

MED DEVICE LIMITED (U.K.) was formed to provide medical device manufacturers world-wide with flexible and well adjusted service packages to assist its clients to comply with EU legislation. In Mid-2009, we started operation in India with strategically located branch office- covering India, as independent Contract Research Organization (CRO) providing quality services to Pharmaceutical, Biotechnology and Medical Device Industry.

MED DEVICES has a vision of serving the Industry By facilitating-

- Complete Clinical Research Services (Phase II to Phase IV) to Pharmaceutical/ Biotechnology/ Medical Device Industry.
- Representation Services (EAR) from our U.K. office, to Medical Device Manufacturers worldwide against mandatory requirement of European Directive.
- MDQMS (Medical Devices Quality Management Services- ISO 13485: 2003)
- Product Certification (CE Marking) of Medical Devices
- US FDA consulting services to Medical Device Manufacturers
- Third party Inspection Services
- Vendor sourcing

We perform clinical trials under strict adherence to ICH-GCP dedicated to a high level of protocol compliance and responsive to critical communication between sponsor, site, IRB, study personnel, and related trial entities.

We are dedicated to offer creative and innovative full service from planning to market authorization, to our customers in pharmaceutical, biotechnology and medical device industry. We focus on maximizing the scientific communication early on in every project to work out the most successful, least risky path for the approval of product. With our experience, communication is the foundation of our entire operations.

We firmly believe in transparency in communication with our clients related to their projects. It is about reinforcing the trust they place in us and we honour that trust. In that spirit of partnership, our team takes on the responsibility of being the sole point of contact, relaying information on a continuous basis through emails, teleconferences and periodic updates.

- To become one of the most preferred partners for conducting clinical research. We will reach this goal through stable and continuous growth based on our customer focus, our expertise, reliability, resources and Quality Clinical Research Services
- Serving Medical Device Industry to support end-to-end statutory & regulatory compliance.

