

Hong Kong Exchanges and Clearing Limited and The Stock Exchange of Hong Kong Limited take no responsibility for the contents of this announcement, make no representation as to its accuracy or completeness and expressly disclaim any liability whatsoever for any loss howsoever arising from or in reliance upon the whole or any part of the contents of this announcement.



Gaush Meditech Ltd

高視醫療科技有限公司

(Incorporated in the Cayman Islands with limited liability)

(Stock Code: 2407)

VOLUNTARY ANNOUNCEMENT
GAUSH TELEON OBTAINS INTERNATIONAL
QUALITY MANAGEMENT SYSTEM CERTIFICATION FOR
MEDICAL DEVICES

This announcement is made by Gaush Meditech Ltd (the “**Company**”, together with its subsidiaries, the “**Group**”) on a voluntary basis, to inform the shareholders and potential investors of the Company regarding the latest business development of the Group.

The board (the “**Board**”) of directors (the “**Director(s)**”) of the Company is pleased to announce that Gaush Teleon Ltd* (高視泰靚醫療科技有限公司) (“**Gaush Teleon**”), a subsidiary of the Company, has recently been officially awarded the ISO 13485:2016/EN ISO 13485:2016 Quality Management System Certification for Medical Devices (Certificate No.: 6214501) by DEKRA.

This certification comprehensively covers the entire process of design, development, production and sale of intraocular lenses used for the treatment of cataract and presbyopia, as well as their supporting intraocular lens implant systems.

DEKRA, founded in 1925, is the world’s leading independent, non-listed expert organization in the field of testing, inspection and certification, serving a wide range of industries.

ISO 13485 is a specialised quality management system standard for the medical device industry, issued by the International Organization for Standardization (“**ISO**”). It focuses on “patient-centric” risk management and full-process quality control, serving as a core compliance benchmark recognised by global medical device regulatory authorities, such as China’s NMPA, EU CE and U.S. FDA. Achieving this certification signifies that an enterprise has established a robust quality management system spanning the entire product lifecycle, including design and development, production, inspection and after-sales service, ensuring compliance with regulatory requirements and a commitment to continuous improvement.

The attainment of this prestigious certification underscores the Company's dedication to enhancing its quality management system, also demonstrates the Company's capabilities in achieving international standards in medical device quality management, risk control and regulatory compliance, laying a solid foundation for the Company to further expand into international markets.

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
Gaush Meditech Ltd
Mr. Gao Tieta
Chairman and Executive Director

Hong Kong, September 29, 2025

As of the date of this announcement, the Board comprises Mr. Gao Tieta as Chairman and executive Director, Mr. Liu Xinwei, Mr. Zhao Xinli, Mr. Zhang Jianjun and Ms. Li Wenqi as executive Directors, Dr. David Guowei Wang as non-executive Director, and Mr. Feng Xin, Mr. Wang Li-Shin and Mr. Chan Fan Shing as independent non-executive Directors.