

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 09/20/10 (Ref # 4349329)

Required Modules	Date Completed	
GCP Introduction	09/13/10	3/3 (100%)
Overview of New Drug Development	09/13/10	5/5 (100%)
International Conference on Harmonisation (ICH): GCP Requirements	09/13/10	4/4 (100%)
FDA Regulated Research and ICH for Investigators	09/13/10	5/5 (100%)
International Conference on Harmonisation - ICH for Investigators	09/13/10	no quiz
Conducting Investigator-Initiated Studies According to FDA Regulations and Good Clinical Practices	09/13/10	1/3 (33%)
Investigator Obligations in FDA-Regulated Clinical Research	09/17/10	4/5 (80%)
Managing Investigational Agents According to GCP Requirements	09/17/10	5/5 (100%)
Conducting Clinical Trials of Medical Devices	09/17/10	3/3 (100%)
Informed Consent-An Ongoing Process	09/17/10	4/4 (100%)
Detection and Evaluation of Adverse Events	09/17/10	4/4 (100%)
Reporting Serious Adverse Events	09/17/10	4/4 (100%)
Audits and Inspections in Clinical Trials	09/18/10	4/5 (80%)
Monitoring of Clinical Trials by Industry Sponsors	09/20/10	8/8 (100%)
Completing the CITI GCP Course	09/20/10	no quiz
Palm Beach Research Center	07/07/06	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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