PharmaRelations Academy

Paid | In-person training

Advanced ICH GCP training

with focus on Sponsor Oversight and upcoming revision (R3).

With this training, PharmaRelations will help you pre-plan and implement the upcoming ICH revision in your daily work and organization. We will train you in the current guidelines and highlight the important changes in the ongoing revision – with a focus on Sponsor Oversight.

AFTER THIS COURSE, YOU WILL HAVE A DEEP UNDERSTANDING OF:

- The Sponsor's responsibilities in Clinical Trials
- How to start building quality into the design of the trials
- How to identify Critical-to-Quality factors
- The differences in E6 R2 and R3 *(draft version)*
- ICH E8(R1)

Read more at:
pharmarelations.com/ich

Full day training
Thursday 25 January 2024
Frösundaviks Allé 1, 169 70 Solna, Sweden

Price model*:
Sponsors | 2500 SEK ex. VAT
Standard fee | 5,000 SEK ex. VAT

* Sponsors: Biotech, Pharma, Medtech companies. Standard fee: CROs, Consultant companies and freelancers.
Terms & conditions available at Pharmarelations.com/ich

Meet the trainer **Anneli Hallmin**

Anneli has more than 30 years within the Pharma industry and 20 of those within Clinical Trials. Annelie has worked in many different roles at both Big Pharma and small Biotech.

- Sr CRA, Project- and Clinical Study Manager at both regional and global level
- GCP Expert and SOP writer
- Dedicated GCP & Process Manager