How to plan risk based approaches for clinical trials

Date : Feb 12, 2018 - 10:00 AM

Event URL : http://www.BostonEventsList.com/events/how-to-plan-risk-based-approaches-for-clinical-trials-feb-2018

Organizer : Netzealous LLC DBA - Compliance4All

Venue : Online Event

Location : 161 Mission Falls Lane, Suite 216,, Mission Falls Ln
Fremont, California, USA, ZIP: 94539
Phone: 8004479407

Ticket Price: One Dial-in One Attendee Price: $150.00
Overview:
You will understand how to identify, evaluate and also how to implement specific risk based techniques for risk management used in clinical trials.

Areas Covered in the Session:
Have explained key risk based process/tools and techniques
Review a risk based approach to protocol design
Understand risk based approach to monitoring / data handling
Review of risk based approaches to QC / QA (Auditing)
Hear best practice of these new risk requirements

Who Will Benefit:
Clinical Development Managers and Personnel
Clinical Research Associates
Clinical Research Archiving and Document Management Personnel
Quality Assurance Managers and Auditors
Clinical Development Managers and Personnel

Speaker Profile:
Dr. Laura Brown , PhD, MBA, Diploma Clinical Sciences, is an independent QA and training consultant in the pharmaceutical industry. She is a managing director with LB Training and

Development Ltd., course director for the M.Sc. in Clinical Research, School of Pharmacy at the University of Cardiff, and course director for M.Sc. Regulatory Affairs, TOPRA. Dr. Brown has 20 years experience running clinical trials and clinical quality assurance in the pharmaceutical industry, and auditing clinical trials internationally.

Event Fee: One Dial-in One Attendee Price: US$150.00

Contact Detail:
Compliance4All DBA NetZealous,
Phone: +1-800-447-9407
Email: support@compliance4All.com

Event Categories :