Workshop on Guidance Document on Clinical Trial Inspection January 31st, 2011, FICCI, New Delhi

The guidance document on Clinical Trial Inspection covers the entire gamut of issues involved in the clinical trial inspection. The scope and extent of the programme covers all clinical trial sites and sponsors/CRO's facilities involved in clinical trial of drugs including the biological and medical device covered under D&C Act.

Federation of Indian Chambers of Commerce & Industry (FICCI) is organizing Workshop on Guidance Document on Clinical Trial Inspection with a purposes to discuss and build awareness on the guidance document on Clinical Trial Inspection for Industry and Investigator.

This workshop will help to develop better understanding of the process and requirements for organization to implement it effectively.

Speakers

Invited
- **Government Regulators***
- **Industry***

Who Should Attend

- **Clinical Research Organization**
- **Investigators**
- **Regulatory professionals**
- **Business Consultants/ Regulatory Advisors**
- **Academics**
- **Hospital Administrators**
- **Government & Regulatory Authority**

Program Highlights

Inaugural Session

Plenary Session

Presentation on Guidance Documents

Technical Sessions

Session I: Planning for Inspection
Session II: Inspection of Trial Sites
Session III: Inspection of CRO/ Sponsor
Session IV: Reporting

Registration

Admission only through prior registration. Limited seating on first-cum-first serve basis

Registration Fees INR 3000/- per delegate

Payment Mode Demand draft/cheque in favor of Federation of Indian Chambers of Commerce & Industry, payable at New Delhi

Postal Address Ms. Ekta Sharma
Clinical Research Divison
Federation of Indian Chambers of Commerce & Industry (FICCI)
Tansen Marg, New Delhi – 110001

Email / Phone clinicalresearch@ficci.com or at 23487305

For more Information please contact- Ms. Geetika Lamba. Research Associate, FICCI; Email :geetika@ficci.com