

Pharmacovigilance

The aim of the course is to provide participants with a basic understanding of pharmacovigilance and the rules and guidelines regulating safety reporting.

Description

- Background
- Definitions
- Laws and regulations in clinical trials
- Sponsor's responsibility
- Laws and regulations post-marketing
- MAH's responsibility

Target group

People needing a basic knowledge of pharmacovigilance, such as Safety Assistants, Clinical Research Managers, Project Managers and Monitors.

Course leader

- Margareta Svensson, MSc, Team Leader Clinical Drug Safety

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.

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