Press release Xintela AB (publ) 556780-3480 2020-06-23



Xintela granted preliminary approval from the European Patent Office for the treatment of brain tumors

Lund, Sweden, June 23, 2020 - Xintela today announces that the European Patent Office (EPO) has issued a preliminary approval ("Intention to grant") for the company's patent application related to antibody treatment of Glioblastoma and other tumors of the brain using the company's target molecule integrin α 10 β 1.

Xintela's patent application (publication number EP3258964) covers the use of integrin α 10 β 1 antibodies for treatment and diagnosis of tumors of the central nervous system (CNS). The present intention to grant is followed by an administrative process leading to the grant of the patent. The patent, which broadens and extends Xintela's fundamental protection in this field, will be valid until 2036.

"We are very pleased with this intention to grant communication which constitutes a validation of the strength and breadth of the company's oncology project, and adds significant value to our oncology business," says Xintela's CEO Evy Lundgren-Åkerlund.

Xintela has previously reported that the company's targeting ADC (Antibody-Drug Conjugate) antibodies have demonstrated killing effects on glioblastoma cells both in cell studies and in an animal model. The results have been published in the journal Cancers (2019, Vol. 11, pg 587). Additionally, in December 2019, Xintela announced that the company's function-blocking antibodies have demonstrated an inhibitory effect on glioblastoma tumors in a preclinical study. The current patent application also protects the target molecule integrin α 10 β 1 and thereto targeting antibodies for treatment of other cancers of the CNS.

This information is such information that Xintela AB (publ) is obligated to publish in compliance with the EU market abuse regulation. The information was provided, through the below contact, for publication on the 23rd of June 2020.

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About Xintela

Xintela is an Advanced Therapy company developing regenerative cell therapies and targeted cancer therapies based on the patented marker technology platform XINMARK®. The platform is built on specific cell surface proteins (integrins) and more than 25 years of research and development. Xintela uses the marker technology to isolate and quality assure stem cells for the treatment of musculoskeletal diseases including osteoarthritis (OA). Studies on horses with OA have demonstrated that the stem cells are safe and that they have a positive effect on cartilage and bone. Xintela has established an in-house GMP-facility for manufacturing of stem cells and is preparing a First in Human clinical study on patients with knee OA. In the oncology program, Xintela develops antibody-based therapies for treatment of aggressive tumors including glioblastoma. Xintela is listed on Nasdaq First North Growth Market Stockholm since 22 March 2016. Xintela's Certified Adviser at Nasdaq First North Growth Market is Erik Penser Bank AB, +46 8-463 80 00, certifiedadviser@penser.se.