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Cantargia announces final outcome of the company's significantly oversubscribed rights issue

12 August 2022 – The final outcome in Cantargia AB's (publ) ("Cantargia" or the "Company") (Nasdaq Stockholm: CANTA) rights issue of approximately SEK 250 million (the "Rights Issue"), is that 63,929,030 shares, corresponding to approximately 95.7 percent of the offered shares, have been subscribed for with the support of subscription rights after the end of the subscription period on 10 August 2022. The remaining 2,866,128 shares were subscribed for without the support of subscription rights, corresponding to approximately 4.3 percent of the offered shares. The Rights Issue was significantly oversubscribed by 44 percent. Thus, the guarantee commitments will not be utilized. Cantargia will receive approximately SEK 250 million before the deduction of transaction costs through the Rights Issue.

This financing further enhances Cantargia's financial position and enables us to continue developing the pipeline and to deliver additional milestone events. We are excited by the progress and recently published ASCO data on our lead program nadulinomab and look forward to providing further updates on nadunolimab and CAN10 as the projects progress. We are grateful for the demonstrable support of our existing shareholders and the significant interest shown from other investors, says Göran Forsberg, CEO of Cantargia.

The Rights Issue comprised a maximum of 66,795,158 shares, of which 63,929,030 shares, corresponding to approximately 95.7 percent of the offered shares, have been subscribed for with the support of subscription rights. The remaining 2,866,128 shares were subscribed for without the support of subscription rights, corresponding to approximately 4.3 percent of the offered shares. Together, subscriptions with the support of subscription rights and subscription applications without the support of subscription rights correspond to 144 percent of the offered shares in the Rights Issue. Thus, the guarantee commitments will not be utilized. Cantargia will receive approximately SEK 250 million before the deduction of transaction costs through the Rights Issue.

As a result of the fully subscribed Rights Issue, Cantargia's share capital will increase by SEK 5,343,612.64 to SEK 13,359,031.60 and the number of shares and votes will increase by 66,795,158 to 166,987,895.

Those who have subscribed for shares without the support of subscription rights will be allotted shares in accordance with the principles set out in the prospectus published on 22 July 2022. Notice of allotment to the persons who have subscribed for shares without the support of subscription rights is expected to be distributed today, 12 August 2022. Subscribed and allotted shares must be paid for in cash in accordance with the instructions on the settlement note sent to the subscriber. Investors who have subscribed through a nominee will be notified of the allotment in accordance with their respective nominee's procedures. Only those who have been allotted shares will be notified.

The last day for trading in paid subscribed shares (BTA) was on 11 August 2022. The new shares subscribed for with the support of subscription rights are expected to be registered with the Swedish Companies Registration Office around 16 August 2022 and expected to start trading on Nasdaq Stockholm as of 17 August 2022. The new shares subscribed for without the support of subscription rights are expected to be registered with the Swedish Companies Registration Office around 18 August 2022 and expected to start trading on Nasdaq Stockholm as of 26 August 2022.

Advisers

The Company has engaged Barclays Bank Ireland PLC and Carnegie Investment Bank AB (publ) as Joint Global Coordinators and Bookrunners in conjunction with the Rights Issue. Advokatfirman Vinge acted as legal counsel to the Company and Baker & McKenzie Advokatbyrå KB acted as legal counsel to the Joint Global Coordinators.

For further information, please contact:

Göran Forsberg, CEO
Mobile: +46 (0)46-275 62 60
E-mail: goran.forsberg@cantargia.com

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The information was submitted for publication, through the agency of the contact person set out above, at 08:00 CEST on 12 August 2022.

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 project, the antibody nadunolimab, is being studied clinically in combination 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 its target IL1RAP and functions by inducing ADCC and 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 multiple ongoing clinical trials. In the phase I/IIa study CANFOUR, first line combination therapy is investigated with standard chemotherapies in patients with PDAC (gemcitabine/nab-paclitaxel) and patients with NSCLC (cisplatin/gemcitabine) (NCT03267316). Positive interim data for the combination therapies show durable responses in 73 patients with PDAC, resulting in median iPFS of 7.2 months and median survival of 12.7 months. Stronger efficacy was also observed in 30 NSCLC patients with median PFS of 6.8 months. A response rate of 53 percent was achieved, with even higher responses in non-squamous NSCLC patients previously treated with pembrolizumab. These results show stronger efficacy than expected from chemotherapy alone. CAN04 is investigated with chemotherapy also in the phase I study CAPAFOUR, with the FOLFIRINOX regimen for first line treatment of metastatic PDAC (NCT04990037), and in two further clinical studies, CESTAFOUR (NCT05116891) and TRIFOUR (NCT05181462), in additional forms of cancer, including biliary tract cancer, colorectal cancer and triple negative breast cancer. CAN04 is also evaluated with the immune checkpoint inhibitor pembrolizumab, with or without chemotherapy, in the phase I study CIRIFOUR (NCT04452214).

Important information

The information in this press release does not contain or constitute an offer to acquire, subscribe or otherwise trade with shares or other securities in Cantargia. No action has been taken and measures will not be taken to permit a public offering in any other jurisdictions besides Sweden.

This press release is not a prospectus according to the definition in Regulation (EU) 2017/2019 (the "**Prospectus Regulation**") and has not been approved by any regulatory authority in any jurisdiction. This press release neither identifies nor pretends to identify risks (direct or indirect) that can be connected to an investment in shares or other securities in Cantargia. Any invitation to the persons concerned to subscribe for shares in Cantargia has only been made through the prospectus (the "**Prospectus**") published by the Company on 22 July 2022. The Prospectus has been approved and registered by the Swedish Financial Supervisory Authority and has been published on the Company's website, www.cantargia.com/en/investors/rights-issue. The approval should not be considered as an endorsement of the Company or as an endorsement of the quality of the securities that are the subject of the Prospectus and does not indicate that the Swedish Financial Supervisory Authority guarantees that the facts in the Prospectus are correct or complete. Investors should make their own assessment as to the suitability of investing in the Company's securities. In order for investors to fully understand the potential risks and benefits associated with a decision to participate in the Rights Issue, any investment decision should only be made based on the information in the Prospectus. Thus, investors are encouraged to review the

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In connection with the Rights Issue, each of the Joint Global Coordinators and Bookrunners and any of their affiliates, acting as investors for their own account, may take up a portion of the shares in the Rights Issue as a principal position and in that capacity may retain, purchase, sell, offer to sell for their own accounts such shares and other securities of the Company or related investments in connection with the Rights Issue or otherwise. Accordingly, references to Rights Issue shares being offered, acquired, placed or otherwise dealt in should be read as including any issue or offer to, or acquisition, placing or dealing by, each of the Joint Global Coordinators and Bookrunners and any of their affiliates acting in such capacity. In addition, each of the Joint Global Coordinators and Bookrunners and any of their affiliates may enter into financing arrangements (including swaps, warrants or contracts for differences) with investors in connection with which each of the Joint Global Coordinators and Bookrunners and any of their respective affiliates may from time to time acquire, hold or dispose of shares. None of the Joint Global Coordinators and Bookrunners intend to disclose the extent of any such investment or transactions otherwise than in accordance with any legal or regulatory obligations to do so.

Each of the Joint Global Coordinators and Bookrunners and their respective affiliates may have engaged in transactions with, and provided various commercial banking, investment banking, financial advisory transactions and services in the ordinary course of their business with the Company and/or its affiliates for which they would have received customary fees and commissions. Each of the Joint Global Coordinators and Bookrunners and their respective affiliates may provide such services to the Company and/or its affiliates in the future.

Information to distributors

Solely for the purposes of the product governance requirements contained within: (a) EU Directive 2014/65/EU on markets in financial instruments, as amended ("**MiFID II**"); (b) Articles 9 and 10 of Commission Delegated Directive (EU) 2017/593 supplementing MiFID II; and (c) local implementing measures (together, the "**MiFID II Product Governance Requirements**"), and disclaiming all and any liability, whether arising in tort, contract or otherwise, which any "manufacturer" (for the purposes of the MiFID II Product Governance Requirements) may otherwise have with respect thereto, the shares of the Company have been subject to a product approval process, which has determined that such shares are: (i) compatible with an end target market of retail investors and investors who meet the criteria of professional clients and eligible counterparties, each as defined in MiFID II; and (ii) eligible for distribution through all distribution channels as are permitted by MiFID II (the "**EU Target Market Assessment**"). Solely for the purposes of each manufacturer's product approval process in the United Kingdom, the target market assessment in respect of the shares in the Company has led to the conclusion that: (i) the target market for such shares is only eligible counterparties, as defined in the FCA Handbook Conduct of Business Sourcebook, and professional clients, as defined in Regulation (EU) No 600/2014 as it forms part of domestic law by virtue of the European Union (Withdrawal) Act 2018 ("**UK MiFIR**"); and (ii) all channels for distribution of such shares to eligible counterparties and professional clients are appropriate (the "**UK Target Market Assessment**") and, together with the EU Target Market Assessment, the "**Target Market Assessment**"). Notwithstanding the Target Market Assessment, Distributors should note that: the price of the shares of the Company may decline and investors could lose all or part of their investment; the shares of the Company offer no guaranteed income and no capital protection; and an investment in the shares in the Company is compatible only with investors who do not need a guaranteed income or capital protection, who (either alone or in conjunction with an appropriate financial or other advisers) are capable of evaluating the merits and risks of such an investment and who have sufficient resources to be able to bear any losses that may result therefrom. The Target Market Assessment is without prejudice to the requirements of any contractual, legal or regulatory selling restrictions in relation to the Rights Issue. Furthermore, it is noted that, notwithstanding the Target Market Assessment, the Joint Global Coordinators will only procure investors who meet the criteria of professional clients and eligible counterparties.

For the avoidance of doubt, the Target Market Assessment does not constitute: (a) an assessment of suitability or appropriateness for the purposes of MiFID II or UK MiFIR; or (b) a recommendation to any investor or group of investors to invest in, or purchase, or take any other action whatsoever with respect to the shares of the Company.

Each distributor is responsible for undertaking its own target market assessment in respect of the shares in Company and determining appropriate distribution channels.

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