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## *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

**Christa menten**

Membership Affiliation and #:  
Employee Number:

For Successfully Completing Training  
on

**June 28, 2018**

**Provided by: Quintiles Global Learning**



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”.*

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# *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

**Christa menten**

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Employee Number:

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

**Module 3: Subject Informed Consent & Protocol  
Compliance**

Congratulations to

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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**

Congratulations to

**Christa menten**

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Employee Number:

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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**  
Congratulations to

**Christa menten**

Membership Affiliation and #:  
Employee Number:

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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  
**Christa menten**

Membership Affiliation and #:  
Employee Number:

For Successfully Completing Training  
on

**July 1, 2018**

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