

Acticor Biotech: extension of the receivership procedure

Paris, France, October 3, 2024 – 6:00 p.m. CEST - ACTICOR BIOTECH (FR0014005OJ5 - ALACT), a clinical stage biopharmaceutical company developing glenzocimab, an innovative drug for the treatment of cardiovascular emergencies, announces that the Paris Commercial Court has today pronounced the extension of the company's receivership proceedings.

As a reminder, the Paris Commercial Court opened receivership proceedings on August 6, 2024. The purpose of these proceedings is to enable the Company to assess all options for pursuing the development of its product, glenzocimab, and its search for financing and partners. This procedure also enables the Company, as announced on August 6, 2024, to finance its operations until January 2025.

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

Contacts

ACTICOR BIOTECH

Gilles AVENARD, MD

CEO and Founder

gilles.avenard@acticor-biotech.com

Sophie BINAY, PhD

General Manager and CSO

Sophie.binay@acticor-biotech.com

NewCap

Mathilde BOHIN

Investor Relations

acticor@newcap.eu

T. : +33 (0)1 44 71 94 95

NewCap

Arthur ROUILLÉ

Media Relations

acticor@newcap.eu

T. : +33 (0)1 44 71 00 15

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