is developing glenzocimab, a humanized monoclonal antibody fragment (fab) targeting the GPVI platelet receptor for the treatment of cardiovascular emergencies and acute thrombotic diseases.

The main clinical indication being evaluated is acute ischemic stroke, due to the strong need for safer treatments, particularly those that do not increase the risk of bleeding, and its high incidence. In three international clinical trials involving over 600 stroke patients, no significant impact on neurological improvement (mRS score at 3 months) was demonstrated, with the exception of a sub-population of patients with intracerebral haemorrhage, where mortality was significantly reduced by a factor of 3 (p=0.035) (Mazighi et al. 2024).

LIBERATE, a Phase 2 clinical trial in the acute phase of myocardial infarction (STEMI), is currently being recruited through an academic partnership with the University of Birmingham (UK). This study aims to demonstrate the efficacy of glenzocimab in reducing the size of myocardial infarction, a critical factor for long-term cardiac function.

In all, more than 800 subjects were included in the clinical trials, over 400 of whom were exposed to glenzocimab without safety concerns.

The use of glenzocimab in thrombotic diseases is covered by 3 patent families, with an expiry date in 2036 for the first family. ACTICOR BIOTECH also has the right to develop a biomarker for stroke patients.

Acticor Biotech is backed by a panel of European and international investors (Mediolanum farmaceutici, Karista, Go Capital, Newton Biocapital, CMS Medical Venture Investment (HK) Limited, A&B (HK) Limited, Anaxago, and the Armesa Foundation) and has been listed on Euronext Growth Paris since November 2021 (ISIN: FR0014005OJ5 - ALACT).

For further information, visit: www.acticor-biotech.com

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