

CITI Collaborative Institutional Training Initiative

CITI Good Clinical Practice Curriculum Completion Report

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CITI Good Clinical Practice Course:

Stage 1. Basic Course Passed on 04/22/10 (Ref # 4325122)

Required Modules	Date Completed	
GCP Introduction	04/19/10	3/3 (100%)
Overview of New Drug Development	04/22/10	5/5 (100%)
International Conference on Harmonisation (ICH): GCP Requirements	04/16/10	4/4 (100%)
FDA Regulated Research and ICH for Investigators	04/19/10	5/5 (100%)
International Conference on Harmonisation - ICH for Investigators	04/20/10	no quiz
Conducting Investigator-Initiated Studies According to FDA Regulations and Good Clinical Practices	04/20/10	3/3 (100%)
Investigator Obligations in FDA-Regulated Clinical Research	04/20/10	5/5 (100%)
Managing Investigational Agents According to GCP Requirements	04/20/10	5/5 (100%)
Conducting Clinical Trials of Medical Devices	04/20/10	3/3 (100%)
Informed Consent-An Ongoing Process	04/22/10	4/4 (100%)
Detection and Evaluation of Adverse Events	04/22/10	3/4 (75%)
Reporting Serious Adverse Events	04/22/10	4/4 (100%)
Audits and Inspections in Clinical Trial	04/22/10	5/5 (100%)
Monitoring of Clinical Trials by Industry Sponsors	04/22/10	8/8 (100%)
Completing the CITI GCP Course	04/22/10	no quiz
Palm Beach Research Center	04/22/10	no quiz

For this Completion Report to be valid, the learner listed above must be affiliated with a CITI participating institution. Falsified information and unauthorized use of the CITI course site is unethical, and may be considered scientific misconduct by your institution.

Paul Braunschweiger Ph.D.
Professor, University of Miami
Director Office of Research Education
CITI Course Coordinator

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