



# Your CRO Partner for Success

## YOUR FIRST CHOICE FOR QUALITY, COST-EFFECTIVE REGULATORY DOCUMENT COLLECTION & MANAGEMENT FOR CLINICAL 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.



## Pharmaceutical Regulatory Document Collection & Management

- Start-up, Maintenance & Closeout for Phase I – IV Pharmaceutical Trials
- Sponsor and Investigator-Initiated Trials
- Confidential Disclosure Agreement & Feasibility Support
- Initial Regulatory Packet Preparation & Dissemination (Paper or Electronic Systems)
- Submission of Template Regulatory Documentation to Central IRB
- Ongoing Regulatory Document Creation, Collection, Review & Maintenance
  - 1572s, Financial Disclosures, CVs, Protocol Agreements, Laboratory & IRB Documentation
- Regulatory Submission
- Maintenance of Sponsor Systems & Databases, Including Initiation & Oversight of Product Shipment
- Tracking, Collating & Follow-up on Patient, Vendor & Study Data
- Study Closure Activities & Closeout Document Collection
- Investigator, Country & Central Regulatory File Housing & Maintenance



**B. McLAUGHLIN ASSOCIATES, INC.**  
A Helping Hand in Clinical Research<sup>SM</sup>

Contact us today to discuss your support needs.

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