

Chapman & Hall/CRC Biostatistics Series

Bayesian Adaptive Methods for Clinical Trials



Scott M. Berry
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CRC Press
Taylor & Francis Group

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Bayesian Adaptive Methods For Clinical Trials

Biostatistics

Rachel Sandford



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Bayesian Adaptive Methods for Clinical Trials Scott M. Berry, Bradley P. Carlin, J. Jack Lee, Peter Muller, 2010-07-19
Already popular in the analysis of medical device trials adaptive Bayesian designs are increasingly being used in drug development for a wide variety of diseases and conditions from Alzheimer's disease and multiple sclerosis to obesity diabetes hepatitis C and HIV Written by leading pioneers of Bayesian clinical trial designs Bayesian Adaptive Clinical Trial Design
Guosheng Yin, 2012-01-03 A balanced treatment of the theories methodologies and design issues involved in clinical trials using statistical methods There has been enormous interest and development in Bayesian adaptive designs especially for early phases of clinical trials However for phase III trials frequentist methods still play a dominant role through controlling type I and type II errors in the hypothesis testing framework From practical perspectives Clinical Trial Design Bayesian and Frequentist Adaptive Methods provides comprehensive coverage of both Bayesian and frequentist approaches to all phases of clinical trial design Before underpinning various adaptive methods the book establishes an overview of the fundamentals of clinical trials as well as a comparison of Bayesian and frequentist statistics Recognizing that clinical trial design is one of the most important and useful skills in the pharmaceutical industry this book provides detailed discussions on a variety of statistical designs their properties and operating characteristics for phase I II and III clinical trials as well as an introduction to phase IV trials Many practical issues and challenges arising in clinical trials are addressed Additional topics of coverage include Risk and benefit analysis for toxicity and efficacy trade offs Bayesian predictive probability trial monitoring Bayesian adaptive randomization Late onset toxicity and response Dose finding in drug combination trials Targeted therapy designs The author utilizes cutting edge clinical trial designs and statistical methods that have been employed at the world's leading medical centers as well as in the pharmaceutical industry The software used throughout the book is freely available on the book's related website equipping readers with the necessary tools for designing clinical trials Clinical Trial Design is an excellent book for courses on the topic at the graduate level The book also serves as a valuable reference for statisticians and biostatisticians in the pharmaceutical industry as well as for researchers and practitioners who design conduct and monitor clinical trials in their everyday work *Design and Analysis of Clinical Trials for Predictive Medicine* Shigeyuki Matsui, Marc Buyse, Richard Simon, 2015-03-19 *Design and Analysis of Clinical Trials for Predictive Medicine* provides statistical guidance on conducting clinical trials for predictive medicine It covers statistical topics relevant to the main clinical research phases for developing molecular diagnostics and therapeutics from identifying molecular biomarkers using DNA microarrays to confirming *Statistical Design and Analysis of Clinical Trials* Weichung Joe Shih, Joseph Aisner, 2015-07-28 *Statistical Design and Analysis of Clinical Trials Principles and Methods* concentrates on the biostatistics component of clinical trials Developed from the authors' courses taught to public health and medical students residents and fellows during the past 15 years the text shows how biostatistics in clinical trials is an integration of many Adaptive Design Methods in Clinical

Trials Shein-Chung Chow, Mark Chang, 2006-11-16 Although adaptive design methods are flexible and useful in clinical research little or no regulatory guidelines are available One of the first books on the topic Adaptive Design Methods in Clinical Trials presents the principles and methodologies in adaptive design and analysis that pertain to adaptations made to trial or statistical procedures

Sample Size Calculations for Clustered and Longitudinal Outcomes in Clinical Research Chul Ahn, Moonseoung Heo, Song Zhang, 2014-12-09 This book explains how to determine sample size for studies with correlated outcomes which are widely implemented in medical epidemiological and behavioral studies For clustered studies the authors provide sample size formulas that account for variable cluster sizes and within cluster correlation For longitudinal studies they present sample size formulas that account for within subject correlation among repeated measurements and various missing data patterns For multiple levels of clustering the authors describe how randomization impacts trial administration analysis and sample size requirement

Design & Analysis of Clinical Trials for Economic Evaluation & Reimbursement Iftekhar Khan, 2015-11-11 Economic evaluation has become an essential component of clinical trial design to show that new treatments and technologies offer value to payers in various healthcare systems Although many books exist that address the theoretical or practical aspects of cost effectiveness analysis this book differentiates itself from the competition by detailing

Noninferiority Testing in Clinical Trials Tie-Hua Ng, 2014-12-01 Take Your NI Trial to the Next Level Reflecting the vast research on noninferiority NI designs from the past 15 years Noninferiority Testing in Clinical Trials Issues and Challenges explains how to choose the NI margin as a small fraction of the therapeutic effect of the active control in a clinical trial Requiring no prior knowledge of NI testing the book is easily accessible to both statisticians and nonstatisticians involved in drug development With over 20 years of experience in this area the author introduces the basic elements of the NI trials one at a time in a logical order He discusses issues with estimating the effect size based on historical placebo control trials of the active control The book covers fundamental concepts related to NI trials such as assay sensitivity constancy assumption discounting and preservation It also describes patient populations three arm trials and the equivalence of three or more groups

Multiregional Clinical Trials for Simultaneous Global New Drug Development Joshua Chen, Hui Quan, 2016-04-21 In a global clinical development strategy multiregional clinical trials MRCTs are vital in the development of innovative medicines Multiregional Clinical Trials for Simultaneous Global New Drug Development presents a comprehensive overview on the current status of conducting MRCTs in clinical development International experts from academia in

Cancer Clinical Trials Stephen L. George, Xiaofei Wang, Herbert Pang, 2016-08-19 Cancer Clinical Trials Current and Controversial Issues in Design and Analysis provides statisticians with an understanding of the critical challenges currently encountered in oncology trials Well known statisticians from academic institutions regulatory and government agencies such as the U S FDA and National Cancer Institute and the pharmaceutical industry share their extensive experiences in cancer clinical trials and present examples taken from actual trials The book

covers topics that are often perplexing and sometimes controversial in cancer clinical trials Most of the issues addressed are also important for clinical trials in other settings After discussing general topics the book focuses on aspects of early and late phase clinical trials It also explores personalized medicine including biomarker based clinical trials adaptive clinical trial designs and dynamic treatment regimes [Advances in Clinical Trial Biostatistics](#) Nancy L. Geller,2003-10-21 From aspects of early trials to complex modeling problems [Advances in Clinical Trial Biostatistics](#) summarizes current methodologies used in the design and analysis of clinical trials Its chapters contributed by internationally renowned methodologists experienced in clinical trials address topics that include Bayesian methods for phase I clinical trials adaptive two stage clinical trials and the design and analysis of cluster randomization trials trials with multiple endpoints and therapeutic equivalence trials Other discussions explore Bayesian reporting methods incorporating compliance in treatment evaluation and statistical issues emerging from clinical trials in HIV infection **Modern Approaches to Clinical Trials Using SAS** Sandeep Menon,Richard C. Zink,2015-12-09 Get the tools you need to use SAS in clinical trial design Unique and multifaceted [Modern Approaches to Clinical Trials Using SAS Classical Adaptive and Bayesian Methods](#) edited by Sandeep M Menon and Richard C Zink thoroughly covers several domains of modern clinical trial design classical group sequential adaptive and Bayesian methods that are applicable to and widely used in various phases of pharmaceutical development Written for biostatisticians pharmacometricians clinical developers and statistical programmers involved in the design analysis and interpretation of clinical trials as well as students in graduate and postgraduate programs in statistics or biostatistics the book touches on a wide variety of topics including dose response and dose escalation designs sequential methods to stop trials early for overwhelming efficacy safety or futility Bayesian designs that incorporate historical data adaptive sample size re estimation adaptive randomization to allocate subjects to more effective treatments and population enrichment designs Methods are illustrated using clinical trials from diverse therapeutic areas including dermatology endocrinology infectious disease neurology oncology and rheumatology Individual chapters are authored by renowned contributors experts and key opinion leaders from the pharmaceutical medical device industry or academia Numerous real world examples and sample SAS code enable users to readily apply novel clinical trial design and analysis methodologies in practice [Group Sequential and Adaptive Methods for Clinical Trials](#) Christopher Jennison,Bruce W. Turnbull,2015-12-08 In clinical trials monitoring accumulating data at regular intervals is essential for balancing ethical and financial considerations against scientific rigor This comprehensive second edition reflects the remarkable evolution in adaptive clinical trial methodology over the past two decades Since publication of the first edition these approaches have transformed from theoretical concepts to widely accepted practices now endorsed by regulatory authorities including formal FDA guidance Features Eleven entirely new chapters including optimal design construction delayed pipeline data accommodation and multiple comparison procedures Eight dedicated chapters on adaptive methods including sample size re assessment seamless Phase II III trials multi arm

multi stage trials and enrichment designs Exact methods for binary and Poisson data with applications to vaccine trials Expanded discussion of binding versus non binding futility boundaries for enhanced decision making A chapter with a thoroughly updated review of Bayesian approaches to sequential trial design and analysis Incorporates response adaptive treatment assignment in group sequential designs Practical computational methods and software guidance for implementing the techniques This definitive resource serves statisticians clinical researchers pharmaceutical scientists and regulatory professionals involved in clinical trial design and analysis The text balances theoretical foundations with practical implementation guidance making it invaluable for both experienced practitioners and those new to adaptive trial methodology With twice the content of the first edition Group Sequential and Adaptive Methods for Clinical Trials Second Edition provides the comprehensive understanding needed to design more efficient ethical and scientifically rigorous clinical trials ultimately accelerating the delivery of effective treatments to patients

A Bayesian Adaptive Design for 2-drug Combination Phase I Clinical Trials with Ordinal Toxicity Outcomes Lu-May Chiang,2007 **Bayesian Adaptive Methods to Incorporate Preclinical Data Into Phase I Clinical Trials** Haiyan Zheng,2019 **Bayesian Statistics 3** J. M. Bernardo,1988 The field of statistics has undergone rapid and wide development during the past two decades and the Bayesian approach to statistics has provided both a general framework and a creative stimulus for all aspects of this development This volume describes the work presented at the Third Valencia International Meeting on Bayesian Statistics the main source of information and communication about the current state of knowledge and research in Bayesian statistics throughout the world The research presented which encompasses both invited papers and selected contributed papers has had a profound effect on the foundations of statistical inference and probability statistical theory and methodology and the applications of statistics in science technology medicine business law and public policy The contributors to this volume form a virtual Who's Who in the area of Bayesian statistics

Bayesian Computational Methods and Applications R. Carter Hill,1996 **Bayesian Adaptive Methods for Phase I Clinical Trials** Ruitao Lin,2017-01-26 **Proceedings of "Current Topics in Biostatistics and Epidemiology"** Daniel Seigel,1982 **Statistical Methods for Dose-Finding Experiments** Sylvie Chevret,2006-05-26 Dose finding experiments define the safe dosage of a drug in development in terms of the quantity given to a patient Statistical methods play a crucial role in identifying optimal dosage Used appropriately these methods provide reliable results and reduce trial duration and costs In practice however dose finding is often done poorly with widely used conventional methods frequently being unreliable leading to inaccurate results However there have been many advances in recent years with new statistical techniques being developed and it is important that these new techniques are utilized correctly Statistical Methods for Dose Finding Experiments reviews the main statistical approaches for dose finding in phase I II clinical trials and presents practical guidance on their correct use Includes an introductory section summarizing the essential concepts in dose finding Contains a section on algorithm based approaches such as the

traditional 3 3 design and a section on model based approaches such as the continual reassessment method Explains fundamental issues such as how to stop trials early and how to cope with delayed or ordinal outcomes Discusses in detail the main websites and software used to implement the methods Features numerous worked examples making use of real data Statistical Methods for Dose Finding Experiments is an important collaboration from the leading experts in the area Primarily aimed at statisticians and clinicians working in clinical trials and medical research there is also much to benefit graduate students of biostatistics

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