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This guide looks at the evolution of remote clinical trial monitoring from a technology perspective. Our goal is to help accelerate the adoption of remote monitoring by providing clinical operations professionals with a concise guide to understanding the technology options available to support remote monitoring.

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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

This guidebook is filled with valuable information on the role and responsibilities of a clinical research coordinator (CRC) and explains the research process from the site and CRC perspective. Topics covered include: identifying the regulations governing clinical research; describing the drug development process; discussing good clinical practices and how to apply them in clinical trials and organizing a clinical practice.

This concise e-book provides clinicians as well as administrative personnel involved in clinical research with an understanding of documentation related to clinical trial monitoring activities at each stage of the study from planning and set up, through conduct and close-out.

Rev. ed. of: Mosby's pocket guide to fetal monitoring / Susan Martin Tucker, Lisa A. Miller, David A. Miller.

The management of clinical data, from its collection during a trial to its extraction for analysis, has become a critical element in the steps to prepare a regulatory submission and to obtain approval to market a treatment. Groundbreaking on its initial publication nearly fourteen years ago, and evolving with the field in each iteration since then,

Condensing the most important topics in all of clinical research in an easy to understand presentation. The 20 percent of what you need to know in order to be 80 percent proficient! The authors who have operated various levels of businesses in the clinical research industry since 2005 believe that more practical information pertaining to clinical research needs to be accessible to individuals who are new to the industry or are curious about entering the rewarding world of clinical trials. This book reads in an easy to understand style and is based on proven methods the authors have developed to train their own employees and students of their various clinical research academies throughout the years. Picking this up and absorbing the information will allow anyone to gain much better insight into the complicated dynamics of clinical research. This practical roadmap is all you will need to get started on your clinical trial journey! In this book you will learn about: Regulations and the history as well as evolution of GCP. Clinical Research Site Operations Monitoring Dynamics and Typical Monitoring Visits CRO Activities Sponsor Level Dynamics Industry Vendors Common Career Opportunities and Employment Roadmaps

A Practical Guide to Managing Clinical Trials is a basic, comprehensive guide to conducting clinical trials. Designed for individuals working in research site operations, this user-friendly reference guides the reader through each step of the clinical trial process from site selection, to site set-up, subject recruitment, study visits, and to study close-out. Topics include staff roles/responsibilities/training, budget and contract review and management, subject study visits, data and document management, event reporting, research ethics, audits and inspections, consent processes, IRB, FDA regulations, and good clinical practices. Each chapter concludes with a review of key points and knowledge application. Unique to this book is "A View from India," a chapter-by-chapter comparison of clinical trial practices in India versus the U.S. Throughout the book and in Chapter 10, readers will glimpse some of the challenges and opportunities in the emerging and growing market of Indian clinical trials.

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