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Ocumention Therapeutics
歐康維視生物

(Incorporated in the Cayman Islands with limited liability)
(Stock code: 1477)

VOLUNTARY ANNOUNCEMENT
PATIENT ENROLLMENT COMPLETED IN THE SECOND
PHASE III CLINICAL TRIAL OF OT-301 IN CHINA

This announcement is made by Ocumention Therapeutics (the “**Company**”, together with its subsidiaries, the “**Group**”) on a voluntary basis to keep the shareholders of the Company and potential investors informed of the latest business updates of the Group.

The board (the “**Board**”) of directors of the Company is pleased to announce that one of the Group’s key drug candidates, OT-301 (NCX 470), a first-in-class, nitric oxide (NO)-donating prostaglandin analog under joint development by Nicox S.A. (“**Nicox**”) and the Group, has completed the enrollment of over 140 patients for its second phase III clinical trial (the “**Denali trial**”) in China recently. The first phase III clinical trial of NCX 470, namely the Mont Blanc trial, was initiated by Nicox in the United States in June 2020, in which the 0.1% dose was selected over the 0.65% dose through an adaptive design, demonstrating robust efficacy and safety in topline results.

The Denali trial is a three-month phase III multi-regional clinical trial evaluating the safety and efficacy of OT-301 (NCX 470) ophthalmic solution, 0.1%, versus the current standard of care, latanoprost ophthalmic solution, 0.005%, for the lowering of intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension. The Denali trial, which includes a long-term safety extension, has enrolled more than 670 patients at approximately 90 clinical sites in the United States and China.

OT-301 (NCX 470) is a new chemical entity invented by Nicox and designed to release both bimatoprost, a United States Food and Drug Administration approved prostaglandin analog, and NO, for the lowering of intraocular pressure in patients with open-angle glaucoma and ocular hypertension. The Group obtained an exclusive license from Nicox to develop, make, have made, import, export and sell OT-301 (NCX 470) in greater China in December 2018, and extended the exclusive right to Korea and 12 countries in Southeast Asia in March 2020.

Cautionary Statement: The Company cannot guarantee that it will ultimately commercialize OT-301 (NCX 470) successfully. 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
Ocumension Therapeutics
Dr. Lian Yong CHEN
Chairman and Non-executive Director

Hong Kong, December 3, 2024

As of the date of this announcement, the Board comprises Mr. Ye LIU and Dr. Zhaopeng HU as executive directors, Dr. Lian Yong CHEN, Dr. Wei LI and Mr. Yanling CAO as non-executive directors, and Mr. Ting Yuk Anthony WU, Mr. Yiran HUANG and Mr. Zhenyu ZHANG as independent non-executive directors.