

TRAINERS & INSTRUCTORS



Ms. Deepali Khandagale, RQAP GCP

Quality Management System professional with 16+ years of extensive and diverse experience in global and domestic Sponsor, CRO, BPO companies for Clinical Research and Medical Affairs. Successfully cleared sponsor and regulatory audits and certifications/renewal of certifications. Hold expertise in managing Quality assurance and quality control activities for clinical programs submitted to regulatory agencies like FDA, EMA, ANVISA, and DCGI. Provided end to end quality support to Phase I-IV trials, BABE and RWE studies and Medical Affairs functions. Successfully established effective Quality Management Systems in all organisations worked for.



Dr. Hitesh Khandagale

Global Clinical Research professional with 18+ years of extensive and diverse experience in global and domestic Sponsor and CRO companies. Successfully handled multiple global and domestic phase I-IV trials and achieved the milestones in the studies much before the timelines committed to the Management by extensive oversight and PM expertise. Worked in various therapeutic areas for global regulatory submissions. Successfully cleared multiple site audits and inspections in the role of Project Manager.



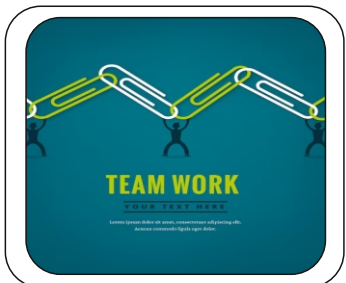
Meenakshi Bagul – Associate Director Data Management

Meenakshi Bagul is having more than 15 years of pharmaceutical industrial experience. Have expertise in End-to-End Clinical Data Management including Startup, Conduct & Close out phases, worked in multiple contract research organization as well as pharmaceutical companies. Handled multiple complex trails like Oncology, Phase I to IV, RWE studies. Also, experienced in areas of Project Management, Resource Management, Team Management. Guiding and managing teams for timely execution, driving greater efficiencies and engagement within the teams.



Ashok Mahangade – Associate Director

Ashok Mahangade having more than 15 years' experience in managing phase I to IV clinical trials across India, Sri Lanka & Bangladesh; worked in large pharmaceuticals like GSK Pharmaceuticals and contract research organization like Aragen Life Sciences, IQVIA, SIRO CLIN pharm. Expertise in managing data collection studies like RWE, claim studies for nutritional supplements and device studies. Managed several independent, regulatory, sponsor & internal audits with no critical findings. Involved in line manager functions, team management, sales & marketing, bid defence, vendor management, trainings activities and project management. Active member of ISCR and was part of training council of ISCR.



Project Managers – having more than 10 years' experience in Clinical Research including more than 6 years of core monitoring experience. worked for both CRO & Pharma companies.

Site Managers- having more than 7 years of Clinical Monitoring experience worked for both CRO & Pharma companies.



DR. MANISH RAJAK – CEO

Manish Rajak having more than 15 years' experience in the pharmaceutical industry encompassing all phase of clinical trials in US, Asia-Pacific, ROW and European countries across multiple therapy areas. Has worked in both contract research organizations, as well as small and large pharmaceutical companies like GSK Pharmaceuticals, PAREXEL International, Novartis, Syneos (previously PharmaNet), Clintec International and IQVIA (previously Quintiles)



SURAJ MADHAVAN – COO

Suraj Madhavan is having more than 14 years' experience in the pharmaceutical industry encompassing all phase of clinical trials in Asia-Pacific and European countries across multiple therapy areas. Has worked in both contract research organizations, as well as small and large pharmaceutical companies like MYLAN Pharmaceuticals, PAREXEL International and IQVIA (previously Quintiles)

And More

***Please note that all sessions would be conducted by Industry Experts in the relevant domain and will be sharing learnings from their own rich experiences to enhance your Knowledge & Skills.**