## How to Read a Clinical Trial Paper: A Lesson in Basic Trial Statistics

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Address correspondence to: Dr. Peter D. R. Higgins 1150 W. Medical Center Drive SPC 5682 Ann Arbor, MI 48109; Tel: 234-647-2964; Fax: 734-763-2535; E-mail: phiggins@umich.edu Abstract: While the number of clinical trials performed yearly is increasing, the application of these results to individual patients is quite difficult. This article reviews key portions of the process of applying research results to clinical practice. The first step involves defining the study population and determining whether these patients are similar to the patients seen in clinical practice in terms of demographics, disease type, and disease severity. The dropout rate should be compared between the different study arms, Design aspects, including randomization and blinding, should be checked for signs of bias. When comparing studies, clinicians should be aware that the outcomes being studied may vary greatly from one study to another, and some outcomes are much more reliable and valuable than others. The definition of clinical response should also be scrutinized, as it may be too lenient. Surrogate outcomes should be viewed cautiously, and their use should be well justified. Clinicians should also note that statistical significance, as defined by a P-value cutoff, may be the result of a large sample size rather than a clinically significant difference. The treatment effect can be estimated by calculating the number needed to treat, which will demonstrate whether changes in clinical practice are worthwhile. Finally, this article discusses some common issues that can arise with figures.

In the last decade, the number of publications dedicated to research in gastroenterology has expanded dramatically. In parallel, the number of clinical trials has been increasing, with more than 18,000 ongoing clinical trials in the United States alone. With this rapid growth in research, clinicians find themselves trying to keep up with wave after wave of studies and trying to determine how these studies apply to their patients. As researchers learn more about disease processes, clinical trial design is also changing and becoming more complex. Medical school and subsequent clinical training provide limited education on how to evaluate these studies studies.

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Principles and Practice of Clinical Trials Steven Piantadosi, Curtis L. Meinert, 2022-07-19 This is a comprehensive major reference work for our SpringerReference program covering clinical trials Although the core of the Work will focus on the design analysis and interpretation of scientific data from clinical trials a broad spectrum of clinical trial application areas will be covered in detail This is an important time to develop such a Work as drug safety and efficacy emphasizes the Clinical Trials process Because of an immense and growing international disease burden pharmaceutical and biotechnology companies continue to develop new drugs Clinical trials have also become extremely globalized in the past 15 years with over 225 000 international trials ongoing at this point in time Principles in Practice of Clinical Trials is truly an interdisciplinary that will be divided into the following areas 1 Clinical Trials Basic Perspectives 2 Regulation and Oversight 3 Basic Trial Designs 4 Advanced Trial Designs 5 Analysis 6 Trial Publication 7 Topics Related Specific Populations and Legal Aspects of Clinical Trials The Work is designed to be comprised of 175 chapters and approximately 2500 pages The Work will be oriented like many of our SpringerReference Handbooks presenting detailed and comprehensive expository chapters on broad subjects The Editors are major figures in the field of clinical trials and both have written textbooks on the topic There

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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 Leger of Hitchcock's **New Method of Teaching Book-keeping** Ira Irvine Hitchcock, 1853 Pharmaceutical Journal; ,1895 Master Dentistry Volume 1 E-Book Paul Coulthard, Philip Sloan, Elizabeth D. Theaker, Anita Sengupta, 2021-05-25 Now in its fourth edition this popular text provides a comprehensive overview of the oral and maxillofacial subjects in dentistry that students will need in order to pass their final exams This invaluable adjunct to exam preparation provides a practical synthesis of core information reflecting real life case scenarios Information is structured to enhance understanding and clinical decision making and a variety of self assessment methods prepare students for success Comprehensively updated the book covers a range of essential topics in the field of contemporary oral and maxillofacial subjects including surgical flap design state of the art surgical techniques zygomatic implants molecular pathology current imaging applications and pain management The text is integrated and evidence based throughout Covers oral and maxillofacial surgery radiology pathology and oral medicine subjects Brand new chapter on facial skin broadens diagnostic ability Range of self assessment tasks to support learning Aligns to dental school curricula globally Concise and easy to follow Designed to support recall for examination purposes Practical guidance on examination preparation and skills Perfect for BDS exam preparation and candidates taking the MFDS MJDF ORE or other post graduate exams

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