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Abbisko Cayman Limited

和譽開曼有限責任公司

(Incorporated in the Cayman Islands with limited liability)

(Stock Code: 2256)

**VOLUNTARY ANNOUNCEMENT
ABBISKO THERAPEUTICS ANNOUNCES IND
CLEARANCE FROM THE NMPA FOR ORAL SMALL
MOLECULE PAN-KRAS INHIBITOR ABSK211**

Abbisko Cayman Limited (the “**Company**”, together with its subsidiaries, the “**Group**”) hereby informs the shareholders and potential investors of the Company of the attached press release that Abbisko Therapeutics Co., Ltd. (“**Abbisko Therapeutics**”), a subsidiary of the Company, announced that the China National Medical Products Administration (“**NMPA**”) has cleared the Investigational New Drug (“**IND**”) application for ABSK211, an oral and highly potent small-molecule pan-KRAS inhibitor with broad alteration coverage, for the treatment of patients with advanced solid tumors harboring KRAS alterations. Prior to this, ABSK211 received IND approval from the U.S. Food and Drug Administration (“**FDA**”) on July 8. Achieving IND clearance in both China and the U.S. marks a key milestone in the global clinical development plan for ABSK211.

This is a voluntary announcement made by the Company. The Group cannot guarantee that ABSK211 will ultimately be successfully marketed. Shareholders and potential investors of the Company are advised to exercise caution when dealing in the shares of the Company.

By order of the Board
Abbisko Cayman Limited
Dr. Xu Yao-Chang
Chairman

Shanghai, 19 August 2026

As at the date of this announcement, the board of directors of the Company comprises Dr. Xu Yao-Chang, Dr. Yu Hongping and Dr. Ji Jing as executive directors; and Dr. Sun Piaoyang, Mr. Sun Hongbin and Ms. Chui Hoi Yam as independent non-executive directors.

Abbisko Therapeutics Announces IND Clearance from the NMPA for Oral Small Molecule pan-KRAS Inhibitor ABSK211

On 19 August 2026, Abbisko Therapeutics Co., Ltd. (“**Abbisko Therapeutics**”) announced that the China National Medical Products Administration (“**NMPA**”) has cleared the Investigational New Drug (“**IND**”) application for ABSK211, an oral and highly potent small-molecule pan-KRAS inhibitor with broad alteration coverage, for the treatment of patients with advanced solid tumors harboring KRAS alterations. Prior to this, ABSK211 received IND approval from the U.S. Food and Drug Administration (“**FDA**”) on July 8. Achieving IND clearance in both China and the U.S. marks a key milestone in the global clinical development plan for ABSK211.

Abbisko Therapeutics will imminently begin an open-label Phase I clinical trial designed to assess the safety, tolerability, efficacy and pharmacokinetics of ABSK211 in patients with advanced solid tumors harboring KRAS alterations.

KRAS is one of the most common oncogenic driver genes in human cancers and is highly prevalent across multiple solid tumor types, including approximately 90% of pancreatic cancers, 35% of colorectal cancers, and 25% of lung cancers. Due to the complexity and diversity of KRAS alteration subtypes, KRAS has long been regarded as one of the most challenging targets in oncology. While several pan-KRAS inhibitors have entered clinical development, there remains a critical need for enhanced potency.

Preclinical studies demonstrated that ABSK211, a novel oral pan-KRAS inhibitor, exhibited broad and potent inhibitory activity across multiple KRAS alterations, including KRAS G12D, G12C, G12S, G12V, G13D, and KRAS wild-type amplification. In vitro studies showed that ABSK211 significantly inhibited proliferation of multiple KRAS-mutant tumor cell lines at sub-nanomolar to nanomolar concentrations, while exhibiting minimal activity in KRAS wild-type cell lines with normal copy number. In vivo studies further demonstrated that ABSK211 induced deep and durable tumor regression across multiple KRAS-altered tumor models, with robust target engagement.

In addition, ABSK211 demonstrated strong potential in combination therapy settings across multiple tumor models. Study results showed that combining ABSK211 with PRMT5 inhibitors, cetuximab, immunotherapy, and chemotherapy further enhanced tumor growth inhibition and improved durability compared to monotherapy, supporting future exploration of combination treatment strategies.

About ABSK211

ABSK211 is a novel investigational oral and highly potent small molecule pan-KRAS inhibitor with broad alteration coverage independently developed by Abbisko Therapeutics. It is designed to target a broad spectrum of KRAS alterations, including common KRAS mutation subtypes and selected amplification settings. Preclinical data demonstrated encouraging anti-tumor activity and favorable selectivity of ABSK211 in both monotherapy and combination treatment settings, supporting its potential to provide a new therapeutic option for patients with KRAS-altered solid tumors. ABSK211 has received IND clearances in both China and the U.S. and has entered global clinical development.

About Abbisko Therapeutics

Founded in 2016, Abbisko Therapeutics (HKEX: 02256) is a pharmaceutical company committed to the research, discovery, and development of innovative and differentiated medicines designed to address unmet medical needs in China and globally.

Adhering to international concepts and standards for drug development, Abbisko primarily centers on small molecules, emphasizing precision oncology and immuno-oncology, with a growing expansion into both non-oncology therapeutic areas and other modalities. Its self-developed Class 1 innovative drug, pimicotinib, has received approval from the China NMPA, while dozens of other candidates are currently in clinical development stages.

Abbisko Therapeutics is committed to creating first-in-class and best-in-class innovative drugs. Through an unwavering pursuit of scientific innovation, the company strives to provide transformative therapies with breakthrough efficacy for patients worldwide.

For more information, please visit www.abbisko.com.

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.