1. Record Nr. UNISA996395449203316

Autore	Perkins William <1558-1602.>
Titolo	A cloud of faithfull vvitnesses, leading to the heauenly Canaan: or, A commentarie vpon the eleuenth chapter to the Hebrewes [[electronic resource]] : Preached in Cambridge, by that godly, and iudicious diuine, M. William Perkins. Long expected and desired; and therefore published at the request of his executors, by William Crashawe, and Thomas Pierson, preachers of Gods Word: who heard him preach it, and wrote it from his mouth
Pubbl/distr/stampa	London, : Printed by William Stansby for Henry Fetherstone and Iohn Parker, 1622
Descrizione fisica	[8], 551, [1] p
Altri autori (Persone)	CrashawWilliam <1572-1626.> PiersonThomas <ca. 1570-1633.>
Lingua di pubblicazione	Inglese
Formato	Materiale a stampa
Livello bibliografico	Monografia
Note generali	Running title reads: A commentarie vpon the eleuenth chapter to the Hebrewes. Reproduction of the original in the Henry E. Huntington Library and Art Gallery.
Sommario/riassunto	eebo-0113

2. Record Nr. UNINA9910818732603321

Autore	Pocock Stuart J.
Titolo	Clinical trials : a practical approach / / Stuart J. Pocock
Pubbl/distr/stampa	Chichester, England : , : John Wiley & Sons, , 1983 ©1983
ISBN	1-118-79391-9 1-118-79392-7
Descrizione fisica	1 online resource (280 p.)
Collana	A Wiley medical publication Clinical trials
Disciplina	615/.7/0724
Soggetti	Clinical trials
Lingua di pubblicazione	Inglese
Formato	Materiale a stampa
Livello bibliografico	Monografia
Note generali	Description based upon print version of record.
Nota di bibliografia	Includes bibliographical references and index.
Nota di contenuto	Cover; Title Page; Contents; Preface; 1. Introduction: The Rationale of Clinical Trials; 1.1 Types of clinical trial; 1.2 Controlled clinical trials and the scientific method; 1.3 An example of a clinical trial for primary breast cancer; 2. The Historical Development of Clinical Trials; 2.1 Clinical trials before 1950; 2.2 Clinical trials since 1950; 2.3 Cancer chemotherapy in the United States; 2.4 Treatment of acute myocardial infarction; 2.5 The pharmaceutical industry; 3. Organization and Planning; 3.1 The protocol; 3.2 Administration, staff and finance; 3.3 Selection of patients 3.4 Treatment schedules 3.5 Evaluation of patient response; 4. The Justification for Randomized Controlled Trials; 4.1 Problems with uncontrolled trials; 4.2 Problems with historical controls; 4.3 Problems with concurrent non-randomized controls; 4.4 Is randomization feasible?; 5. Methods of Randomization; 5.1 Patient registration; 5.2 Preparing the randomization list; 5.3 Stratified randomization; 5.4 Unequal randomization; 6. Blinding and Placebos; 6.1 The justification for double-blind trials; 6.2 The conduct of double-blind trials; 6.3 When is blinding feasible?; 7. Ethical Issues 7.1 Medical progress and individual patient care 7.2 Informed patient consent; 8. Crossover Trials; 8.1 Within-patient comparisons; 8.2 The two-period crossover design; 8.3 The analysis and interpretation of crossover trials; 8.4 Multi-period crossover designs; 9. The Size of a Clinical Trial; 9.1 Statistical methods for determining trial size; 9.2 The

realistic assessment of trial size; 9.3 The inadequacy of small trials; 9.4 Multi-centre trials; 9.5 The number of treatments and factorial designs; 10. Monitoring Trial Progress; 10.1 Reasons for monitoring; 10.2 Interim analyses
10.3 Repeated significance testing: group sequential designs 10.4 Continuous sequential designs; 11. Forms and Data Management; 11.1 Form design; 11.2 Data management; 11.3 The use of computers; 12. Protocol Deviations; 12.1 Ineligible patients; 12.2 Non-compliance and incomplete evaluation; 12.3 Inclusion of withdrawals in analysis; 13. Basic Principles of Statistical Analysis; 13.1 Describing the data; 13.2 Significance tests; 13.3 Estimation and confidence limits; 14. Further Aspects of Data Analysis; 14.1 Prognostic factors; 14.2 The analysis of survival data; 14.3 Multiplicity of data
15. Publication and Interpretation of Findings 15.1 Trial reports and their critical evaluation; 15.2 An excess of false-positives; 15.3 Combining evidence and overall strategy; References; Index

Sommario/riassunto

This comprehensive, unified text on the principles and practice of clinical trials presents a detailed account of how to conduct the trials. It describes the design, analysis, and interpretation of clinical trials in a non-technical manner and provides a general perspective on their historical development, current status, and future strategy. Features examples derived from the author's personal experience.

3.

Record Nr.	UNIORUON00062561
Autore	MALM, William P.
Titolo	Japanese music and musical instruments / William Malm
Pubbl/distr/stampa	Toko, : Tuttle, 1959
Descrizione fisica	299 p. : ill. ; 27 cm
Classificazione	GIA IX H
Soggetti	MUSICA GIAPPONESE
Lingua di pubblicazione	Inglese
Formato	Materiale a stampa
Livello bibliografico	Monografia