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#### Recombinant DNA LAB - Making Paper Plasmids

Standard: 5c Objective:

To understand how genes can be inserted into another DNA - "recombine"

To conceptualize "restriction enzymes" and recognition of specific sites

Students will model the process of using restriction enzymes and plasmids to form recombinant DNA.

The "Recombinant DNA Lab" activity can help you see how genes may be manipulated for genetic research, namely, gene cloning/genetic engineering. This activity will help you conceptualize the mechanics involved in cutting and ligating DNAs into a plasmid vector with "sticky ends" of complementary DNA base pairs.

#### **Background Information**

Recombinant DNA technology is one of the new techniques of biotechnology. Biotechnology uses living organisms to carry out chemical processes or to produce substances, combining biology with chemistry and science with industry. Biotechnology includes the field of **genetic engineering**, which is the science of directly manipulating genes to produce recombinant DNA. Genetic engineering techniques are used in a variety of industries, in agriculture, in basic research, and in medicine.

Through genetic engineering, there is great potential for the development of useful products. Researchers have already developed sources of interferon, human growth hormone, insulin, and resistant fruits and vegetables. A gene coding for a particular protein is transferred into a host organism. The host organism reproduces and the population produces the desired protein in volume. For example, the gene that codes for the production of human insulin has been inserted into the common bacterium E. coli. Then the bacteria can be grown in huge containers and large amounts of insulin can be collected.

As an introduction to recombinant DNA technology, the exercise that follows illustrates on paper some of the steps of recombinant DNA experiments done in laboratories.

#### Steps to Recombination

- Scientists must first identify the gene that codes for the production of the protein they want to manufacture.
- 2. Next scientists must isolate the desired gene. Restriction enzymes from bacterial cells are important in this step. Each restriction enzyme recognizes and cleaves (cuts) a very specific sequence of DNA called a restriction site. Some restriction enzymes make a staggered cut of the DNA producing sticky ends or single strands of unpaired nucleotide bases capable of binding with complementary sticky ends. By using restriction enzymes that will cut the DNA on either side of the gene, the desired gene can be clipped out of the DNA strand.



3. Once the desired gene has been isolated scientists must then insert the gene into a plasmid. Plasmids are small circular pieces of DNA found naturally in most bacteria. The plasmid must be removed from a bacterium and cleaved with the same restriction enzyme used to isolate the gene. This way the sticky ends of the plasmid will match those of the gene.

# **Recombinant Paper Plasmids Lab Answers**

**Michael Henson** 

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## **Table of Contents Recombinant Paper Plasmids Lab Answers**

- 1. Understanding the eBook Recombinant Paper Plasmids Lab Answers
  - The Rise of Digital Reading Recombinant Paper Plasmids Lab Answers
  - Advantages of eBooks Over Traditional Books
- 2. Identifying Recombinant Paper Plasmids Lab Answers
  - 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 Recombinant Paper Plasmids Lab Answers
  - User-Friendly Interface
- 4. Exploring eBook Recommendations from Recombinant Paper Plasmids Lab Answers
  - Personalized Recommendations
  - Recombinant Paper Plasmids Lab Answers User Reviews and Ratings
  - Recombinant Paper Plasmids Lab Answers and Bestseller Lists

- 5. Accessing Recombinant Paper Plasmids Lab Answers Free and Paid eBooks
  - o Recombinant Paper Plasmids Lab Answers Public Domain eBooks
  - Recombinant Paper Plasmids Lab Answers eBook Subscription Services
  - Recombinant Paper Plasmids Lab Answers Budget-Friendly Options
- 6. Navigating Recombinant Paper Plasmids Lab Answers eBook Formats
  - ∘ ePub, PDF, MOBI, and More
  - Recombinant Paper Plasmids Lab Answers Compatibility with Devices
  - Recombinant Paper Plasmids Lab Answers Enhanced eBook Features
- 7. Enhancing Your Reading Experience
  - Adjustable Fonts and Text Sizes of Recombinant Paper Plasmids Lab Answers
  - Highlighting and Note-Taking Recombinant Paper Plasmids Lab Answers
  - Interactive Elements Recombinant Paper Plasmids Lab Answers
- 8. Staying Engaged with Recombinant Paper Plasmids Lab Answers
  - Joining Online Reading Communities
  - Participating in Virtual Book Clubs
  - $\circ\,$  Following Authors and Publishers Recombinant Paper Plasmids Lab Answers
- 9. Balancing eBooks and Physical Books Recombinant Paper Plasmids Lab Answers
  - Benefits of a Digital Library
  - Creating a Diverse Reading Collection Recombinant Paper Plasmids Lab Answers
- 10. Overcoming Reading Challenges
  - Dealing with Digital Eye Strain
  - Minimizing Distractions
  - Managing Screen Time
- 11. Cultivating a Reading Routine Recombinant Paper Plasmids Lab Answers
  - Setting Reading Goals Recombinant Paper Plasmids Lab Answers
  - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Recombinant Paper Plasmids Lab Answers
  - Fact-Checking eBook Content of Recombinant Paper Plasmids Lab Answers
  - 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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