

Translating Experience Into Reality

December 6, 2008 8:30am – 4:30pm
ITC Hotel Windsor Manor, Bangalore

Introduction to the Conference

The theme of this conference is “translating experience into reality”. Conducting world class clinical trials in India is a unique opportunity, with unique challenges. This conference, jointly conducted by Indian Society of Clinical Research (ISCR) and Indian Association of Statistics in Clinical Trials (IASCT), focuses on communicating the experience, triumphs and despairs of established luminaries in the field of clinical research in India.

About ISCR and IASCT

Indian Society for Clinical Research is an association of clinical research professionals registered under the Societies Registration Act (1860). The Society brings together all those who are engaged in clinical research activities in India and provides a forum for exchange of information and learning. ISCR aims to build awareness of clinical research as a specialty in India and to facilitate its growth in the country while helping to evolve the highest standards of quality and ethics.

IASCT is a professional network of Statisticians in the field of Clinical Research in India. Founded in 2007, the primary objectives are to enhance awareness about the role of statistics in clinical trials, to promote biostatistics and statistical programming in clinical research and to enable professional development of statisticians and statistical programmers.



Keynote Speakers



Dr. Ramakrishna holds the Intellectual Property Rights Chair [IPR Chair] of the Ministry of Human Resources and Development at the NLSIU and is the Coordinator of the Centre of Intellectual Property Rights Research and Advocacy [CIPRA] at the NLSIU.

Dr. Ramakrishna, on behest of an international body, has recently completed a survey across investigators, hospitals, pharmaceutical companies and has submitted a report to the Ministry of Health, funded by an international body, on legal and regulatory framework for Clinical Trials, Data Exclusivity and Data Protection in India.



Dr. Chaitanya Dutt is Director on the board of Torrent Pharmaceuticals.

Dr. Dutt holds an MD in Medicine. He practiced as a consulting physician before joining the company in 1982. Since then he has been associated with the Company. His rich experience spans areas of Pharma R&D, clinical research, manufacturing, quality assurance, etc. He is one of the key professionals in the top management team of the Company. He has been instrumental in setting up the Torrent Research Centre (TRC), the research wing of the Company. Under his prudent guidance and leadership, TRC has achieved tremendous progress in the areas of discovery research as well as development work on formulations.

Featured Speakers

Aparna Raychaudhuri, Ph.D. - Aparna, a Biostatistician, has about twenty years of experience in statistics in academia and industry in India and abroad. She has held senior management experience in Abbot Laboratories, GSK, Centocor, Reliance Clinical Research Services and ClinRX.

Mandar Oak, V.P – SIRO Clinpharm, Mumbai .He looks after selection, implementation and rollout of IT systems used in clinical trial process. In his clinical research career of seven years, he has played various roles in Medical Affairs, Data Management, IT and Statistical teams in CRO as well as in the Pharma industry.

Mala Srivastava - Partner at Nextvel Consulting. Mala has about sixteen years of experience in the pharmaceutical industry with about eleven years in clinical research. She has held leadership positions in various organizations in India and abroad.

Agnes Shiewe, Ph.D., - Director, Clinical Research Ecron Acunova, Frankfurt. Dr. Schiewe has been with ECRON since 1993. She has about 20 years and completed over 60 studies spanning US, Europe and Asia.

Vishwanath Mahesh Iyer – Head of Biostatistics and Statistical Reporting for the Oncology group at Novartis, India. He has a Masters in Statistics from the University of Connecticut, and is currently pursuing his Ph.D at Temple University, Philadelphia. Mahesh Iyer has been involved with analyzing and reporting clinical trials since 1998, and has worked at Boehringer-Ingelheim, BMS .



Topics

- Legal and Regulatory framework in conducting studies in india
- Breeding drugs to set-up a study; Medical and Statistical Outlook
- Managing project through project Leadership; monitoring, logistics and management
- Managing data to report

Registration Fees

The Registration fees per person is Rs. 1800, to be paid either by Cheque or DD drawn in favour of "Indian Society for Clinical Research" and mailed to: Shobha Radhakrishnan, Pharm Olam International (India) Pvt Ltd, 217, 4th cross, 19th Main, Koramangala, Bangalore - 560 095

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Shamiq Hussain
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Translating Experience into Reality Registration Form

Kindly fill this form and send by e-mail or fax before the 24th of November 2008 to:

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shobha.radhakrishnan@pharm-olam.com

or

N V Ramamurthy
ramamurthyv@yahoo.co.in

Fax Number: 080 4110 5220

Title (Dr/ Mr/ Ms/ Mrs)	
Name (In Capital Letters)	
Designation	
Name of the Organization	
Mailing Address	
City	
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Pincode	
Bank Name/Cheque/DD No/Date/Amount	
Meal Preference (Veg/ Non.Veg)	

Date: _____

