

14-3 Human Molecular Genetics

Watson and Crick took the first step in making genetics a molecular science when they discovered the double-helical structure of DNA in 1953. Today, the transformation they started is complete. The exploration of human genes is now a major scientific undertaking. Biologists can now read, analyze, and even change the molecular code of genes.

Human DNA Analysis

The roughly 6 billion base pairs you carry in your DNA are a bit like an encyclopedia with thousands of volumes. In principle, biologists would like to know everything the volumes contain, but as a practical matter there isn't enough time to read all of them. Nonetheless, if you've used an encyclopedia you've already learned one of the ways to handle huge amounts of information—you find a way to look up only what you need. In an encyclopedia, you can use an index or an alphabetical list of articles. As you might suspect, biologists search the volumes of the human genome using sequences of DNA bases.

Testing for Alleles. If two prospective parents suspect they might be carrying recessive alleles for a genetic disorder such as cystic fibrosis (CF) or Tay-Sachs disease, how could they find out for sure? Because the Tay-Sachs and CF alleles have slightly different DNA sequences from their normal counterparts, a variety of genetic tests have been developed that can spot these differences. Sometimes these genetic tests use labeled DNA probes. These are specific DNA base sequences that detect the complementary base sequences found in disease-causing alleles. Other tests search for changes in restriction enzyme cutting sites. Tests also detect differences between the lengths of normal and abnormal alleles.

Genetic tests are now available for hundreds of disorders, making it possible to determine whether prospective parents risk passing such alleles to their children. In an increasing number of such cases, DNA testing can pinpoint the exact genetic basis of a disorder, making it possible to develop more effective treatment for individuals affected by genetic disease.

► **Figure 14-17** This laboratory worker is preparing a report on DNA evidence. The inset shows vials of DNA lying on a printout of a DNA analysis chart.

Guide for Reading



Key Concepts

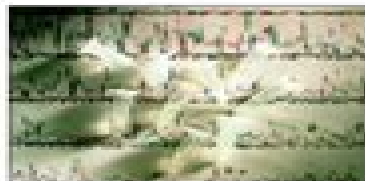
- What is the goal of the Human Genome Project?
- What is gene therapy?

Vocabulary

DNA fingerprinting

Reading Strategy:

Finding Main Ideas. As you read, find evidence to support the following statement: The influence of human molecular genetics on society is growing rapidly.



Section 14-3

1 FOCUS

Objectives

- 14.3.1 Summarize** methods of human DNA analysis.
- 14.3.2 State** the goal of the Human Genome Project.
- 14.3.3 Describe** how researchers are attempting to cure genetic disorders.

Guide for Reading

Vocabulary Preview

Invite student volunteers to describe what a fingerprint is and what it is used for. Then, challenge them to make inferences about what DNA fingerprinting is. List the inferences on the board, and narrow them down to two or three. Revisit this list throughout the section, if necessary, until the class correctly defines the term.

Reading Strategy

Instruct students to write the statement from the text onto a sheet of paper. As they read, students should list under the statement the evidence that supports it.

2 INSTRUCT

Human DNA Analysis

Build Science Skills

Designing Experiments Challenge students to write the steps in a protocol in which they test for the allele of a gene that causes a genetic disorder using restriction enzymes and gel electrophoresis. Students who need an extra challenge can list the steps to testing for alleles using a labeled DNA probe. Encourage students to use what they learned in Chapter 13. They might also wish to use the Internet to find additional resources.



SECTION RESOURCES

Print:

- **Teaching Resources**, Section Review 14-3, Chapter 14 Real-World Lab
- **Reading and Study Workbook A**, Section 14-3
- **Adapted Reading and Study Workbook B**, Section 14-3
- **Issues and Decision Making**, Issues and Decisions 9, 10, 11, 12
- **Biotechnology Manual**, Labs 2, 11, 12; Concepts 2, 3, 4, 6

- **Lesson Plans**, Section 14-3

Technology:

- **iText**, Section 14-3
- **Animated Biological Concepts Videotape Library**, 30
- **Transparencies Plus**, Section 14-3

Answers For 14 3 Human Molecular Genetics

**Joseph Jankovic, John C.
Mazziotta, Scott L. Pomeroy**



Answers For 14 3 Human Molecular Genetics:

An Introduction to Human Molecular Genetics Jack J. Pasternak, 2005-06-03 An Introduction to Human Molecular Genetics Second Edition Jack J Pasternak The Second Edition of this internationally acclaimed text expands its coverage of the molecular genetics of inherited human diseases with the latest research findings and discoveries Using a unique systems based approach the text offers readers a thorough explanation of the gene discovery process and how defective genes are linked to inherited disease states in major organ and tissue systems All the latest developments in functional genomics proteomics and microarray technology have been thoroughly incorporated into the text The first part of the text introduces readers to the fundamentals of cytogenetics and Mendelian genetics Next techniques and strategies for gene manipulation mapping and isolation are examined Readers will particularly appreciate the text s exceptionally thorough and clear explanation of genetic mapping The final part features unique coverage of the molecular genetics of distinct biological systems covering muscle neurological eye cancer and mitochondrial disorders Throughout the text helpful figures and diagrams illustrate and clarify complex material Readers familiar with the first edition will recognize the text s same lucid and engaging style and will find a wealth of new and expanded material that brings them fully up to date with a current understanding of the field including New chapters on complex genetic disorders genomic imprinting and human population genetics Expanded and fully revised section on clinical genetics covering diagnostic testing molecular screening and various treatments This text is targeted at upper level undergraduate students graduate students and medical students It is also an excellent reference for researchers and physicians who need a clinically relevant reference for the molecular genetics of inherited human diseases

Human Molecular Genetics Tom Strachan, Andrew Read, 2018-03-29 Human Molecular Genetics is an established and class proven textbook for upper level undergraduates and graduate students which provides an authoritative and integrated approach to the molecular aspects of human genetics While maintaining the hallmark features of previous editions the Fourth Edition has been completely updated It includes new Key Concepts at the beginning of each chapter and annotated further reading at the conclusion of each chapter to help readers navigate the wealth of information in this subject The text has been restructured so genomic technologies are integrated throughout and next generation sequencing is included Genetic testing screening approaches to therapy personalized medicine and disease models have been brought together in one section Coverage of cell biology including stem cells and cell therapy studying gene function and structure comparative genomics model organisms noncoding RNAs and their functions and epigenetics have all been expanded

SBAs for the MRCOG Part 1 Babu Karavadra, Richard A. Haines, Medha Sule, 2024-06-25 SBAs for the MRCOG Part 1 is a question and answer style revision guide designed to help candidates prepare for Papers 1 and 2 of the MRCOG Part 1 examination Mapped to the exam syllabus the book contains 500 single best answer SBA questions carefully written to reflect fundamental areas of the curriculum along with explanatory answers based on the most recent

Green top Guidelines from the Royal College of Obstetricians and Gynaecologists RCOG and journal articles from The Obstetrician and Gynaecologist TOG The questions test knowledge of the basic sciences as well as considerations relevant to day to day clinical practice to help candidates to understand the wider context of their learning Alongside the Q As for the individual curriculum areas the book contains two 100 question mock papers to help candidates feel fully prepared for the real exam Providing a thorough assessment of the key topics and expert guidance this is an essential resource for obstetrics and gynaecology trainees looking to maximise their exam success

Molecular Biology of the Cell 6E - The Problems Book John Wilson,Tim Hunt,2014-11-21 The Problems Book helps students appreciate the ways in which experiments and simple calculations can lead to an understanding of how cells work by introducing the experimental foundation of cell and molecular biology Each chapter reviews key terms tests for understanding basic concepts and poses research based problems The Problems Book has been

Combinatorial Pattern Matching Ricardo Baeza-Yates,Edgar Chávez,Maxime Crochemore,2007-03-06 The refereed proceedings of the 14th Annual Symposium on Combinatorial Pattern Matching CPM 2003 held in Morelia Michoacán Mexico in June 2003 The 28 revised full papers presented were carefully reviewed and selected from 57 submissions The papers are devoted to current theoretical and computational aspects of searching and matching strings and more complicated patterns such as trees regular expressions graphs point sets and arrays Among the application fields addressed are computational biology bioinformatics genomics the Web data compression coding multimedia information retrieval pattern recognition and computer vision

Bradley and Daroff's Neurology in Clinical Practice - E-Book Joseph Jankovic,John C. Mazziotta,Scott L. Pomeroy,2021-03-23 A practical dynamic resource for practicing neurologists clinicians and trainees Bradley and Daroff's Neurology in Clinical Practice Eighth Edition offers a straightforward style evidence based information and robust interactive content supplemented by treatment algorithms and images to keep you up to date with all that's current in this fast changing field This two volume set is ideal for daily reference featuring a unique organization by presenting symptom sign and by specific disease entities allowing you to access content in ways that mirror how you practice More than 150 expert contributors led by Drs Joseph Jankovic John C Mazziotta Scott L Pomeroy and Nancy J Newman provide up to date guidance that equips you to effectively diagnose and manage the full range of neurological disorders Covers all aspects of today's neurology in an easy to read clinically relevant manner Allows for easy searches through an intuitive organization by both symptom and grouping of diseases Features new and expanded content on movement disorders genetic and immunologic disorders tropical neurology neuro ophthalmology and neuro otology palliative care pediatric neurology and new and emerging therapies Offers even more detailed videos that depict how neurological disorders manifest including EEG and seizures deep brain stimulation for PD and tremor sleep disorders movement disorders ocular oscillations EMG evaluation cranial neuropathies and disorders of upper and lower motor neurons as well as other neurologic signs Enhanced eBook version included with purchase Your enhanced eBook allows you to access all of the text

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Applications of Statistical and Machine Learning Methods in Bioinformatics Jaroslaw Meller,Wieslaw Nowak,2007 Statistical and machine learning approaches play an increasingly important role in biomedical research In the absence of fundamental first principle based models or because of the computational complexity of such models statistical and machine learning approaches are being used to identify interesting structures in the data e g patterns in gene expression profiles correlate these patterns and other input attributes with e g medically relevant outcomes and to develop predictors that can generalize from known data and make predictions for new data instances Examples of important applications include structural bioinformatics in which one of the goals is to predict elements of protein structure from amino acid sequence or microarray gene expression profiling in which the goal is to discover interesting patterns in gene expression data and correlate them with clinically relevant phenotypes This volume includes papers submitted to the BIT 2005 workshop on the Applications of Machine and Statistical Learning Methods in Bioinformatics that took place in September 2005 in Torun Poland Animal Biotechnology Ashish S. Verma,Anchal Singh,2020-06-09 Animal Biotechnology Models in Discovery and Translation Second Edition provides a helpful guide to anyone seeking a thorough review of animal biotechnology and its application to human disease and welfare This updated edition covers vital fundamentals including animal cell cultures genome sequencing analysis epigenetics and animal models gene expression and ethics and safety concerns along with in depth examples of implications for human health and prospects for the future New chapters cover animal biotechnology as applied to various disease types and research areas including in vitro fertilization human embryonic stem cell research

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