

Basic Principles of Clinical Research and Methodology



Institute of Clinical Research (India)

**D-41, Defence Colony
New Delhi**



Basic Principles Of Clinical Research And Methodology 1st Edition

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Basic Principles Of Clinical Research And Methodology 1st Edition:

Basic Principles of Clinical Research and Methodology SK Gupta, 2007-05-30 **A Textbook of Clinical Research and Pharmacovigilance** KPR Chowdary, 2025-06-01 This book describes all concepts practices methods and regulatory guidelines related to clinical research clinical trials and pharmacovigilance in a simple lucid and easily understandable manner and covers the entire syllabus prescribed by Pharmacy Council of India PCI New Delhi for Pharm D and M Pharm courses The book provides a comprehensive knowledge of various aspects such as drug development and approval process pharmacological and toxicological approaches and methods pharmaceutical dosage form approaches for drug development clinical approaches and clinical trials phases types designs and statistical tests of clinical trials regulatory aspects GCP as per ICH WHO ICMR Schedule Y and regulatory environment in US Europe and India in 20 chapters Special emphasis is given to Pharmacovigilance methods and Pharmacovigilance programme of India PvPI The book provides a comprehensive knowledge of all aspects of clinical research clinical trials GCP guidelines and Pharmacovigilance as per the requirements of clinical research industry and personnel The subject is presented in a simple lucid and easily understandable way in logical flow for the benefit of pharmacy students as well as industry persons Latest practices and regulatory guidelines are included and hence the book provides updated knowledge This book is ideal for Pharm D M Pharm and PhD students of Pharmacy and also for research personnel involved in clinical research Contents 1 Drug Discovery Development and Approval Process An Overview 2 Approaches to Drug Discovery Pharmacological and Toxicological 3 Drug Characterization Preformulation and Dosage Form Development 4 The Investigational New Drug IND Application and New Drug Application NDA 5 Clinical Development of Drugs Introduction and Evolution of Clinical Research 6 Clinical Research Methodology Phases Types Designs and Statistical Concepts of Clinical Trials 7 Clinical Trials Research in India Clinical Trial Phases Process Documentation and Regulations 8 Methods of Post Marketing Surveillance PMS 9 Abbreviated New Drug Application ANDA Submissions 10 Guidelines and Principles of Good Clinical Practices ICH WHO 11 Comparison of Clinical Trial Regulations in India Europe and USA 12 Challenges in the Implementation of GCP Guidelines 13 Ethical Guidelines in Clinical Research 14 Composition Role and Responsibilities of Institutional Ethics Committee IEC in Clinical Trials 15 Regulatory Environment in US India and Europe 16 Role and Responsibilities of Clinical Trial Personnel as per GCP 17 Designing of Clinical Study Documents and Informed Consent Process 18 Data Management in Clinical Research 19 Safety Monitoring in Clinical Trials 20 Pharmacovigilance *ESSENTIALS OF BIOSTATISTICS AND RESEARCH METHODOLOGY* INDRANIL SAHA, BOBBY PAUL, 2023-04-14 THIS TEXTBOOK IS A COMPREHENSIVE USER FRIENDLY AND EASY TO READ RESOURCE ON BIOSTATISTICS AND RESEARCH METHODOLOGY IT IS MEANT FOR UNDERGRADUATE AND POSTGRADUATE MEDICAL STUDENTS AND ALLIED BIOMEDICAL SCIENCES HEALTH RESEARCHERS RESEARCH SUPERVISORS AND FACULTY MEMBERS MAY FIND IT USEFUL AS A REFERENCE BOOK Fundamentals of Clinical

Trials Lawrence M. Friedman, Curt D. Furberg, David L. DeMets, 2010-09-09 The clinical trial is the most definitive tool for evaluation of the applicability of clinical research. It represents a key research activity with the potential to improve the quality of health care and control costs through careful comparison of alternative treatments. It has been called on many occasions the gold standard against which all other clinical research is measured. Although many clinical trials are of high quality, a careful reader of the medical literature will notice that a large number have deficiencies in design, conduct, analysis, presentation, and/or interpretation of results. Improvements have occurred over the past few decades, but too many trials are still conducted without adequate attention to its fundamental principles. Certainly, numerous studies could have been upgraded if the authors had had a better understanding of the fundamentals. Since the publication of the first edition of this book, a large number of other texts on clinical trials have appeared, most of which are indicated here. Several of them, however, discuss only specific issues involved in clinical trials. Additionally, many are no longer current. The purpose of this fourth edition is to update areas in which major progress has been made since the publication of the third edition. We have revised most chapters considerably and added one on ethical issues.

Bayesian Analysis with R for Drug Development
Harry Yang, Steven Novick, 2019-06-26 Drug development is an iterative process. The recent publications of regulatory guidelines further entail a lifecycle approach. Blending data from disparate sources, the Bayesian approach provides a flexible framework for drug development. Despite its advantages, the uptake of Bayesian methodologies is lagging behind in the field of pharmaceutical development. Written specifically for pharmaceutical practitioners, *Bayesian Analysis with R for Drug Development: Concepts, Algorithms, and Case Studies* describes a wide range of Bayesian applications to problems throughout pre-clinical, clinical, and Chemistry, Manufacturing, and Control (CMC) development. Authored by two seasoned statisticians in the pharmaceutical industry, the book provides detailed Bayesian solutions to a broad array of pharmaceutical problems. Features: Provides a single source of information on Bayesian statistics for drug development. Covers a wide spectrum of pre-clinical, clinical, and CMC topics. Demonstrates proper Bayesian applications using real-life examples. Includes easy-to-follow R code with Bayesian Markov Chain Monte Carlo performed in both JAGS and Stan. Bayesian software platforms. Offers sufficient background for each problem and detailed description of solutions suitable for practitioners with limited Bayesian knowledge. Harry Yang, Ph.D., is Senior Director and Head of Statistical Sciences at AstraZeneca. He has 24 years of experience across all aspects of drug research and development and extensive global regulatory experiences. He has published 6 statistical books, 15 book chapters, and over 90 peer-reviewed papers on diverse scientific and statistical subjects, including 15 joint statistical works with Dr. Novick. He is a frequent invited speaker at national and international conferences. He also developed statistical courses and conducted training at the FDA and USP, as well as Peking University. Steven Novick, Ph.D., is Director of Statistical Sciences at AstraZeneca. He has extensively contributed statistical methods to the biopharmaceutical literature. Novick is a skilled Bayesian computer programmer and is frequently invited to speak at

conferences having developed and taught courses in several areas including drug combination analysis and Bayesian methods in clinical areas Novick served on IPAC RS and has chaired several national statistical conferences *Current Catalog* National Library of Medicine (U.S.),1980 First multi year cumulation covers six years 1965 70 **Principles and Practice of Clinical Research** John I. Gallin, Frederick P Ognibene, Laura Lee Johnson, 2017-11-17 Principles and Practice of Clinical Research Fourth Edition has been thoroughly revised to provide a comprehensive look at both the fundamental principles and expanding practice of clinical research New to this edition of this highly regarded reference authors have focused on examples that broadly reflect clinical research on a global scale while including a discussion of international regulations studies and implications In addition to key topics such as bioethics clinical outcome data cultural diversity protocol guidelines and omic platforms this edition contains new chapters devoted to electronic health records and information resources for clinical researchers as well as the many opportunities associated with big data Covering a vast number of topics and practical advice for both novice and advanced clinical investigators this book is a highly relevant and essential resource for all those involved in conducting research Features input from experts in the field dedicated to translating scientific research from bench to bedside and back Provides expanded coverage of global clinical research Contains hands on practical suggestions illustrations and examples throughout Includes new chapters on the international regulation of drugs and biologics the emergence of the important role of comparative effectiveness research and how to identify clinical risks and manage patient safety in a clinical research setting *Principles of Research Methodology* Phyllis G. Supino, Jeffrey S. Borer, 2012-06-22 Principles of Research Methodology A Guide for Clinical Investigators is the definitive comprehensive guide to understanding and performing clinical research Designed for medical students physicians basic scientists involved in translational research and other health professionals this indispensable reference also addresses the unique challenges and demands of clinical research and offers clear guidance in becoming a more successful member of a medical research team and critical reader of the medical research literature The book covers the entire research process beginning with the conception of the research problem to publication of findings Principles of Research Methodology A Guide for Clinical Investigators comprehensively and concisely presents concepts in a manner that is relevant and engaging to read The text combines theory and practical application to familiarize the reader with the logic of research design and hypothesis construction the importance of research planning the ethical basis of human subjects research the basics of writing a clinical research protocol and scientific paper the logic and techniques of data generation and management and the fundamentals and implications of various sampling techniques and alternative statistical methodologies Organized in thirteen easy to read chapters the text emphasizes the importance of clearly defined research questions and well constructed hypothesis reinforced throughout the various chapters for informing methods and in guiding data interpretation Written by prominent medical scientists and methodologists who have extensive personal experience in biomedical investigation and in teaching

key aspects of research methodology to medical students physicians and other health professionals the authors expertly integrate theory with examples and employ language that is clear and useful for a general medical audience A major contribution to the methodology literature Principles of Research Methodology A Guide for Clinical Investigators is an authoritative resource for all individuals who perform research plan to perform it or wish to understand it better

Handbook of Research Methods in Clinical Psychology Philip C. Kendall, James N. Butcher, Grayson N. Holmbeck, 1999-01-25 Revising a successful handbook which covers all major areas of clinical research this edition covers all the relevant aspects of and tactics for conducting clinical research American Book Publishing Record ,2006

Canadian Medical Association Journal Canadian Medical Association, 1923 Drug Intelligence & Clinical Pharmacy ,1981 The United States Catalog Mary Burnham, Carol Hurd, 1928 **International Record of Medicine and General Practice Clinics** Edward Swift Dunster, Frank Pierce Foster, James Bradbridge Hunter, Charles Eucharist de Medicis Sajous, Gregory Stragnell, Henry J. Klaunberg, Félix Martí-Ibáñez, 1900 British Medical Journal ,1930 **Subject Guide to Books in Print** ,1997 **International Books in Print** ,1997 Stroke Moira A. Banks, 1986 Book Review Index ,2003 Vols 8 10 of the 1965 1984 master cumulation constitute a title index Pharmaceutical Journal ,1921

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