Medical devices, medicines, foodstuffs

Quality and Regulatory Training

This training is for you

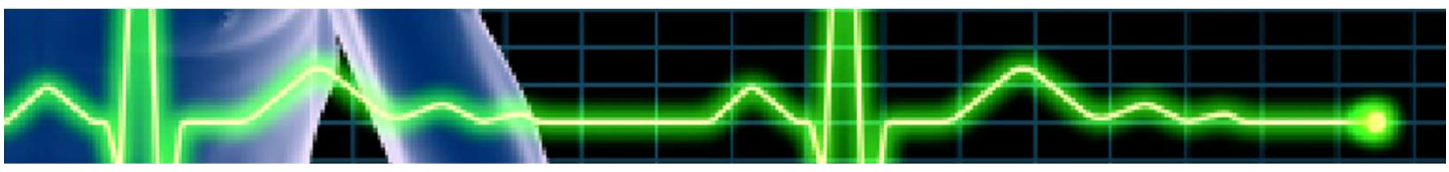
- whose research can produce results that can developed to products like medical devices, foodstuffs and medicines
- who are working in TUTLI-projects
- who are planning to apply TUTLI-funding
- who are interested in working in regulated environment

Time and Place:
Part 1 – 23.1.2019 at 12.15-15.00 @Frost Club / Tellus
Part 2 – 24.1.2019 at 9.15-16.30 @Frost Club / Tellus

Trainers: Jani Hopia and Nina Vartiainen / Kasve Oy
Organized by: Innovation Services

Registration to: https://bit.ly/2qcJdNP at latest 16th January
OBS! You can participate to both parts or choose those sessions which interest you

More info: maarit.jokela@oulu.fi



Program

23rd Jan at 12.15-15.00 @Frost Club

Part 1: Learn the basics from quality management and regulatory requirements

- Why quality assurance is important in companies?
- How quality and regulation should be taken into account in R&D phase?

24th Jan @Frost Club

Part 2: Quality management and regulatory requirements when you develop

9.15 – 11.00 Medical devices (MD)
- Legislation and regulations
- Essential requirements, registration and market entry
- From MD directive to MD regulation: transition period in practice

12.00 – 14.00 Medicinal products
- Surveillance of medicines in Finland and Europe
- Medicine life cycle and key operators in Europe
- Market authorization in Europe

14.30 – 16.30 Foodstuffs and dietary supplements
- Development process, claims and trends
- Self monitoring and safety
- Import/export regulations