

DNA replication

DNA Polymerase

→ Arthur Kornberg
→ 5' → 3' only

→ functions without RNA and DNA primer in test tube.

Okazaki fragments

Replication fork is asymmetrical.

Main reason → ATP requirement for DNA helicase

The 3' nucleotide has a triphosphate, that can be broken to release energy. In 5' end if there is triphosphate, where does the primer sit? → Why 5' → 3'

Fact: only one polymerase acts / two? X

Looping of lagging strand in replisome helps in coordinated movement of two strands

Joining of Okazaki Fragments

Why RNA as primer! (and not DNA)

RNAse H → 5' exonuclease

→ does it recognise 2' OH RNA

→ Yes

DNA polymerase III → elongation.

DNA polymerase I → filling of gaps.

↳ does this remove the primers?

↳ how can it sit without something

Fidelity of replication

→ does not allow incorrect base pairs

→ can recognise unpaired end.

→ can also recognise RNAs due to steric hindrance

Tautomers & incorrect base pairing

Structural isomers of chemical compounds that readily interconvert.

Thymine (enol), Cytosine (imino)

Transient → unpairs → recognised again

↳ as DNA pol moves slower than this change.

How does it slide back? → source of energy is ATP.

we should print pictures of DNA pol III.

CT on Monday HEHE