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ASCENTAGE PHARMA GROUP INTERNATIONAL

亞盛醫藥集團

(Incorporated in the Cayman Islands with limited liability)

(Stock Code: 6855)

VOLUNTARY ANNOUNCEMENT

ASCENTAGE PHARMA ANNOUNCES INCLUSION OF LISAFTOCLAX IN CHINA'S 2026 NATIONAL REIMBURSEMENT DRUG LIST AND SUCCESSFUL RENEWAL OF COVERAGE FOR OLVEREMBATINIB

Ascentage Pharma Group International (the "**Company**" or "**Ascentage Pharma**") is pleased to announce that its two core assets are covered by China's 2026 National Reimbursement Drug List (NRDL). Specifically:

- Lisaftoclax has been included in the 2026 NRDL for the first time through negotiation for the treatment of adult patients with chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL) who have previously received at least one systemic therapy including a Bruton's tyrosine kinase (BTK) inhibitor.

- Olverembatinib has successfully renewed its NRDL coverage through renegotiation for the treatment of (1) adult patients with chronic myeloid leukemia in chronic phase (CML-CP) who are resistant and/or intolerant to first – and second-generation tyrosine kinase inhibitors (TKIs); and (2) adult patients with CML-CP or accelerated phase (CML-AP) harboring the T315I mutation.

As of now, all approved indications for both of Ascentage Pharma's novel drugs are covered by the NRDL, providing more patients with greater access to innovative treatment options. The updated NRDL will officially take effect on January 1, 2027.

Lisaftoclax is a novel, oral, Bcl-2 selective inhibitor independently developed by Ascentage Pharma to treat patients with malignancies by selectively blocking the anti-apoptotic protein Bcl-2 and hence restoring the normal apoptosis process in cancer cells. Approved in China in July 2025, Lisaftoclax became the first domestically developed, original Bcl-2 inhibitor to receive marketing approval in the country. Its inclusion in the updated NRDL is expected to significantly improve treatment accessibility for patients with CLL/SLL in China, enabling more patients to gain timely access to new treatment options.

Olverembatinib is a novel, oral, third-generation BCR-ABL inhibitor independently developed by Ascentage Pharma, with support from China's National Major New Drug Innovation Program. It is jointly commercialized in China by Ascentage Pharma and Innovent Biologics, Inc. Approved in China in November 2021, Olverembatinib became the first third-generation BCR-ABL inhibitor to receive marketing approval in the country. The successful renewal of its NRDL coverage will continue to benefit more patients with CML in China by further improving the affordability and accessibility of long-term treatment.

By order of the Board
Ascentage Pharma Group International
Dr. Yang Dajun
Chairman and Executive Director

Suzhou, People's Republic of China, November 16, 2026

As at the date of this announcement, the Board comprises Dr. Yang Dajun as Chairman and executive Director, Dr. Wang Shaomeng and Dr. Lu Simon Dazhong[Note1] as non-executive Directors, and Mr. Ye Changqing, Mr. Ren Wei, Dr. David Sidransky[Note2], Ms. Marina S. Bozilenko, Dr. Debra Yu and Dr. Marc E. Lippman, MD as independent non-executive Directors.

Notes:

1. *Dr. Lu Simon Dazhong satisfy the independence requirements of the U.S. Securities and Exchange Commission and Nasdaq corporate governance requirements*

2. *Dr. David Sidransky is the Lead Independent Non-Executive Director of the Company.*