

Clinical Study Report Writing

The aim of the course is to teach participants how to write clinical study reports including synopses and appendices in such a way that they fulfil the regulatory requirements.

Description

The course includes a comprehensive survey of ICH E3 with tips on how to avoid the most common problems, how to simplify the writing process and how to structure quality control and review. The course will also briefly discuss when a complete clinical study report is not necessary and what to include in a report written under such circumstances.

- ICH E3
- Data presentation
- Quality control & review
- Preparation of appendices
- "Abbreviated clinical study reports"

Target group

The course is designed for anybody who is planning to or already has started to write clinical study reports. The course will also benefit anybody involved in review and quality control of clinical study reports and other medical documents. The course will be of most value to those who already have some experience in clinical development.

Course leader

- Cecilia Falkenberg, PhD, Team Leader Medical Writing

Cost

The cost for the course is EURO 375 + VAT when applying one month prior to the date of the course. An application after that date costs EURO 400 + VAT. The price includes morning and afternoon coffee as well as lunch.

Please check next occasion on our web page www.tfsacademy.com or e-mail us at tfs.academy@trialformsupport.com