

# Antengene Announces Participation in the H.C. Wainwright 26<sup>th</sup> Annual Global Investment Conference

Shanghai and Hong Kong, PRC, September 5, 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 its
participation in the H.C. Wainwright 26th Annual Global Investment
Conference. The Company will share the latest advancements in R&D
during a Fireside Chat.

#### **Fireside Chat:**

September 10<sup>th</sup>, 2024 (Tuesday), 4:30 pm - 5:00 pm (US Eastern Time)

September 11<sup>th</sup>, 2024 (Wednesday), 4:30 am - 5:00 am (Beijing Time)

**Location:** Holmes I - 4<sup>th</sup> Floor, Lotte New York Palace Hotel

#### **Webcasting Link for the Fireside Chat:**

https://journey.ct.events/view/48a63922-9bbf-487b-8152bf12ba2f890e

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



### 1x1 Meetings:

September 9<sup>th</sup> to 11<sup>th</sup>, 2024 (Monday to Wednesday)

**Location:** Lotte New York Palace Hotel

#### **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 29 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 and Australia.

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## **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.

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