

Ting

Statistics for Biology and Health

Naitee Ting
Editor

Dose Finding in Drug Development

Dose Finding in Drug Development



 **Springer**

Dose Finding In Drug Development Dose Finding In Drug Development

J. Venitz, W. Sittner



Dose Finding In Drug Development Dose Finding In Drug Development:

Dose Finding in Drug Development Naitee Ting, 2006-12-29 This book emphasizes dose selection issues from a statistical point of view. It presents statistical applications in the design and analysis of dose response studies. The importance of this subject can be found from the International Conference on Harmonization ICH E4 Guidance document. Establishing the dose response relationship is one of the most important activities in developing a new drug. A clinical development program for a new drug can be broadly divided into four phases, namely Phases I, II, III, and IV. Phase I clinical trials are designed to study the clinical pharmacology. Information gained from these studies will help in designing Phase II studies. Dose response relationships are usually studied in Phase II. Phase III clinical trials are large scale long term studies. These studies serve to confirm findings from Phases I and II. Results obtained from Phases I, II, and III clinical trials would then be documented and submitted to regulatory agencies for drug approval. In the United States, reviews from Food and Drug Administration (FDA) review these documents and make a decision to approve or to reject this New Drug Application (NDA). If the new drug is approved, then Phase IV studies can be started. Phase IV clinical trials are also known as postmarketing studies.

Dose Finding and Beyond in Biopharmaceutical Development Jingjing Ye, Ding-Geng Chen, Wen Zhou, Qiqi Deng, Joseph C. Cappelleri, 2024-10-28 This book covers topics in 2 parts: 1. Review of FDA Guidance 2. Novel Designs and Analyses. While covering basic principles of dose finding, this book details advancements made in drug development. Finding the right dose is one of the most important objectives in new drug development. In Phase I clinical development, one of the objectives is to escalate test doses from low to high. The low doses should be safe, then escalate up to the maximally tolerable dose (MTD). Phase II clinical trials then lower test doses to the minimal efficacious dose (MinED). Dose range of a study drug can be thought of as the doses between MinED and MTD. From this dose range, one or a few doses are selected for Phase III confirmation. In practice, dose finding is a very difficult in every phase of clinical development for new drugs. The editors brought distinguished researchers and practitioners in biopharmaceuticals and universities to discuss the statistical procedures, useful methods, and their novel applications in dose finding. The chapters in the book present emerging topics in dose finding and related interdisciplinary areas. This timely book is a valuable resource to stimulate the development of this growing and exciting field in drug development.

Appropriate Dose Selection - How to Optimize Clinical Drug Development J. Venitz, W. Sittner, 2007-03-06 Optimal dose individualization has become more important in improving clinical efficacy and safety given the variability in drug response, e.g. due to concurrent illnesses or co-medications. Therefore, the role of optimal dose finding in early clinical drug development so as to maximize successful clinical use is emphasized. The continued use of biomarkers based on the known pharmacology of the drug and/or biology of the underlying disease along with exposure response evaluation throughout all phases of drug development can quantitatively integrate clinical pharmacology knowledge, provide early proof of concept, and help in rational dose selection and rational drug product

labeling for clinical use **Dose Finding In Drug Development** Ed Naitee Ting, 2008-12-01 *Appropriate Dose Selection - How to Optimize Clinical Drug Development* J. Venitz, W. Sittner, 2009-09-02 Optimal dose individualization has become more important in improving clinical efficacy and safety This is due in part to the variability in drug response Therefore the role of optimal dose finding in early clinical drug development so as to maximize successful clinical use is emphasized This book reviews innovative methods tools and examples of rational drug development strategies particularly for novel oncological agents **Handbook of Methods for Designing, Monitoring, and Analyzing Dose-Finding Trials** John O'Quigley, Alexia Iasonos, Björn Bornkamp, 2017-04-27 Handbook of Methods for Designing Monitoring and Analyzing Dose Finding Trials gives a thorough presentation of state of the art methods for early phase clinical trials The methodology of clinical trials has advanced greatly over the last 20 years and arguably nowhere greater than that of early phase studies The need to accelerate drug development in a rapidly evolving context of targeted therapies immunotherapy combination treatments and complex group structures has provided the stimulus to these advances Typically we deal with very small samples sequential methods that need to be efficient while at the same time adhering to ethical principles due to the involvement of human subjects Statistical inference is difficult since the standard techniques of maximum likelihood do not usually apply as a result of model misspecification and parameter estimates lying on the boundary of the parameter space Bayesian methods play an important part in overcoming these difficulties but nonetheless require special consideration in this particular context The purpose of this handbook is to provide an expanded summary of the field as it stands and also through discussion provide insights into the thinking of leaders in the field as to the potential developments of the years ahead With this goal in mind we present An introduction to the field for graduate students and novices A basis for more established researchers from which to build A collection of material for an advanced course in early phase clinical trials A comprehensive guide to available methodology for practicing statisticians on the design and analysis of dose finding experiments An extensive guide for the multiple comparison and modeling MCP Mod dose finding approach adaptive two stage designs for dose finding as well as dose time response models and multiple testing in the context of confirmatory dose finding studies John O Quigley is a professor of mathematics and research director at the French National Institute for Health and Medical Research based at the Faculty of Mathematics University Pierre and Marie Curie in Paris France He is author of Proportional Hazards Regression and has published extensively in the field of dose finding Alexia Iasonos is an associate attending biostatistician at the Memorial Sloan Kettering Cancer Center in New York She has over one hundred publications in the leading statistical and clinical journals on the methodology and design of early phase clinical trials Dr Iasonos has wide experience in the actual implementation of model based early phase trials and has given courses in scientific meetings internationally Bj rn Bornkamp is a statistical methodologist at Novartis in Basel Switzerland researching and implementing dose finding designs in Phase II clinical trials He is one of the co developers of the MCP Mod methodology for dose finding

and main author of the DoseFinding R package He has published numerous papers on dose finding nonlinear models and Bayesian statistics and in 2013 won the Royal Statistical Society award for statistical excellence in the pharmaceutical industry

Design and Analysis of Dose-finding Studies in Drug Development Corine Baayen,2015

Dose Finding by the Continual Reassessment Method Ying Kuen Cheung,2011-03-29 As clinicians begin to realize the important role of dose finding in the drug development process there is an increasing openness to novel methods proposed in the past two decades In particular the Continual Reassessment Method CRM and its variations have drawn much attention in the medical community though it has yet to become a commonplace tool To overcome the status quo in phase I clinical trials statisticians must be able to design trials using the CRM in a timely and reproducible manner A self contained theoretical framework of the CRM for researchers and graduate students who set out to learn and do research in the CRM and dose finding methods in general Dose Finding by the Continual Reassessment Method features Real clinical trial examples that illustrate the methods and techniques throughout the book Detailed calibration techniques that enable biostatisticians to design a CRM in timely manner Limitations of the CRM are outlined to aid in correct use of method This book supplies practical efficient dose finding methods based on cutting edge statistical research More than just a cookbook it provides full unified coverage of the CRM in addition to step by step guidelines to automation and parameterization of the methods used on a regular basis A detailed exposition of the calibration of the CRM for applied statisticians working with dose finding in phase I trials the book focuses on the R package dform for the CRM and its major variants The author recognizes clinicians skepticism of model based designs and addresses their concerns that the time professional and computational resources necessary for accurate model based designs can be major bottlenecks to the widespread use of appropriate dose finding methods in phase I practice The theoretically and empirically based methods in Dose Finding by the Continual Reassessment Method will lessen the statistician s burden and encourage the continuing development and implementation of model based dose finding methods

Desing and Analysis of Dose-finding Studies in Drug Development Corine Baayen,2015

Statistical Issues in Drug Development Stephen S. Senn,2008-02-28 Drug development is the process of finding and producingtherapeutically useful pharmaceuticals turning them into safe andeffective medicine and producing reliable information regardingthe appropriate dosage and dosing intervals With regulatoryauthorities demanding increasingly higher standards in suchdevelopments statistics has become an intrinsic and criticalement in the design and conduct of drug development programmes Statistical Issues in Drug Development presents anessential and thought provoking guide to the statistical issues andcontroversies involved in drug development This highly readable second edition has been updated toinclude Comprehensive coverage of the design and interpretation ofclinical trials Expanded sections on missing data equivalence meta analysisand dose finding An examination of both Bayesian and frequentist methods A new chapter on pharmacogenomics and expanded coverage ofpharmaco epidemiology and pharmaco economics Coverage of the ICH

guidelines in particular ICH E9 Statistical Principles for Clinical Trials It is hoped that the book will stimulate dialogue between statisticians and life scientists working within the pharmaceutical industry The accessible and wide ranging coverage make it essential reading for both statisticians and non statisticians working in the pharmaceutical industry regulatory bodies and medical research institutes There is also much to benefit undergraduate and postgraduate students whose courses include a medical statistics component

Guide to Paediatric Drug Development and Clinical Research K. Rose, J. N. van den Anker, 2010-05-21 Children in the developed world have never enjoyed better medical care mortality has decreased and many fatal diseases of the past can today be prevented or even cured However the current practice of pharmacotherapy in children does not reflect existing scientific knowledge and has come under scrutiny by paediatricians pharmacists and regulatory authorities In order to advance the development of medicines tailored to paediatric needs US and EU legislators have taken action and the WHO has initiated a global paediatric campaign This book gives an overview over the worldwide activities that increasingly include children in the development of new medicines Triggered by both a better understanding of how the child's body develops as well as recent legislation in the USA and in Europe this comprises dosing ethics age appropriate pharmaceutical forms and clinical trials to name just a few aspects A wide spectrum of readers will profit from this book including paediatricians pharmacists general practitioners and health care professionals involved in child care and paediatric research clinical trial personnel patient advocacy groups ethics committees politicians parents and interested lay persons

Dose Finding by the Continual Reassessment Method Ying Kuen Cheung, 2011-03-29 This book presents the continual reassessment method CRM as a tool for dose finding studies With a focus on the implementation and practice of the CRM and its variations it explains how the CRM may be calibrated and extended to suit common clinical settings The book includes examples of real clinical trials data to illustrate the calibration techniques and shows how R can be used to carry out the techniques It reviews the literature related methodology and theoretical properties of the CRM It also explores alternatives for situations where the CRM fails

A Dose Finding Method in Joint Modeling of Efficacy and Safety Endpoints in Phase II Studies Aiyang Tao, 2010 Determination of appropriate doses to advance into Phase III is one of the most challenging and important decisions made during drug development Selecting a dose too high may result in unacceptable safety problems while a too low dose may lead to ineffective drugs Proper estimation of such dose response profiles for relevant safety and efficacy endpoints allows the reliable evaluation of the risk benefit profile of a drug at the end of Phase II as well as the selection of appropriate doses to be brought into confirmatory Phase III trials This dissertation will address how to select doses in Phase II trials by combining information about the efficacy and safety in a joint model setting The methods we present in the dissertation may play a key role in drug development programs and are often the gate keeper for large confirmatory Phase III trials with greater chance of successful approval The dose selection when both safety and efficacy are represented by continuous responses is discussed in Part I of the dissertation while Part II addresses the

methodology when the safety and efficacy are mixed type responses Both scenarios involve joint modeling of safety and efficacy endpoints The methodology will focus on the following 1 Joint modeling approaches 2 Model selection 3 Identification of minimum effective dose MED and maximum safety dose MSD 4 Selection of optimal dose s for the Phase III program

Encyclopedia of Biopharmaceutical Statistics - Four Volume Set Shein-Chung Chow, 2018-09-03 Since the publication of the first edition in 2000 there has been an explosive growth of literature in biopharmaceutical research and development of new medicines This encyclopedia 1 provides a comprehensive and unified presentation of designs and analyses used at different stages of the drug development process 2 gives a well balanced summary of current regulatory requirements and 3 describes recently developed statistical methods in the pharmaceutical sciences Features of the Fourth Edition 1 78 new and revised entries have been added for a total of 308 chapters and a fourth volume has been added to encompass the increased number of chapters 2 Revised and updated entries reflect changes and recent developments in regulatory requirements for the drug review approval process and statistical designs and methodologies 3 Additional topics include multiple stage adaptive trial design in clinical research translational medicine design and analysis of biosimilar drug development big data analytics and real world evidence for clinical research and development 4 A table of contents organized by stages of biopharmaceutical development provides easy access to relevant topics About the Editor Shein Chung Chow Ph D is currently an Associate Director Office of Biostatistics U S Food and Drug Administration FDA Dr Chow is an Adjunct Professor at Duke University School of Medicine as well as Adjunct Professor at Duke NUS Singapore and North Carolina State University Dr Chow is the Editor in Chief of the Journal of Biopharmaceutical Statistics and the Chapman Hall CRC Biostatistics Book Series and the author of 28 books and over 300 methodology papers He was elected Fellow of the American Statistical Association in 1995

Modern Dose-Finding Designs for Cancer Phase I Trials: Drug Combinations and Molecularly Targeted Agents Akihiro Hirakawa, Hiroyuki Sato, Takashi Daimon, Shigeyuki Matsui, 2018-01-30 This book deals with advanced methods for adaptive phase I dose finding clinical trials for combination of two agents and molecularly targeted agents MTAs in oncology It provides not only methodological aspects of the dose finding methods but also software implementations and practical considerations in applying these complex methods to real cancer clinical trials Thus the book aims to furnish researchers in biostatistics and statistical science with a good summary of recent developments of adaptive dose finding methods as well as providing practitioners in biostatistics and clinical investigators with advanced materials for designing conducting monitoring and analyzing adaptive dose finding trials The topics in the book are mainly related to cancer clinical trials but many of those topics are potentially applicable or can be extended to trials for other diseases The focus is mainly on model based dose finding methods for two kinds of phase I trials One is clinical trials with combinations of two agents Development of dose finding methods for two agent combination trials requires reasonable models that can adequately capture joint toxicity probabilities for two agents taking into consideration

possible interactions of the two agents on toxicity probability such as synergistic or antagonistic effects Another is clinical trials for evaluating both efficacy and toxicity outcomes in single and two agent combination trials These methods are often applied to the phase I trials including MTAs because the toxicity and efficacy for a MTA does not monotonically increase with dose but the efficacy often increases initially with the dose and then plateaus Successful software implementations for several dose finding methods are introduced in the book and their operating characteristics in practice are discussed Recent advance of the adaptive dose finding methods in drug developments are also provided

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

Model-Assisted Bayesian Designs for Dose Finding and Optimization Ying Yuan, Ruitao Lin, J. Jack Lee, 2022-11-11 Bayesian adaptive designs provide a critical approach to improve the efficiency and success of drug development that has been embraced by the US Food and Drug Administration FDA This is particularly important for early phase trials as they form the basis for the development and success of subsequent phase II and III trials The objective of this book is to describe the state of the art model assisted designs to facilitate and accelerate the use of novel adaptive designs for early phase clinical trials Model assisted designs possess avant garde features where superiority meets simplicity Model assisted designs enjoy exceptional performance comparable to more complicated model based adaptive designs yet their decision rules often can be pre tabulated and included in the protocol making implementation as simple as conventional algorithm based designs An example is the Bayesian optimal interval BOIN design the first dose finding design to receive the fit for purpose designation from the FDA This designation underscores the regulatory agency s support of the use of the novel adaptive design to improve drug development Features Represents the

first book to provide comprehensive coverage of model assisted designs for various types of dose finding and optimization clinical trials Describes the up to date theory and practice for model assisted designs Presents many practical challenges issues and solutions arising from early phase clinical trials Illustrates with many real trial applications Offers numerous tips and guidance on designing dose finding and optimization trials Provides step by step illustrations of using software to design trials Develops a companion website www.trialdesign.org to provide freely available easy to use software to assist learning and implementing model assisted designs Written by internationally recognized research leaders who pioneered model assisted designs from the University of Texas MD Anderson Cancer Center this book shows how model assisted designs can greatly improve the efficiency and simplify the design conduct and optimization of early phase dose finding trials It should therefore be a very useful practical reference for biostatisticians clinicians working in clinical trials and drug regulatory professionals as well as graduate students of biostatistics Novel model assisted designs showcase the new KISS principle Keep it simple and smart

Wiley Encyclopedia of Clinical Trials Lisa Marie Sullivan, Joseph Massaro, 2008 Here you ll find more than 500 entries from the world s leading experts in the field on the basic concepts methodologies and applications in clinical trials The range of topics includes basic statistical concepts design and analysis of clinical trials ethics regulatory issues and methodologies for clinical data management and analysis

Statistical Issues in Drug Development Stephen S. Senn, 2021-05-25 Statistical Issues in Drug Development The revised third edition of Statistical Issues in Drug Development delivers an insightful treatment of the intersection between statistics and the life sciences The book offers readers new discussions of crucial topics including cluster randomization historical controls responder analysis studies in children post hoc tests estimands publication bias the replication crisis and many more This work presents the major statistical issues in drug development in a way that is accessible and comprehensible to life scientists working in the field and takes pains not to gloss over significant disagreements in the field of statistics while encouraging communication between the statistical and life sciences disciplines In addition to new material on topics like invalid inversion severity random effects in network meta analysis and explained variation readers will benefit from the inclusion of A thorough introduction to basic topics in drug development and statistics including the role played by statistics in drug development An exploration of the four views of statistics in drug development including the historical methodological technical and professional An examination of debatable and controversial topics in drug development including the allocation of treatments to patients in clinical trials baselines and covariate information and the measurement of treatment effects Perfect for life scientists and other professionals working in the field of drug development Statistical Issues in Drug Development is the ideal resource for anyone seeking a one stop reference to enhance their understanding of the use of statistics during drug development

Dose Optimization in Drug Development Rajesh Krishna, 2006-05 This reference provides a concise overview of the key principles in dose selection and optimization and demonstrates applicability to recent successful new drug

applications Compiling key issues and current research on safety efficacy and clinical pharmacology and PK PD this volume critically highlights the multidisciplinary nature of drug development and spans the fields of pharmacokinetics clinical pharmacology biostatistics and experimental medicine

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Table of Contents Dose Finding In Drug Development Dose Finding In Drug Development

1. Understanding the eBook Dose Finding In Drug Development Dose Finding In Drug Development
 - The Rise of Digital Reading Dose Finding In Drug Development Dose Finding In Drug Development
 - Advantages of eBooks Over Traditional Books
2. Identifying Dose Finding In Drug Development Dose Finding In Drug Development
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Dose Finding In Drug Development Dose Finding In Drug Development
 - User-Friendly Interface
4. Exploring eBook Recommendations from Dose Finding In Drug Development Dose Finding In Drug Development
 - Personalized Recommendations
 - Dose Finding In Drug Development Dose Finding In Drug Development User Reviews and Ratings

- Dose Finding In Drug Development Dose Finding In Drug Development and Bestseller Lists
- 5. Accessing Dose Finding In Drug Development Dose Finding In Drug Development Free and Paid eBooks
 - Dose Finding In Drug Development Dose Finding In Drug Development Public Domain eBooks
 - Dose Finding In Drug Development Dose Finding In Drug Development eBook Subscription Services
 - Dose Finding In Drug Development Dose Finding In Drug Development Budget-Friendly Options
- 6. Navigating Dose Finding In Drug Development Dose Finding In Drug Development eBook Formats
 - ePub, PDF, MOBI, and More
 - Dose Finding In Drug Development Dose Finding In Drug Development Compatibility with Devices
 - Dose Finding In Drug Development Dose Finding In Drug Development Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Dose Finding In Drug Development Dose Finding In Drug Development
 - Highlighting and Note-Taking Dose Finding In Drug Development Dose Finding In Drug Development
 - Interactive Elements Dose Finding In Drug Development Dose Finding In Drug Development
- 8. Staying Engaged with Dose Finding In Drug Development Dose Finding In Drug Development
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Dose Finding In Drug Development Dose Finding In Drug Development
- 9. Balancing eBooks and Physical Books Dose Finding In Drug Development Dose Finding In Drug Development
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Dose Finding In Drug Development Dose Finding In Drug Development
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Dose Finding In Drug Development Dose Finding In Drug Development
 - Setting Reading Goals Dose Finding In Drug Development Dose Finding In Drug Development
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Dose Finding In Drug Development Dose Finding In Drug Development
 - Fact-Checking eBook Content of Dose Finding In Drug Development Dose Finding In Drug Development
 - Distinguishing Credible Sources

13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
14. Embracing eBook Trends
 - Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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