

DNA Replication Webquest

Answer all questions in a **DIFFERENT COLOR** and **IN YOUR OWN WORDS**.

Watch the animations and answer these questions:

http://www.wiley.com/college/pratt/0471393878/student/animations/dna_replication/index.html

1. What is DNA replication? **The process where an entire double-stranded DNA is copied to produce a second, identical DNA double helix.**
2. How does DNA replicate? (highlight one) A. Conservatively **B. Semi-conservatively** C. Dispersively
3. Semi-conservative means **When each resulting DNA double helix retains one strand of the original DNA**
4. Watch section 3 in its entirety. Then define the function of the following players in DNA replication..... If needed, go back and watch it again.
 - a. Polymerase - **extends the DNA chain**
 - b. Helicase - **unwinds the DNA double helix into two individually strands**
 - c. Ligase - **links the short DNA chains**
 - d. Primase - **creates RNA primers**
 - e. SSB - **prevents reannealing**
5. What is the first step of DNA replication? **The two double helix strands are unwound and separated from each other by the helicase enzyme.**
6. How does the DNA get unwind and stay unwind? **Its get unwinded by the Helicase.**
7. What is the replication fork? **The point where the DNA is separated into single strands, and where new DNA will be synthesized. The area where the parental DNA strands are copied to create two daughter strands.**
8. In what direction can DNA polymerase make new strand of DNA? **It starts at the 3' end of the RNA primer and creates a new strand in the 5' to 3' order.**
9. In other words, DNA polymerase adds nucleotides to the '3 end of the DNA strand.
10. What is needed before DNA polymerase can start building a new DNA strand? **A template is always needed. They can't start making a DNA chain from scratch, only from the 3' end of the primer.**
11. Contrast the leading and lagging strands. **The leading strand is single DNA strand and is replicated in the 3' to 5' direction. The lagging strand one of two strands of the DNA at the replication fork and needs a slight delay. It is also synthesized as a series of Okazaki fragments.**
12. The short pieces of DNA formed on the lagging strand are called... **Okazaki fragments.**

Recombinant Dna Webquest Answers

Xiang Xie



Recombinant Dna Webquest Answers:

Recombinant DNA and Biotechnology Helen Kreuzer, Adrienne Massey, 1996 Since the last edition was published more European legislation has been incorporated into the law of the United Kingdom and the third edition contains a full account of the 1992 regulations implementing European directives The Treaty of Amst **DNA Science** David A. Micklos, Greg A. Freyer, 1990 This laboratory text combines the theory practice and applications of recombinant DNA technology into one articulated package Unlike super texts that can only be sampled by even the most ambitious instructor or student DNA Science is designed to be read from cover to cover The eight text chapters are written in a semi journalistic style and adopt a historical perspective to explain where DNA science has come from and where it is going Combining the unique perspectives of both a research biologist and a science writer the topical treatment integrates up to the minute examples drawn directly from the research literature Extensively tested by thousands of high school and college teachers and students in 25 states and Canada the ten laboratory experiments cover the basic techniques of gene isolation and analysis The experiments engender systematic repetition to build student confidence and mastery of techniques Extensive prelab notes at the beginning of each experiment explain how to schedule and prepare and flowcharts and icons make the protocols easy to follow The laboratory course is completely supported by quality assured Carolina Biological Supply Company products from bulk reagents to reusable reagent systems to single use kits satisfying a range of teaching applications Truly a first course in recombinant DNA technology the laboratory sequence presupposes no prior experience on the part of the instructor or student Structured to follow directly from an introduction to principles of biology the experiments are equally appropriate for the advanced high school student and the beginning college student The book can be used as the first course in a molecularbiology sequence be integrated as a genetics DNA structure component of a general biology course or be used as a unit within a microbiology or genetics course The text is suitable for introducing recombinant DNA in science and society courses An Introduction to Recombinant DNA Alan E. H. Emery, 1984-11-27 After presenting a brief historical review this introduction to recombinant DNA technology deals with the essentials of the technology and the light it has shed on the molecular basis of various genetic disorders as well as common diseases such as diabetes coronary artery disease and cancer The applications of the technology in prenatal diagnosis the synthesis of vaccines and other medically important products and treatments through gene therapy are also reviewed Some broader applications with respect to human evolution and various agricultural commercial and industrial uses are also discussed The final chapters of the book examine the biohazards ethical and legal problems raised by the technology and discuss possible future developments *Recombinant DNA Technology* Keya Chaudhuri, 2013-01-01 Recombinant DNA Technology is focussed on the current state of knowledge on the recombinant DNA technology and its applications The book will provide comprehensive knowledge on the principles and concepts of recombinant DNA technology or genetic engineering protein expression of cloned genes PCR amplification of

DNA RFLP AFLP and DNA fingerprinting and finally the most recent siRNA technology It can be used by post graduate students studying and teachers teaching in the area of Molecular Biology Biotechnology Genetics Microbiology Life Science Pharmacy Agriculture and Basic Medical Sciences Guidelines for Research Involving Recombinant DNA Molecules National Institutes of Health (U.S.),1976 **Recombinant DNA Technology** Aleš Prokop, **Recombinant DNA and Biotechnology** Helen Kreuzer,1996 **Recombinant DNA and Genetic Experimentation** Joan Morgan,W. J. Whelan,2013-10-22 Recombinant DNA and Genetic Experimentation contains papers from the Proceedings of a Conference on Recombinant DNA held in London on April 1 4 1979 This books reviews recombinant DNA research and discusses advances in the application of recombinant DNA research and the regulations affecting such research Part 1 of the book deals with recombinant DNA techniques that are useful in the biological perspective These techniques include tests for rare gene exchanger and laboratory genetic manipulations Part 2 addresses the achievements of recombinant DNA research such as the detection of homologous sequences and progress made in the research of animal viruses Part 3 discusses the practical benefits of recombinant DNA research covering topics such as the production of valuable proteins in alternate biological hosts These proteins are shown as being valuable to society besides being scientific curiosities An important presentation is Part 4 of the symposium which discusses the guidelines and legislations affecting recombinant DNA research such as prior restraint prohibitions risks and approval of the conduct of such experiments Part 5 concerns a review of the basic assumptions made in the symposium while Part 6 tackles the question of what options are left open in the international arena in the medical field and in the eyes of the public This collection of papers can prove beneficial for molecular biologists DNA researchers molecular geneticists ecologists and endocrinologists and pharmacologists **Recombinant DNA Technology and Applications** Aleš Prokop,1991 **Recombinant DNA Technical Bulletin** ,1977 Recombinant DNA James D. Watson,1992 *Recombinant DNA Technology* Sardul Singh Sandhu,2010-06-30 The objective of the book is to introduce the basic principle and techniques used to make Recombinant DNA The book commences with an introduction to different tools used for Gene cloning The final chapters cover the application of Recombinant Technology on current research and provide an inside look on Human Genome Project Ribozyme Technology Antisense technology DNA sequencing Protein Engineering Transgenic technology and development of vaccines It features summary of chapter in the form of flow charts highlighting the key points The book also includes an appendix which provides in depth descriptions of protocols which cover the basic aspects of Molecular biology and glossary defining nearly all the possible terms mentioned in the book The purpose of this book is to provide an insight on theoretical aspects of Recombinant DNA manipulation with special emphasis on different procedures to create chimeric molecules using examples from actual experimental works The book has been designed for under graduates post graduates and technicians who wish to know and use the principles and techniques of Recombinant DNA Technology **Recombinant DNA [Deoxyribonucleic Acid] and genetic experimentation** ,1979

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Recombinant DNA Principles and Methodologies James Greene,2021-12-16 This comprehensive yet balanced work emphasizes the principles and rationale underlying recombinant DNA methodology while furnishing a general understanding of the experimental protocols suggesting flexible approaches to resolving particular molecular necessities that are easily adaptable to readers specific applications Features summary tables presenting at a glance information on practices of recombinant DNA methodologies Recombinant DNA Principles and Methodologies discusses basic and advanced topics requisite to the employment of recombinant DNA technology such as plasmid biology nucleic acid biochemistry restriction enzymes cloning strategies gel electrophoresis southern and northern blotting preparation of probes phage lambda biology cosmids and genome analysis cloned gene expression polymerase chain reaction conventional and automated DNA sequencing site directed mutagenesis and more Elucidating the material with over 2250 edifying references equations

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