Clinical Research Coordinator Handbook, 2009, 157 pages, Deborrah Norris, 0937548707, 9780937548707, Plexus Pub., 2009

In this fully revised and expanded fourth edition of the essential reference for clinical research coordinators, Deborrah Norris provides expanded coverage of CRC duties and regulatory requirements, including new sections on investigator responsibilities, data clarification, and adverse event reporting. The book s five appendices include a directory of CRC resources, updated forms and checklists, state regulatory requirements and contact information, conversion charts and tables, and glossary.

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The GCP Harmonization Handbook , Donald E. Maynard, Robert S. Dewoskin, May 31, 1996, Medical, 486 pages. This book is designed to present those involved in clinical research with the similarities and the important differences between the GCP regulations of the United States.

Institutional review board (IRB) reference book protecting the people who volunteer to participate in research, Michele K. Russell-Einhorn, Thomas Puglisi, 2001, Institutional review boards (Medicine), 680 pages.

Intelligent Drug Development Trials and Errors in Clinical Research, Michael Tansey, 2014, Medical, 220 pages. Clinical research is heavily regulated and involves coordination of numerous pharmaceutical-related disciplines. Each individual trial involves contractual, regulatory, and.

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