



# Your CRO Partner for Success

## YOUR FIRST CHOICE FOR QUALITY, COST-EFFECTIVE STUDY MANAGEMENT



*Our team has the experience and industry knowledge required for successful study management.*

BMA can help you meet your corporate and product development goals by supporting many aspects of your clinical trials. We allow you to focus on other critical functions, effectively getting more out of your internal resources.

### “ 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.



## Study Management

- **In-House CRA Responsibilities**
  - Feasibility & Site Selection Activities
  - Creation, Collection & Review of Regulatory Documentation
  - Negotiation of Contracts & Patient Budgets
  - Maintenance of Sponsor Systems & Databases
  - Drug & Biologics Temperature Monitoring
  - Study Closure Activities
  - Regulatory Document Audit & FDA Submission Support
  - Site Consultation & Support
  - Safety Information & Correspondence Distribution
  - Patient Tracking & Post-Study Follow-Up
  - Provide Support to Field Monitors
- **Clinical Study Manager Responsibilities**
  - Study Budget Creation, Review & Maintenance
  - Informed Consent Creation, Review & Maintenance
  - Investigator & CRA Meetings
  - Study Team Management
  - Recruitment & Enrollment Tracking
  - IP, Device & Study Supply Management
  - Study Timeline Planning & Adherence
  - Provide Direction to Field Monitors



**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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