Antengene Corporation Limited
德琪醫藥有限公司

(Incorporated in the Cayman Islands with limited liability)

(Stock Code: 6996)

VOLUNTARY ANNOUNCEMENT

APPROVAL OF IND APPLICATION IN CHINA FOR A PHASE 3 CLINICAL TRIAL OF ATG-010 (SELINEXOR)

Antengene Corporation Limited (the "**Company**", together with its subsidiaries, the "**Group**") hereby informs the shareholders and potential investors of the Company of the attached press release in respect of the approval of Investigational New Drug ("**IND**") application in China for a Phase 3 clinical trial of ATG-010 (selinexor) in combination with bortezomib and dexamethasone (SVd) for the treatment of relapsed/refractory multiple myeloma ("**rrMM**") .

This is a voluntary announcement made by the Company. The Group cannot guarantee that ATG-010 (selinexor) will ultimately be successfully developed and marketed. Shareholders and potential investors of the Company are advised to exercise caution when dealing in the shares of the Company.

By order of the Board
Antengene Corporation Limited
Dr. Jay Mei
Chairman

Hong Kong, December 18, 2020

As at the date of this announcement, the board of directors of the Company comprises Dr. Jay Mei, Mr. John F. Chin and Mr. Yiteng Liu as executive directors; Mr. Xubo Hu, Mr. Zhen Li and Mr. Yanling Cao as non-executive directors; and Mr. Mark J. Alles, Ms. Jing Qian and Mr. Sheng Tang as independent non-executive directors.

Antengene Announces Approval of IND Application in China for a Phase 3 Clinical Trial of ATG-010 (Selinexor) in Combination with Bortezomib and Dexamethasone (SVd) for the Treatment of rrMM

Shanghai and Hong Kong, PRC, December 18, 2020 – Antengene Corporation Limited ("**Antengene**", SEHK: 6996.HK), a leading innovative biopharmaceutical company dedicated to discovering, developing and commercializing global first-in-class and/or best-in class therapeutics in hematology and oncology, announced that the National Medical Products Administration (NMPA) has approved the Investigational New Drug (IND) application for ATG-010 (selinexor), an oral Selective Inhibitor of Nuclear Export compound, in combination with bortezomib and dexamethasone for the treatment of patients with relapsed/refractory multiple myeloma (rrMM) in China.

The trial is a Phase 3 randomized, controlled, open-label, multicenter clinical trial, aiming to evaluate the efficacy and safety of ATG-010, bortezomib and dexamethasone (SVd) regimen against bortezomib and dexamethasone (Vd) regimen in Chinese adult patients with rrMM who have received one to three prior lines of therapy. A total of 150 patients will be randomized in a 2:1 ratio to receive SVd or Vd treatment.

ATG-010 is a first-in-class and only-in-class oral selective inhibitor of nuclear export (SINE) and the first and only drug approved by the Food and Drug Administration (FDA) for use in both relapsed/refractory multiple myeloma and diffuse large B-cell lymphoma. In December 2020, National Comprehensive Cancer Network (NCCN®) added three different ATG-010 combination regimens to its Clinical Practice Guidelines in Oncology (NCCN® Guidelines) for previously treated multiple myeloma, including SVd, SDd and SPd. In China, Antengene is conducting a Phase 2 registrational clinical trial of ATG-010 for rrMM (MARCH).

"The NMPA approval of BENCH trial demonstrates our ability to efficiently execute, and marks a great start of Antegene's first Phase 3 registrational trial to validate SVd regimen's efficacy and safety (as evidenced in the global BOSTON trial) in Chinese population." said Dr. Jay Mei, Founder, Chairman and CEO of Antengene. "Since becoming a public company, our clear focus has been on advancing the clinical development of ATG-010. We will initiate immediately our Phase 3 trial of ATG-010 for rrMM patients in China and believe the unique and novel MoA of ATG-010 will provide physicians new treatment options for more oncology indications."

About ATG-010 (selinexor, XPOVIO®)

ATG-010 (selinexor, XPOVIO®) is a first-in-class and only-in-class oral selective inhibitor of nuclear export compound, developed by Antengene and Karyopharm Therapeutics Inc. (NASDAQ: KPTI). In July 2019, the US Food and Drug Administration (FDA) approved ATG-010 in combination with low-dose dexamethasone for the treatment of relapsed/refractory multiple myeloma (rrMM) and in June 2020 approved ATG-010 as a single-agent for the treatment of relapsed/refractory diffuse large B-cell lymphoma (rrDLBCL). ATG-010 is so far the first and only oral SINE compound approved by the FDA. ATG-010 is also being evaluated in several other mid-and later-phase clinical trials across multiple solid tumor indications, including liposarcoma and endometrial cancer. In November 2020, at the Connective Tissue Oncology Society 2020

Annual Meeting (CTOS 2020), Antengene's partner, Karyopharm Therapeutics, presented positive results from the Phase 3 randomized, double blind, placebo controlled, cross-over SEAL study evaluating single agent, oral ATG-010 versus matching placebo in patients with liposarcoma. Karyopharm also recently announced that the ongoing Phase 3 SIENDO study of ATG-010 in patients with endometrial cancer passed planned interim futility analysis and that Data and Safety Monitoring Board (DSMB) recommended the study should proceed as planned without any modifications. Top-line SIENDO study results are expected in the second half of 2021.

Antengene is conducting two Phase 2 registrational clinical trials of ATG-010 in China for relapsed refractory multiple myeloma (MARCH) and for relapsed refractory diffuse large B-cell lymphoma (SEARCH), and has initiated clinical trials for high prevalence cancer types in the Asia Pacific region including peripheral T-cell lymphoma and NK/T-cell lymphoma (TOUCH) and KRAS-mutant non-small cell lung cancer (TRUMP).

About Antengene

Antengene is a leading clinical-stage Asia-Pacific biopharmaceutical company focused on innovative oncology medicines. Antengene aims to provide the most advanced anti-cancer drugs to patients in China, the Asia Pacific Region and around the world. Since its establishment, Antengene has built a pipeline of 12 clinical and pre-clinical stage assets, obtained 11 investigational new drug approvals and has 9 ongoing cross-regional clinical trials in Asia Pacific. The vision of Antengene is to "Treat Patients Beyond Borders". Antengene aims to address significant unmet medical needs by discovering, developing and commercializing first-in-class/ best-in-class therapeutics.

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.

* XPOVIO® is a registered trademark of Karyopharm Therapeutics Inc.;

NCCN® is a registered trademark of National Comprehensive Cancer Network;

SVd: selinexor, bortezomib and dexamethasone;

SDd: selinexor, daratumumab and dexamethasone;

SPd: selinexor, pomalidomide and dexamethasone.