

# CITI Collaborative Institutional Training Initiative

## CITI Good Clinical Practice Curriculum Completion Report Printed on 11/2/2011

**Learner:** Novlett Lawrence (username: Novlett)

**Institution:** Palm Beach Research Center

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**CITI Good Clinical Practice Course:**

### Stage 1. Basic Course Passed on 11/01/11 (Ref # 6935075)

Required Modules	Date Completed	
GCP Introduction	11/01/11	2/3 (67%)
Overview of New Drug Development	11/01/11	5/5 (100%)
ICH Overview	11/01/11	4/4 (100%)
FDA Regulated Research and ICH for Investigators	11/01/11	4/5 (80%)
ICH - Comparison Between ICH GCP E6 and U.S. FDA Regulations	11/01/11	no quiz
Conducting Investigator-Initiated Studies According to FDA Regulations and Good Clinical Practices	11/01/11	3/3 (100%)
Investigator Obligations in FDA-Regulated Clinical Research	11/01/11	4/5 (80%)
Managing Investigational Agents According to GCP Requirements	11/01/11	5/5 (100%)
Conducting Clinical Trials of Medical Devices	11/01/11	3/3 (100%)
Informed Consent-An Ongoing Process	11/01/11	4/4 (100%)
Detection and Evaluation of Adverse Events	11/01/11	4/4 (100%)
Reporting Serious Adverse Events	11/01/11	3/4 (75%)
Audits and Inspections in Clinical Trials	11/01/11	4/5 (80%)
Monitoring of Clinical Trials by Industry Sponsors	11/01/11	8/8 (100%)
Completing the CITI GCP Course	11/01/11	no quiz
Palm Beach Research Center	11/01/11	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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