SITE SUPPORT

imarc
WE'LL EARN YOUR APPROVAL.

SITE SUPPORT THAT BRINGS YOU CLEAN DATA ... FASTER

We go beyond filling the gaps. Our team continuously looks for opportunities to enhance efficiency, improve communication and coordination and ultimately, work smarter. When IMARC's site support team helps extract and report critical study and safety data faster, monitors can verify accuracy and provide clean data for analysis sooner.

EXPEDITE DATA REPORTING AND MAINTAIN COMPLIANCE

Our clinical research site support staff will become part of the site team and work with them to ensure accurate data are reported in a timely manner. This allows your team to provide clean data for analysis efficiently and maintain compliance so your study will make it to the finish line faster.

Reach the Finish Line Faster

Your Site. Our Eyes. Is your study data being held hostage by busy sites that cause unnecessary delays in reporting and analysis? Are your research coordinators too overwhelmed with essential tasks like providing patient care and managing administrative duties to keep up with data entry? IMARC is here to help.

Analyze … Extract … Report

IMARC provides comprehensive site support that starts with assessing your site and processes. Then our team will work with yours to develop solutions for more effective coordination and deliver results on time, on budget.

IMARC's site support includes:

- Assisting with remote review of electronic medical records
- Providing remote data entry support
- Call center support
- On-site data and imaging collection support

See the Whole Picture With Our Full Range of Services

Our site support services can stand alone or complement any of the full range of services we provide, including:

- **Monitoring**
  Providing critical thinking, practical experience and full oversight to your trial
- **Auditing**
  An efficient, cost-effective way to identify compliance issues before the FDA does so you can approach your inspection with confidence
- **Training**
  Comprehensive, cost-effective and convenient training programs to ensure your team understands the regulations and how to apply them using critical thinking
- **Project Management**
  Coordinating with team members and vendors to prevent delays and keep your project moving forward
- **Safety Management**
  Providing additional layers of oversight with the Data Safety Monitoring Board and Clinical Events Committee
- **Consulting**
  Helping your team handle quality system development, compliance concerns or other procedural issues

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To learn more, contact John E. Lehmann, Director of Business Development at 440.801.1540
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