

MED DEVICES help you to develop and implement cost-effective strategies and programs to meet the Regulatory requirements for your specific Pharmaceutical/ Biotechnology/ Medical Device's business objectives.

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

Customer satisfaction is the essential driver in our daily business. Each client is unique and different. MED DEVICE customizes an individual solution for each one.

MED DEVICES also maintains an up-to-date Directory of Medical Devices Manufacturers in the world with clear indication –whether the manufacturer has been audited by us or not.

