

Hong Kong International Medical Device Regulatory Forum

Navigating the Future of Innovation and AI in Medical Device Regulation

28 May 2025

- Date** : 28 May 2025 (Wednesday)
- Time** : 8:30 am – 9:00 am | Registration
9:00 am – 12:45 pm | Forum and Panel Discussions
- Venue** : Charles K. Kao Auditorium
1/F, Building 10W, Hong Kong Science Park,
Shatin, Hong Kong
- Language** : English or Putonghua
(simultaneous interpretation will be provided)
- Registration** : <https://invitation.hk/HKIMDRF2025.html>
- Deadline** : 19 May 2025
- Note** : Free admission.
Limited seats are available.
Prior registration is required.



Register Now!

Featured Speakers and Moderators



Dr. Ronald LAM Man-kin, JP

Director of Health,
Department of Health,
The Government of the
Hong Kong Special Administrative Region,
The People's Republic of China



Prof. Philip CHIU

Dean,
Faculty of Medicine,
CUHK

National and Overseas



Mr. LIU Bin

Director,
Greater Bay Area Center for
Medical Device Evaluation
and Inspection,
National Medical
Products Administration,
The People's Republic of China



Dr. Muralitharan PARAMASUA

Chief Executive of
Medical Device Authority,
Ministry of Health Malaysia



Prof. Peter HEGYI

Chairperson,
Basic and Clinical
Translational Sciences,
Academia Europea



Engr. Sameer ALHAMDAN

Acting Director of Regulatory Affairs,
Medical Devices Sector,
Saudi Food and Drug Authority

Local



Prof. Samuel AU

Director,
Multi-Scale
Medical Robotics Center,
CUHK



Prof. Jason CHAN

Assistant Dean and Chairman,
Department of Otorhinolaryngology,
Head and Neck Surgery,
Faculty of Medicine,
CUHK



Ms. Rachel FAN

Chief Executive Officer,
GenieBiome Limited



Prof. KWOK Ka Wai

Department of Mechanical and
Automation Engineering,
Faculty of Engineering,
CUHK



Dr. Jerry WANG

Co-Founder,
Chief Technology Officer
and Chief Operating Officer,
Cornerstone Robotics



Dr. WONG Chi Hong, Ambrose

Principal Medical &
Health Officer (Medical Device),
Department of Health,
The Government of the
Hong Kong Special Administrative Region,
The People's Republic of China

Why Join?

- Gain **global regulatory insights** e.g. NMPA, EMA, on medical devices
- Network with **regulators, industry leaders, and experts** worldwide
- Foster **collaborations** with innovators, manufacturers, and distributors
- Learn from **real-world cases studies** on regulatory challenges and solutions
- Explore **post-market surveillance, clinical evaluations, and QMS** best practices
- Enhance expertise in **regulatory affairs, R&D and quality assurance**

Who Should Attend?



Medical device
manufacturers,
importers,
and distributors



Investors
and venture
capitalists



Regulatory
consultants



Healthcare
professionals



AI and
medtech
innovators



Academic
researchers

Enquiries

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