Antengene Unveils Its First Australian Office to Continue Expanding Global Presence

Antengene recently unveiled a new office space situated in Melbourne, Australia's most sought-after CBD, taking another important step in the company's continued global expansion. The new office offers an appealing environment that will help Antengene to attract more top talents, and further accelerate its expansion in the APAC region as well as its committed efforts to bring high quality and affordable innovative medicines to patients with life-threatening conditions including hematologic malignancies and solid tumors in Asia Pacific region and around the world.

Members of Antengene's Australia team at work in the newly opened office

Occupying a space of 250 square meters in a Grade A office building in Melbourne's CBD that has an impressive 4.5-star NABERS energy rating and a 5.5-star NABERS water rating, the new office provides the fast-growing Australian team a superb work environment boasting a bird view of the downtown CBD area. In addition, the office has adopted a modern design with a state-of-the-art video conferencing system to facilitate both effective communications within the local team and close collaborations with teams and partners across the globe.

A state-of-the-art video conferencing system

In March 2022, XPOVIO® (selinexor) was approved by the Australia Therapeutic Goods Administration (TGA) for the treatment of patients with relapsed/refractory multiple myeloma (R/R MM) or triple-refractory R/R MM. In September 2022, XPOVIO® was included into the Pharmaceutical Benefits Scheme (PBS) for the treatment of patients with penta-refractory R/R MM. While making steady progress with the commercialization of Antengene's lead asset, XPOVIO®, in Australia and the wider Asia Pacific region, the Australian team continued to grow at a rapid rate to a total of fifteen members in a variety of functions covering Commercial, Finance, Medical Affairs, Clinical and Business Development. The office will enable closer collaborations with teams in other APAC markets and provide the extra space needed to support the team's continued expansion.

At present, **Antengene is conducting 4 clinical trials in Australia with the company's four drug candidates, including ATG-018, ATG-101, ATG-037, and ATG-017,** exploring these novel agents in a range of solid tumors and hematological malignancies**.** It is worth noting that ATG-037 is the first oral available, small molecule CD73 inhibitor entering clinical

development in China and the wider Asia Pacific region. ATG-101, the first PD-L1/4-1BB bispecific antibody approved to enter clinical stage in Australia, is currently being evaluated in clinical setting in Australia, China, and the U.S.

Thomas Karalis, Antengene's Corporate Vice President, Head of Asia Pacific Region

Mr. Thomas Karalis, Antengene's Corporate Vice President, Head of Asia Pacific Region, commented on the new office, "Our first dedicated Antengene office in the Asia Pacific Region is an inspiring place to work and has been embraced by our growing Antengene team. Its opening marks a significant milestone for Antengene, as it not only meets our current business needs but will also support our future growth in the APAC region and globally."

上海市长宁区中山西路 1065 号 SOHO 中山广场 B 座 1206-1209 室
Suite 1206-1209, Building B, SOHO Plaza, 1065 West Zhongshan Road, Shanghai 200051, China
Tel: (86) 021 3250 1095
Fax: (86) 021 3250 1062
www.antengene.com

Michele Robbins, Antengene's Senior Director of Commercialisation for ANZ

Ms. Michele Robbins, Antengene's Senior Director of Commercialisation for ANZ, commented, "We have hired fantastic, capable people since starting the business in Australia a little over two years ago. Our new, open-plan Melbourne office enables us to continue the rapid growth of the business achieved to date, with a collaborative space to continue to drive teamwork and innovation."

Dr. Tamara Etto, Antengene's Head of Medical Affairs, APAC

Dr. Tamara Etto, Antengene's Head of Medical Affairs, APAC, commented on Antengene's plans for growth, "Antengene's focus for the team in Australia is to align with our teams in China and across APAC and China advancing the clinical development and commercialization of XPOVIO®, a first-in-class/best-in-class therapy for patients with R/R MM.

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We are excited to have a growing Australian team and look to the future of Antengene with eagerness to positively impact patient lives."

About Antengene

Antengene Corporation Limited (**"Antengene"** , SEHK: 6996.HK) is a leading commercial-stage R&D-driven global biopharmaceutical company focused on the discovery, development, manufacturing and commercialization of innovative first-in-class/best-in-class therapeutics for the treatment of hematologic malignancies and solid tumors, in realizing its vision of **"Treating Patients Beyond Borders"** .

Since 2017, Antengene has built a broad and expanding pipeline of 15 clinical and preclinical assets, of which 10 are global rights assets, and 5 came with rights for Asia Pacific markets including the Greater China region. To date, Antengene has obtained 24 investigational new drug (IND) approvals in the U.S. and Asia, and submitted 6 new drug applications (NDAs) in multiple Asia Pacific markets, with the NDA for XPOVIO® (selinexor) already approved in mainland China, South Korea, Singapore and Australia.

Forward-looking statements

The forward-looking statements made in this article relate only to the events or information as of the date on which the statements are made in this article. Except as required by law, we undertake no obligation to update or revise publicly any forward-looking statements, whether as a result of new information, future events or otherwise, after the date on which the statements are made or to reflect the occurrence of unanticipated events. You should read this article completely and with the understanding that our actual future results or performance may be materially different from what we expect. In this article, statements of, or references to, our intentions or those of any of our Directors or our Company are made as of the date of this article. Any of these intentions

may alter in light of future development. For a further discussion of these and other factors that could cause future results to differ materially from any forward-looking statement, see the section titled "Risk Factors" in our periodic reports filed with the Hong Kong Stock Exchange and the other risks and uncertainties described in the Company's Annual Report for year-end December 31, 2021, and subsequent filings with the Hong Kong Stock Exchange.

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