DH and NMPA organise seminar on “GBA Medical Device Regulations and Registration Process Training” for local medical device trade (with photos)

The Department of Health (DH) and the Guangdong-Hong Kong-Macao Greater Bay Area (GBA) Center for Medical Device Evaluation and Inspection of the National Medical Products Administration (NMPA) today (May 27) organised a seminar on "GBA Medical Device Regulations and Registration Process Training" in Hong Kong. The seminar aims to help the local industry tap into the Mainland market and inject new impetus into healthcare innovation, co-operation and development in Mainland China and Hong Kong.

The Senior Engineer of the Integrated Affairs (Quality Management) Division and a number of reviewers from the Center were keynote speakers at the seminar. They explained to the local industry the major regulations governing medical devices on the Mainland and elaborated on the process and requirements for registering medical devices (including active, passive and in-vitro diagnostic medical devices) on the Mainland. This information will help medical device manufacturers and agents better prepare for and successfully enter the Mainland market.

The half-day seminar was attended by over 130 representatives of the local medical device industry.

The DH expresses its sincere gratitude to the NMPA for its unfailing support in co-organising the seminar for the Hong Kong trade, which was well received by the participants for its highly useful content.

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China is a vital market and manufacturing hub for the global medical device and pharmaceutical industries. In the future, the DH will continue to organise similar activities for the industry, leveraging Hong Kong's unique advantages and reinforcing connectivity with the Mainland and the world. The DH is committed to aligning with the nation's direction of high-standard opening up and promoting healthcare innovation and development.


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2025年5月27日