



WE'LL EARN YOUR APPROVAL.

# MONITORING

## 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 entrusted.

## Always Watching. Always Thinking. Always Advancing.

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

### CLINICALLY SOUND SITE EXPERIENCE

- ICU
- CCU
- Post-Cardiac Care
- Post-Anesthesia
- Neurology
- Medical / Surgical
- Public Health
- Podiatry
- Oncology
- Psychiatry
- Women's Health

- **Manage site protocol-related issues**
- **Assist with handling site data queries**
- **Develop systems for submitting accurate site data in a timely fashion**
- **Provide GCP training and mentoring for site research coordinators or replacements**
- **Alert site team members to pressing issues**

## 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.



## Your Site. *Our Eyes.*

Assessments • Initiations • Periodic Visits • Close-Outs • FDA Preps

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

## 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.

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### 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.

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### 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, implement strategies that prevent reoccurrence, mentor and conduct comprehensive record reviews.

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### 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.” 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.

## 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 site study team.



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

22560 Lunn Road, Strongsville, Ohio 44149 • **tel** 440.801.1540 • **fax** 440.801.1542  
**info@imarcresearch.com** • **imarcresearch.com**