

AP Biology Practice Exam

Section I: Multiple-Choice Questions

Complete one hundred multiple-choice questions in 80 minutes, worth 60 percent of the total score.

BIOLOGY

SECTION I

Time—80 minutes

100 Questions

Directions: Each of the questions or incomplete statements below is followed by five suggested answers or completions. Select the one that is best in each case.

1. During the evolutionary process, which of the following is the correct sequence of events?
 - A. Change in phenotype—change in genotype—speciation—selection
 - B. Speciation—change in genotype—selection—change in phenotype
 - C. Speciation—selection—change in phenotype—change in genotype
 - D. Change in genotype—change in phenotype—selection—speciation
 - E. Selection—speciation—change in phenotype—change in genotype
2. A lenticel is to a plant as a _____ is to an insect.
 - A. stoma
 - B. spiracle
 - C. trachea
 - D. mouth
 - E. siphon

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The Publishers Weekly ,2008 **Guide to College Visits** Princeton Review,2007-04-10 A practical handbook covers everything needed to plan a successful visit to campuses across the country with information on nearby accommodations for every budget the best and worst times to visit itineraries and campus tours **Developing and Validating Test Items** Thomas M. Haladyna,Michael C. Rodriguez,2013-07-18 Since test items are the building blocks of any test learning how to develop and validate test items has always been critical to the teaching learning process As they grow in importance and use testing programs increasingly supplement the use of selected response multiple choice items with constructed response formats This trend is expected to continue As a result a new item writing book is needed one that provides comprehensive coverage of both types of items and of the validity theory underlying them This book is an outgrowth of the author s previous

book *Developing and Validating Multiple Choice Test Items 3e* Haladyna 2004 That book achieved distinction as the leading source of guidance on creating and validating selected response test items Like its predecessor the content of this new book is based on both an extensive review of the literature and on its author's long experience in the testing field It is very timely in this era of burgeoning testing programs especially when these items are delivered in a computer based environment Key features include Comprehensive and Flexible No other book so thoroughly covers the field of test item development and its various applications Focus on Validity Validity the most important consideration in testing is stressed throughout and is based on the Standards for Educational and Psychological Testing currently under revision by AERA APA and NCME Illustrative Examples The book presents various selected and constructed response formats and uses many examples to illustrate correct and incorrect ways of writing items Strategies for training item writers and developing large numbers of items using algorithms and other item generating methods are also presented Based on Theory and Research A comprehensive review and synthesis of existing research runs throughout the book and complements the expertise of its authors

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strengthen your understanding of key concepts Media Resources Lists encourage you to develop a study plan to master the important content in each chapter

Innovative Strategies, Statistical Solutions and Simulations for Modern Clinical Trials Mark Chang, John Balser, Jim Roach, Robin Bliss, 2019-03-20 This is truly an outstanding book It brings together all of the latest research in clinical trials methodology and how it can be applied to drug development Chang et al provide applications to industry supported trials This will allow statisticians in the industry community to take these methods seriously Jay Herson Johns Hopkins University The pharmaceutical industry s approach to drug discovery and development has rapidly transformed in the last decade from the more traditional Research and Development R D approach to a more innovative approach in which strategies are employed to compress and optimize the clinical development plan and associated timelines However these strategies are generally being considered on an individual trial basis and not as part of a fully integrated overall development program Such optimization at the trial level is somewhat near sighted and does not ensure cost time or development efficiency of the overall program This book seeks to address this imbalance by establishing a statistical framework for overall global clinical development optimization and providing tactics and techniques to support such optimization including clinical trial simulations Provides a statistical framework for achieve global optimization in each phase of the drug development process Describes specific techniques to support optimization including adaptive designs precision medicine survival endpoints dose finding and multiple testing Gives practical approaches to handling missing data in clinical trials using SAS Looks at key controversial issues from both a clinical and statistical perspective Presents a generous number of case studies from multiple therapeutic areas that help motivate and illustrate the statistical methods introduced in the book Puts great emphasis on software implementation of the statistical methods with multiple examples of software code both SAS and R It is important for statisticians to possess a deep knowledge of the drug development process beyond statistical considerations For these reasons this book incorporates both statistical and clinical medical perspectives

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