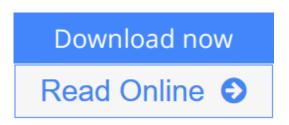


## Clinical Trials and Human Research: A Practical Guide to Regulatory Compliance

By Fay A. Rozovsky, Rodney K. Adams



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This easy-to-read reference book provides a practical approach for dealing with the legal and regulatory compliance issues involved in human research. Covering a broad range of topics, such as consent, confidentiality, subject recruitment and selection, the role of the investigator and Institutional Review Board, it offers timely and useful strategies for achieving regulatory compliance while reducing liability. In addition, insurance, quality management, accreditation, and risk management are topics examined in the book. The practical insights found in this volume are not found in other books on the subject. *Clinical Trials and Human Research* is a practical tool to help anyone involved in clinical research.

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#### Praise for Clinical Trials and Human Research

"I would recommend the book for anyone new to the Institutional Review Board process, regardless of their clinical or administrative background or responsibilities. It is a valuable resource to orient risk managers, clinicians, and administrators to the clinical trial process and pitfalls."

— Deborah Boyd, M.S., R.H.I.A., C.P.H.R.M., senior risk management consultant, Zurich North America, Atlanta, Georgia

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#### About the Author

**Fay A. Rozovsky**, Fay A. Rozovsky, J.D., M.P.H. is an affiliate associate professor in the Department of Legal Medicine at Virginia Commonwealth University's School of Medicine. Ms. Rozovsky has served as the administrator of an Institutional Review Board and is a member of human research committees in the United States and Canada.

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