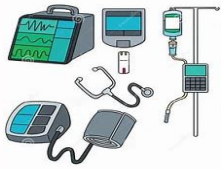
Regulatory Requirements for Medical Device Clinical Trials



SingHealth Research Integrity, Compliance and Ethics (RICE) and ARQon (Asia Regulatory & Quality Consultancy For Medical Device & Drug) will jointly host a webinar to share on the regulatory requirements of conducting Medical Device Clinical Trials.

Scope of Webinar

What is a Medical Device? Device, Software, AI

Risk Control? Classification, Design Risk

HSA Regulatory Requirements

Clinical Research Material (CRM) & Import License

Adverse Events Reporting

Speaker Details:

Ms May Ng

MSc (Biomed. Eng), Post Grad (Medtech Manufacturing), BSc (Biochem & Microbio)

Global Regulatory Director, ARQon

- ISO13485 lead auditor trained. MDSAP, CE MDD/MDR/IVDD/IVDR trained.
- ARQon Group 8 years, MedtechBOSS 3 years, IMDS Group 3 years
- Biosensors International Group, 4 years
- Singapore Health Science Authority (HSA),10 years

Event Details:

Date: 20 July 2022 (WED)
Time: 12:00 PM – 1.30 PM
Location: Zoom Webinar
Registration is required.

Click here to register →
https://ihis.zoom.us/webinar/register/WN_vaRI2HrOQmKi8dKoq40JPg
***Kindly register using your SingHealth corporate email (not personal email)**

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Organised by:
SingHealth Research Integrity, Compliance and Ethics (RICE)
For enquiries, please email oric@singhealth.com.sg