CLINICAL TRIALS

Design, Conduct, and Analysis

SECOND EDITION

CURTIS L. MEINERT

Jørn Olsen, Rodolfo Saracci, Dimitrios Trichopoulos

ClinicalTrials Curtis L. Meinert PhD, 2012-02-17 First published in 1986 this landmark text is the definitive guide to clinical trials written by one of the leading experts in the field This fully updated second edition continues to be the most authoritative reference text on randomized clinical trials It contains a wealth of practical information on the design conduct and analysis of both single center and multicenter trials No other book on clinical trials offers as much detail on such issues as sample size calculation stratification and randomization data systems design development of consent forms publication policies preparation of funding requests and reporting procedures While the basics of design conduct and analysis of clinical trials remain the same there have been significant changes since the first edition of Clinical Trials was published two decades ago In this new edition the author discusses the refinements and improvements made to methods and procedures changes in the policies and guidelines underlying trials as well as requirements for registration of trials He also discusses current practices for data sharing for gender representation for treatment effects monitoring and for ethical standards of clinical trials The importance of the randomized controlled trial has grown significantly over time and they are now the cornerstone of all evidence based medicine Still rich in tables checklists charts and other resources for the trialist the second edition of Clinical Trials is an indispensable reference for clinicians biostaticians epidemiologists and anyone involved in the design and Clinical Trials Curtis L. Meinert, Susan Tonascia, 1986 The definitive reference work on implementation of a clinical trial clinical trials this book presents a wealth of detailed practical information on the design conduct and analysis of both single center and multicenter trials No other book on clinical trials offers as much detail as Meinert does on such issues as sample size calculation stratification and randomization data systems design consent form development publication policies preparation of funding requests and reporting procedures ClinicalTrials Curtis L. Meinert, 2012-03-27 The classic definitive guide to the design conduct and analysis of randomized clinical trials **Biostatistics and Epidemiology** Sylvia Wassertheil-Smoller, Jordan Smoller, 2015-02-05 Since the publication of the first edition Biostatistics and Epidemiology has attracted loyal readers from across specialty areas in the biomedical community Not only does this textbook teach foundations of epidemiological design and statistical methods but it also includes topics applicable to new areas of research Areas covered in the fourth edition include a new chapter on risk prediction risk reclassification and evaluation of biomarkers new material on propensity analyses and a vastly expanded chapter on genetic epidemiology which is particularly relevant to those who wish to understand the epidemiological and statistical aspects of scientific articles in this rapidly advancing field Biostatistics and Epidemiology was written to be accessible for readers without backgrounds in mathematics It provides clear explanations of underlying principles as well as practical guidelines of how to do it and how to interpret it Key features include a philosophical and logical explanation at the beginning of the book subsections that can stand alone or serve as reference cross referencing recommended reading and appendices covering sample calculations for various statistics in the

Clinical Trials Steven Piantadosi, 2024-04-03 Comprehensive resource presenting methods essential in planning text designing conducting analyzing and interpreting clinical trials The Fourth Edition of Clinical Trials builds on the text s reputation as a straightforward detailed and authoritative presentation of quantitative methods for clinical trials discussing principles of design for various types of clinical trials and elements of planning the experiment assembling a study cohort assessing data and reporting results Each chapter contains an introduction and summary to reinforce key points Discussion questions stimulate critical thinking and help readers understand how they can apply their newfound knowledge Written by a highly qualified author with significant experience in the field the Fourth Edition of Clinical Trials approaches the topic with Problems that may arise during a trial and accompanying common sense solutions Design alternatives for addressing many questions in the rapeutic development Statistical principles with new and provocative topics such as generalizing results operating characteristics trial issues during the COVID 19 pandemic and more Alternative medicine ethics middle development comparative studies adaptive designs and clinical trials using point of care data Revamped exercise sets updated and extensive references new material on endpoints and the developmental pipeline and revisions of numerous sections tables and figures Standing out due to its accessible and broad coverage of statistical design methods which are the building blocks of clinical trials and medical research Clinical Trials is an essential learning aid on the subject for undergraduate and graduate clinical trials courses Thomas' Hematopoietic Cell Transplantation, 2 Volume Set Stephen J. Forman, Robert S. Negrin, Joseph H. Antin, Frederick R. Appelbaum, 2016-12-27 Fully revised for the fifth edition this outstanding reference on bone marrow transplantation is an essential field leading resource Extensive coverage of the field from the scientific basis for stem cell transplantation to the future direction of research Combines the knowledge and expertise of over 170 international specialists across 106 chapters Includes new chapters addressing basic science experiments in stem cell biology immunology and tolerance Contains expanded content on the benefits and challenges of transplantation and analysis of the impact of new therapies to help clinical decision making Includes a fully searchable Wiley Digital Edition with downloadable figures linked references and more References for this new edition are online only accessible via the Wiley Digital Edition code printed inside the front cover or at www wiley com go forman hematopoietic

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Organized for quick access to key topics and replete with practical examples the book describes a variety of research designs and statistical methods and explains how to choose the best design for a particular project Research implementation including regulatory issues and grant writing is also covered The book opens with a section on the basics of research design discussing the many ways in which studies can be organized executed and evaluated The second section is devoted to statistics and explains how to choose the correct statistical approach and reviews the varieties of data types descriptive and inferential statistics methods for demonstrating associations hypothesis testing and prediction specialized methods and considerations in epidemiological studies and measure construction The third section covers implementation including how to develop a grant application step by step the project budget and the nuts and bolts of the timely and successful completion of a research project and documentation of findings procedural manuals and case report forms collecting managing and securing data operational structure and ongoing monitoring and evaluation and ethical and regulatory concerns in research with human subjects With a concise presentation of the essentials for successful research the Handbook for Clinical Research is a valuable addition to the library of any student research professional or clinician interested in expanding the knowledge base of his or her field Key Features Delivers the essential elements practical insights and trade secrets for ensuring successful research design analysis and implementation Presents the nuts and bolts of statistical analysis Organized for quick access to a wealth of information Replete with practical examples of successful research designs from single case designs to meta analysis and how to achieve them Addresses research implementation including regulatory issues and grant writing

Handbook of Drug Abuse Prevention Zili Sloboda, William J. Bukoski, 2007-08-06 There are a number of reasons why this book is important First there is no one source that summarizes what we know about the prevention of substance abuse from the research field so the book serves as a repository of accumulated knowledge on prevention theory intervention design and development and prevention research methodology Second as an evolving field prevention science has only begun to assert itself in both the arenas of practice and policy The formation of the Society for Prevention Research in 1991 was the first recognition that a science of prevention existed and required a separate forum to present the rapidly growing content of the field for dis cussion and review Finally there is a need to establish a baseline a reference point against which progress in the field of prevention science can be assessed This book serves all of these purposes The idea for this book grew from the observation in the early 1990s that after decades of attempts to develop effective interventions to prevent drug use among children and adolescents in the United States that we were finally having success particularly in addressing the initiation of use These successes are the result of research that has provided a better understanding of the factors and processes associated with the onset of substance use

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