



# Your CRO Partner for Success

## YOUR FIRST CHOICE FOR QUALITY, COST-EFFECTIVE ON-SITE MONITORING SERVICES



Ensuring that your trial has quality data, is compliant with all applicable regulations, and meets all timelines is a must for success.

Allow BMA field monitors to work with you and your clinical sites to help meet corporate and study timelines by providing monitoring support throughout all phases of a clinical trial. Our professional, industry experienced monitors will support your sites from feasibility through study closeout, as well as assist with audit preparation and submission support.

### “ 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.



## On-Site Monitoring Activities

- Certified Field Monitors
  - ACRP
  - SoCRA
- Site Feasibility, Qualification & Selection Support
- Initiation, Interim & Closeout Visits
- Source Documentation Review & Data Verification
- Data Discrepancy/Query Support
- Regulatory Compliance Support & Review
- IRB Documentation Review
- Site Preparation for FDA & Sponsor Audits
- Participation & Presentation at Investigator Meetings
- Enrollment Review
- Identification & Addressing of Protocol Deviations & Violations
- Study Supply Inventory
- Reporting of Adverse Events & Serious Adverse Events

Contact us today to discuss your support needs.



1540 McDaniel Drive  
West Chester, PA 19380  
Tel: 610.430.1847

16515 South 40th Street, #125  
Phoenix, AZ 85048  
Tel. 480.659.9120

Info@bmclaughlin.com

www.bmclaughlin.com