Intermediate Professional Study Coordinator Training Workshop

21st – 23rd September 2011
Sunway Medical Centre, Kuala Lumpur

Emerged

HIGHLIGHTS

- ✓ ICF taking role play
- ✓ Hands-on biosamples packing
- ✓ CRF completion

we teach you 'head-to-toe' Clinical Trial Management!

MORE LECTURES

- ❖ Archiving process
- ❖ PSV/SIV
- ❖ Tips to be a good SC
- ❖ Clinical pathway of SC

www.info-kinetics.com

LOOK UP OUR WEBSITE FOR FINALIZED LIST OF SPEAKERS AND WORKSHOP SCHEDULE

Registration Form
21st - 23rd September 2011
Sunway Medical Centre, Kuala Lumpur

The organizers are committed to equality of educational opportunity and do not discriminate against registrations. Seats are limited to 30 based on a first come first served basis. Photocopies of this form are acceptable for registration purpose. Confirmation of your registration will be sent out once payment has been processed.

Title : **Full Name(Print) :** _____

Preferred name (name tag) : _____

Address (for certificate sending) : _____

_____

Organization: _____ **Designation in organization :** _____

Contact No. : _____ **Email Address :** _____

Dietary remarks : _____

Learning objectives

A successful clinical trial is supported by a well trained and knowledgeable study team working hand in hand. Study coordinator or research nurse plays an important role in the study team, being the left and right hand of the principal investigator. This workshop aims to prepare and build up individual capabilities in new technology for a research position at any phase of Good Clinical Practice (GCP) trials with minimal supervision after the completion of the training.

Who should attend

If your work involves 'head-to-toe' clinical trial management, this is a workshop that you can't miss. Study coordinators and research nurses planning or currently engaged in clinical trials, clinicians who wish to know better about the operation of clinical trial will draw direct and immediate benefits. Attendants are preferably GCP trained and have at least assisted/conducted 2 clinical trials.

Faculty

Speakers comprise experts whom have valuable experiences from clinical research and pharmaceutical industries. Take this opportunity to learn from people with real life testimony on how to conduct quality trials. A mixture of lectures and hands-on application in small breakout groups maximize learning benefits.

Workshop registration fees

Registration fee is RM1200 per person. The fees will cover workshop materials, lunch and refreshment breaks during the workshop. We reserve the right to cancel the workshop without any liability other than full refund of the registration fees.

Accommodation

Participants are required to make their own arrangement, if needed.

Cancellation

Refunds (RM 50 processing fee) will be granted to requests received in writing 14 days prior to the start of the workshop. No refunds will be granted after this date.

How to register

You may register by filling up the registration slip attached and forward back to us via mail or fax. Registration will not be processed without payment. Please send cheque/bank draft/postal order of **RM1200** payable to '**Info Kinetics Sdn. Bhd**.'. A confirmation of your registration will be sent to you upon processed.

Closing date for registration

09th September 2011. Places are limited to 30 only to ensure that participants derive maximum benefit of working in small groups.

Kindly forward registration and enquiries to:

Contact Person: Ms. Patricia Khoo / Mr. Toh
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Organisers

Association of Clinical Research Professionals

Clinical Research Centre
Ministry of Health

Info Kinetics Sdn. Bhd.