



TRAINING

Never Stop Learning

Your Site. Our Eyes. A well-trained staff is key to the success and integrity of your clinical study. That's why IMARC University offers comprehensive training that can be customized to meet the needs of your team or individual staff members. Our training includes cost-effective in-person training, teleconferencing, and convenient online courses available 24/7.

Train ... Prepare ... Accelerate

IMARC University's training programs have been developed to ensure clinical researchers understand the regulations as well as best practices and how to apply them using critical-thinking skills.

Our training staff brings a broad range of perspectives, with backgrounds ranging from nursing, engineering, medical technology, laboratory research, and more, to their roles as research professionals. Many have worked in the research industry for years as research coordinators, monitors, auditors, and project managers, and bring a variety of experiences to training programs.

TRAINING
is in Our DNA.

TRAINING SESSIONS

Expand your regulatory knowledge of study operations with these courses:

- Introduction to Research Coordinating
- Monitoring "Musts"
- Understanding Clinical Research Sites (for Project Managers & Monitors)
- Clinical Site Management
- Human Subject Protection
- Monitoring Device Studies
- Device vs. Drugs
- Monitoring Team Roles for Global Studies
- Warning Letters... and the Implications for You

TRAINING TECHNIQUES

IMARC training solutions are designed to provide you with a better understanding of your specific research setting and how it functions. These informative, interactive and inspirational solutions include:

- Didactic Presentations
- Group Activities
- Case Study Discussions
- Warning Letter Reviews & Discussions
- Games & Quizzes
- Virtual Study Experiences



Online Training, Courses on Your Schedule

Whether your team members need to prepare for new roles in clinical research or just need a refresher course in the regulations, IMARC University offers affordable, convenient online courses for individuals and teams. Learners can access their courses online anytime and complete them at their own pace.

IMARC University's online courses prepare individuals for a variety of roles within the clinical research field. With a well-trained staff, your team will be using a consistent process that meets the most stringent standards of patient safety and compliance.



In-Person Training Solutions

Need to onboard an entire team or provide them with specialized training in specific areas? IMARC University delivers practical, experience-based training from clinical research professionals working in the trenches.

Training can be customized to meet the unique needs (and schedules!) of teams of all sizes. We will bring experienced, dynamic trainers to your team or desired location, or host groups at our training facility in Strongsville, Ohio.



Remote Training, Live via Web Conferencing

When interactive training is a must, remote web conference training offers an alternative to in-person methods to allow for questions in real time, from anywhere in the world.

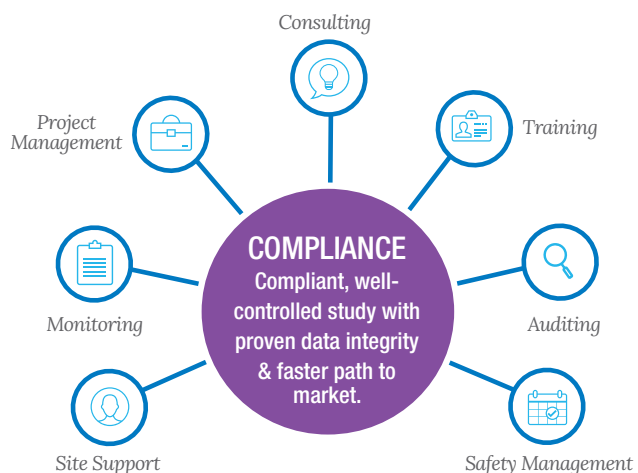
Bring your global team together for remote training, covering a wide variety of clinical research topics.

TRAINING SOLUTIONS FOR ANY ROLE

Our training is ideal for clinical research professionals in a variety of roles at the Sponsor, CRO and Site levels, including:

- Monitors
- Research Coordinators
- Investigators
- Project Managers
- Auditors
- IRB Staff
- Other Sponsor and Vendor Staff Members

PARTNERS IN *ensuring compliance*



TRAINING FOR TRIALS WORLDWIDE

- Human Subject Protection
- GCP Compliance
- Introduction to Research Coordinating
- Monitoring 101
- GCP, JGCP and ICH Guidelines
- FDA Regulations: 21 CFR 11, 50, 54, 56, 312 and 812



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