

Cantargia publishes preclinical data on the potential of CAN10 in systemic sclerosis in a leading scientific journal

Cantargia (Cantargia AB; Nasdaq Stockholm: CANTA) today reported publication of preclinical results obtained with CAN10 in systemic sclerosis in one of the leading rheumatology journals, 'Annals of the Rheumatic Diseases'. CAN10 reduces both lung and skin fibrosis in multiple preclinical models of systemic sclerosis. The results are strengthened by the target of CAN10, IL1RAP and its signaling systems (IL-1, IL-33, and IL-36) being upregulated and disease-promoting in systemic sclerosis patient skin. CAN10 is an antibody in phase I clinical development.

"We are really pleased that the important results supporting CAN10 development in systemic sclerosis is recognized in a leading scientific journal. With phase I clinical studies ongoing, we are already now preparing for phase II clinical development in this life-threatening disease" said Göran Forsberg, CEO of Cantargia.

Systemic sclerosis is a life-threatening autoimmune disease resulting in fibrosis in the skin, lung, and other internal organs. Patients often have a severely impacted quality of life and there are no effective treatments today. Systemic sclerosis is one of the lead indications in the CAN10 development program.

The published data demonstrate that the target for CAN10, IL1RAP, and the IL1RAP-dependent signaling molecules IL-1, IL-33 and IL-36, are upregulated in skin from systemic sclerosis patients and that IL-1, IL-33 and IL-36 have profibrotic effects on skin fibroblasts from systemic sclerosis patients, which can be reduced by CAN10. Moreover, therapeutic treatment with a surrogate of CAN10 (mCAN10) in three different preclinical models of systemic sclerosis potently reduced both skin and lung fibrosis. Gene expression analysis indicated a broad mode of action of mCAN10, which normalized the expression of a majority of the genes commonly dysregulated in systemic sclerosis. In summary, the published data show that CAN10 targets central processes important for systemic sclerosis and that CAN10 provides a novel and promising opportunity to treat this disease. The publication in this high-impact journal (impact factor 27.4) reflects the scientific significance of the data.

This work was performed in collaboration with a world-leading research group headed by Prof. Dr. Jörg Distler at the Heinrich-Heine University, Düsseldorf, Germany. Key data from these studies were recently presented as a poster at the Systemic Sclerosis World Congress March 14-16, 2024.

"Systemic sclerosis patients have a very high need for new treatments. The published data provide strong evidence that signaling via IL1RAP regulates disease development in systemic sclerosis. The use of both patient samples and three different preclinical models strengthens the

translational aspect of the data and indicates an effect of CAN10 on key signaling pathways in this disease. I am very much looking forward to following the clinical development of CAN10, which has the potential of becoming a novel, targeted treatment for systemic sclerosis patients.” said Prof. Dr. Jörg Distler.

The article, titled “Combined inhibition of IL-1, IL-33 and IL-36 signaling by targeting IL1RAP ameliorates skin and lung fibrosis in preclinical models of Systemic Sclerosis”, is authored by Grönberg *et al.* and is available via this [link](#).

For further information, please contact

Göran Forsberg, CEO

Telephone: +46 (0)46-275 62 60

E-mail: goran.forsberg@cantargia.com

About Cantargia

Cantargia AB (publ), reg. no. 556791-6019, is a biotechnology company that develops antibody-based treatments for life-threatening diseases and has established a platform based on the protein IL1RAP, involved in a number of cancer forms and inflammatory diseases. The main program, the antibody nadunolimab (CAN04), is being studied clinically primarily in combination with chemotherapy with a focus on pancreatic cancer, non-small cell lung cancer and triple-negative breast cancer. Positive interim data for the combinations indicate stronger efficacy than would be expected from chemotherapy alone. Cantargia’s second development program, the antibody CAN10, blocks signaling via IL1RAP in a different manner than nadunolimab and addresses treatment of serious autoimmune/inflammatory diseases, with initial focus on systemic sclerosis and myocarditis.

Cantargia is listed on Nasdaq Stockholm (ticker: CANTA). More information about Cantargia is available at www.cantargia.com.

About CAN10

The CAN10 antibody binds strongly to its target IL1RAP and has a unique capability to simultaneously inhibit signaling via IL-1, IL-33 and IL-36. Inhibition of these signals can be of significant value in the treatment of several inflammatory or autoimmune diseases. The initial focus of CAN10 will be on two severe diseases: myocarditis and systemic sclerosis. In preclinical *in vivo* models of myocarditis, a CAN10 surrogate antibody significantly reduced the development of inflammation and fibrosis, and significantly counteracted the deterioration of the cardiac function. The CAN10 surrogate also inhibited disease development in models of systemic sclerosis, psoriasis, psoriatic arthritis, atherosclerosis and peritonitis. A clinical phase I study, investigating CAN10 in healthy volunteers and psoriasis patients, is ongoing. Up to 80 subjects may be included in the trial, the first clinical data set shows good safety. Additional data from the trial are expected continuously during 2024.



PRESS RELEASE
10 April 2024 08:45:00 CEST

Attachments

Cantargia publishes preclinical data on the potential of CAN10 in systemic sclerosis in a leading scientific journal