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content since the last edition, Clinical Decision Support, 2nd Edition explores the crucial new motivating factors poised to accelerate Clinical Decision Support (CDS) adoption. This book is mostly focused on the US perspective because of initiatives driving EHR adoption, the articulation of

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'meaningful use', and new policy attention in process including the Office of the National Coordinator for Health Information Technology (ONC) and the Center for Medicare and Medicaid Services (CMS). A few chapters focus on the broader international perspective. Clinical

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savings, as well as the still substantial roadblocks needed to be overcome by policy-makers, clinicians, and clinical informatics experts, the field is poised anew on the brink of broad adoption. Clinical Decision Support, 2nd Edition provides an updated and pragmatic view of the methodological processes

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and implementation considerations. This book also considers advanced technologies and architectures, standards, and cooperative activities needed on a societal basis for truly large-scale adoption. At least 40% updated, and seven new chapters since the previous edition, with the

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new and revised content focused on new opportunities and challenges for clinical decision support at point of care, given changes in science, technology, regulatory policy, and healthcare finance Informs healthcare leaders and planners, health IT system developers, healthcare IT organization

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leaders and staff, clinical informatics professionals and researchers, and clinicians with an interest in the role of technology in shaping healthcare of the future

Clinical trials are the engine of progress in the development of new drugs and devices for the detection,

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thinking about cancer biology. A poorly done trial does little to advance the field or guide clinical practice, consumes precious clinical and financial resources and challenges the validity of the ethical contract between investigators and the volunteers who willingly give their time and effort to

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benefit future patients. With chapters written by oncologists, researchers, biostatisticians, clinical research administrators, and industry and FDA representatives, *Oncology Clinical Trials*, provides a comprehensive guide for both early-career and senior oncology investigators into the

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successful design, conduct and analysis of an oncology clinical trial. Oncology Clinical Trials covers how to formulate a study question, selecting a study population, study design of Phase I, II, and III trials, toxicity monitoring, data analysis and reporting, use of genomics, cost-

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effectiveness analysis, systemic review and meta-analysis, and many other issues. Many examples of real-life flaws in clinical trials that have been reported in the literature are included throughout. The book discusses clinical trials from start to finish focusing on real-life examples in

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the development, design and analysis of clinical trials. *Oncology Clinical Trials* features: A systematic guide to all aspects of the design, conduct, analysis, and reporting of clinical trials in oncology Contributions from oncologists, researchers, biostatisticians, clinical research

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administrators, and industry and FDA representatives Hot topics in oncology trials including multi-arm trials, meta-analysis and adaptive design, use of genomics, and cost-effectiveness analysis Real-life examples from reported clinical trials included throughout

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This open access book comprehensively covers the fundamentals of clinical data science, focusing on data collection, modelling and clinical applications. Topics covered in the first section on data collection include: data sources, data at scale (big data), data stewardship

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(FAIR data) and related privacy concerns. Aspects of predictive modelling using techniques such as classification, regression or clustering, and prediction model validation will be covered in the second section. The third section covers aspects of (mobile) clinical decision support

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systems, operational excellence and value-based healthcare.

Fundamentals of Clinical Data Science is an essential resource for healthcare professionals and IT consultants intending to develop and refine their skills in personalized medicine, using solutions based on large datasets from

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chapters have been thoroughly revised, updated, and made more user-friendly.

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therapists, as a cornerstone, a component, or an adjunct of their work. In particular, research studies are identifying the Buddhist practice of mindfulness—a non-judgmental self-observation that promotes

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compliance with ACGME standards. The list goes on and on. Unfortunately, new coordinators are not often given the resources they need to effectively perform these tasks. They are left on their own to navigate complicated standards, overcome scheduling

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challenges, and master documentation requirements. Failure to quickly learn and carry out these duties puts the program's accreditation at risk. And because standards and expectations are constantly changing, even the most experienced coordinators often find

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Clinical research monitoring is a vital aspect of Good Clinical Practice (GCP). Its principles are

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straightforward: they are aimed at protecting those subjects that participate in the trial, and their goal is to provide reliable data that will contribute to the safety and efficacy of the intervention under study, i.e. to support the health of future subjects.

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However, the practical implementation of these major goals is complicated. Various mishaps have happened in recent history, and an extensive set of international rules and regulations have emerged. This book gives a thorough survey of the ethical and legal aspects

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of clinical research and provides a detailed guideline for implementing these aspects into the practice of studying investigational medicinal products in humans, in the European context. It can be used as a study aid for starting monitors, a reference guide

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for more experienced monitors, and anyone else involved in clinical research. Contents: The Past Medicinal Products: The Development Process Clinical Trials: Design Aspects The Rules and the Regs The Ethical Pillars of Clinical Research The

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Development; Investigational Medicinal
Products (IMPs) Review: Key Features:
Current textbooks are US
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The third edition of the bestselling *Clinical Trials in Oncology* provides a concise, nontechnical, and thoroughly up-to-date review of methods and issues related to cancer clinical trials.

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clinical trials issues, and issues specific to Phases I, II, and III. New sections cover innovations in Phase I designs, randomized Phase II designs, and overcoming the challenges of array data. Although this book focuses on cancer trials, the same issues and

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concepts are important in any clinical setting. As always, the authors use clear, lucid prose and a multitude of real-world examples to convey the principles of successful trials without the need for a strong statistics or mathematics background. Armed with

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chapter concludes with a review of key points and knowledge application.

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It is superior to
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exist in those designs.

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of using the randomized control design for a specific intervention and disease may differ, the basic fundamentals still apply in developing the study protocol and

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operational procedures.

These fundamentals still
apply in developing the
study protocol and
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These fundamentals include
identifying the specific

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questions to be tested and appropriate outcome measures, determining an adequate sample size, specifying the randomization procedure, detailing the intervention

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organizational structure. This text is structured to address the fundamentals as the protocol for a clinical trial is being developed. A chapter is devoted to

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each of the critical areas of a protocol to aid the clinical trial researcher. The fundamentals described in this text are based on sound scientific methodology, statistical

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including cardiology,
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diabetes, osteoporosis,
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steering committee responsible for developing the protocol and as members of data and safety monitoring committees. The fundamentals were proposed in the first edition

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published in 1981 and have not changed substantially in the later editions.

However, the number of examples illustrating the fundamentals has greatly expanded base on the

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collective experience of the authors. This text is intended for the clinical researcher who is interested in designing a clinical trial and developing a protocol. It

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and the implications for the care and treatment of patients. The test uses numerous examples of published clinical trials from a variety of medical disciplines to meaningfully

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illustrate the fundamentals. Technical design issues such as sample size are considered but the technical details have been suppressed as much as possible through

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Data sharing can accelerate new discoveries by avoiding duplicative trials, stimulating new ideas for research, and enabling the maximal scientific knowledge and benefits to be gained from the efforts of clinical

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trial participants and investigators. At the same time, sharing clinical trial data presents risks, burdens, and challenges. These include the need to protect the privacy and honor the consent of clinical trial participants; safeguard the

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legitimate economic interests of sponsors; and guard against invalid secondary analyses, which could undermine trust in clinical trials or otherwise harm public health. Sharing Clinical Trial Data presents activities and strategies for the

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responsible sharing of clinical trial data. With the goal of increasing scientific knowledge to lead to better therapies for patients, this book identifies guiding principles and makes recommendations to maximize the benefits and minimize

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access to data, and future knowledge and infrastructure needs. Responsible sharing of clinical trial data will allow other investigators to replicate published findings and carry out additional analyses, strengthen the evidence

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and reviews the ethical and legal standards for conducting it. It also considers problems with the interpretation and application of these standards and conduct, concluding that while children should not be excluded from potentially beneficial clinical

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