

Instructions:

1. You have 1.0 hours to complete this test.
2. Calculators will not be used.
3. Answers must include units of measurement e.g. mg, ml, tab
4. To receive full marks all calculations must be shown
5. Please circle the correct answer for each question where applicable.

* note: for all of the following questions, you can assume that all of the tablets are scored.

Part A: Conversions

Convert the following (show all your work):

1. 1 g = 1000 mg

2. 1.5 g = 1500 mg

3. 0.05 mg = 50 mcg

4. 0.25 g = 250 mg

5. 15,000 mg = 15 g

6. 100 mcg = 0.1 mg

7. 400 mg = 0.4 g

Practice Test For Drug Calculations Cma

M Carnoy



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Medical Assisting Exam Review for CMA, RMA & CMAS Certification Helen Houser,Janet Sesser,2020-06-08 This Fourth Edition of Medical Assisting Exam Review for CMA RMA CMAS Certification focuses on the critical most current components of the MA and MAS curricula making it an indispensable tool for recent graduates practicing medical assistants medical administrative specialists and medical administrative assistants preparing to sit for any recognized national certification exams

Principles of Research Design and Drug Literature Evaluation Rajender R. Aparasu,John P. Bentley,2014-03-07 Principles of Research Design and Drug Literature Evaluation is a unique resource that provides a balanced approach covering critical elements of clinical research biostatistical principles and scientific literature evaluation techniques for evidence based medicine This accessible text provides comprehensive course content that meets and exceeds the curriculum standards set by the Accreditation Council for Pharmacy Education ACPE Written by expert authors specializing in pharmacy practice and research this valuable text will provide pharmacy students and practitioners with a thorough understanding of the principles and practices

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The Clinical Practice of Drug Information Michael Gabay,2015-03-09 This resource will educate students and pharmacists on traditional drug information topics while providing an extensive background on more recent practice areas This is a user friendly text with multiple examples that can be used in education and training as well as clinical practice Each chapter includes learning objectives key terms example

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the general welfare and well being of a country's citizens. Financial infrastructure and policy reform can play a significant role in optimizing existing healthcare programs. *Health Economics and Healthcare Reform Breakthroughs in Research and Practice* is a comprehensive source of academic material on the importance of economic structures and policy reform initiatives in modern healthcare systems. Highlighting a range of pertinent topics such as clinical costing, patient engagement, and e-health, this book is ideally designed for medical practitioners, researchers, professionals, and students interested in the optimization of healthcare delivery. Clinical Pharmacy Education, Practice and Research Dixon Thomas, 2018-11-23

Clinical Pharmacy Education Practice and Research offers readers a solid foundation in clinical pharmacy and related sciences through contributions by 83 leading experts in the field from 25 countries. This book stresses educational approaches that empower pharmacists with patient care and research competencies. The learning objectives and writing style of the book focus on clarifying the concepts comprehensively for a pharmacist from regular patient counseling to pharmacogenomics practice. It covers all interesting topics a pharmacist should know. This book serves as a basis to standardize and coordinate learning to practice, explaining basics and using self-learning strategies through online resources or other advanced texts. With an educational approach, it guides pharmacy students and pharmacists to learn quickly and apply. *Clinical Pharmacy Education Practice and Research* provides an essential foundation for pharmacy students and pharmacists globally. Covers the core information needed for pharmacy practice courses. Includes multiple case studies and practical situations with 70% focused on practical clinical pharmacology knowledge. Designed for educational settings but also useful as a refresher for advanced students and researchers. *Quantitative Drug Safety and Benefit Risk Evaluation* William Wang, Melvin Munsaka, James Buchanan, Judy Li, 2021-12-30

Quantitative Methodologies and Process for Safety Monitoring and Ongoing Benefit Risk Evaluation provides a comprehensive coverage on safety monitoring methodologies covering both global trends and regional initiatives. Pharmacovigilance has traditionally focused on the handling of individual adverse event reports; however, recently there has been a shift towards aggregate analysis to better understand the scope of product risks. Written to be accessible not only to statisticians but also to safety scientists with a quantitative interest, this book aims to bridge the gap in knowledge between medical and statistical fields, creating a truly multi-disciplinary approach that is very much needed for 21st century safety evaluation.

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