Certificate of Completion

Introduction to the Clinical Drug Development Process: ICH Good Clinical Practice for Clinical Trial Sites

Module 1: Course Overview & Overview of ICH Good Clinical Practice

Congratulations to

Caroline Dejonckheere

Membership Affiliation and #:
Employee Number:

For Successfully Completing Training
on

July 10, 2018

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QUINTILES

This document must be retained by the licensee for a period of four years after the course concludes

This ICH E6 GCP investigator Site Training meets the Minimum Criteria for ICH GCP investigator Site Personnel Training identified by TransCelerate BioPharma as necessary to enable mutual recognition of GCP training among trial sponsors".

Certificate of Completion

Introduction to the Clinical Drug Development Process: ICH Good Clinical Practice for Clinical Trial Sites

Module 2: Clinical Investigator Obligations & Qualifications, Resources, IRBs/IECs

Congratulations to

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Module 3: Subject Informed Consent & Protocol Compliance

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Introduction to the Clinical Drug Development Process: ICH Good Clinical Practice for Clinical Trial Sites

Module 4: Investigational Product, Randomization, and Unblinding & Source Documents and Case Report Form Completion

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Introduction to the Clinical Drug Development Process: ICH Good Clinical Practice for Clinical Trial Sites

Module 5: Safety Reporting, Financial Disclosure, & Study Closeout, Trial Termination, and Record Retention

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Introduction to the Clinical Drug Development Process: ICH Good Clinical Practice for Clinical Trial Sites

Module 6: Non-compliance, Scientific Misconduct and Fraud & Monitoring and Preparing for Audits and Inspections

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Introduction to the Clinical Drug Development Process: ICH Good Clinical Practice for Clinical Trial Sites

Module 7: The European Clinical Trial Directives

Congratulations to

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