



XPOVIO® Receives Reimbursement Approval in South Korea for a Second Multiple Myeloma Indication

- *XPOVIO® is the first XPO1 inhibitor approved for reimbursement by South Korea's National Health Insurance Service (NHIS) for the treatment of adult patients with multiple myeloma (MM).*
- *In South Korea, XPOVIO® has been approved for three indications across MM and diffuse large B-cell lymphoma (DLBCL), and two of these approved indications have been included in the national reimbursement scheme.*

Shanghai and Hong Kong, PRC, March 2, 2026 — Antengene Corporation Limited (“**Antengene**” , SEHK: 6996.HK) , a leading innovative, commercial-stage global biotech company dedicated to discovering, developing and commercializing first-in-class and/or best-in-class medicines for autoimmune disease, solid tumors and hematological malignancies indications, announced that **South Korea's National Health Insurance Service (NHIS) has approved the reimbursement of XPOVIO® (selinexor) in combination with bortezomib and dexamethasone for the treatment of adult patients with multiple myeloma (MM) after one prior therapy. The reimbursement has taken effect on March 1, 2026. This marks the second XPOVIO® indication to**



be approved for reimbursement in South Korea.

As Antengene continues to expand its footprint across the Asia Pacific region, the Company remains committed to improving patient access to its innovative therapies. **To date, XPOVIO® has been approved in South Korea for three indications in MM and diffuse large B-cell lymphoma (DLBCL), both major hematological malignancies. Two of these indications have been approved for reimbursement,** including XPOVIO® in combination with dexamethasone for the treatment of adult patients with relapsed or refractory MM (R/R MM) who have received at least four prior therapies, as well as the newly reimbursed indication. With expanded reimbursement coverage, XPOVIO® is expected to benefit a broader patient population and further contribute to the management of hematological malignancies in South Korea.

With a novel mechanism of action, XPOVIO® is the world's first approved orally-available, selective XPO1 inhibitor. XPOVIO® has already been approved in ten countries and regions in APAC, and has been included in the national insurance schemes in five of these markets (the mainland of China, Taiwan market, Australia, Singapore and South Korea). Moving forward, Antengene will continue to pursue broader access for XPOVIO® across APAC markets.



About Antengene

Antengene Corporation Limited (“**Antengene**” , SEHK: 6996.HK) is a global, R&D-driven, commercial-stage biotech company focused on developing first-in-class/best-in-class therapeutics for diseases with significant unmet medical needs. Its pipeline spans from preclinical to commercial stages and includes several in-house discovered programs, including ATG-022 (CLDN18.2 ADC), ATG-037 (oral CD73 inhibitor), ATG-101 (PD-L1 × 4-1BB bispecific antibody), and ATG-042 (oral PRMT5-MTA inhibitor).

Antengene has also developed AnTenGager™, a proprietary T cell engager 2.0 platform featuring “2+1” bivalent binding for low expressing targets, steric hindrance masking, and proprietary CD3 sequences with fast on/off kinetics to minimize cytokine release syndrome (CRS) and enhance efficacy. These characteristics support the platform’s broad applicability across autoimmune disease, solid tumors and hematological malignancies, with programs targeting CD19 x CD3 (ATG-201 for B cell-related autoimmune diseases), CDH6 x CD3 (ATG-106 for ovarian cancer and kidney cancer), ALPPL2 x CD3 (ATG-112 for gynecologic tumors and non-small cell lung cancer), LY6G6D x CD3 (ATG-110 for microsatellite-stable colorectal cancer), GPRC5D x CD3 (ATG-021



for multiple myeloma), LILRB4 x CD3 (ATG-102 for acute myeloid leukemia and chronic myelomonocytic leukemia) and FLT3 x CD3 (ATG-107 for acute myeloid leukemia).

To date, Antengene has obtained 32 investigational new drug (IND) approvals in the U.S. and Asia, and obtained new drug application (NDA) approvals in 10 Asia Pacific markets. Its lead commercial asset, XPOVIO® (selinexor), is approved in the Mainland of China, Taiwan China, Hong Kong China, Macau China, South Korea, Singapore, Malaysia, Thailand, Indonesia and Australia, and has been included in the national insurance schemes in five of these markets (Mainland of China, Taiwan China, Australia, South Korea and Singapore).

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, 2024, and the documents subsequently submitted to the Hong Kong Stock Exchange.