



## **Antengene to Present Results from Two Late-Stage Studies of Selinexor Signaling Potential Clinical Breakthrough at ASH 2024**

Shanghai and Hong Kong, PRC, November 6, 2024 — Antengene Corporation Limited ( “**Antengene**” , SEHK: 6996.HK), a leading innovative, commercial-stage global biopharmaceutical company dedicated to discovering, developing and commercializing first-in-class and/or best-in-class medicines for cancer, today announced that **it will present the latest data from two clinical studies of selinexor in two Posters at the 2024 American Society of Hematology Annual Meeting (ASH 2024), taking place on December 7-10, 2024, in San Diego, CA, the United States.**

### **Details on the Posters:**

**Title:** Weekly Selinexor, Bortizomib and Dexamethasone (SVd) Versus Twice Weekly Bortizomib and Dexamethasone (Vd) in Chinese Patients with Relapsed and Refractory Multiple Myeloma (RRMM): Primary Analysis of Phase III Bench Study

**Publication Number:** 4748

**Session:** 654. Multiple Myeloma: Pharmacologic Therapies: Poster III

**Date:** Monday, December 9, 2024



**Time:** 6:00 PM - 8:00 PM (Eastern time)

7:00 AM - 9:00 AM, December 10, 2024 (Beijing time)

**First Author:** Dr. Jin Lu (Peking University People's Hospital)

**Corresponding Author:** Dr. Jian Hou (Ren Ji Hospital, Shanghai Jiao Tong University School of Medicine)

**Title:** Selinexor Combined with Tislelizumab in Patients with Relapsed or Refractory Extranodal NK/T-Cell Lymphoma (R/R ENKTL): Preliminary Results of Arm C, from a Multicenter, Single-Arm, Phase I/II Study, Touch

**Publication Number:** 4448

**Session:** 625. T Cell, NK Cell, or NK/T Cell Lymphomas: Clinical and Epidemiological: Poster III

**Date:** Monday, December 9, 2024

**Time:** 6:00 PM - 8:00 PM (Eastern time)

7:00 AM - 9:00 AM, December 10, 2024 (Beijing time)

**First Author:** Dr. Rong Tao (Fudan University Shanghai Cancer Center)

**Corresponding Author:** Dr. Huiqiang Huang (Sun Yat-Sen University Cancer Center)

### **About XPOVIO® (selinexor)**

XPOVIO® is the world's first approved orally-available, selective inhibitor of the nuclear export protein XPO1. **It offers a novel**



**mechanism of action, synergistic effects in combination regimens, fast onset of action, and durable responses.**

By blocking the nuclear export protein XPO1, XPOVIO® can promote the intranuclear accumulation and activation of tumor suppressor proteins and growth regulating proteins, and down-regulate the levels of multiple oncogenic proteins. XPOVIO® delivers its antitumor effects through three mechanistic pathways: 1) exerting antitumor effects by inducing the intranuclear accumulation of tumor suppressor proteins; 2) reducing the level of oncogenic proteins in the cytoplasm by inducing the intranuclear accumulation of oncogenic mRNAs; 3) restoring hormone sensitivity by activating the glucocorticoid receptors (GR) pathway. To utilize its unique mechanism of actions, XPOVIO® is being evaluated for use in multiple combination regimens in a range of indications. At present, Antengene is conducting multiple clinical studies of XPOVIO® in the mainland of China for the treatment of relapsed/refractory hematologic malignancies and solid tumors (3 of these studies are being jointly conducted by Antengene and Karyopharm Therapeutics Inc. [Nasdaq:KPTI]).



## 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 pipeline of 9 oncology assets at various stages going from clinical to commercial, including 6 with global rights, and 3 with rights for the APAC region. To date, Antengene has obtained 31 investigational new drug (IND) approvals in the U.S. and Asia, and submitted 10 new drug applications (NDAs) in multiple Asia Pacific markets, with the NDA for XPOVIO® (selinexor) already approved in Mainland of China, Taiwan China, Hong Kong China, Macau China, South Korea, Singapore, Malaysia, Thailand 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, please see the other risks and uncertainties described in the Company's Annual Report for the year ended December 31, 2023, and the documents subsequently submitted to the Hong Kong Stock Exchange.