

CITI Collaborative Institutional Training Initiative

CITI Good Clinical Practice Curriculum Completion Report Printed on 11/1/2011

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Institution: Palm Beach Research Center

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CITI Good Clinical Practice Course:

Stage 1. Basic Course Passed on 10/28/11 (Ref # 6940972)

Required Modules	Date Completed	
GCP Introduction	10/27/11	3/3 (100%)
Overview of New Drug Development	10/27/11	5/5 (100%)
ICH Overview	10/27/11	4/4 (100%)
FDA Regulated Research and ICH for Investigators	10/27/11	5/5 (100%)
ICH - Comparison Between ICH GCP E6 and U.S. FDA Regulations	10/27/11	no quiz
Conducting Investigator-Initiated Studies According to FDA Regulations and Good Clinical Practices	10/27/11	3/3 (100%)
Investigator Obligations in FDA-Regulated Clinical Research	10/27/11	3/5 (60%)
Managing Investigational Agents According to GCP Requirements	10/27/11	4/5 (80%)
Conducting Clinical Trials of Medical Devices	10/28/11	3/3 (100%)
Informed Consent-An Ongoing Process	10/28/11	4/4 (100%)
Detection and Evaluation of Adverse Events	10/28/11	4/4 (100%)
Reporting Serious Adverse Events	10/28/11	4/4 (100%)
Audits and Inspections in Clinical Trials	10/28/11	5/5 (100%)
Monitoring of Clinical Trials by Industry Sponsors	10/28/11	7/8 (88%)
Completing the CITI GCP Course	10/28/11	no quiz
Palm Beach Research Center	10/28/11	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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