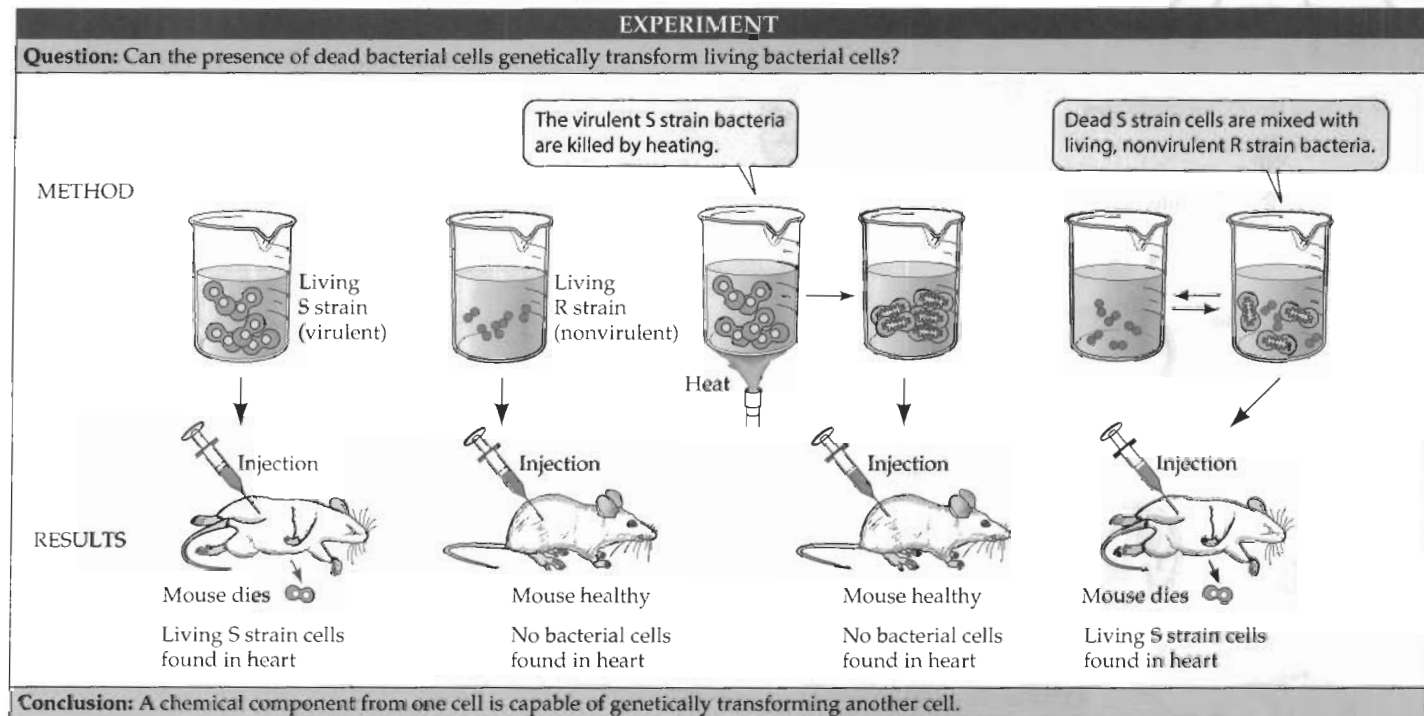


# 11

## DNA and Its Role in Heredity



### 11.1 Genetic Transformation of Nonvirulent Pneumococci (Page 214)

## EXPERIMENT

**Question:** Which component of a bacteriophage—DNA or protein—is the hereditary material that enters a bacterial cell to direct the assembly of new virus particles?

## Experiment 1

- 1a** T2 phage are grown in a medium containing  $^{32}\text{P}$  (P is an element in DNA but not in proteins).

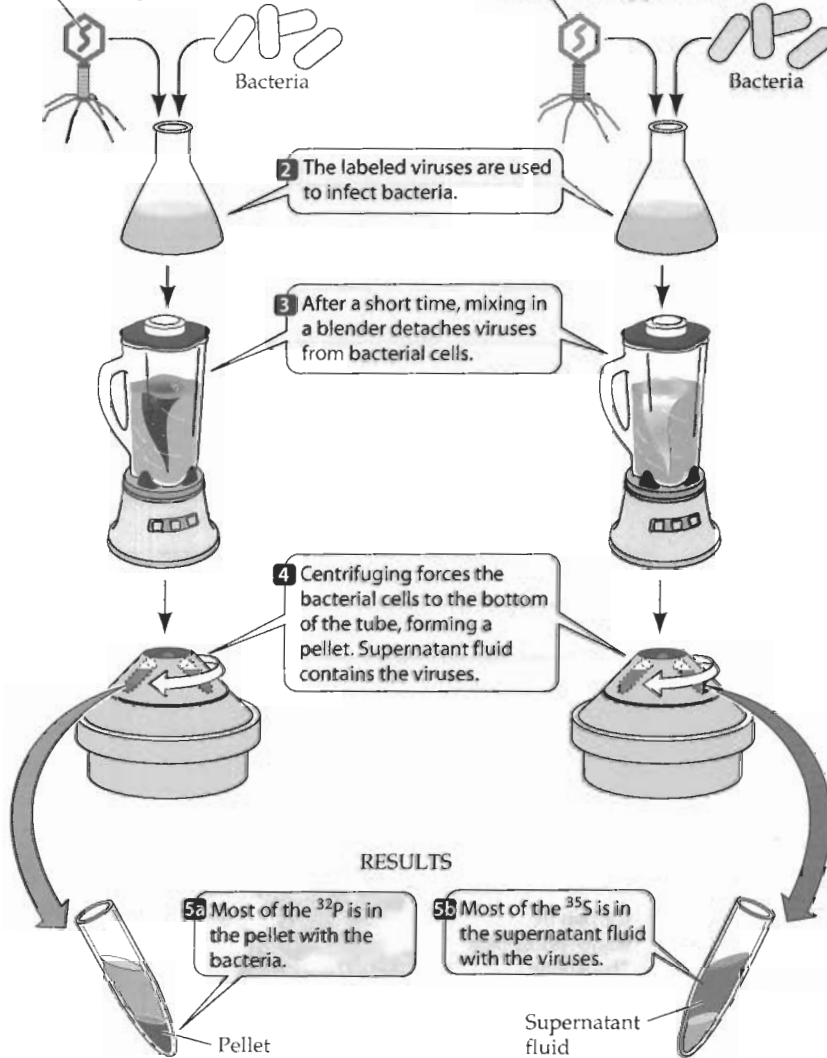
## Experiment 2

- 1b** T2 phage are grown in a medium containing  $^{35}\text{S}$  (S is an element in proteins but not in DNA).

$^{32}\text{P}$ -containing DNA

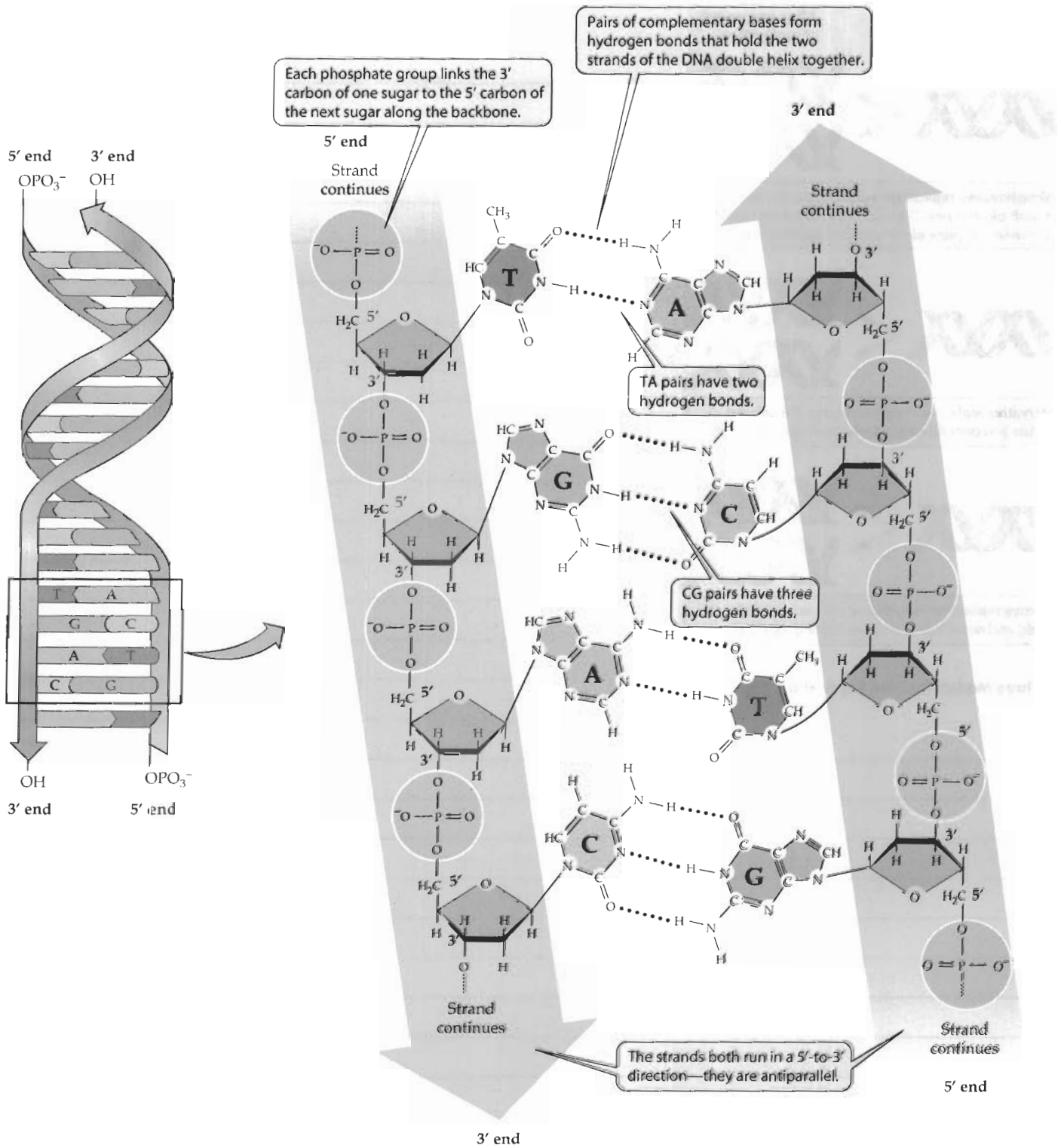
METHOD

$^{35}\text{S}$ -containing protein coats



**Conclusion:** DNA, not protein, enters bacterial cells and directs the assembly of new viruses.

### 11.3 The Hershey-Chase Experiment (Page 216)



### 11.7 Base Pairing in DNA Is Complementary (Page 219)

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## EXPERIMENT

**Question:** Does DNA replicate semiconservatively, or by some other mechanism?

## METHOD

1 Grow bacteria in  $^{15}\text{N}$  (heavy) medium.

2 Transfer some bacteria to  $^{14}\text{N}$  (light) medium; bacterial growth continues.

3 Before the bacteria reproduce the first time in the light medium, all DNA (parental) is heavy.

Sample at 0 minutes

Sample after 20 minutes

Sample after 40 minutes

4 Samples are taken after 0 minutes, 20 minutes (after one round of replication), and 40 minutes (two rounds of replication).

## RESULTS

$^{14}\text{N}/^{14}\text{N}$  (light) DNA

$^{14}\text{N}/^{15}\text{N}$  (intermediate) DNA

$^{15}\text{N}/^{15}\text{N}$  (heavy) DNA

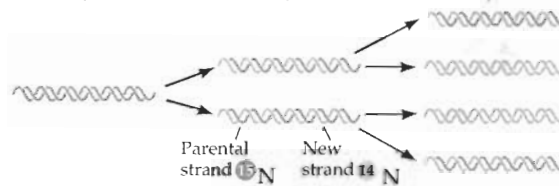
Parental  
(all heavy)

First generation  
(intermediate)

Second generation  
(half are all light)

5 If each strand served as a template for a new strand, DNA of the first generation would be of an intermediate density, and half the DNA from the second generation would be intermediate and half light. This is what was in fact observed.

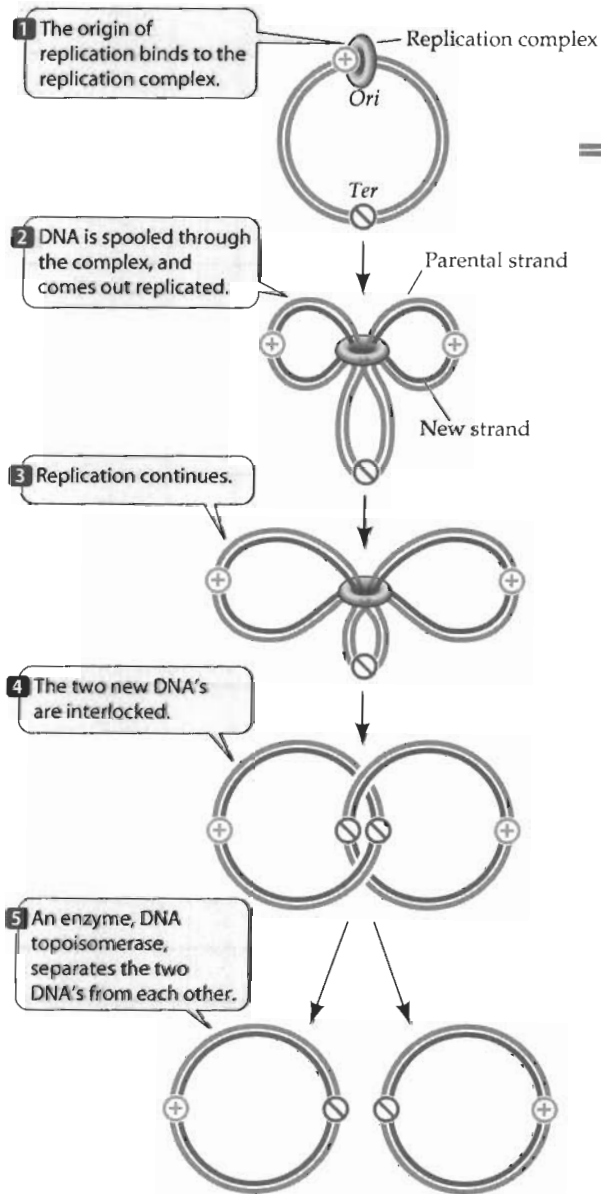
## INTERPRETATION



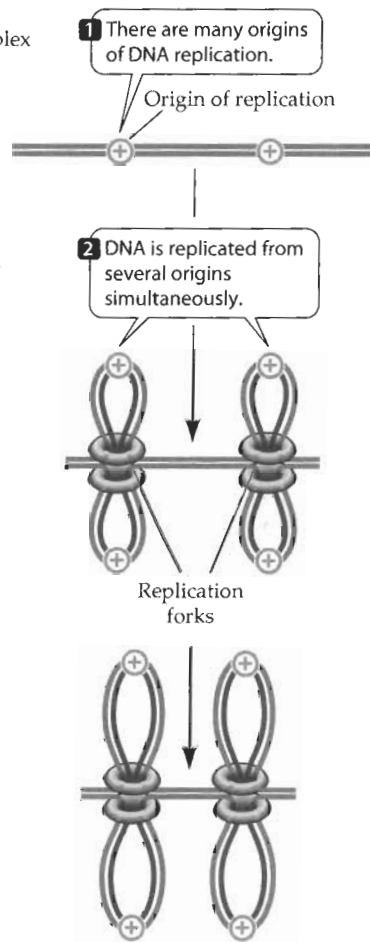
**Conclusion:** DNA replication is semiconservative.

## 11.9 The Meselson-Stahl Experiment (Page 221)

(a) Circular chromosome

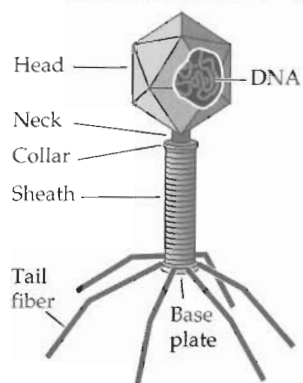


(b) Linear chromosome

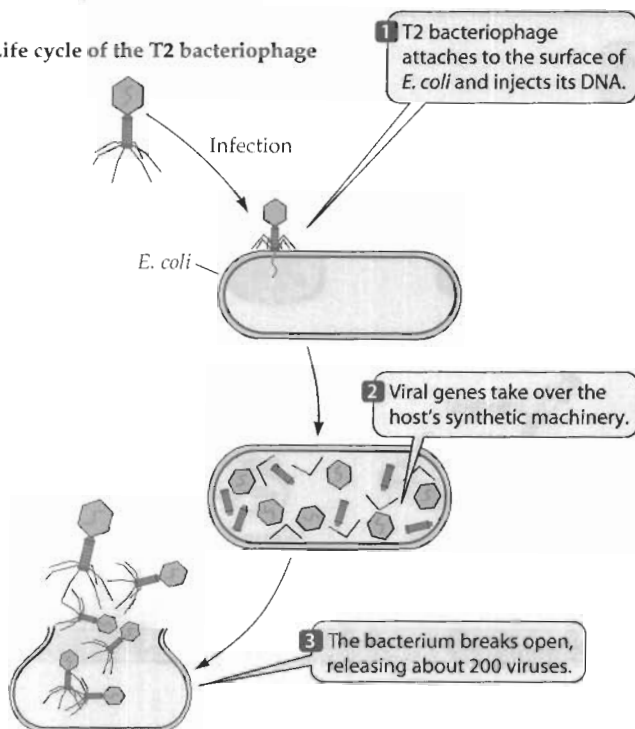


### 11.12 Replication in Small Circular and Large Linear Chromosomes (Page 224)

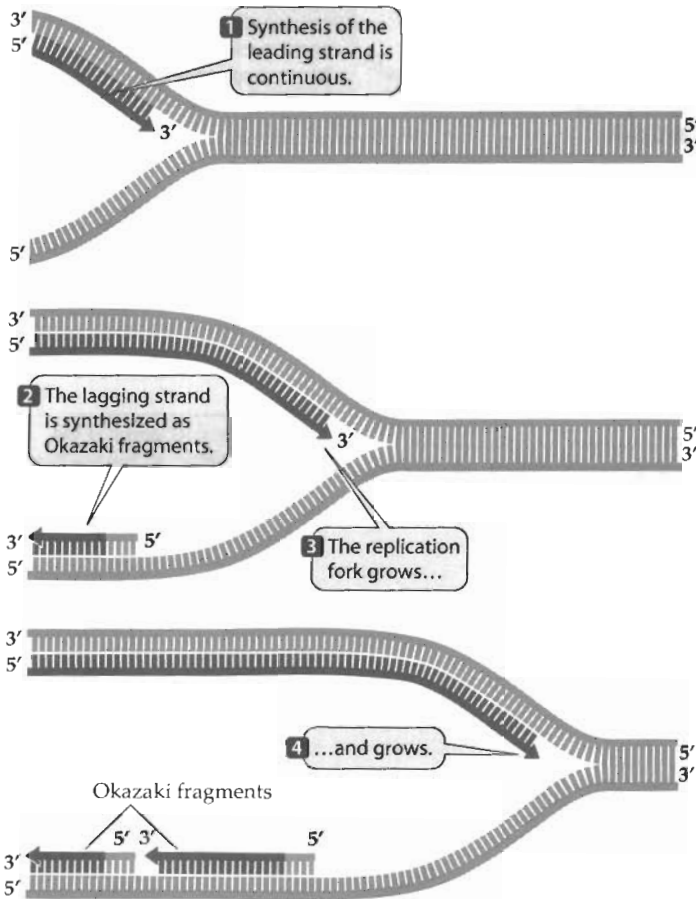
## (a) The virus: T2 bacteriophage



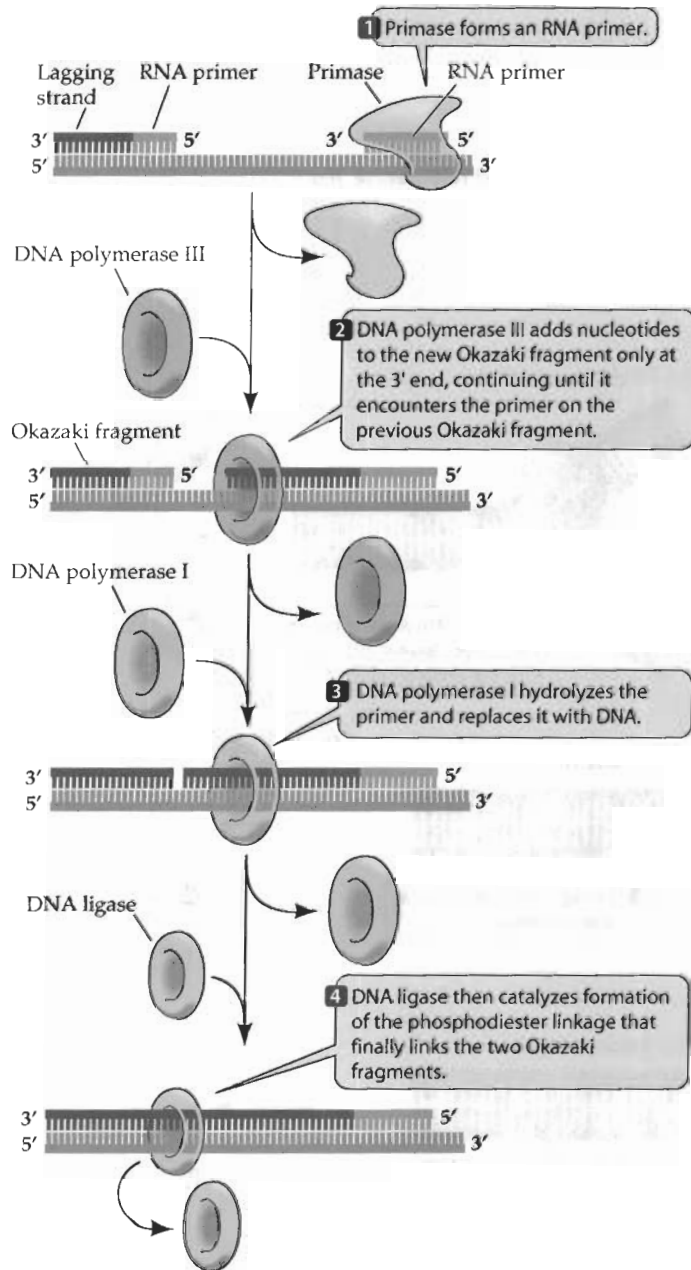
## (b) Life cycle of the T2 bacteriophage



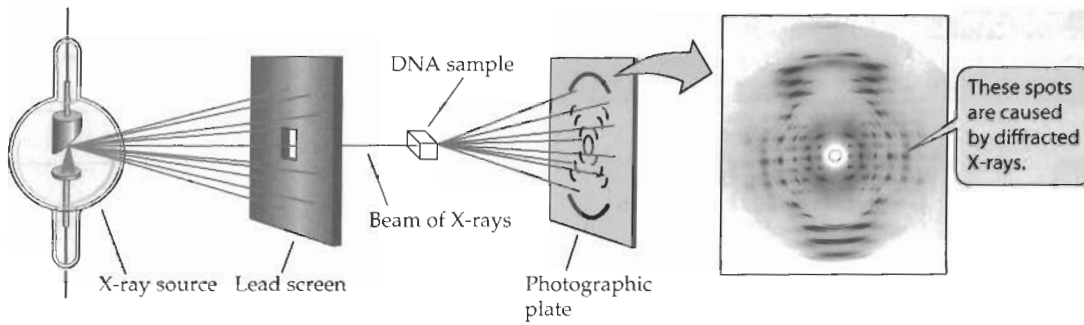
## 11.2 T2 and the Bacteriophage Reproduction Cycle (Page 215)



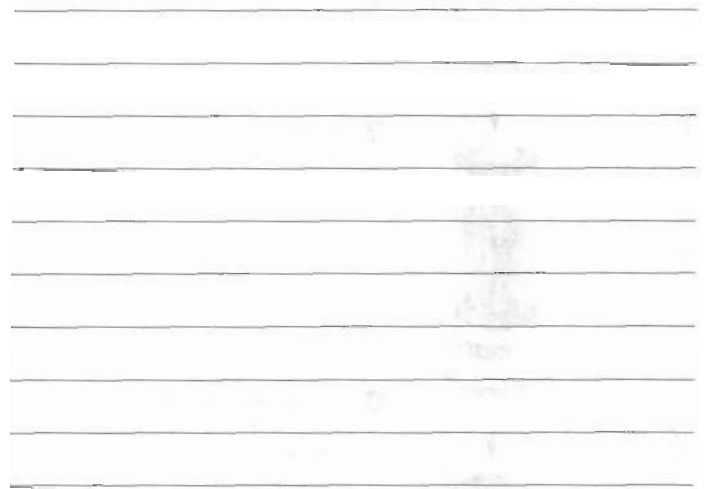
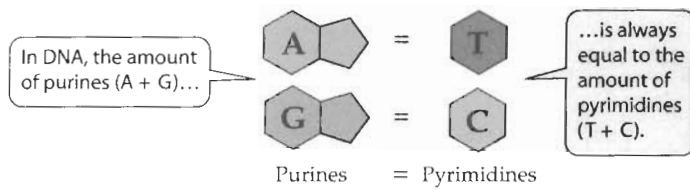
11.16 The Two New Strands Form in Different Ways (Page 225)



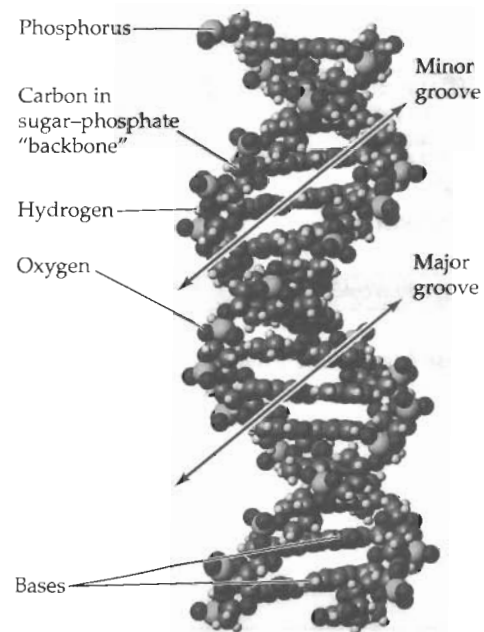
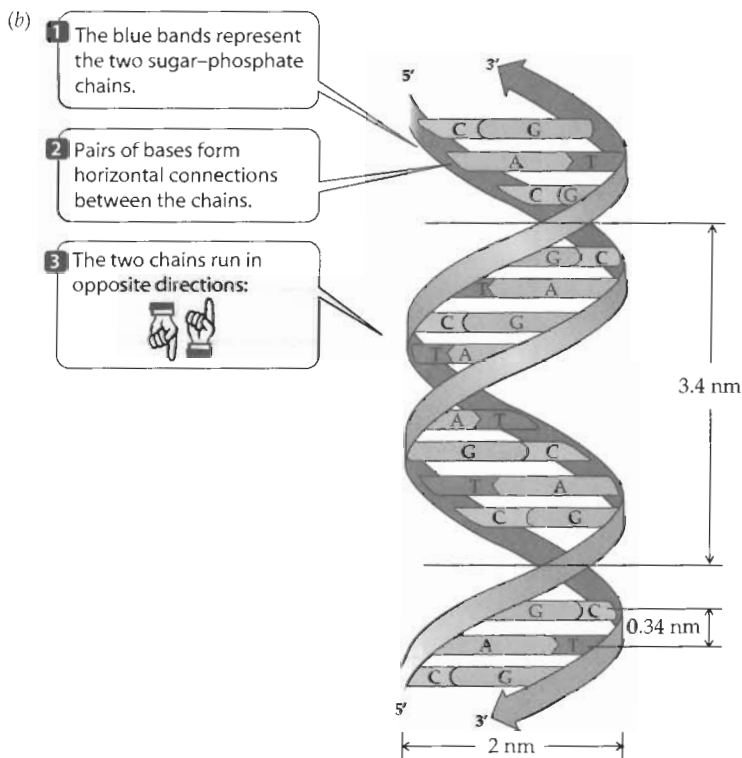
11.17 The Lagging Strand Story (Page 226)



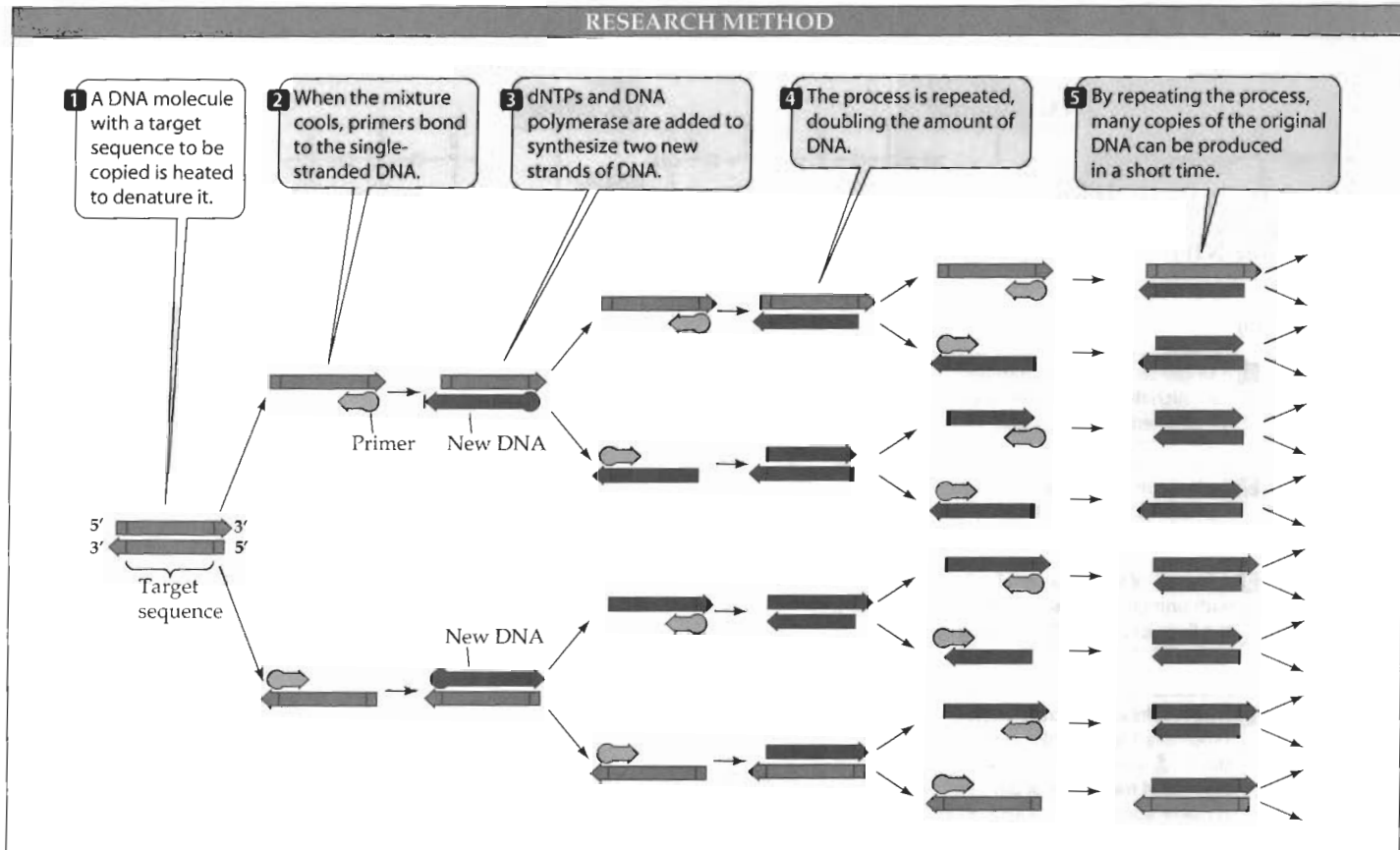
#### 11.4 X-Ray Crystallography Revealed the Basic Helical Structure of the DNA Molecule (Page 217)



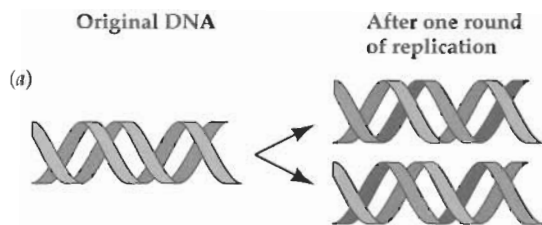
#### 11.5 Chargaff's Rule (Page 217)



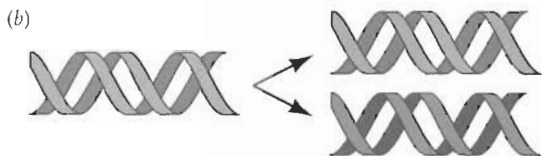
#### 11.6 DNA Is a Double Helix (Page 218)



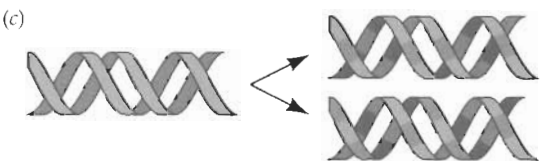
11.20 The Polymerase Chain Reaction (Page 229)



Semiconservative replication would produce molecules with both old and new DNA, but each molecule would contain one complete old strand and one new one.



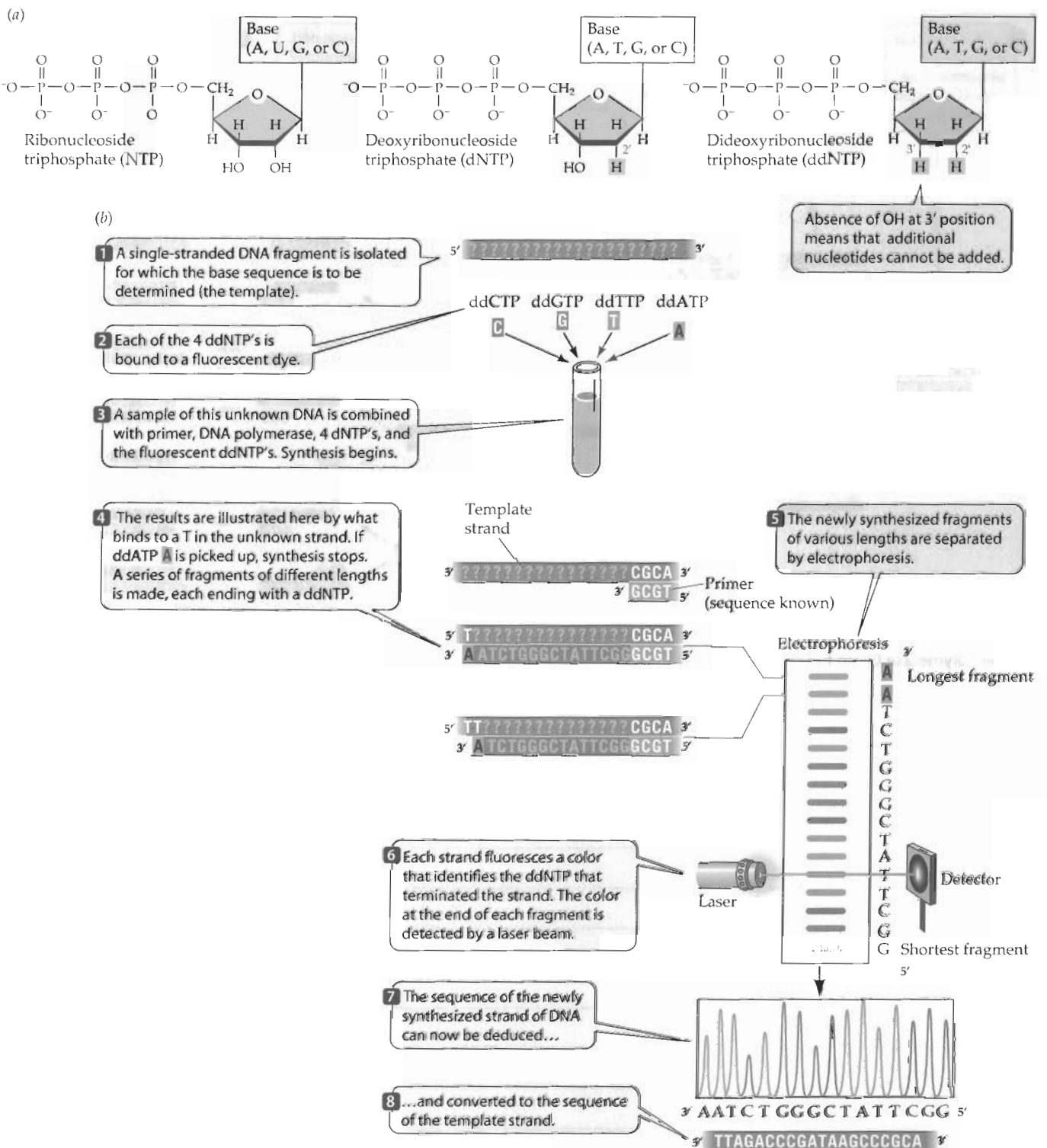
Conservative replication would preserve the original molecule and generate an entirely new molecule.



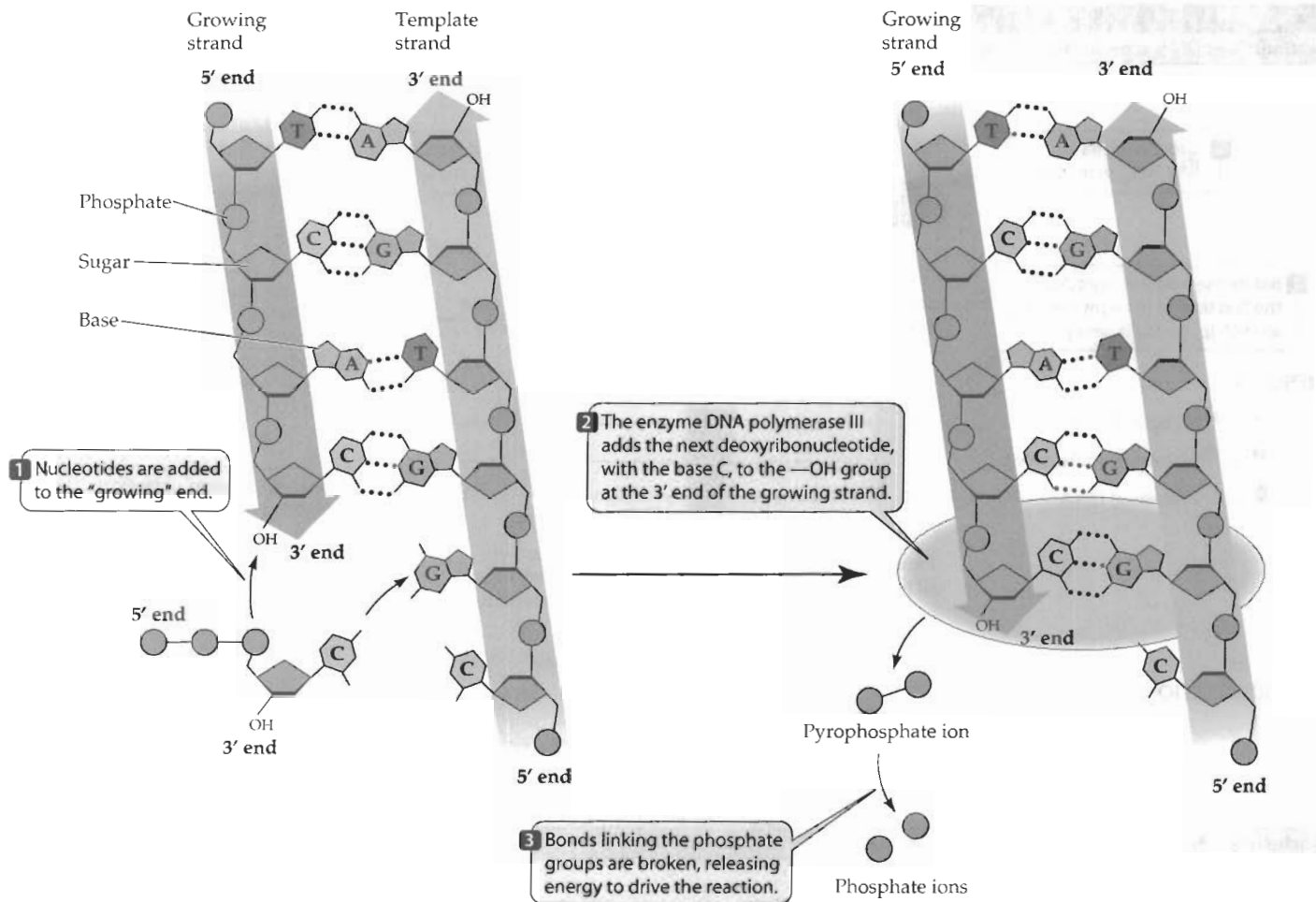
Dispersive replication would produce two molecules with old and new DNA interspersed along each strand.

### 11.8 Three Models for DNA Replication (Page 220)

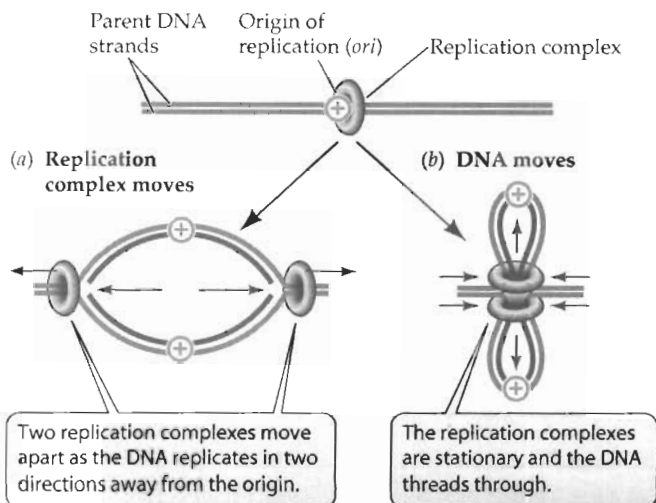
## RESEARCH METHOD



## 11.21 Sequencing DNA (Page 230)

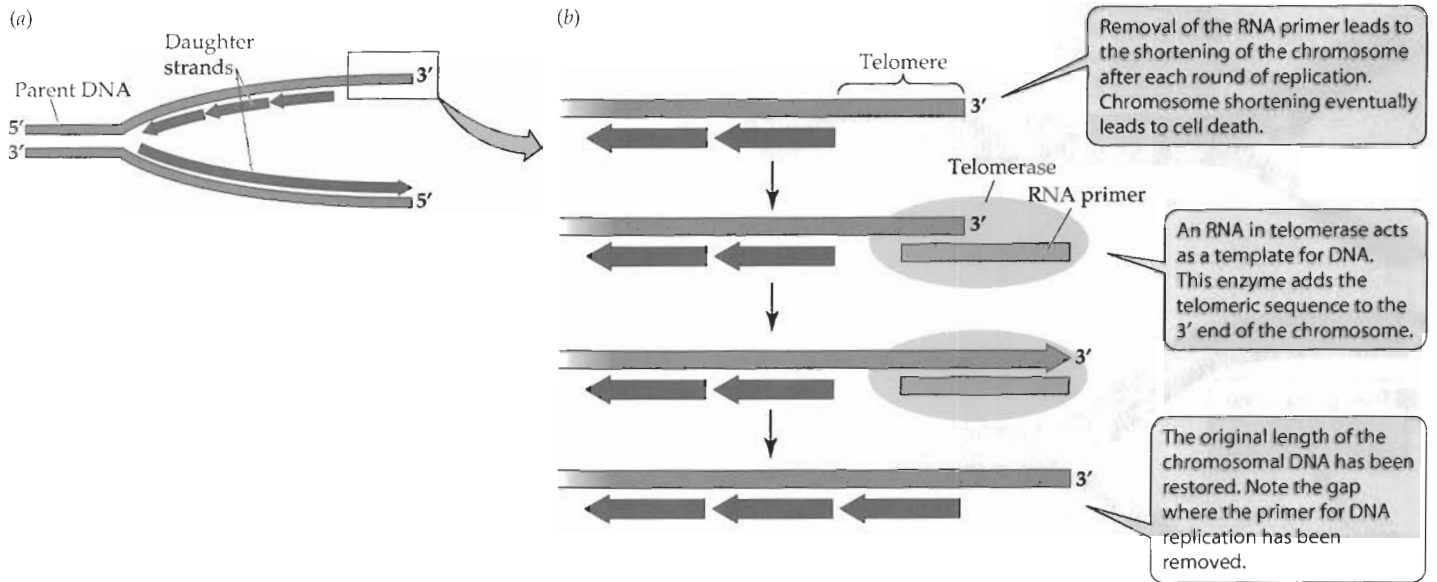


### 11.10 Each New DNA Strand Grows from its 5' End to its 3' End (Page 223)

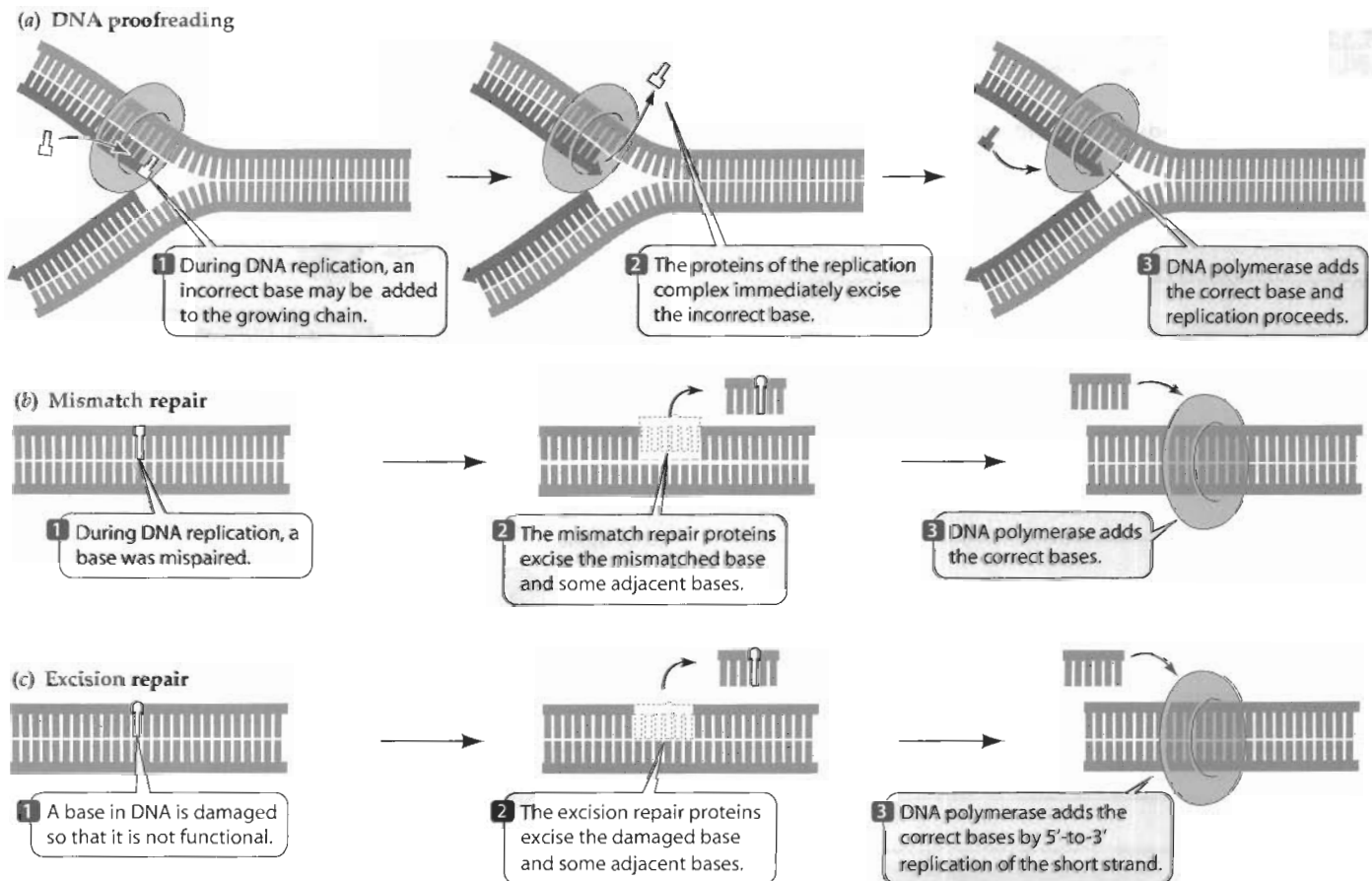


## 11.11 Two Views of DNA Replication (Page 223)

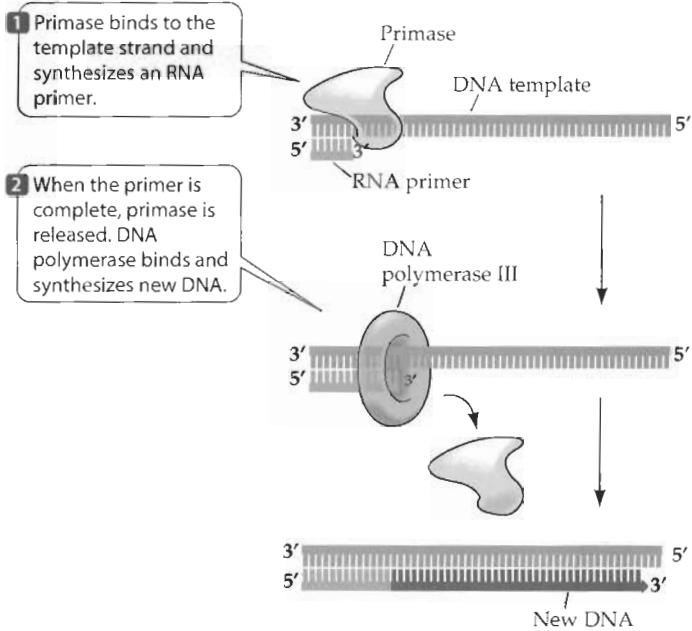
This image shows a single sheet of white paper with horizontal blue or grey ruling lines. The lines are evenly spaced and run across the width of the page. There are approximately 20 lines visible. The paper appears slightly aged or off-white.



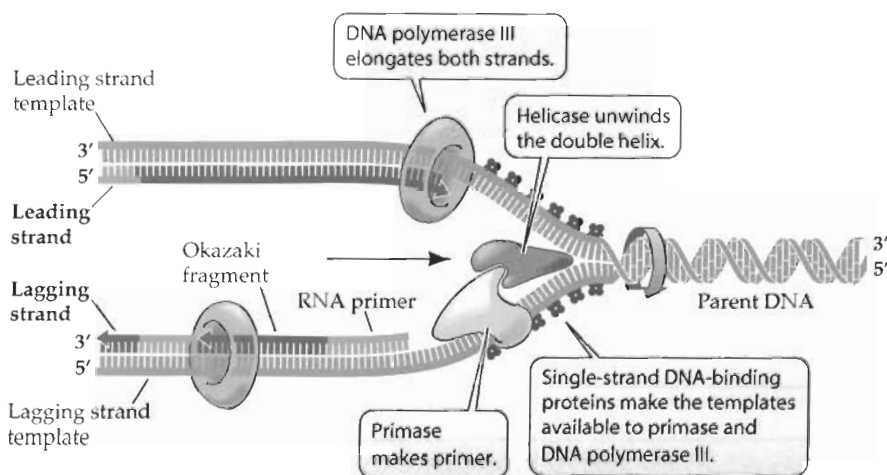
## 11.18 Telomeres and Telomerase (Page 227)



## 11.19 DNA Repair Mechanisms (Page 228)



#### 11.14 No DNA Forms without a Primer (Page 225)



#### 11.15 Many Proteins Collaborate at the Replication Fork (Page 225)