



**A Helping Hand in Clinical Research<sup>SM</sup>**

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



## **PARTNERSHIP FOR SUCCESS**

At BMA, we strive to develop a true partnership with our clients and bring over 18 years of proven excellence in working with study teams, sites and sponsor selected vendors and CROs.

## **COST-EFFECTIVE SOLUTIONS**

Understanding your need to stay within budget, we provide our expertise at highly competitive rates, firmly believing that quality should never be sacrificed in order to keep costs in line. Whether your outsourcing requirements are for the full life cycle of a clinical trial or just a specific need, BMA is here to provide a flexible solution.

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BMA is a WBENC certified CRO providing expertise in clinical trial Study Management, Cold Chain Management, Data Management, Auditing and Clinical Monitoring. Our 18 years of success and growth have resulted from an unchanged approach that has stood the test of time - provide quality, commitment, excellence, value and personalized, flexible service to our clients to support their clinical trial needs. Our commitment to our customers and their projects, as well as the consistent quality of our work is the foundation of our company.

## SUPPORT SOLUTIONS THAT WORK FOR YOU

## Innovative Expertise & Flexible Solutions to Meet Your Needs

**REPUTATION** 18 Years of building a reputation...not simply relying on one

**DIVERSITY** WBENC Certified woman owned, small business and diverse supplier

**GROWTH** Growth driven by repeat business and word of mouth referrals

**EXPERTISE** Cross-functionally trained, experienced and dedicated staff with an average 7 years at BMA and 11 years in industry

### BROAD THERAPEUTIC EXPERIENCE

Biologics	Infectious Disease
Cardiovascular	Musculoskeletal Disease
CNS	Nephrology
Dermatology	Oncology
Endocrinology	Respiratory
Gastroenterology	Urology
Hematology	Women's Health

### CUSTOMIZABLE SERVICES

#### Study Management

- In-House CRAs & Clinical Study Managers
- Negotiation of Clinical Study Contracts & Patient Budgets
- Informed Consent Creation & Review
- Maintenance of Sponsor Systems & Databases
- Creation, Collection & Review of Regulatory Documentation
- Regulatory File Audit & FDA Submission Support

#### Global Data Management

- Full Service & Niche Data Management
- Clinical Scientist Support
- Data Discrepancy/Query & Data Form Collection (Paper & eCRF)
- eCRF User Support & Site Training
- Patient Listing & Narrative Review
- SAE, Lab Normal Range & Lesion Cleaning

#### Cold Chain Management

- Monitoring of Drugs, Devices & Biologics from Manufacture to Subject Administration
- Temperature Deviation Identification, Follow-up, Quarantine & Corrective Action

#### On-Site Monitoring

- Certified Field Monitors
- Site Feasibility, Qualification & Selection
- Initiation, Interim & Closeout Visits
- Source Document Review & Data Verification