PROJECT MANAGEMENT

imarc
WE'LL EARN YOUR APPROVAL.

PROJECT MANAGEMENT SERVICES

Our project management services are designed to accommodate your team's most pressing needs. We can be involved from start to finish or anywhere in between, depending on your goals and your in-house resources.

Some of our project management services include:

- Project planning
- Conducting ongoing risk analysis
- Qualifying and managing vendors
- Managing study start-up processes
- Overseeing monitoring to ensure study compliance
- Strategizing to meet enrollment goals
- Ensuring timely data submission and resolution of data queries
- Coordinating data locks, analyses and study reports
- Supervising study closeout activities, including preparation of regulatory submissions

Keep Your Trial on Track

Are your studies experiencing delays due to limited internal resources or inefficiencies? Our team understands, and we're here to help.

Plan...Execute...Streamline

IMARC has extensive expertise managing a wide range of clinical research trials in the United States and across the world. Having a project manager gives you a single point of contact for your study, allowing you to communicate more efficiently with vendors, team members and site staff across multiple locations. Our highly trained staff will work with yours to develop a strategy that helps you achieve your objectives within your time frame. Then we'll build a plan to tackle key milestones and assess progress throughout the trial. During the process, we will identify opportunities to enhance efficiency without compromising integrity.

The result? Fewer delays and data integrity beyond reproach so you can bring your device to market faster.

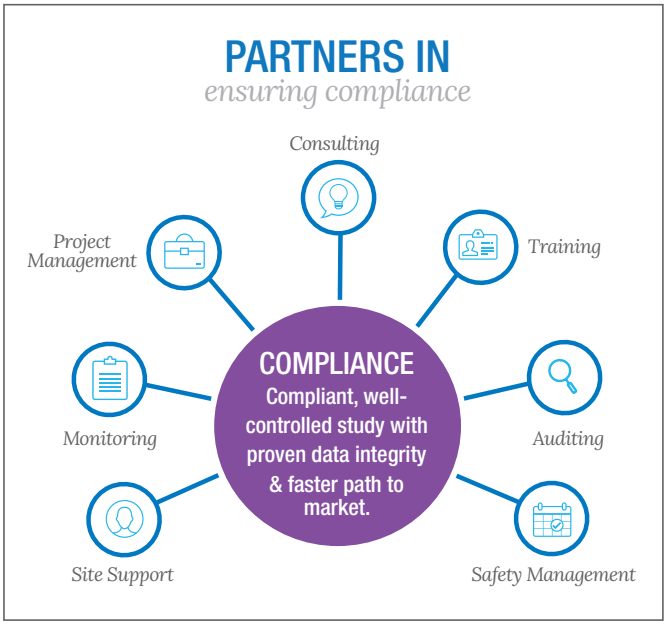
See the Whole Picture With Our Full Range of Services

Our project management 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
- **Safety Management**
  Providing additional layers of oversight with the Data Safety Monitoring Board and Clinical Events Committee
- **Site Support**
  Handling paperwork, data entry and other site responsibilities so on-site staff can focus on subject enrollment and patient care
- **Consulting**
  Helping your team handle quality system development, compliance concerns or other procedural issues
- **Training**
  Comprehensive, cost-effective and convenient training programs to ensure your team understands the regulations and how to apply them using critical thinking

An Extension of Your Team

Our project managers collaborate with your existing team or assemble new teammates, aligning with your company's objectives. IMARC employs true leaders that undergo extensive training in clinical research processes, enabling them to identify and quickly address risks. IMARC project managers have the skills and experience needed to support the study team to guide your study to the finish line faster.



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