

Your CRO for Success



YOUR FIRST CHOICE FOR QUALITY, COST-EFFECTIVE REGULATORY DOCUMENT MANAGEMENT FOR MEDICAL DEVICE TRIALS



Maximize your internal resources by using BMA for regulatory document collection, review, and management.

Timely collection of accurate regulatory documentation throughout the life of your trial is vital to meeting study timelines. Our staff will work with your team, investigative sites, and vendors to ensure that all required regulatory, financial, and IRB documentation is promptly collected, reviewed, and managed.

“ Get the most out of your study and resources by partnering with BMA ”

Our goal is to provide world class support services that keep your projects on time, within budget and compliant to all regulatory authorities.



Medical Device Regulatory Document Collection & Management

- Confidential Disclosure Agreement & Feasibility Support
- Initial Regulatory Packet Preparation and Dissemination (Paper or Electronic Systems)
- Regulatory Document Creation, Collection, Review & Maintenance
- Investigator Agreements, Financial Disclosures, CVs & IRB Documentation
- Patient Enrollment Tracking
- Regulatory Support for IDE Submissions
- Investigational Device Shipment, Tracking & Reconciliation at End of Study
- Maintenance of Sponsor Systems & Databases
- Study Closure Activities & Document Collection
- Investigator, Country & Central Regulatory File Housing & Maintenance



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A Helping Hand in Clinical ResearchSM

Contact us today to discuss your support needs.

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