10 CIEC years

INVITATION / TRAINING

GOOD CLINICAL PRACTICE

GCP Advanced Training (for Experienced Investigators and Research Teams)

Modern Risk Management in Clinical Trials

AGENDA

15:00 Welcome and introduction

15:10 Update on GCP requirements for investigators under the ICH-GCP Addendum

15:30 Exercise and joint discussion: How to handle risks in a clinical trial

16:00 Relevance of the Informed Consent Process for minimisation of the risk of low patient recruitment and retention

16:15 Exercise and joint discussion: How to improve the Informed Consent Process at the site?

16:45 Break

17:00 Essential tools in clinical trial organisation

17:15 Exercise and joint discussion: How to better organise the clinical trial at my site?

17:45 Collection, assessment and reporting obligations for safety data from a site's perspective

18:00 Exercise and joint discussion: How to improve safety data collection and assessment?

18:30 What did I learn? Final Multiple Choice Test

19:00 End of training

DATE & TIME

Wednesday: **13 June 2018**
3:00 p.m. - **7:00** p.m.
Followed by a cocktail

VENUE

Centre Hospitalier Luxembourg (CHL)

Amphithéâtre
4, rue Barblé
L-1210 Luxembourg

INFORMATION & REGISTRATION

Registration: **gcptraining.lih.lu**

As the number of seats is limited, we will accept registrations on a first come first served basis

CONTACT

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TRAINER

Ingrid Klingmann

MD, PhD Chairman, European Forum for Good Clinical Practice (EFGCP)

LANGUAGE

English

ORGANIZED BY:

LIH - Clinical and Epidemiological Investigation Center in collaboration with **EFGCP**

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