

MED DEVICES recognizes the need for well-trained clinical researchers to bridge the gap between the basic researcher's bench and the patient's bedside. MED DEVICES is committed to result oriented, effective, dedicated management of clinical research training in India.

MED DEVICES will work with a clinical study center to provide project specific guidance. MED DEVICES will travel to your clinical center for providing consulting and/ or training to prepare yourself for Clinical Research Trials. Our training sessions are always available at your disposal or let us work with you to customize training schedule that best suits your needs.

To meet the demand of our clients for Clinical Research Professionals, MED DEVICES selects eligible candidates & trains them in:

- Clinical Trial Management (Principles & Practices)
- Pharmacology / Therapeutics
- Pharmacovigilance
- Clinical Data Management
- Regulatory Affairs
- Medical Writing

The training objectives are:

- Obtain a clear understanding of Good Clinical Practice and Standard Operating Procedure for Clinical Research and Clinical Data Management
- Provide trained professionals to CROs, Clinical Investigators at Hospitals'/Medical Site conducting trials.
- Provide an understanding of the use of the Electronic Data Capture (EDC) system and Project Monitoring in the Clinical Trials process.
- Enable participation in design, conduct and management of global clinical trials, conducted at multi-centric sites in India.
- Acquire an understanding of evolving regulatory processes, standards and practice of ICH-GCP in the conduct of different therapeutic trials and in the preparation of submissions to be made to regulatory authorities in India.

