

## CIP Level 1 study guide questions and answers

291). Define corrosion

✓ Ans: The deterioration of a substance, usually a metal from a reaction with its environment

292). What is passivation?

✓ Ans: A layer of oxides formed on the surface of a metal that provides corrosion protection.

293). What is an example of passivation?

✓ Ans: Stainless Steel

294). What are the elements of a corrosion cell?

✓ Ans: Anode, Cathode, Metallic Pathway, Electrolyte

295). Describe what happens at the anode.

✓ Ans: The metal dissolves into the electrolyte

296). What is the function of the electrolyte?

✓ Ans: Allows the passage of ions to connect the corrosion cell

297). What is the function of the metallic pathway?

✓ Ans: Allows the flow of electrons to connect the corrosion cell

298). What is the galvanic series?

✓ Ans: A listing of metals in order of reactivity (more or less noble) in seawater at 25 C

# Cip Certification Study Guide

**Jill Spealman**



## **Cip Certification Study Guide:**

**Cip Exam Study Guide** Pramod M. Lad, 2019-02-03 **PRODUCT DESCRIPTION** This study guide provides one tool for the preparation and study for the CIP examination. It is a companion book to the CIP Exam Workbook. The sequence of chapters in the study guide follows the same sequence as in the CIP exam workbook, and the flow of ideas in each chapter is concordant with the sequence of questions in the workbook. It is recommended that the two books be studied together for the most effective exam preparation. The study guide is organized in distinct chapters, each of which covers one aspect of the regulations or guidances. The study material is designed to instruct on core information relevant to the examination. However, it is hoped that the study guide can also function as an IRB Handbook. The study guide is therefore designed not only to prepare for the CIP examination but also to educate IRB professionals and Clinical Research Coordinators on matters which arise frequently in IRB administration. The Institutional Review Board (IRB) is responsible for the review of a wide variety of clinical research. As the complexity of clinical research has grown over the years, the duties and responsibilities of the IRB have grown increasingly complex. This complex environment demands that the IRB be staffed and managed by professionals. As a part of affirming the professionalism of IRB staff administrators and directors, the Public Responsibility in Research and Medicine (PRIM R) provides an important forum for education and affirmation of ethical standards for the performance and management of clinical research. An important component of this program is the certification exam known as the CIP Certified IRB Professional. This examination, which is offered twice a year, covers a wide range of regulatory topics. The book addresses the key issues in federal regulations outlined in statutes including Title 45 part 4 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 CIP exam covers material based not only on these regulations but also on guidances issued by OHRP and the FDA. Special attention has been devoted to material covered in these guidances. Also addressed are interactions of the IRB with other committees in the institutional environment. **National Library of**

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Certified IRB Professional This examination which is offered twice a year covers a wide range of regulatory topics This workbook provides one tool for the preparation and study for the CIP examination The book addresses the key issues in 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 CIP exam covers material based not only on these regulations but also on guidances issued by OHRP and the FDA Special attention has been devoted to material covered in these guidances Also addressed are interactions of the IRB with other committees in the institutional environment Some of the material also covers ICH guidelines for clinical trial management 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 offering linguistically ingenious choices An answer key is provided The workbook is therefore designed not only to prepare for the CIP examination but also to educate IRB professionals on matters which arise frequently in IRB administration Current Catalog ,1991 First multi year cumulation covers six years 1965 70 *CIP Exam Workbook* Pramod Lad,2018-06-10 The Institutional Review Board IRB is responsible for the review of a wide variety of clinical research As the complexity of clinical research has grown over the years the duties and responsibilities of the IRB have grown increasingly complex This complex environment demands that the IRB be staffed and managed by professionals As a part of affirming the professionalism of IRB staff administrators and directors the Public Responsibility in Research and Medicine PRIM R provides an important forum for education and affirmation of ethical standards for the performance and management of clinical research An important component of this program is the certification exam known as the CIP Certified IRB Professional This examination which is offered twice a year covers a wide range of regulatory topics This workbook provides one tool for the preparation and study for the CIP examination The book addresses the key issues in federal regulations outlined in statutes including Title 45 part 4 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 CIP exam covers material based not only on these regulations but also on guidances issued by OHRP and the FDA Special attention has been devoted to material covered in these guidances Also addressed are interactions of the IRB with other committees in the institutional environment Some of the material also covers ICH guidelines for clinical trial management 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

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