

## BioInvent to expand pipeline with new development programs in solid cancer

- Aim is to add three clinical programs in solid cancer starting H1 2019
- BioInvent Capital Markets Day in Stockholm on December 10 to provide information on existing programs and rationale for new programs
- Company to engage in intensive investor meeting program in the U.S., Europe and Israel in October-November

**Lund, Sweden – October 16, 2018** – BioInvent International AB (OMXS: BINV) today announced its intention to initiate three new clinical programs in solid cancer. In addition to the previously disclosed programs (BI-1206 + rituximab – projected topline results H1 2020; and the Transgene and Pfizer collaborations), and subject to successful preclinical results and sufficient financial resources, BioInvent intends to advance the following programs in solid cancer: anti FcγRIIB antibody/anti-PD1 antibody – projected start phase I/Ila in H1 2019; BI-1607, an anti FcγRIIB antibody/check point inhibitor – projected start phase I proof of concept trial in H2 2019; BI-1808 (anti-"EmergingTNFRS" antibody) +/- anti-PD1 antibody – projected start phase I in H1 2020.

On December 10, BioInvent has the pleasure to invite investors, financial analysts and media to attend the Company's Capital Markets Day at the IVA Conference Center, Grev Turegatan 16 in Stockholm between 2 pm and 4 pm. BioInvent has made a concerted effort over the past months to reframe its equity story to better bring out its value drivers – the strength of its discovery platform, the scientific and clinical rationale of targeting Fc gamma receptors in cancer immunotherapy and its leadership in that space, the substantial unmet medical need of the cancer indications it is pursuing, and the associated commercial opportunity.

Speakers at the Capital Markets Day include Martin Welschof, CEO, Björn Frendéus, Chief Scientific Officer and Andres McAllister, Chief Medical Officer. The presentations will give an overview of BioInvent's current and planned cancer immunotherapy programs, and the company's strategy to create shareholder value. The Company looks forward to engaging with all its stakeholders.

To attend the Capital Markets Day, please register by e-mail to stefan.ericsson@bioinvent.com or by phone +46 46 286 85 54 no later than December 6. Following the event, a video recording will be published on BioInvent's website.

During the months of October and November, BioInvent's management will undertake an intensive program of investor meetings across the U.S., Europe, and Israel. As part of this program, BioInvent will host a Key Opinion Leader (KOL) event in New York on November 6. Details of this event, which will be webcast live and archived on BioInvent's website, will be announced shortly. At the KOL event, BioInvent's management and an invited scientific key opinion leader will engage with the U.S. investor community to present the Company's scientific and clinical value drivers.

## **About BioInvent**

BioInvent International AB (OMXS: BINV) is focused on the discovery and development of novel and first-in-class immuno-modulatory antibodies to treat cancer. The Company's lead program BI-1206 is currently in Phase I/II for non-Hodgkin lymphoma and chronic lymphatic leukemia. BioInvent's preclinical portfolio is focused on targeting key immune suppressive cells and pathways of the tumor microenvironment, including regulatory T cells, tumor-associated myeloid cells and mechanisms of antibody drug-resistance. The Company has a strategic research collaboration with Pfizer Inc., and partnerships with Transgene, Bayer Pharma, Daiichi Sankyo, and Mitsubishi Tanabe Pharma. BioInvent generates near term revenues from its fully integrated manufacturing unit producing antibodies for third parties for research through to late-stage clinical trials. More information is available at www.bioinvent.se.

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The press release contains statements about the future, consisting of subjective assumptions and forecasts for future scenarios. Predictions for the future only apply as the date they are made and are, by their very nature, in the same way as research and development work in the biotech segment, associated with risk and uncertainty. With this in mind, the actual outcome may deviate significantly from the scenarios described in this press release.