MONITORING

imarc
WE'LL EARN YOUR APPROVAL.

Hit Your Marks with IMARC

Reaching benchmarks isn't enough. It is how you reach them that matters. With IMARC Monitors protecting your interests from Day One, your data integrity is ensured, compliance requirements enforced and patient safety protected.

Always Watching. Always Thinking. Always Advancing.

Monitoring sites starts with understanding how to manage them. IMARC has extensive experience doing both. Are your studies suffering setbacks from slow enrollment or delayed submission of critical data? Do costly delays or non-compliant sites keep pushing you further away from approval? There are solutions — and IMARC has them. Whether from the start, or during an ongoing study, our highly trained monitors are qualified to take the burden off you and act as site liaisons to help:

- ☐ Manage site compliance and complete source data verification on time and on budget
- ☐ Assist with handling data queries and ensure sites maintain proper regulatory and product records
- ☐ Develop systems for submitting accurate data in a timely fashion
- ☐ Provide GCP training and mentoring for research coordinators and investigators
- ☐ Alert site team members to pressing issues
- ☐ Assure protection of human subjects

Constantly Monitoring Your Situation

IMARC believes monitoring means having eyes everywhere. Whether it is communicating with upper management, following through on issues and maintenance, tracking trends during studies, or creating customized monitoring tools to boost performance, IMARC stays on top of it for you.

THERAPEUTIC EXPERTISE

- Wound Care
- Cardiovascular
- Peripheral Vascular
- Neurovascular
- Orthopedic
- Gastrointestinal
- Urological
- Urogynecological
- Women's Health

YOUR SITE. *OUR EYES.*

- Site and Investigator Assessments
- Site Initiation Visits
- Periodic Visits
- Remote Monitoring
- Closeout Visits
- FDA Inspections Preparation

BUILD... CULTIVATE... THINK.

These three words should serve as the foundation for your next study. They will with IMARC. Our approach to monitoring extends beyond just working from a checklist. IMARC operates from experience and eliminates barriers to compliance.

IMARC Monitors BUILD Confidence, Trust and Relationships

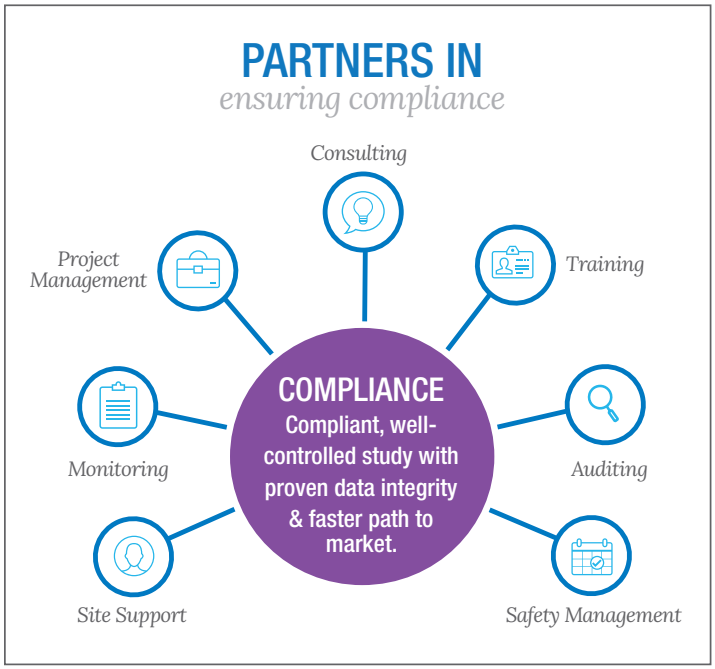
Creating partnerships powered by open communication and healthy dynamics is how IMARC helps you navigate through the complexities of clinical studies toward your goals. We see ourselves as a true extension of your study team and care about the success of your study as much as you do.

IMARC Monitors CULTIVATE Compliance, Commitment and Creativity

Sites monitored from the start by IMARC have never received a warning letter. The reasons: We work diligently to identify non-compliance issues early, proactively problem-solve and implement strategies that prevent recurrence.

STRATEGICALLY POSITIONED

Monitors can be wherever you need them to be – in the U.S. or as Consultants at an international location – to support your study team.



IMARC Monitors THINK Strategically, Logically and Effectively

Our monitors not only embrace data verification, they push it "off the chart" during chart reviews by challenging conventional thinking to verify more than "X=X" and "Y=Y." IMARC monitors use critical thinking skills to see the bigger picture to ensure your study data is accurate and has integrity. Extensive training in the FDA's BIMO process enables us to identify, report and fix vulnerabilities quickly in clinical studies to keep you thinking and looking ahead.

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