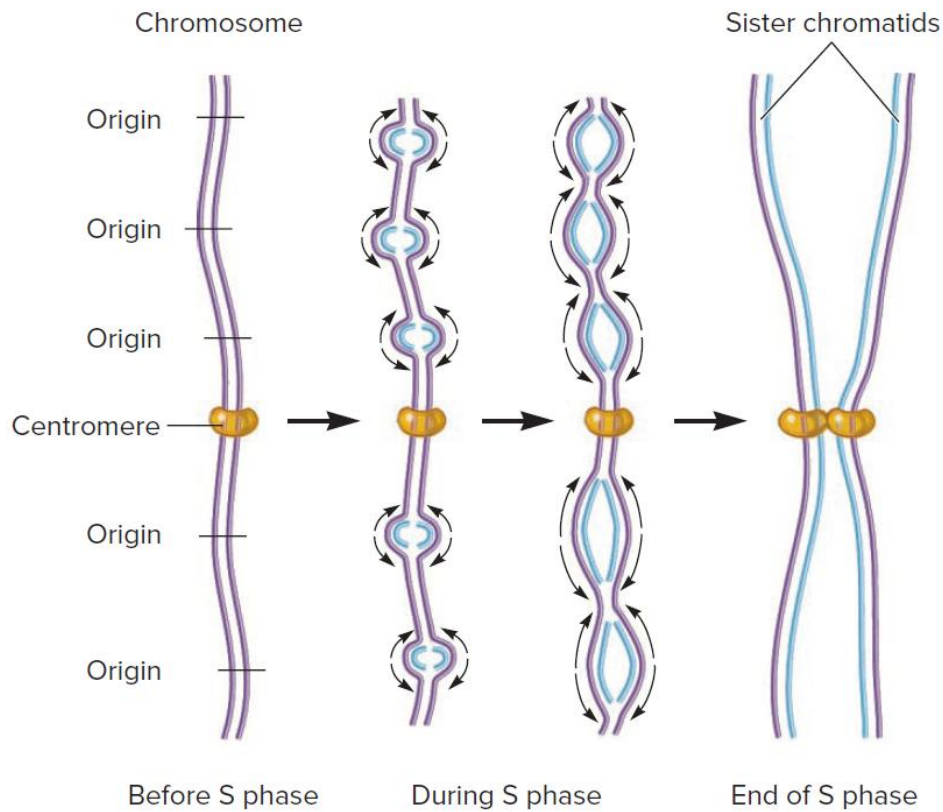
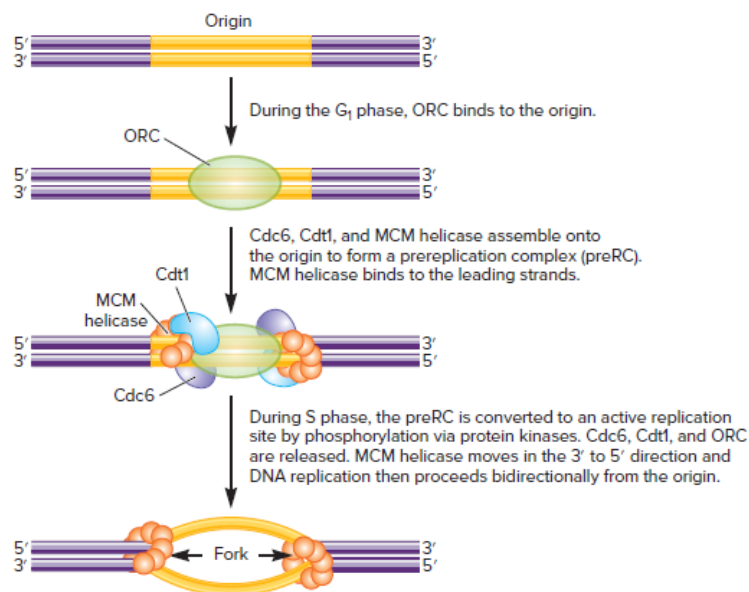


## La replication chez les eucaryotes TD 3



**FIGURE 11.19 The replication of eukaryotic chromosomes.** Replication begins from multiple origins of replication, and the replication forks move bidirectionally to replicate the DNA. Eventually, all of the replication forks will merge. The net result is two sister chromatids attached to each other at the centromere.

**CONCEPT CHECK:** Why do eukaryotes need multiple origins of replication?



11.5 EUKARYOTIC DNA REPLICATION

**FIGURE 11.20** The formation of a prereplication complex in eukaryotes. This is a simplified model; more proteins are involved in this process than are shown here.

The molecular features of eukaryotic origins of replication may have some similarities to the origins found in bacteria. At the molecular level, eukaryotic origins of replication have been extensively studied in the yeast *Saccharomyces cerevisiae*. In this organism, several replication origins have been identified and sequenced. They have been named **ARS elements** (for **autonomously replicating sequence**). ARS elements, which are about 50 bp in length, are

### Eukaryotes Have Many Different DNA Polymerases

Eukaryotes have many types of DNA polymerases. For example, mammalian cells have well over a dozen different DNA polymerases (**Table 11.4**). Four of these, designated  $\alpha$  (alpha),  $\epsilon$  (epsilon),  $\delta$  (delta), and  $\gamma$  (gamma), have the primary function of replicating DNA. DNA polymerase  $\gamma$  functions in the mitochondria to repli-

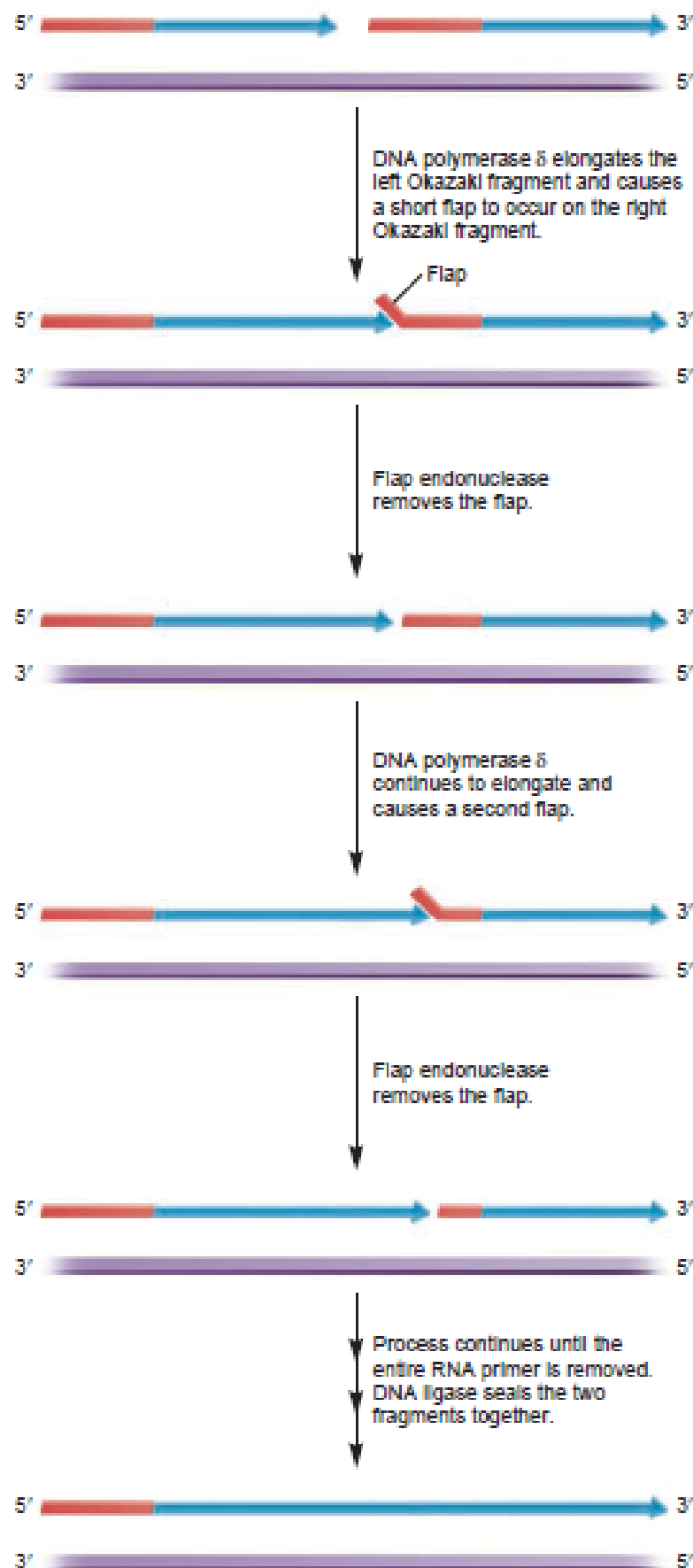
**TABLE 11.4**

#### Eukaryotic DNA Polymerases

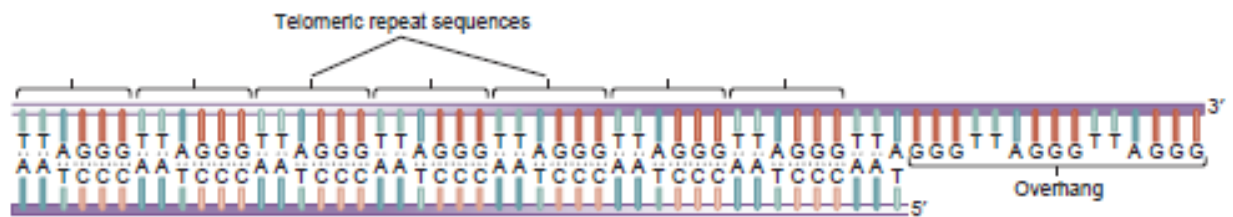
Polymerase Types*	Function
$\alpha$	Initiates DNA replication in conjunction with primase
$\epsilon$	Replication of the leading strand during S phase
$\delta$	Replication of the lagging strand during S phase
$\gamma$	Replication of mitochondrial DNA
$\eta$ , $\kappa$ , $\iota$ , $\xi$ (lesion-replicating polymerases)	Replication of damaged DNA
$\alpha$ , $\beta$ , $\delta$ , $\epsilon$ , $\sigma$ , $\lambda$ , $\mu$ , $\varphi$ , $\theta$ , $\eta$	DNA repair or other functions†

\*The designations are those of mammalian enzymes.

†Many DNA polymerases have dual functions. For example, DNA polymerases  $\alpha$ ,  $\delta$ , and  $\epsilon$  are involved in the replication of normal DNA and also play a role in DNA repair. In cells of the immune system, certain genes that encode antibodies (i.e., immunoglobulin genes) undergo a phenomenon known as hypermutation. This increases the variation in the kinds of antibodies the cells can make. Certain polymerases in this list, such as  $\eta$ , may play a role in hypermutation of immunoglobulin genes. DNA polymerase  $\sigma$  may play a role in sister chromatid cohesion, a topic discussed in Chapter 10.



**FIGURE 11.23** Removal of an RNA primer by flap endonuclease.

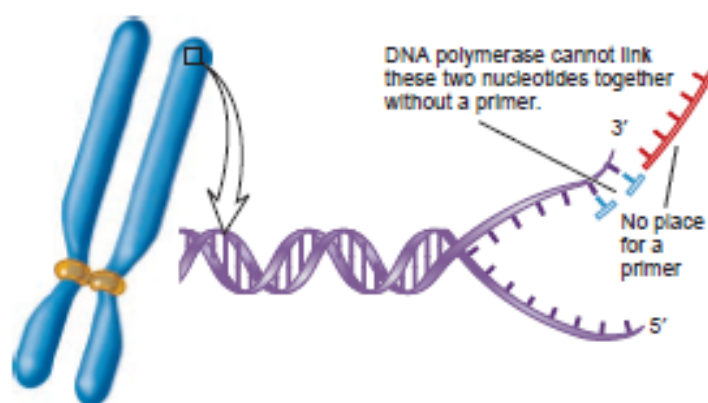


**FIGURE 11.24** General structure of telomeric sequences. The telomere DNA consists of a tandemly repeated sequence and a 12- to 16-nucleotide overhang.

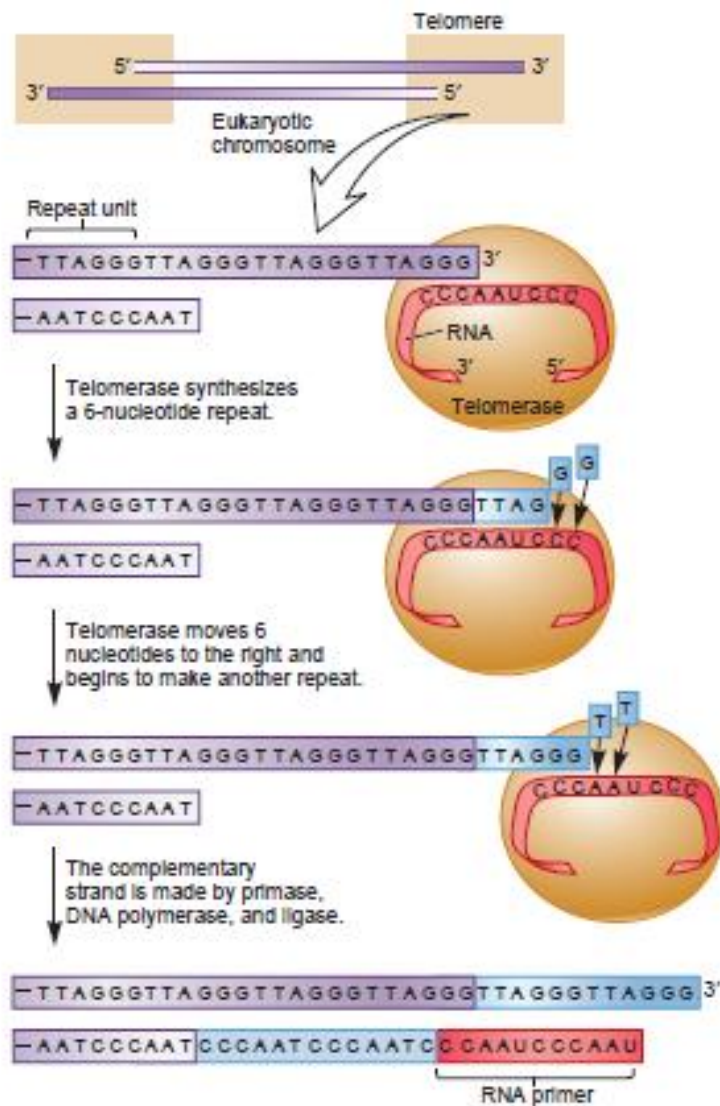
**TABLE 11.5**

**Telomeric Repeat Sequences Within Selected Organisms**

Group	Examples	Telomeric Repeat Sequence
Mammals	Humans	TTAGGG
Slime molds	<i>Physarum, Didymium</i>	TTAGGG
	<i>Dictyostelium</i>	$AG_{(1-4)}$
Filamentous fungi	<i>Neurospora</i>	TTAGGG
Budding yeast	<i>Saccharomyces cerevisiae</i>	$TG_{(1-3)}$
Ciliates	<i>Tetrahymena</i>	TTGGGG
	<i>Paramecium</i>	TTGGG(TG)
	<i>Euplotes</i>	TTTTGGGG
Higher plants	<i>Arabidopsis</i>	TTTAGGG



**FIGURE 11.25** The replication problem at the ends of linear chromosomes. DNA polymerase cannot synthesize a DNA strand that is complementary to the 3' end because a primer cannot be made upstream from this site.



**FIGURE 11.26 The enzymatic action of telomerase.**

A short, three-nucleotide segment of RNA within telomerase causes it to bind to the 3' overhang. The adjacent part of the RNA is used as a template to make a short, six-nucleotide repeat of DNA. After the repeat is made, telomerase moves six nucleotides to the right and then synthesizes another repeat. This process is repeated many times to lengthen the top strand shown in this figure. The bottom strand is made by DNA polymerase, using an RNA primer at the end of the chromosome that is complementary to the telomeric repeat sequence in the top strand. DNA polymerase fills in the region, which is sealed by ligase.

Bonne lecture mes grands

