

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

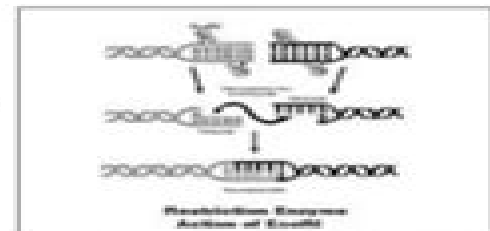
Recombinant DNA technology is one of the new techniques of biotechnology. Biotechnology uses living organisms to carry out chemical processes or to produce substances, combining biology with chemistry and science with industry. Biotechnology includes the field of **genetic engineering**, which is the 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

1. 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 Dna Paper Lab Answers

Anthony S. Fauci



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Table of Contents Recombinant Dna Paper Lab Answers

1. Understanding the eBook Recombinant Dna Paper Lab Answers
 - The Rise of Digital Reading Recombinant Dna Paper Lab Answers
 - Advantages of eBooks Over Traditional Books
2. Identifying Recombinant Dna Paper 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 Dna Paper Lab Answers
 - User-Friendly Interface
4. Exploring eBook Recommendations from Recombinant Dna Paper Lab Answers

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

- Fact-Checking eBook Content of Recombinant Dna Paper 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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