

COLLABORATIVE INSTITUTIONAL TRAINING INITIATIVE (CITI PROGRAM)

COMPLETION REPORT - PART 1 OF 2 COURSEWORK REQUIREMENTS*

* NOTE: Scores on this Requirements Report reflect quiz completions at the time all requirements for the course were met. See list below for details. See separate Transcript Report for more recent quiz scores, including those on optional (supplemental) course elements.

- **Name:** Shicheng Guo (ID: 6805118)
- **Institution Affiliation:** Marshfield Clinic Research Institute (ID: 781)
- **Institution Email:** Guo.Shicheng@marshfieldresearch.org
- **Institution Unit:** Center for Human Genetics
- **Phone:** 2816855882

- **Curriculum Group:** Good Clinical Practice Course for Clinical Trials Involving Drugs (ICH focus)
- **Course Learner Group:** GCP for Clinical Trials with Investigational Drugs and Biologics (ICH Focus)
- **Stage:** Stage 2 - GCP Refresher

- **Record ID:** 25419310
- **Completion Date:** 09-Dec-2019
- **Expiration Date:** 08-Dec-2021
- **Minimum Passing:** 80
- **Reported Score*:** 86

REQUIRED AND ELECTIVE MODULES ONLY	DATE COMPLETED	SCORE
GCP Refresher - International Council for Harmonisation (ICH): GCP Requirements (ID: 16779)	09-Dec-2019	5/5 (100%)
GCP Refresher - Investigator's Responsibilities and GCP (ID: 16780)	09-Dec-2019	4/5 (80%)
GCP Refresher - Informed Consent (ID: 16781)	09-Dec-2019	5/5 (100%)
GCP Refresher - Safety Management (ID: 16782)	09-Dec-2019	5/5 (100%)
GCP Refresher - Investigational Product (Drug) Management (ID: 16783)	09-Dec-2019	5/5 (100%)
GCP Refresher - Audits, Inspection, and Monitoring of Research Studies (ID: 16784)	09-Dec-2019	4/5 (80%)
GCP Refresher - Sponsor Responsibilities and GCP (ID: 16785)	09-Dec-2019	2/5 (40%)

For this Report to be valid, the learner identified above must have had a valid affiliation with the CITI Program subscribing institution identified above or have been a paid Independent Learner.

Verify at: www.citiprogram.org/verify/?k3fee036d-c169-4048-8770-94723d765ce0-25419310

Collaborative Institutional Training Initiative (CITI Program)

Email: support@citiprogram.org

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COLLABORATIVE INSTITUTIONAL TRAINING INITIATIVE (CITI PROGRAM)

COMPLETION REPORT - PART 2 OF 2 COURSEWORK TRANSCRIPT**

** NOTE: Scores on this Transcript Report reflect the most current quiz completions, including quizzes on optional (supplemental) elements of the course. See list below for details. See separate Requirements Report for the reported scores at the time all requirements for the course were met.

- **Name:** Shicheng Guo (ID: 6805118)
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- **Curriculum Group:** Good Clinical Practice Course for Clinical Trials Involving Drugs (ICH focus)
- **Course Learner Group:** GCP for Clinical Trials with Investigational Drugs and Biologics (ICH Focus)
- **Stage:** Stage 2 - GCP Refresher
- **Description:** This GCP training contains all of the attested CITI Program modules from the **GCP ICH Refresher Version 2**. 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.

- **Record ID:** 25419310
- **Report Date:** 09-Dec-2019
- **Current Score**:** 86

REQUIRED, ELECTIVE, AND SUPPLEMENTAL MODULES	MOST RECENT	SCORE
GCP Refresher - International Council for Harmonisation (ICH): GCP Requirements (ID: 16779)	09-Dec-2019	5/5 (100%)
GCP Refresher - Investigator's Responsibilities and GCP (ID: 16780)	09-Dec-2019	4/5 (80%)
GCP Refresher - Informed Consent (ID: 16781)	09-Dec-2019	5/5 (100%)
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