

Services at a Glance

- 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

