

EUROPEAN MEDICAL DEVICE LEADERSHIP: ADVANCED TRAINING COURSE



An advanced course organized through a collaborative initiative by the Academy for Patient Centred Excellence and Innovation in Regulatory Sciences and The Organisation for Professionals in Regulatory Affairs (TOPRA)

Venue: Malta Life Sciences Park, San Gwann

Date: 29 - 30 October 2019

Early bird registration till 15 September 2019	€1250
Trio Delegation till 15 September 2019	€2750
Late registration	€1500

Advocating medical device safety

Do you want to...

- ... be a **champion** in taking the medical device industry and science to the next higher level?
- ... be part of an **enthusiastic** forum and share knowledge and ideas?
- ... **embrace** the challenges ahead in line with the EU Medical Device regulations?
- ... meet the **experts** in the field?

Benefits of the course

- * Understand regulatory sciences of medical devices
- * Transform regulations into real world practice
- * Discuss the emerging science of medical devices
- * Sustain a culture of excellence in medical device intelligence
- * Form partnerships between stakeholders
- * Establish a legacy of knowledge on medical devices

Course participants will be able to:

- ◇ Classify medical devices
- ◇ Appreciate conformity assessment
- ◇ Incorporate quality into medical devices
- ◇ Prepare and plan for EU Medical Devices Regulations
- ◇ Understand post-marketing surveillance devices
- ◇ Implement field safety corrective action
- ◇ Interpret device labelling
- ◇ Market in a timely, safe and effective rewarding manner

Are you...

- ... interested in helping people **live better** through the use of safe medical devices?
- ... responsible for having **affordable and accessible** medical devices to ensure superior patient care?
- ... concerned with medical device **marketing or acquisition**?
- ... facilitating the **use** of medical devices?
- ... enjoying a satisfying **career** involving medical devices?

**REGISTER
NOW**

Register at:

<https://forms.gle/MscCTRw4mZoDHL2C7>

Further queries may be forwarded to:

academy.medicinesauthority@gov.mt

Who should participate?

- * Distributors, importers, manufacturers, retailers
- * Healthcare professionals
- * Patient advocates: carers, users, managers
- * Scientists, technologists and technicians with interest in medical devices
- * Regulatory professionals and other personnel
- * Professionals from the legal field
- * Procurement personnel
- * Stakeholders with an interest in medical devices industry and business

Courses organised by the **Academy for Patient Centred Excellence and Innovation in Regulatory Sciences**

- * Establish innovative active learning environments
- * Stimulate intellectual curiosity to novel heights
- * Challenge scientific boundaries for excellence
- * Sustain advanced research in emerging sciences
- * Foster sharing of knowledge through partnerships and networking
- * Promote leadership to be nationally recognised and world-renowned
- * Serve national, European and international populations



Certificate of participation is awarded to successful participants