

## CITI Collaborative Institutional Training Initiative

### CITI Good Clinical Practice Curriculum Completion Report

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**Learner:** Kristen Neely (username: KristenNeely)

**Institution:** Palm Beach Research Center

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**CITI Good Clinical Practice Course:**

#### Stage 1. Basic Course Passed on 05/20/08 (Ref # 1627172)

Required Modules	Date Completed	
GCP Introduction	05/20/08	no quiz
Overview of New Drug Development	05/20/08	6/6 (100%)
International Conference on Harmonisation (ICH): GCP Requirements	05/20/08	5/5 (100%)
FDA Regulated Research and ICH for Investigators	05/20/08	5/5 (100%)
International Conference on Harmonisation - ICH for Investigators	05/20/08	no quiz
Conducting Investigator-Initiated Studies According to FDA Regulations and Good Clinical Practices	05/20/08	4/4 (100%)
Investigator Obligations in FDA-Regulated Clinical Research	05/20/08	5/5 (100%)
Managing Investigational Agents According to GCP Requirements	05/20/08	5/5 (100%)
Conducting Clinical Trials of Medical Devices	05/20/08	6/6 (100%)
Informed Consent-An Ongoing Process	05/20/08	4/4 (100%)
Detection and Evaluation of Adverse Events	05/20/08	7/7 (100%)
Reporting Serious Adverse Events	05/20/08	6/6 (100%)
Audits and Inspections in Clinical Trial	05/20/08	8/8 (100%)
Monitoring of Clinical Trials by Industry Sponsors	05/20/08	10/10 (100%)
Completing the CITI GCP Course	05/20/08	no quiz
Palm Beach Research Center	04/09/06	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

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