Organized by: NUS Department of Biomedical Engineering Faculty of Engineering | RAPS Regulatory Affairs Professionals Society

EU Medical Device Legislation- RELOADED

Seminar Overview

- Background
  - Moving from Directive to Regulation
- In vitro diagnostic regulation
- Medical Device Regulation
- Retained and new requirements

:: NUS LT 6 ::
:: 9am to 12pm ::
:: Sat, 4th March 2017 ::
:: Tea break and lunch provided ::

Target audience

- GCMDRA students
- All Executives and Professionals in the IVD and MD industries
- Registration fee : $120 (50% discount for GCMDRA alumni)

Mr Rod Ruston, Director of Priory Analysts

Rod has a career which spans manufacturing, design, quality assurance and regulatory affairs. He has been working with the EU medical device regulatory from its introduction in the early 90's. He was first Chair of the EU Notified Bodies Group and also chaired the RAPS Regulatory Affairs Certification Board

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http://bioeng.nus.edu.sg/edu/MDRA/EUseminar