CETERA
ACADEMIC CONSULTING RESEARCH ORGANIZATION


# CLINICAL RESEARCH

## WHAT'S IT ALL ABOUT?

2022 EDITION

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

## WHEN:

21 & 22 November 2022 | Online

## TARGET AUDIENCE:

Investigators, pharmacists, study nurses and study coordinators with zero to 5 years of experience in clinical research and anyone who would like to learn the fundamentals of Good Clinical Practice and extend their knowledge in clinical research methodologies and procedures.

**COURSE DIRECTORS**
Inês Zimbarra Cabrita, PhD
Francisca Patuleia Figueiras, PhD

**SCIENTIFIC COMMITTEE**
Fausto J. Pinto, MD, PhD, FESC, FACC, FSCAI, FASE
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Catarina Sousa, MD, PhD
Cristina Valente, PharmD

## DESCRIPTION:

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 performance.

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.

**ACCREDITATIONS**

ATIVIDADE CREDITADA PELA ORDEM DOS FARMACÊUTICOS
00180/9/2020
1,95 CDP

ORGANIZED BY
CETERA, a Portuguese Academic CRO, an Autonomous department of the Association for Research and Development of the Faculty of Medicine, based at the Lisbon Academic Medical Centre (CAML).

## PROGRAM | Main Topics

### DAY 1

9am – 5pm

Session 1. Types of Studies and Research Design
Session 2. Regulatory Aspects
Session 3. Clinical Study: the first contact with investigators and study team
Session 4. Patient-centered Research
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

IN COLLABORATION WITH

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