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 Alleies 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 affeles have slightly different DNA sequences from their normal counterparts, a variety of genetic tests have been developed that can soot those 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 affelies.

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 printend of a DNA analysis chart.

Guide for Reading



May Concepts

- . What is the goal of the Human Corrognee Project?
- What is gone therapy?

DINA Singerprinting

Reading Strategy: Finding Main Ideas Acyon read, find evidence to support. the following statement: The influence of human molecular. genetics on society is growing suppositly.





SECTION RESOURCES

- . 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

Legion Plans, Section 14-3.

Technology:

- Ffext, Section 14-3.
- Animated Biological Concepts Videotope Library, 30
- Transparencies Plus, Section 14–3.

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

levite student volunteers to describe what a fingerprint is and what it is used for. Then, challenge them tomake 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 gelelectrophoresis. 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.

Answers For 14 3 Human Molecular Genetics

Arturo Cuomo

Answers For 14 3 Human Molecular Genetics:

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 **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 figures and references from the book on a variety of devices Central Nervous System Diseases—Advances in Research and Treatment: 2013 Edition ,2013-06-21 Central Nervous System Diseases Advances in Research and Treatment 2013 Edition is a ScholarlyEditions book that delivers timely authoritative and comprehensive information about Brain Diseases The editors have built Central Nervous System Diseases Advances in Research and Treatment 2013 Edition on the vast information databases of ScholarlyNews You can expect the information about Brain Diseases in this book to be deeper than what you can access anywhere else as well as consistently reliable

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Biology ,2004 Molecular Biology of the Cell ,2007 Craniofacial Development, Growth and Evolution Murray Clyde Meikle,2002 Over the past twenty years craniofacial biology has been revolutionized by major developments in our understanding of the cellular molecular and genetic mechanisms underlying embryonic development Many of these advances have been based on animal models most notably the fruitfly Drosophila the chick and the mouse Since these developmental processes have been highly conserved during evolution this information is relevant not only to understanding normal human development but also to understanding how genetic mutations produce particular malformations or inherited diseases This book incorporates these discoveries into traditional morphological description of craniofacial development and in a form accessible to clinicians with an interest in the head and neck book cover

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