



Press release

Cantargia AB
556791-6019
10 August 2021

Invitation to presentation of Cantargia's report for the second quarter of 2021 on August 19 at 3.00 p.m. CET

Cantargia AB will publish the company's report for the period April – June 2021 on Thursday, August 19, 2021, at 08:30 a.m. CET.

In conjunction to the report, Cantargia invites investors, analysts, and media to an audiocast with teleconference (in English) on August 19, at 3:00 p.m. CET, where Göran Forsberg, CEO, and Bengt Jöndell, CFO, will present Cantargia and comment on the quarterly report for the period April – June 2021, followed by a Q&A-session.

The conference call can be followed at: <https://tv.streamfabriken.com/cantargia-q2-2021>.

To attend through telephone, please dial-in at one of the numbers below:

SE: +46 850558369
UK: +44 3333009274
US: +1 6319131422 PIN: 40478324#

The webcast will also be available on demand on Cantargia's corporate website: www.cantargia.com

For further information, please contact

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About Cantargia

Cantargia AB (publ), reg. no. 556791-6019, is a biotechnology company that develops antibody-based treatments for life-threatening diseases. The basis for this is the protein IL1RAP that is involved in a number of diseases and where Cantargia has established a platform. The main project, the antibody CAN04, is being studied clinically as combination therapy with chemotherapy or immune therapy with a primary focus on non-small cell lung cancer and pancreatic cancer. Positive interim data from the combination with chemotherapy indicate stronger efficacy than would be expected from chemotherapy alone. Cantargia's second project, the antibody CAN10, 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 nadunolimab (CAN04)

The antibody CAN04 binds strongly to the target IL1RAP and functions both through ADCC as well as blocking IL-1 α and IL-1 β signaling. Thereby, CAN04 can counteract the contribution of the IL-1 system to the immune suppressive tumor microenvironment and development of resistance to chemotherapy. CAN04 is investigated in three ongoing clinical trials. In the first phase I/IIa-study, CANFOUR, first line combination therapy is investigated using two different standard chemotherapies in patients with NSCLC (gemcitabine/cisplatin) and patients with PDAC (gemcitabine/nab-paclitaxel), as well as monotherapy in late stage patients (<https://clinicaltrials.gov/ct2/show/NCT03267316>). Phase I monotherapy data from 22 patients were presented at ASCO 2019 and showed good safety with infusion-related reaction being the most common side effect. In addition, the biomarkers IL-6 and CRP decreased during treatment. Positive interim data from the combination therapies show durable responses or pseudoprogression in patients with PDAC, resulting in iPFS of 7.8 months, and also a higher response rate of patients with NSCLC, compared to chemotherapy alone. A phase I study, CAPAFOUR, was initiated in H1 2021 and will investigate CAN04 in combination with the chemotherapy regimen FOLFIRINOX for first line treatment of metastatic PDAC. A phase I study, CIRIFOUR, is also currently investigating CAN04 in combination with an immune checkpoint inhibitor and was started H2 2020 (<https://clinicaltrials.gov/ct2/show/NCT04452214>). Additional clinical combination studies are planned to start during 2021.