

# **STERILE PROCESSING FINAL**

## **EXAM STUDY GUIDE**

### **What is Central Service ?**

Centralized distribution of supplies to all customers in the hospital.

### **What is the Central Service One Way Work Flow ?**

- Decontamination (soiled)
- Preparation, Packaging, Sterilization (clean)
- Sterile Storage (sterile)

### **What is the purpose of a One Way Work Flow ?**

To prevent cross contamination.

### **Decontamination**

Physically or chemically render instruments or equipment that may be contaminated with harmful micro organisms to make it safe to handle.

### **Negative air flow**

Air flows into a room from other surrounding areas, pressure is less than other adjacent areas.

### **Positive Air Pressure**

Maintains air flow out of a room, this helps with keeping all dust particles and microorganisms out.

### **Standard Precautions**

Treat everyone and everything as infected /contaminated.

### **Job Description**

Define and communicate job duties and requirements in organization.

### **Nosocomial Infection**

Hospital Acquired Infection.

**Textile Packaging Materials should be held at a room temperature of \_\_\_\_\_ and a relative humidity of \_\_\_\_\_ for a minimum of \_\_\_\_\_ prior to sterilization.**

-64-72 degrees F

-35-70%

-2 Hours

### **Superheating**

When steam is at a temperature which exceeds that of saturated steam at the same pressure.

**Linen packaging material is limited to a size of \_\_\_\_\_,not weigh more than \_\_\_\_\_, and maximum density must not exceed \_\_\_\_\_.**

-12x12x20

-12 lbs

-7.2 lbs

### **What is a wet pack?**

Containers with moisture after the sterilization process is complete.

# Study Guide For Sterile Processing Tech

**Feroz Jameel, John W. Skoug, Robert R.  
Nesbitt**



## **Study Guide For Sterile Processing Tech:**

*Surgical Tech Certification Study Guide* Pasquale De Marco, 2025-04-28 Embark on an educational journey into the realm of surgical technology with this comprehensive guide meticulously crafted for aspiring and practicing surgical technologists. Within these pages you will find an in depth exploration of the intricate procedures and techniques that ensure the safety and well being of patients undergoing surgery. As you delve into the chapters you will gain a thorough understanding of the surgical environment encompassing the specialized equipment instrumentation and sterilization techniques that uphold the highest standards of asepsis. Moreover you will delve into the nuances of preoperative care encompassing patient assessment preparation and informed consent ensuring that patients are adequately prepared for surgery. Furthermore this guide unveils the intricacies of intraoperative care guiding you through the meticulous steps of scrubbing gowning draping and positioning the patient. You will master the art of monitoring the patient's vital signs anticipating potential complications and assisting the surgeon with precision and dexterity. Beyond the operating room this book explores the multifaceted role of the surgical technologist in postoperative care emphasizing the importance of pain management wound care and patient education. It also delves into the realm of surgical emergencies equipping you with the knowledge and skills to respond effectively to unforeseen circumstances. This comprehensive resource extends its reach to encompass legal and ethical considerations in surgical practice ensuring that you navigate the complexities of patient rights informed consent confidentiality and medical malpractice with integrity and professionalism. Additionally it explores the avenues for professional development highlighting the significance of continuing education certification and networking in advancing your career as a surgical technologist. Enrich your knowledge and elevate your skills with this indispensable guide empowering you to excel in your role as a surgical technologist and contribute to the delivery of exceptional patient care. If you like this book write a review on google books.

*Critical Care Study Guide* Gerard J. Criner, Gilbert E. D'Alonzo, 2013-03-09 Critical care medicine is a dynamic and exciting arena where complex pathophysiologic states requiring extensive knowledge and clinical acumen are commonly found. Caring for critically ill patients requires an extensive knowledge of basic pathophysiology as well as awareness of the appropriate diagnostic tests and therapeutic interventions. Because this knowledge base crosses many different disciplines introduction to caring for the intensive care patient while exciting may also be intimidating. This textbook is designed toward making the dynamic environment of the critical care unit understandable and the approach to the patient both logical and successful. The book contains three components: 1 description of the procedural tasks commonly performed for the critically ill patient; 2 explanation of the most common pathophysiologic states encountered; and 3 description of specific disease entities with details of their differential diagnosis, diagnostic strategy and therapeutic plan. This book also uses several educational approaches that we have found useful in our own teaching sessions and prior textbooks. Clinical cases introduce chapters and highlight chapter segments to emphasize clinical relevancy. As an additional study aid margin notes highlight

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Study Guide to Accompany Maternal Child Nursing Care, Second Edition Karen A. Piotrowski,Anne Rath Rentfro,2002

*Development of Biopharmaceutical Drug-Device Products* Feroz Jameel,John W. Skoug,Robert R. Nesbitt,2020-03-13 The biotechnology biopharmaceutical sector has tremendously grown which led to the invention of engineered antibodies such as Antibody Drug Conjugates ADCs Bispecific T cell engager BITES Dual Variable Domain DVD antibodies and fusion proteins that are currently being used as therapeutic agents for immunology oncology and other disease conditions Regulatory agencies have raised the bar for the development and manufacture of antibody based products expecting to see the use of Quality by Design QbD elements demonstrating an in depth understanding of product and process based on sound science Drug delivery systems have become an increasingly important part of the therapy and most biopharmaceuticals for self administration are being marketed as combination products A survey of the market indicates that there is a strong need for a new book that will provide one stop shopping for the latest information and knowledge of the scientific and engineering advances made over the last few years in the area of biopharmaceutical product development The new book entitled *Development of Biopharmaceutical Drug Device Products* is a reference text for scientists and engineers in the biopharmaceutical industry academia or regulatory agencies With insightful chapters from experts in the field this new book reviews first principles covers recent technological advancements and provides case studies and regulatory strategies relating to the development and manufacture of antibody based products It covers topics such as the importance of early preformulation studies during drug discovery to influence molecular selection for development formulation strategies for new modalities and the analytical techniques used to characterize them It also addresses important considerations for later stage development such as the development of robust formulations and processes including process engineering and modeling of manufacturing unit operations the design of analytical comparability studies and characterization of primary containers pre filled syringes and vials Finally the latter half of the book reviews key considerations to ensure the development and approval of a patient centered delivery system design This involves the evolving regulatory framework with perspectives from both the

US and EU industry experts the role of international standards design control risk management human factors and its importance in the product development and regulatory approval process as well as review of the risk based approach to bridging between devices used in clinical trials and the to be marketed device Finally case studies are provided throughout The typical readership would have biology and or engineering degrees and would include researchers scientific leaders industry specialists and technology developers working in the biopharmaceutical field     *NASA Scientific and Technical Reports* United States. National Aeronautics and Space Administration Scientific and Technical Information Division,1967



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