

- extremely accurate
- proofreads
- does not distinguish b/w DNA & RNA (!)
- very fast reaction due to processive nature

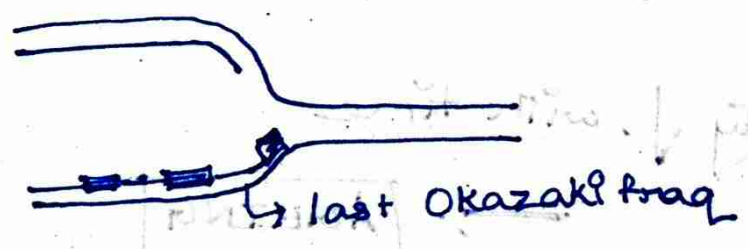
Replication at chromosomal ends:

- Repetitive telomere sequence (TTAGGG)
- Protects chromosome ends from damage
  - telomeric proteins

Why repetition? Why proteins?

Shelterin — protects telomere end — as the end is free and the end is perceived by DNA polymerase as a strand break (exonucleated)

End Replication Problem:



Primer removed → place not filled → bases lost and unreplicated

Replication cycles → chromosome gets shorter & shorter

# Telomerase

→ ribonucleoprotein

TERT - Telomerase reverse transcriptase

TER - Telomerase RNA

→ RNA dependent DNA polymerase

In telomerase → complementary RNA seq.



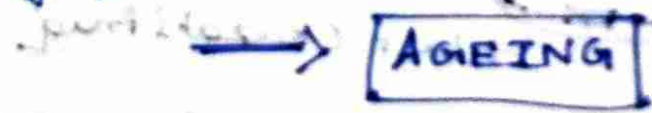
Telomere lengthened by several 1000 bases



After extension → shortening does not affect because of -15~20 bases

① Why TTAGGG

Telomerase activity ↓ with time



① Frequency of telomerase activity w.r.t. rep. cycle.

① Half-life of telomerase

# Regulation of replication

Cell cycle — blah blah blah

## Helicase — Loading and activation

helix → straight

→ Rate determining state

Loading — Binding of Helicase to ORC

Activation — Separation of strands.

MCM 2-7  
+ Cdt1

ORC

→ Complex

→ Cdt1 sep. through ATP.

Origin Recog. Complex

YAY NO REMEMBERING

CDK phosphorylates helicase → Activation

G<sub>1</sub> — loading  
(Act. inhibited)

G<sub>2</sub>, S, M

→ Activation  
(loading inhibited)