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**VOLUNTARY ANNOUNCEMENT
CLOVER INITIATES 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 of the first participants 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 enrolling up to 420 older adults (60-85 years) in Australia, and the participants will be randomized to receive either SCB-1022 (RSV + hMPV), SCB-1033 (RSV + hMPV + PIV3) or placebo. The study will assess safety, reactogenicity and immunogenicity.

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, January 12, 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.