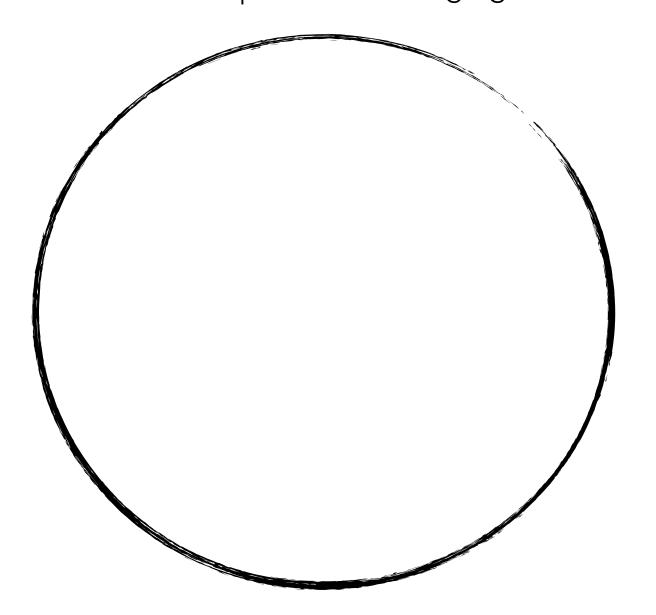


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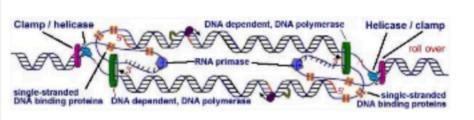
## course web site

**Group work**: outline the molecular activities (proteins) and processes involved in the replication and segregation of a circular chromosome.



Here is a circular DNA molecule; given what you know about the properties of DNA polymerase, draw and label a picture of the molecule halfway through its replication. Indicate the replication origin(s) and fork(s).

There is a mutation that inactivates a **single** protein. DNA synthesis begins and proceeds in both directions around the DNA molecule, but then slows and stops. Which protein is most likely to have been inactived by the mutation?

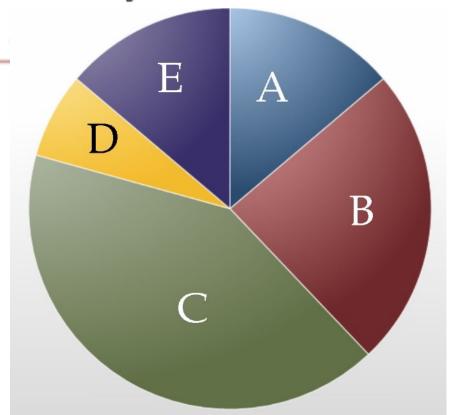


- A: Replication origin binding protein
- B: type I topoisomerase
- C: DNA-dependent, DNApolymerase

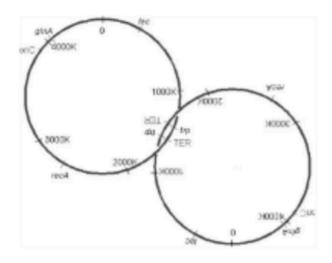
explain why the other choices cannot be correct

what inactivating mutations in the other activities do?

D: DNA ligase



There is a mutation that inactivates a single enzymatic activity. DNA synthesis begins and proceeds to completion, but the two DNA molecules remain intertwined. Which protein is most likely to be inactive?

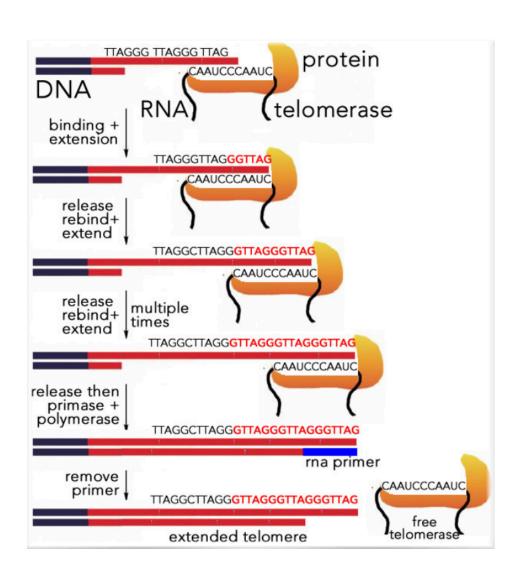


- A. Origin binding protein
- B. type I topoisomerase
- C. DNA-dependent, DNA-polymerase
- D. type II topoisomerase.
- \_ . .

**Group work**: outline the molecular activities (proteins) and processes involved in the replication and segregation of a linear chromosome.

Linear DNA molecules (telomeres)

## Linear DNA molecules (telomeres)



## **Questions to answer**

During DNA/RNA synthesis what is the average ratio of productive to unproductive interactions between an incoming nucleotide and the polymerase?

Why do you need to denature (melt) the DNA double-helix to copy it?

How would DNA replication change if H-bonds were as strong as covalent bonds?

How would evolution be impacted if DNA were totally stable and DNA replication was error-free?

What would be the impact of mutations that altered the proof-reading function of the DNA polymerase complex?

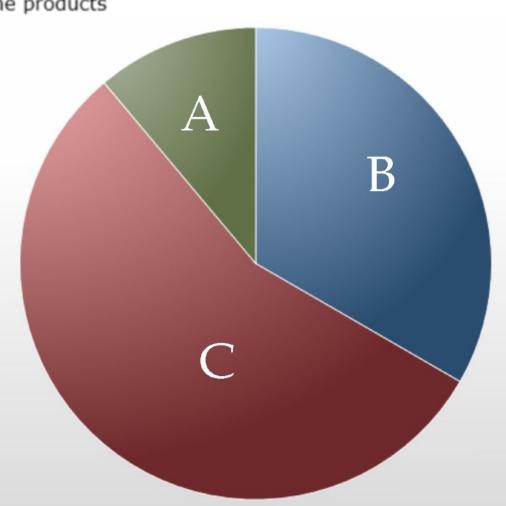
How might mutations in the genes encoding the clamp/clamp-loader system influence DNA replication?

Mutations DNA occur, how are they recognized?

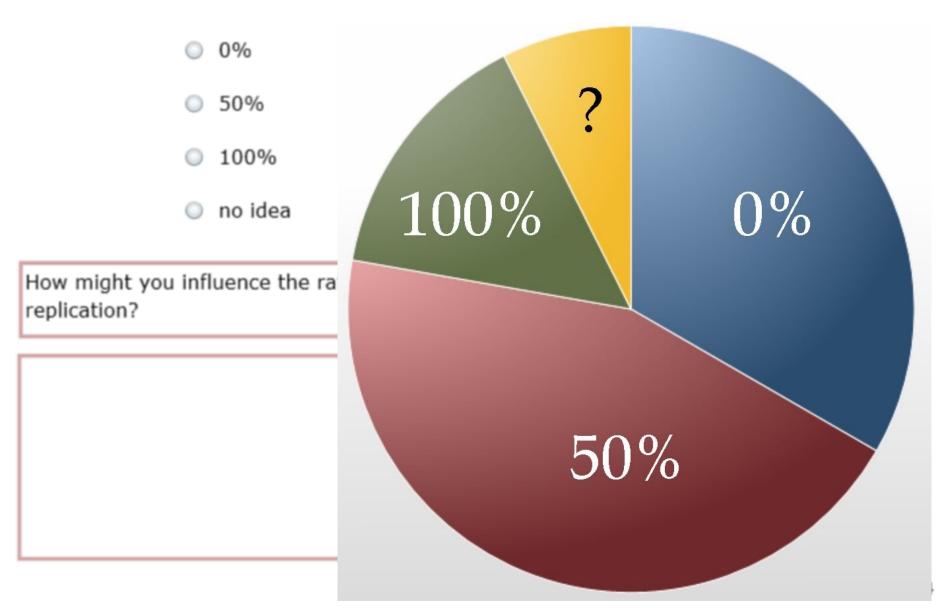
- there are proteins that know what the right sequence should be
- they change the structure of the DNA molecule
- they encode non-functional gene products
- no idea

What does it mean to be recognized?

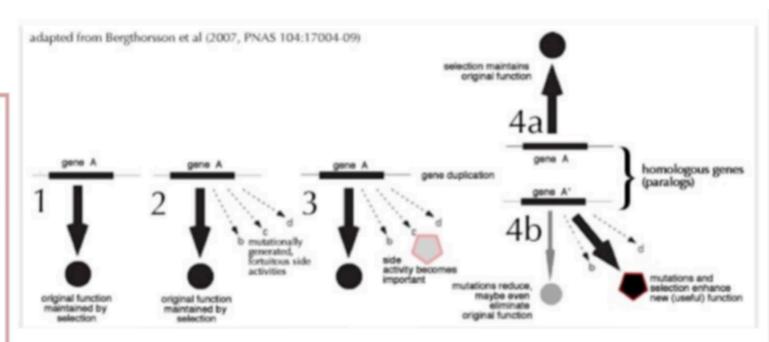
Why are A and C wrong?



During DNA replication mistakes can be made; if they were repaired randomly, what would be the probablilty that they would be repaired correctly



What is the advantage of duplicating genes, a opposed to developing new genetic activities de novo.



If you had to speculate, how would a repeat disease lead to changes in gene product functions?

would the effect be positive, negative, or neutral?

