

ASSOCIATE CLINICAL RESEARCH

Elective: Study Monitoring

“Develop expertise in study monitoring to ensure protocol compliance, data integrity, and participant safety throughout the clinical trial lifecycle.”



WHY JOIN THIS PROGRAM?

- ✓ Builds Core Skills In Study Monitoring With Global Standards
- ✓ Guidance From Clinical Project Managers & Quality Professionals

WHAT YOU WILL LEARN



ICH-GCP E6 (R3) & Global And Local Regulatory Guidelines



Hands-On Exposure To eTMF, EDMS, EDC, And Remote Monitoring Tools



Monitoring Visits, SDV, ALCOA+ And Protocol Compliance

REGISTRATION
FEE
Rs. 500/-

LIMITED SEATS AVAILABLE !

WHY SHOULD YOU JOIN?

- Develop strong monitoring & quality assurance skills
- Ensure compliance with GCP & SOPs
- Protect patient safety & data reliability
- Advance your career in clinical research

WHO SHOULD JOIN?



Life Science Graduates (B.Pharm, MBBS, BDS, BSc, etc.), Healthcare Professionals, Aspiring Clinical Research Professional

ASSOCIATE CLINICAL RESEARCH

Elective: Site Management

“Master the art of site coordination and trial management to ensure successful clinical research from start to finish.”



WHY JOIN THIS PROGRAM?

- ✓ Industry Relevant Skills For Clinical Trial Site Management
- ✓ Hands-On Site Coordination & Compliance Training

WHAT YOU WILL LEARN



Site Initiation & Feasibility



Study Start-up & Regulatory Approvals



Investigator & Site Staff Coordination



Trial Implementation & Oversight



Site Performance & Issue Resolution

REGISTRATION
FEE
Rs. 500/-

LIMITED SEATS AVAILABLE !

WHY SHOULD YOU JOIN?

- Build end-to-end study management expertise
- Ensure regulatory compliance & site readiness
- Improve communication & team collaboration
- Drive study success through effective coordination

WHO SHOULD JOIN?



Life Science Graduates (B.Pharm, MBBS, BDS, BSc, etc.), Healthcare Professionals, Aspiring Clinical Research Professional