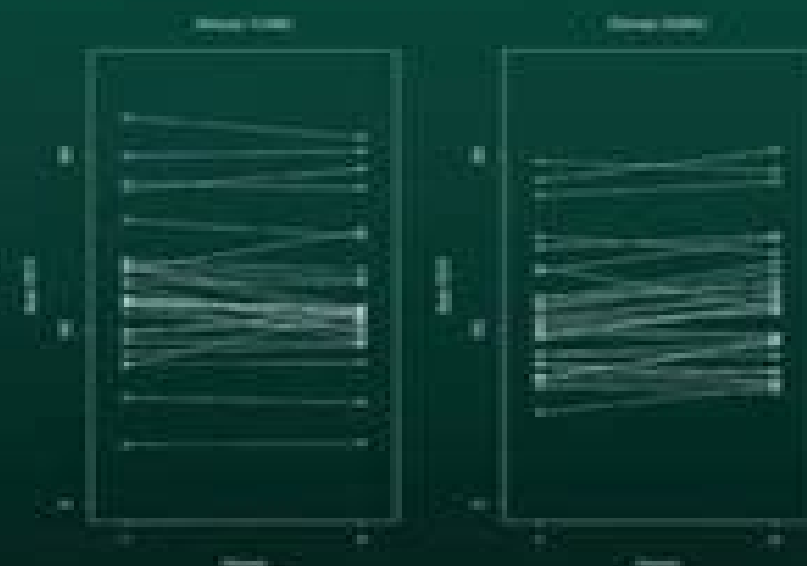


Design and Analysis of Cross-Over Trials

Third Edition



Byron Jones
Michael G. Kenward

Design And Analysis Of Cross Over Trials

Lisa Doreen Fairchild



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Cross-over Trials in Clinical Research Stephen S. Senn, 2003-07-25 Cross over trials are an important class of design used in the pharmaceutical industry and medical research and their use continues to grow Cross over Trials in Clinical Research Second Edition has been fully updated to include the latest methodology used in the design and analysis of cross over trials It includes more background material greater coverage of important statistical techniques including Bayesian methods and discussion of analysis using a number of statistical software packages Comprehensive coverage of the design and analysis of cross over trials Each technique is carefully explained and the mathematics is kept to a minimum Features many real and original examples taken from the author's vast experience Includes discussion of analysis using SAS S Plus and GenStat StatXact and Excel Written in a style suitable for statisticians and physicians alike Computer programs to accompany the examples in the book can be downloaded from the Web Primarily aimed at statisticians and researchers working in the pharmaceutical industry the book will also appeal to physicians involved in clinical research and students of medical statistics

The Design and Analysis of Crossover Trials with Many Periods Michael Edward Smith, 1994

Design and Analysis of Cross-Over Trials, Second Edition Byron Jones, Michael G. Kenward, 2003-03-12 The first edition of Design and Analysis of Cross Over Trials quickly became the standard reference on the subject and has remained so for more than 12 years In that time however the use of cross over trials has grown rapidly particularly in the pharmaceutical arena and researchers have made a number of advances in both the theory and methods applicable to these trials Completely revised and updated the long awaited second edition of this classic text retains its predecessor's careful balance of theory and practice while incorporating new approaches more data sets and a broader scope Enhancements in the second edition include A new chapter on bioequivalence Recently developed methods for analyzing longitudinal continuous and categorical data Real world examples using the SAS system A comprehensive catalog of designs datasets and SAS programs available on a companion Web site at www.crcpress.com The authors exposition gives a clear unified account of the design and analysis of cross over trials from a statistical perspective along with their methodological underpinnings With SAS programs and a thorough treatment of design issues Design and Analysis of Cross Over Trials Second Edition sets a new standard for texts in this area and undoubtedly will be of direct practical value for years to come

Optimal Crossover Designs Mausumi Bose, Aloke Dey, 2009 This monograph presents a comprehensive and up to date account of the developments in optimality aspects of crossover designs Crossover designs are immensely useful in various areas of human investigation including agriculture animal nutrition clinical trials pharmaceutical studies biological assays weather modification experiments sensory evaluation of food products and learning experiments Research on the optimality aspects of crossover designs has developed only in the last three decades and it has now emerged as a potential field for further investigation This book is the first comprehensive treatise on this subject It covers optimal crossover designs at length by consolidating vast amounts of

material from the literature and includes many recent and deep results It is expected that this book will not only provide a one stop reference for the available results but also encourage further research in this area of substantial practical relevance

Clinical Trials Handbook Shayne Cox Gad,2009-06-17 Best practices for conducting effective and safe clinical trials Clinical trials are arguably the most important steps in proving drug effectiveness and safety for public use They require intensive planning and organization and involve a wide range of disciplines data management biostatistics pharmacology toxicology modeling and simulation regulatory monitoring ethics and particular issues for given disease areas Clinical Trials Handbook provides a comprehensive and thorough reference on the basics and practices of clinical trials With contributions from a range of international authors the book takes the reader through each trial phase technique and issue Chapters cover every key aspect of preparing and conducting clinical trials including Interdisciplinary topics that have to be coordinated for a successful clinical trial Data management and adverse event reporting systems Biostatistics pharmacology and toxicology Modeling and simulation Regulatory monitoring and ethics Particular issues for given disease areas cardiology oncology cognitive dementia dermatology neuroscience and more With unique information on such current issues as adverse event reporting AER systems adaptive trial designs and crossover trial designs Clinical Trials Handbook will be a ready reference for pharmaceutical scientists statisticians researchers and the many other professionals involved in drug development

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Biostatistics in Clinical Trials Carol K. Redmond, Theodore Colton, 2001-04-25 The second volume in the Wiley reference series in Biostatistics Featuring articles from the prestigious Encyclopedia of Biostatistics many of which have been fully revised and updated to include recent developments Biostatistics in Clinical Trials also includes up to 25% newly commissioned material reflecting the latest thinking in Bayesian methods Benefit risk assessment Cost effectiveness Ethics Fraud With exceptional contributions from leading experts in academia government and industry Biostatistics in Clinical Trials has been designed to complement existing texts by providing extensive up to date coverage and introducing the reader to the research literature Offering comprehensive coverage of all aspects of clinical trials Biostatistics in Clinical Trials Includes concise definitions and introductions to numerous concepts found in current literature Discusses the software and textbooks available Uses extensive cross references helping to facilitate further research and enabling the reader to locate

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major forum for the interplay of statistics and computer sciences with contributions from many well known scientists all over the world The scientific programme of COMPSTAT 96 covers all aspects of this interplay from user experiences and evaluation of software through the development and implementation of new statistical ideas All papers presented belong to one of the three following categories Statistical methods preferable new ones that require a substantial use of computing Computer environments tools and software useful in statistics Applications of computational statistics in areas of substantial interest environment health industry biometrics etc

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