

GOOD CLINICAL PRACTICE WORKSHOP

8-10 March 2010

Institut Jantung Negara
Kuala Lumpur

A three-day course designed to provide individuals with an in-depth understanding of the clinical research process, the roles and responsibilities of key players as well as regulatory requirements. The course consists of lecture and exercises, and upon completion of this course, each participant will have a thorough knowledge and understanding of the International Conference on Harmonization of Good Clinical Practice (ICH-GCP) guidelines. There will be a formal assessment at the end of the workshop, leading to certification in GCP.

Day 1

Overview of Clinical Research in Malaysia
Principles of ICH/GCP and Malaysian GCP
Role of IRB / Ethics Committee

Day 2

Role and responsibilities of Investigator
Role and responsibilities of Sponsor
Role and responsibilities of Study Monitors
Inspection and Auditing of Clinical Trials

Day 3

Role of Regulatory Authority
Good Laboratory Practice
Assessment

Who Should Attend

- Clinicians
- Study Coordinators
- Pharmacists
- Nurses
- All interested Allied-Health Professionals

REGISTRATION FORM

Name

Department

Organization Name and Address :

Tel : ----- Fax : -----

Email : -----

REGISTRATION FEE : RM800

Payable to :

INSTITUT JANTUNG NEGARA SDN BHD

For more information, please contact :

Secretariat, GCP 01/2010
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