MEDICINA TAL ESCOLAR DE LIS

LINICAL RESEARCH:

WHATS IT ALL ABOUT?



...a focused training for busy investigators and study teams

Next Edition

October 11 and 12, 2018

October 15 and 16, 2018

PORTUGUESE VERSION

COURSE DIRECTORS:

Inês Zimbarra Cabrita, Ph.D Francisca Patuleia Figueiras, Ph.D.

SCIENTIFIC COMMITTEE:

Fausto J. Pinto, MD, Ph.D, FESC, FACC, FSCAI, FASE Dulce Brito, MD, Ph.D, FESC Joaquim Ferreira, MD, Ph.D Catarina Sousa, MD Cristina Valente, Pharm D.





for Continuing Medical Education 12 CME points

For English Edition only

Health Care Professionals (doctors, allied health professionals, nurses, study Clinical Research and anyone who would like to learn the fundamentals of Good Clinical Practice and extend their knowledge in Clinical Research methodology and procedures.

Running clinical studies is a complex task that requires several skills. Skills ranging from Good Clinical Practice over all applicable regulations up to operational aspects on how to carry out clinical studies. Having a high trained and specialized study team conducting clinical research is the main key to achieving success in recruitment objectives and high standards of quality and

This interactive 2-day course will provide you a comprehensive knowledge on the practical aspects of clinical studies, essential to reach the highest quality of data whilst ensuring the study participants' safety and well-being and that your professional knowledge is optimized.

PROGRAM (Main Topics)

DAY 1

9am - 5pm

Session 1, Types of Studies and Research Design

Session 2. Regulatory Aspects

Session 3. Patient-Centered Research

Session 4. Clinical Study: the first contact with investigators and study team

Session 5. Interactive Workshop

DAY 2

9am - 5pm

Session 6. The Physician as Clinician and Principal Investigator

Session 7. Clinical Study Ongoing Activities & Stakeholders

Session 8, Audits and Inspections

Session 9. Pharmacovigilance Session 10. Interactive Workshop













