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Voluntary Announcement

Clover Completes Enrollment in Australian Phase 2 Clinical Trial for RSV + hMPV ± PIV3 Respiratory Combination Vaccine Candidates

This announcement is made by the board (the “**Board**”) of directors (the “**Directors**”) of Clover Biopharmaceuticals, Ltd. (the “**Company**” or “**Clover**”, together with its subsidiaries, the “**Group**”) on a voluntary basis to inform the shareholders of the Company and potential investors on the latest business development of the Group.

The Company is pleased to announce that enrollment has been completed in a Phase 2 clinical trial in Australia evaluating SCB-1022 (RSV + hMPV) and SCB-1033 (RSV + hMPV + PIV3) protein-based vaccine candidates based on prefusion-stabilized F (PreF)-Trimer subunit vaccine antigens utilizing Clover’s validated Trimer-Tag vaccine technology platform.

The ongoing Phase 2 trial for Clover’s combination vaccine candidates is a randomized, observer-blinded, multi-center study with 420 older adults (60-85 years) enrolled in Australia, and the participants have been randomized to receive either SCB-1022 (RSV + hMPV), SCB-1033 (RSV + hMPV + PIV3) or placebo. The study is assessing safety, reactogenicity and immunogenicity, and initial results are expected in Q3-2026.

Shareholders of the Company and potential investors are advised to exercise caution when dealing in the shares of the Company.

By order of the Board
Clover Biopharmaceuticals, Ltd.
Dr. Peng LIANG
Chairman of the Board

Shanghai, PRC, March 30, 2026

As of the date of this announcement, the Board comprises Dr. Peng LIANG and Mr. Joshua G LIANG as executive Directors; Dr. Xiaodong WANG and Dr. Donna Marie AMBROSINO as non-executive Directors; and Dr. Xiaobin WU, Mr. Xiang LIAO, Mr. Jeffrey FARROW and Mr. Thomas LEGGETT as independent non-executive Directors.