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

Julia Starrett

Membership Affiliation and #:
Employee Number:

For Successfully Completing Training
on

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Provided by: Quintiles Global Learning

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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".