

CITI Collaborative Institutional Training Initiative

CITI Good Clinical Practice Curriculum Completion Report Printed on 11/4/2011

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Institution: Palm Beach Research Center

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CITI Good Clinical Practice Course:

Stage 1. Basic Course Passed on 11/04/11 (Ref # 6971198)

Required Modules	Date Completed	
GCP Introduction	11/02/11	3/3 (100%)
Overview of New Drug Development	11/03/11	5/5 (100%)
ICH Overview	11/03/11	4/4 (100%)
FDA Regulated Research and ICH for Investigators	11/03/11	5/5 (100%)
ICH - Comparison Between ICH GCP E6 and U.S. FDA Regulations	11/03/11	no quiz
Conducting Investigator-Initiated Studies According to FDA Regulations and Good Clinical Practices	11/03/11	3/3 (100%)
Investigator Obligations in FDA-Regulated Clinical Research	11/03/11	5/5 (100%)
Managing Investigational Agents According to GCP Requirements	11/04/11	5/5 (100%)
Conducting Clinical Trials of Medical Devices	11/04/11	3/3 (100%)
Informed Consent-An Ongoing Process	11/04/11	4/4 (100%)
Detection and Evaluation of Adverse Events	11/04/11	4/4 (100%)
Reporting Serious Adverse Events	11/04/11	4/4 (100%)
Audits and Inspections in Clinical Trials	11/04/11	5/5 (100%)
Monitoring of Clinical Trials by Industry Sponsors	11/04/11	8/8 (100%)
Completing the CITI GCP Course	11/04/11	no quiz
Palm Beach Research Center	11/04/11	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.
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Director Office of Research Education
CITI Course Coordinator