Hands on Workshop on
Clinical Trial Management and Recruitment Planning
&
New Perspectives of Pharmacovigilance

ScientiaBio announces two workshops for clinical research, healthcare, medical and life science professionals at Singapore.

Workshop 1: Clinical Trial Management and Recruitment Planning
Date: 26th and 27th July, 2012
Venue: Concorde Hotel, Singapore
(Please find the details of this workshop on Page 2)

Workshop 2: New Perspectives of Pharmacovigilance
Date: 28th July, 2012
Venue: Concorde Hotel, Singapore
(Please find the details of this workshop on Page 3)

Brief Background of ScientiaBio's workshop on Clinical Research

- We have conducted 14 hands on workshops across India and other countries on various topics of Clinical Research in last 2 years.
- This is our second annual workshop at Singapore.
- Trainers of the workshops have more than 10+ years of industry experience in the relevant domain and have international training experience.
- Our workshop put more focus on case studies, takes up the examples in areas where professional need more focus.
- We had participants from all major, minor pharmaceuticals, healthcare organization including Pfizer, Novartis, Sanofi, BMS, AstraZeneca and Abbott in our earlier workshops.

Following is couple of sample feedback of our Clinical Research Workshop

- *"Very informative workshop and the trainers engaged the participants" –* Raja Norazireen Raja Ahmad, Research Scientist, Inno Biologics
- *"Quite well organized with most relevant info on stats espy for the non-statistician. Fundamentals were touched upon with a lot of clarity and case studies were very helpful"- Dr Sonica, Associate Director, CPCD, Clinsys*
- *"Good methodology adopted by trainer, very engaging"-* Ms *Ching Chee Juin, Medical Research Associate, Bayer Healthcare*
- *"Thank you for the excellent workshop you organized on Pharmacovigilance. It was an eye opener to us in many aspects " - Ms Geeta, Assistant General Manager – Pharmacovigilance & Medico-regulatory Affairs, Ipca Laboratories*
- *"Good workshop, excellent learning methodology by trainers" – Dr Chirag, Medical Advisor, AstraZeneca*

For more feedback, kindly login http://www.scientiabio.com/UpComingEvents.aspx

Workshop on
Clinical Trial Management and Recruitment Planning

Clinical Trial recruitment is always a very crucial step. In past we have witnessed that lack of proper recruitment plan leads to unsuccessful trial result.

It is also very important to properly conduct the trial for a successful end result.

This hand on workshop will help you to plan and manage the clinical trial in most efficient ways. It will be based on case studies discussion based.

Date: 26th and 27th July, 2012

Venue: Concorde Hotel, Singapore

Workshop Topics

Trial Management (26th July)

- **Trial phases**
- **Trail phase breakdown structure**
- **Preparing a trial management plan**
- **Identifying operational risks at various phases**
- **Preparation of a risk mitigation plan**
- **Execution and monitoring**

Recruitment Planning (27th July)

- **Determination of study population and distribution**
- **Understanding the variance of distribution for different Phases of trials**
- **Laying the groundwork for preparing a recruitment plan**
- **Key factors impacting successful recruitment**
- **Connecting the recruitment plan to the product business**
- **Managing recruitment to minimize screen failures**

Trainers' Profile

Dr Nagendran M V

Dr Nagendran has 30 years of clinical experience, has founded two clinical research organization. He had been principle investigator for 14 Global Trials. He had over sighted more than 30 global trails. He had worked with clients across the globe including USA, Europe, Israel, and Australia. He has vast experience in training for various clinical research activities. He has more than 20 publications in international journal and conferences.

Dr Nagendran has MD degree in medicine from Kasturba Medical College, Bangalore. He also has diploma degree in statistics from London School of Economics.

Workshop on
New Perspectives of Pharmacovigilance

In the current drug development scene the role of pharmacovigilance is becoming increasingly important. The workshop will help you to:

- Learn how to develop a risk management plan
- Prepare for an audit and inspection
- Learn how to handle AE reporting in post marketing surveillance studies
- Be part of an event where the topics are discussed in a clear and comprehensive way

Date: 28th July, 2012

Venue: Concorde Hotel, Singapore

Workshop Topics

- Pharmacovigilance regulations in different Asian countries
- Pharmacovigilance regulations in the US and in Europe
- Pharmacovigilance in Asia
- Preparing for audit and inspection
- Best practices in risk benefit analysis
- Meeting global safety requirements

Trainers' Profile

Prof Jean-Paul Deslypere

Prof Jean-Paul Deslypere currently CEO of Aesculape Pte Ltd and of Proclin Therapeutic Research Pte Ltd , two Asian based, full service CROs. Prof Jean-Paul got his medical degree from the University of Gent in Belgium in 1977. After graduation he obtained his PhD in Endocrinology and Metabolic Diseases in 1984 and his Specialty in Internal medicine/Endocrinology in 1987.

Between 1997 to 2010, he had worked with multiple CRO at vice president or regional director post at Singapore. In 2002, he joined some of Singapore public sector organizations as Research Director, director and adjunct professor.

Jean-Paul Deslypere is Chairman of the CEPM (Council on Education in Pharmaceutical Medicine) . He is Advisory Board Member of the IMI Pharmatrain and of the IMI EHR4CR projects. Recently he received a Honorary Fellowship from the Faculty of Pharmaceutical Medicine (Royal College of Physicians UK), for his outstanding contributions to Pharmaceutical Medicine.

General Information of the Workshop

Who should attend?

This training program is ideal for:

- Individual who is working in Clinical Research, Pharmaceutical, Healthcare medical and related industry or academics.
- Student/research scholar/faculty who work in various fields of life sciences, medical sciences and pharmaceutical sciences.

Registration Fees:

SGD 480* for any one day
(any one day of clinical trial recruitment/planning workshop or PV workshop)
SGD 900* for any two days of the workshop
SGD 1260* for both the workshops (all three days)
*Group discounts (10% for group of 3 to 4, 15% for group for 5 or more) will be available, amount inclusive of tax
10% discount for govt officials and representatives from academics

Registration fees include training material, workshop participation certificate, two refreshments and lunch.

Registration Process / Query

✉ **Fill up the registration form and send us in email to admin@scientiabio.com with subject line "Clinical Research Workshop –Singapore", write to this same email if for any query.**

☎ **Call us at 0091 99453 18216 / 0091 77608 79299**

Visit our website www.scientiabio.com to find out more details.

Payment Method

Bank transfer the registration fee directly to the following bank account:
Account Number: 016905009418
Account name: SCIENTIABIO
Swift code: ICICINBBNRI
Bank :ICICI Bank
Branch: C M H Road, Indira Nagar, Bangalore, India
Payment Message: Registration fees for scientific training

Payment can also be also made via credit card, write to us at admin@scientiabio.com for your secure payment gateway link.

We don't accept bank cheques.

Terms and Condition

- **Payment Confirmation Notifications** will be sent to your email address, once your payment is processed successfully.
- **Refund Policy:** Registrants who cannot attend, and do not send a substitute, are entitled to a refund of 50% if a request is received in writing on or before July 5th, 2012. Registrants are liable for their full fees after that date.

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