Asia Regulatory & Quality Consultancy for Medical Devices & Drugs

A consultancy firm for medical devices & drug companies. We assist our clients in areas of regulatory & quality from ***product development technical file*** *to* ***product registration submission*** *to attain market approval in ASIA, ASEAN, EU, US & the Rest of the World.*

Team from ex-regulator in Singapore and trainer for Medical Device Authorities in ASEAN & ASIA.

SERVICES

- Registration
- Development
- Quality Management System
- Process Validation
- Distribution
- Clinical Trials
- Regulatory Management System
- Training

Please visit us at **Hall 16, Booth G58-05**

www.arqon.com

PROGRAMME (15 & 16 Nov 2016)

10 – 11 AM **Regulatory Strategy for Product Registration:**
Least regulated countries vs most regulated countries globally?

11 – 12PM **Marketing strategy & Distributor search in Asia**

2 – 3 PM **Global Regulatory Changes & Impact in next 3 to 5 Years:**
EU MDR/IVDR, Canada MDSAP, ISO13485:2016, ASEAN MDD

3 – 4 PM **Clinical Evaluation/trial:** Europe & Asia

4 – 5 PM **Medical Device Regulatory Control & Biocompatibility lab in Malaysia**

Speakers

Ray Soh

Shikharesh Das

Daniel Shoukier & May Ng

Joalin Lin & Daniel Shoukier

Medical Device Authority & MOH

INVITATION

RSVP to

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☐ **YES**, I will attend on

15th ☐10AM ☐11AM ☐2PM ☐3PM ☐4PM

16th ☐10AM ☐11AM ☐2PM ☐3PM ☐4PM

Name

Email

Company