

Chapter 14 The Human Genome

Making Karyotypes

Introduction

Several human genetic disorders are caused by extra, missing, or damaged chromosomes. In order to study these disorders, cells from a person are grown with a chemical that stops cell division at the metaphase stage. During metaphase, a chromosome exists as two chromatids attached at the centromere.

The cells are stained to reveal banding patterns and placed on glass slides. The chromosomes are observed under the microscope, where they are counted, checked for abnormalities, and photographed. The photograph is then enlarged, and the images of the chromosomes are individually cut out. The chromosomes are identified and arranged in homologous pairs. The arrangement of homologous pairs is called a karyotype. In this investigation, you will use a sketch of chromosomes to make a karyotype. You will also examine the karyotype to determine the presence of any chromosomal abnormalities.

Problem

How can chromosomes be observed?

Pre-Lab Discussion

Read the entire investigation. Then work with a partner to answer the following questions.

1. What clues to the presence of certain genetic disorders can be seen in a karyotype?

2. Why might a laboratory worker attempting to diagnose a genetic disorder prefer to work with photographs of chromosomes rather than the chromosomes themselves?

3. Why would it be much more difficult to construct a karyotype of unstained chromosomes?

4. Which pair of chromosomes can contain two very different chromosomes and still be considered normal? Explain your answer.

5. How do autosomes differ from sex chromosomes?

Making Karyotypes Chapter 14 The Human Genome

Albert A Gayle



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Table of Contents Making Karyotypes Chapter 14 The Human Genome

1. Understanding the eBook Making Karyotypes Chapter 14 The Human Genome
 - The Rise of Digital Reading Making Karyotypes Chapter 14 The Human Genome
 - Advantages of eBooks Over Traditional Books
2. Identifying Making Karyotypes Chapter 14 The Human Genome
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Making Karyotypes Chapter 14 The Human Genome
 - User-Friendly Interface
4. Exploring eBook Recommendations from Making Karyotypes Chapter 14 The Human Genome

- Personalized Recommendations
- Making Karyotypes Chapter 14 The Human Genome User Reviews and Ratings
- Making Karyotypes Chapter 14 The Human Genome and Bestseller Lists
- 5. Accessing Making Karyotypes Chapter 14 The Human Genome Free and Paid eBooks
 - Making Karyotypes Chapter 14 The Human Genome Public Domain eBooks
 - Making Karyotypes Chapter 14 The Human Genome eBook Subscription Services
 - Making Karyotypes Chapter 14 The Human Genome Budget-Friendly Options
- 6. Navigating Making Karyotypes Chapter 14 The Human Genome eBook Formats
 - ePub, PDF, MOBI, and More
 - Making Karyotypes Chapter 14 The Human Genome Compatibility with Devices
 - Making Karyotypes Chapter 14 The Human Genome Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Making Karyotypes Chapter 14 The Human Genome
 - Highlighting and Note-Taking Making Karyotypes Chapter 14 The Human Genome
 - Interactive Elements Making Karyotypes Chapter 14 The Human Genome
- 8. Staying Engaged with Making Karyotypes Chapter 14 The Human Genome
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Making Karyotypes Chapter 14 The Human Genome
- 9. Balancing eBooks and Physical Books Making Karyotypes Chapter 14 The Human Genome
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Making Karyotypes Chapter 14 The Human Genome
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Making Karyotypes Chapter 14 The Human Genome
 - Setting Reading Goals Making Karyotypes Chapter 14 The Human Genome
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Making Karyotypes Chapter 14 The Human Genome

- Fact-Checking eBook Content of Making Karyotypes Chapter 14 The Human Genome
 - Distinguishing Credible Sources
13. Promoting Lifelong Learning
- Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
14. Embracing eBook Trends
- Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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