

CITI Collaborative Institutional Training Initiative

CITI Good Clinical Practice Curriculum Completion Report

Printed on 12/20/2010

Learner: Shirley Zahn (username: SZahn)

Institution: Palm Beach Research Center

Contact Information Email: shirley@PalmBeachResearch.com

CITI Good Clinical Practice Course:

Stage 1. Basic Course Passed on 12/20/10 (Ref # 3772261)

Required Modules	Date Completed	
GCP Introduction	12/20/10	3/3 (100%)
Overview of New Drug Development	12/20/10	5/5 (100%)
International Conference on Harmonisation (ICH): GCP Requirements	12/20/10	4/4 (100%)
FDA Regulated Research and ICH for Investigators	12/20/10	5/5 (100%)
International Conference on Harmonisation - ICH for Investigators	12/20/10	no quiz
Conducting Investigator-Initiated Studies According to FDA Regulations and Good Clinical Practices	12/20/10	3/3 (100%)
Investigator Obligations in FDA-Regulated Clinical Research	12/20/10	5/5 (100%)
Managing Investigational Agents According to GCP Requirements	12/20/10	5/5 (100%)
Conducting Clinical Trials of Medical Devices	12/20/10	3/3 (100%)
Informed Consent-An Ongoing Process	12/20/10	4/4 (100%)
Detection and Evaluation of Adverse Events	12/20/10	4/4 (100%)
Reporting Serious Adverse Events	12/20/10	4/4 (100%)
Audits and Inspections in Clinical Trials	12/20/10	5/5 (100%)
Monitoring of Clinical Trials by Industry Sponsors	12/20/10	8/8 (100%)
Completing the CITI GCP Course	12/20/10	no quiz
Palm Beach Research Center	04/30/10	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

