

## **Methods and applications of sample size calculation and recalculation in clinical trials**

Type de contenu : Texte

Type de médiation : sans médiation

Type de support : Volume

Titre(s) : Methods and applications of sample size calculation and recalculation in clinical trials / Meinhard Kieser

Auteur(s) : Kieser, Meinhard

Editeur, producteur : Cham, Switzerland : Springer [2020]

Description matérielle : 1 vol. (xiv-396 p.) ; 25 cm

Collection : Springer series in pharmaceutical statistics

ISBN : 978-3-030-49530-5

Appartient à la collection : Springer series in pharmaceutical statistics 2016-.... Cham Springer

Classification décimale Dewey : 610.724 23

Note sur les bibliographies et les index : Bibliogr. p. 371-390 (414 réf.).Index.

Résumé ou extrait : This book provides an extensive overview of the principles and methods of sample size calculation and recalculation in clinical trials. Appropriate calculation of the required sample size is crucial for the success of clinical trials. At the same time, a sample size that is too small or too large is problematic due to ethical, scientific, and economic reasons. Therefore, state-of-the art methods are required when planning clinical trials. Part I describes a general framework for deriving sample size calculation procedures. This enables an understanding of the common principles underlying the numerous methods presented in the following chapters. Part II addresses the fixed sample size design, where the required sample size is determined in the planning stage and is not changed afterwards. It covers sample size calculation methods for superiority, non-inferiority, and equivalence trials, as well as comparisons between two and more than two groups. A wide range of further topics is discussed, including sample size calculation for multiple comparisons, safety assessment, and multi-regional trials. There is often some uncertainty about the assumptions to be made when calculating the sample size upfront. Part III presents methods that allow to modify the initially specified sample size based on new information that becomes available during the ongoing trial. Blinded sample size recalculation procedures for internal pilot study designs are considered, as well as methods for sample size reassessment in adaptive designs that use unblinded data from interim analyses. The application is illustrated using numerous clinical trial

examples, and software code implementing the methods is provided. The book offers theoretical background and practical advice for biostatisticians and clinicians from the pharmaceutical industry and academia who are involved in clinical trials. Covering basic as well as more advanced and recently developed methods, it is suitable for beginners, experienced applied statisticians, and practitioners. To gain maximum benefit, readers should be familiar with introductory statistics. The content of this book has been successfully used for courses on the topic

Sujet - Nom commun : Études cliniques

Médecine clinique

Statistique médicale