## 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**

Ricardo Baeza-Yates, Edgar Chávez, Maxime Crochemore

## **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 SBAs for the MRCOG Part 1 Babu Karavadra, Richard A. 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Derry, 2015-01-01 Canadian historian Margaret Derry examines the evolution of modern animal breeding from the invention of improved breeding methods in 18th century England to the application of molecular genetics in the 1980s and 1990s 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 biosensors enteric diseases biopharming organ transplantation tuberculosis neurodegenerative disorders and more 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 authoritative informed and relevant The content of Central Nervous System Diseases Advances in Research and Treatment 2013 Edition has been produced by the world's leading scientists engineers analysts research institutions and companies All of the content is from peer reviewed sources and all of it is written assembled and edited by the editors at ScholarlyEditions and available exclusively from us You now have a source you can cite with authority confidence and credibility More information is available at http www ScholarlyEditions com

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