23 - 25 APRIL 2018

**ADAMSON HOTEL. KG BARU, KUALA LUMPUR**

# GOOD CLINICAL PRACTICE WORKSHOP


JOINTLY ORGANIZED BY CLINICAL RESEARCH CENTER &
PERSATUAN PASCASISWAZAH HOSPITAL KUALA LUMPUR

## INFO

### REGISTRATION FEE: RM 450.00

FEES INCLUDE COURSE MATERIALS, REFRESHMENTS AND LUNCH

CANCELLATION
ANY CANCELLATION MUST BE MADE IN WRITING TO THE ORGANIZERS. FULL REFUND FOR CANCELLATIONS 14 DAYS PRIOR TO WORKSHOP, 50% REFUND FOR CANCELLATIONS 7 DAYS PRIOR TO WORKSHOP. NO REFUNDS FOR CANCELLATIONS LESS THAN 7 DAYS OR NO SHOW.

WE RESERVE THE RIGHT TO CHANGE THE DATE(S) OR SPEAKER(S) FOR THIS COURSE IF DEEMED FIT WITHOUT PRIOR NOTICE. WE FURTHER RESERVE THE RIGHT TO CANCEL THE COURSE WITHOUT LIABILITY OTHER THAN RETURNING THE COURSE FEE

THE SECRETARIAT WILL NOTIFY REGISTRATION CONFIRMATION VIA EMAIL, UPON WHICH PAYMENT MUST BE MADE WITHIN 7 DAYS

BANK DETAILS: CIMB ISLAMIC BANK BERHAD (KG BARU)
BANK ACC NO: 86-0005640-3
ACC NAME: PERSATUAN PERUBATAN PASCASISWAZAH
PAYMENT METHOD: ATM TRANSFER / ONLINE TRANSFER (SEND A COPY OF THE PAYMENT RECEIPT TO OUR EMAIL)

## ABOUT GCP

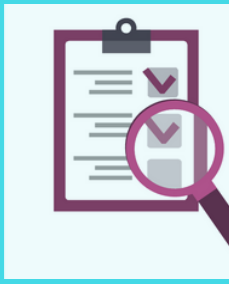
Good Clinical Practice (GCP) is a set of rules & regulations provided by ICH; an international body that regulates clinical trials involving human subjects. It is standard for the design, conduct, performance, monitoring, auditing, recording, analysis and reporting of clinical trials that provides assurance that the data & reported results are credible & accurate, as well as the rights, integrity and confidentiality of trial subjects are protected.

GCP is particularly aimed for Healthcare Professionals such as Clinicians, Nurses, Pharmacists and Allied Health Professionals

It is also beneficial for Research Associates & Study Coordinators, Biomedical & Research Scientists, Statisticians & Database managers, Experienced research personnel who are interested in updating their knowledge regarding GCP

## PROGRAMME

Day 1
Clinical Research in Malaysia
Overview of ICH/GCP
ICH GCP / Malaysian GCP compared
Overview of Ethics of Clinical Research, Ethical Principles & Requirements
Independant Ethics Committee
Ethical Problems in Clinical Research
Clinical Trial Protocol & Investigator's Brochure
Informed Consent



Day 2
Investigator's Resposibility
* Group work
Sponsor's Responsibility
Adverse Events
* Group work

Day 3
GCP Inspection & Regulatory Aspects of Clinical Trial in Malaysia
GCP Examination

## REGISTRATION

NAME: PROF/ DR/ MR/ MS (PLEASE ENTER IN BLOCK LETTERS AS PER IC)

_____

IC/ PASSPORT NO: (PLEASE ATTACH A COPY)

_____

DEPARTMENT: _____
INSTITUTION: _____
DESIGNATION: _____
CONTACT ADDRESS: _____
_____
_____
TEL: _____ (OFFICE) _____ (HP) _____ (FAX)
EMAIL: _____
SIGNATURE: _____

MENU REQUEST: VEGETARIAN / NON-VEGETARIAN

SPONSORED BY: ____ (GOVERNMENT) ____ (COMPANY): PLEASE STATE DETAILS BELOW
____ (SELF)

* FULL ATTENDANCE IS COMPULSORY. CERTIFICATES WILL ONLY BE AWARDED TO PARTICIPANTS UPON PASSING THE MCQ ASSESSMENT (AS PER REQUIREMENTS)

**Secretariat:**
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