

SOCRA CCRP Certification Exam Study Guide **2024 Update.**

Nuremberg Code - Correct answer the first set of principles outlining professional ethics for clinical research.

Nuremberg Code elements - Correct answer 1. Voluntary informed consent

2. Research benefits society
3. Should be based on prior animal work
4. Avoid suffering and injury
5. Research where death and disabling injury is expected shouldn't be conducted
6. Risks should be justified
7. Proper preparations and adequate facilities
8. Conducted by scientifically qualified
9. Subjects can withdraw
10. Research must end the study if there is probable because that continuing would lead to injury, disability, or death.

Timeline of Historical Events - Correct answer Nuremberg Code first (1947). Declaration of Helsinki second (1964). Belmont Report third (1979).

Belmont Report Principles and Application - Correct answer there are 3:

1. Respect for persons = informed consent
2. Beneficence = risk/benefit analysis
3. Justice = appropriate selection of patients

Language Level Recommended for Informed Consent - Correct answer 6th-8th grade

8 Basic Elements of informed Consent - Correct answer 1. Statement explaining the study involves research.

2. Description of risks or discomforts.
3. Description of benefits.
4. Alternative treatments/procedures.
5. Confidentiality.
6. Compensation for involvement and/or injury.
7. Who to contact.
8. Voluntary and can discontinue

Additional Elements of Informed Consent - Correct answer 1. Unforeseeable risks to subject.

2. Participation can be terminated by investigator.
3. Any additional costs.
4. Consequences of the subject's decision to withdraw
5. Significant new findings will be shared
6. approx. # of subjects in the study.

Differences between short and long informed consent form - Correct answer Long = the standard consent form.

Short = document states that the elements have been presented orally to and understood by the subject or LAR.

ICF Monitoring Considerations - Correct answer 1. Subject has signed most recent IRB approved version

2. Subject signature is present in addition to subject name

Socra Certification Study Guide

Karen E. Woodin



Socra Certification Study Guide:

Ccrp Exam Study Guide Pramod Lad, 2019-03-24 This is a companion volume to the CCRP EXAM WORKBOOK. The sequence of chapters is the same in both books to facilitate parallel review. The study guide provides the didactic material while the exam workbook provides test questions pertaining to it. For maximum effectiveness in exam preparation, the two volumes should be studied together. Clinical research management, including the management of clinical trials, is a complex activity involving several different individuals with varying educational and professional backgrounds. Research investigators, clinical research coordinators, research nurses, monitors, IRB staff, regulatory personnel, to name a few, all play an important role in clinical trial and clinical research management. The Society of Clinical Research Associates (SOCRA) provides an important forum for the education and training of clinical research professionals. A significant component of this training is the certification exam, which results in the CCRP Certified Clinical Research Professional designation. This designation is particularly important to clinical research coordinators and research nurses who provide the main site-associated support for clinical trial and clinical research management. The certification serves as an important milestone in career development and can assist clinical research coordinators in careers in both academic and teaching hospitals, CROs, as well as within the pharmaceutical industry. The examination evaluates knowledge, understanding, and application of the conduct of clinical research and clinical trials involving humans. It tests the familiarity with the International Conference on Harmonisation Guideline for Good Clinical Practice (ICH GCP), ICH Clinical Safety Data Management Definitions and Standards for Expedited Reporting (E2A), the United States Code of Federal Regulations (CFR), and the ethical principles that guide clinical research consistent with the principles of the Nuremberg Code, the Belmont Report, and the Declaration of Helsinki. This study guide provides one tool for the preparation and study for the CCRP examination. The book addresses the key issues in ICH GCP, federal regulations outlined in statutes including Title 45, part 46, Protection of Human Subjects; Title 21, part 50, Protection of Human Subjects; Title 21, part 56, Institutional Review Boards; Title 21, part 54, Financial Disclosures by Clinical Investigators. Also addressed are key FDA statutes involved in the regulation of clinical trials: Title 21, part 312, Investigational New Drug Application; Title 21, part 812, Investigational Device Exemptions; and Title 21, part 11, Electronic Records and Electronic Signatures. The CCRP exam covers material based not only on these regulations but also on guidances issued by OHRP and the FDA. The study guide is organized in distinct chapters, each of which covers one aspect of the regulations or guidances. The chapters are deliberately designed to instruct on core materials. The study guide is therefore designed not only to prepare for the CCRP examination but also to educate clinical research professionals, particularly clinical research coordinators and research nurses, on matters which arise frequently in clinical research management and administration.

Ccrp Exam Workbook Pramod Lad, 2018-10-09 Clinical research management, including the management of clinical trials, is a complex activity involving several different individuals with varying educational and professional backgrounds.

Research investigators clinical research coordinators research nurses monitors IRB staff regulatory personnel to name a few all play an important role in clinical trial and clinical research management The Society of Clinical Research Associates SOCRA provides an important forum for the education and training of clinical research professionals A significant component of this training is the certification exam which results in the CCRP Certified Clinical Research Professional designation This designation is particularly important to clinical research coordinators and research nurses who provide the main site associated support for clinical trial and clinical research management The certification serves as an important milestone in career development and can assist clinical research coordinators in careers in both academic and teaching hospitals CROS as well as within the pharmaceutical industry The examination evaluates knowledge understanding and application of the conduct of clinical research and clinical trials involving humans It tests the familiarity with the International Conference on Harmonisation Guideline for Good Clinical Practice E6 ICH GCP ICH Clinical Safety Data Management Definitions and Standards for Expedited Reporting E2A the United States Code of Federal Regulations CFR and the ethical principles that guide clinical research consistent with the principles of the Nuremberg Code the Belmont Report and the Declaration of Helsinki This workbook provides one tool for the preparation and study for the CCRP examination The book addresses the key issues in in ICH GCP federal regulations outlined in statutes including Title 45 part 46 Protection of Human Subjects Title 21 part 50 Protection of Human Subjects Title 21 part 56 Institutional review Boards Title 21 part 54 Financial Disclosures by Clinical Investigators Also addressed are key FDA statutes involved in the regulation of clinical trials Title 21 part 312 Investigational New Drug Application Title 21 part 812 Investigational Device Exemptions and Title 21 part 11 Electronic Records and Electronic Signatures The CCRP exam covers material based not only on these regulations but also on guidances issued by OHRP and the FDA The workbook is organized in distinct chapters each of which covers one aspect of the regulations or guidances The multiple choice questions are deliberately designed to instruct on core materials rather than offering linguistically ingenious choices The workbook is therefore designed not only to prepare for the CCRP examination but also to educate clinical research professionals particularly clinical research coordinators and research nurses on matters which arise frequently in clinical research management and administration

The CRC's Guide to Coordinating Clinical Research Karen E. Woodin, 2004 This guidebook is filled with valuable information on the role and responsibilities of a clinical research coordinator CRC and explains the research process from the site and CRC perspective Topics covered include identifying the regulations governing clinical research describing the drug development process discussing good clinical practices and how to apply them in clinical trials and organizing a clinical practice

Responsible Research Philip A. Cola, Christine Pierre, 2006 The editors of U Hospitals of Cleveland and Rx Trials Inc offer a guide to the practical and ethical issues in the conduct of clinical research coordinators that places the topic in broad international perspective by including approaches from the European Union Japan Canada and the United States Thirteen chapters discuss

ethics and human subjects protection responsible conduct the informed consent process pediatric informed consent and assent study implementation and start up recruitment and retention of research subjects documentation quality assurance in clinical trials communication education and training and future trends in professionalization Distributed in the US by BookMasters Annotation 2006 Book News Inc Portland OR booknews.com **The Stroke Center Handbook** Marilyn M. Rymer,Debbie Summers,Pooja Khatri,2014-03-12 Since publication of the first edition of this book new treatments have become available in acute intervention for stroke and new evidence has been uncovered regarding prevention and neurorehabilitation Designed for the entire team at any stroke center including physicians nurses therapists and administrators The Stroke Center Handbook Orga *The Sourcebook for Clinical Research* Natasha Martien,Jeff Nelligan,2018-08-01 A single trial is complex with numerous regulations administrative processes medical procedures deadlines and specific protocol instructions to follow And yet there has existed no single volume comprehensive clinical research reference manual for investigators medical institutions and national and international research personnel to keep on the shelf as a ready reference to navigate through trial complexities and ensure compliance with U S Federal Regulations and ICH GCP until The Sourcebook for Clinical Research An actionable step by step guide through beginning to advanced topics in clinical research with forms templates and checklists to download from a companion website so that study teams will be compliant and will find all the necessary tools within this book Additionally the authors developed Display Posters for Adverse Events Plus Reporting and Medicare Coverage Analysis that can be purchased separately here <https://www.elsevier.com/books-and-journals/book-companion/9780128162422> order display posters Moreover The Sourcebook for Clinical Research contains clear information and guidance on the newest changes in the industry to keep seasoned investigators and staff current and compliant in addition to providing detailed information regarding the most complex topics This book serves as a quick actionable off the shelf resource to keep by your side at the medical clinic Makes vital trial conduct information easy to understand and instructs on how to practically apply current Federal regulations and Good Clinical Practice ICH GCP Offers extensive guidance that is crucial for guaranteeing compliance to clinical research regulations during each step of the clinical research process Provides up to date and extensive coverage of beginning to advanced topics and step by step actions to take during exceptional circumstances including compassionate use emergency use human subjects protections for vulnerable populations and federal audits Furnishes a detailed clinical research Glossary and a comprehensive Appendix containing ready to use forms templates and checklists for clinical trial personnel to download and begin using immediately Written for the fast paced clinic environment with action steps and forms in the book to respond to a research subject s needs urgently and compliantly A Guide to Patient Recruitment and Retention Diana L. Anderson (Ph.D.),2004 Patient recruitment and retention are clearly complex and challenging components of the clinical trials process In the industry s foremost resource A Guide to Patient Recruitment and Retention the authors provide a wealth of practical advice and

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