

## 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 05/30/10 (Ref # 4257994)

Required Modules	Date Completed	
GCP Introduction	05/28/10	3/3 (100%)
Overview of New Drug Development	04/03/09	5/6 (83%)
International Conference on Harmonisation (ICH): GCP Requirements	04/03/09	4/5 (80%)
FDA Regulated Research and ICH for Investigators	04/03/09	5/5 (100%)
International Conference on Harmonisation - ICH for Investigators	05/28/10	no quiz
Conducting Investigator-Initiated Studies According to FDA Regulations and Good Clinical Practices	05/29/10	3/3 (100%)
Investigator Obligations in FDA-Regulated Clinical Research	05/29/10	5/5 (100%)
Managing Investigational Agents According to GCP Requirements	05/29/10	5/5 (100%)
Conducting Clinical Trials of Medical Devices	05/30/10	3/3 (100%)
Informed Consent-An Ongoing Process	05/30/10	4/4 (100%)
Detection and Evaluation of Adverse Events	05/30/10	4/4 (100%)
Reporting Serious Adverse Events	05/30/10	4/4 (100%)
Audits and Inspections in Clinical Trial	05/30/10	5/5 (100%)
Monitoring of Clinical Trials by Industry Sponsors	05/30/10	8/8 (100%)
Completing the CITI GCP Course	05/30/10	no quiz
Palm Beach Research Center	04/10/10	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

[Return](#)