



CLINICAL TRIAL OUTSOURCING SOLUTIONS

- PROJECT MANAGEMENT
- SITE MONITORING
- MEDICAL MONITORING
- SITE SELECTION & STARTUP
- DATA REVIEW
- FINAL REPORT
- DOCUMENT MANAGEMENT
- VENDOR MANAGEMENT & PAYMENT
- STUDY STAFFING
- INVESTIGATOR RECRUITMENT

CREATING OPPORTUNITIES FROM CHANGE

Change in the BioPharma R&D market is creating significant opportunities for those who can adapt. Research is no longer done by a single firm, but is often a distributed global multi-partner effort. Escalating costs of R&D and increased demand for accelerated new product development are pushing BioPharma firms to outsource projects to trusted partners. At SterlingBio we help you to create opportunity from industry changes. We help you to accelerate clinical trial projects through a combination of our top talent and through our unique technology for Clinical Trial eDocument Management, SureTrial™.

By facilitating close team alliances and communications throughout the clinical trial study lifecycle, our team works with your team to drive cohesive alliances and to deliver quality products on time and under budget. Convert today's change into opportunity by leveraging our flexible team and technology for your next clinical trial. The gains in cost savings, accelerated development cycles and product quality from outsourcing will provide you with new opportunities for success.

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Delivering Sterling Results