Program

Clinical regulatory (Susan Hommerson)
Thursday, September 24, 14.00-17.00; Room: CASCADE 2.21, obligatory for all PhD’s and PostDocs

This course is an initiative by e/MTIC (Eindhoven MedTech Innovation Center) and organized by SMPE/e (School of Medical Physics and Engineering). The course is obligatory for all e/MTIC related PhD’s and PostDocs. In this half day course, the ethical and regulatory framework for clinical trials will be presented.

Course contents
The fundamentals of Good Clinical Practice (GCP) and the Dutch regulatory framework including the role and the responsibilities of the researchers involved will be discussed. Furthermore, the process workflow within e/MTIC studies and the respective institutes and where to find support documentation (templates) will be highlighted.

The course will consist of an introduction in GCP and legislation mandatory for all PhDs, post-docs and master students involved in clinical research.

Course objective
After the course, you will have an understanding of the regulatory framework and basic knowledge of the first steps to set-up a clinical trial. In addition, you will know where to find the necessary required information and support within TU/e and e/MTIC related institutions.

Course information
- Date: Thursday, September 26, 2020
- Time: 14-17 hrs
- Location: TU/e, CASCADE 2.21
- Costs: free of charge, with a no show, € 50 will be charged (a presentation list will be circulated)
- This course is obligatory for all e/MTIC PhD’s and PostDocs.
- To register, click here.

Course coordination
- Course contents: Susan Hommerson, Coordinator medical & medical device research at TU/e, s.m.hommerson@tue.nl
- Course registration: Annebee Langenuizen, Course coordinator SMPEe/ courses, smpee@tue.nl