The CRA's Guide to Monitoring Clinical Research, Third Edition

Karen E. Woodin, Ph.D., John C. Schneider

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The CRA's Guide to Monitoring Clinical Research, now in its third edition, continues to be a key resource for both novice and experienced CRAs seeking to learn more about the field of monitoring or to better understand their roles and responsibilities as the industry becomes more global and technologically focused. With helpful tips and strategies, checklists, personal experiences, key takeaways and exercises, plus new chapters on clinical trial roles and responsibilities, monitoring for device and biologic trials, globalization of studies, EDC and more, *The CRA's Guide* is a must-have training and educational tool that you ll refer to again and again.

Topics include: -A comprehensive review of CRA roles and responsibilities
-Understanding regulations and GCPs
-Study initiation and monitoring plans
-Recruiting and retaining study subjects -The informed consent process
-Conducting adverse event and safety monitoring
-Preparing for audits and detecting fraud
-The future outlook
-Job descriptions and current academic programs
-Devices and Biologics
-Managing Multi-national Trials
-IRBs and Data Safety Monitoring Boards
-Exercises with Answers

Recommended for: -Novice and experienced CRAs
-Health professionals interested in pursuing a career as a study monitor
-Instructors conducting training and educational programs

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