Bio.10-Q2W4-Qs.Bank-Bacteria and Viruses

Multiple Choice

Identify the choice that best completes the statement or answers the question.

 1.	Which of the bacteria is the cause of pneumoni a. staphylococci	a? c.	Treponema pallidum
2	D. FICKEUSIA	a.	streptococcus pneumoniae
 2.	a farming	C	the food industry
	b. the medical industry	d.	all of these
3.	A structure in some bacteria that is resistant to	adve	erse environmental factors is a(n)
 0.	a. prophage	с.	autotroph
	b. endospore	d.	coccus
 4.	Which of the following is NOT an evolutionary	y ada	aptation in bacteria?
	a. They reproduce rapidly.		-
	b. They have a high rate of mutation.		
	c. They cannot exist under adverse conditions	5.	
_	d. They can utilize substances harmful to othe	er or	ganisms.
 5.	Which of the following processes brings about	an e	exchange of genetic information between bacterial cells?
	a. Dinary lission b mutualism	c. d	replication
6	$\Lambda(n)$ is a virus that infacts a hactorial cal	u. 11	replication
 0.	a endospore	п. С	plasmid
	b. decomposer	d.	bacteriophage
7.	Viruses are found in .		
	a. air	c.	soil
	b. water	d.	all of these
 8.	Viruses are		
	a. producers	c.	parasites
	b. consumers	d.	decomposers
 9.	Penicillin kills bacteria by		
	a. consuming them		
	b. causing noises to develop in their cell wans		
	d. depriving them of nutrients		
10	The name streptococcus tells you that the bacte	ria a	are arranged as
 10.	a. pairs of round cells	с.	groups of spirals
	b. long chains of round cells	d.	chains of rods
 11.	Cyanobacteria are		
	a. eubacterial heterotrophs	c.	salt-loving archaebacteria
	b. photosynthetic autotrophs	d.	chemosynthetic autotrophs

Completion *Complete each statement.*



Figure 18-1

- 12. The virus attaching to a host cell is shown in stage ______ of Figure 18-1.
- of Figure 18-1. 13. New virus particles are assembled in stage _____
- 14. Virus injecting its nucleic acid into the host cell is shown in stage ______ of Figure 18-1.
- 15. The host cell breaks open, and the new virus particles are released, as shown in stage _____ of Figure 18-1.
- 16. The host DNA is destroyed, and the viral genes are copied as shown in stage ______ of Figure 18-1.

Matching

Match each item with the correct statement below.

- binary fission a. conjugation f. b. reverse transcriptase g. toxin c. bacteriophage
- d. nitrogen fixation

- h. virus
- i. host cell

- e. endospore
- 17. enzyme injected into a host cell, which copies viral RNA into DNA
- 18. cell in which a virus reproduces
- 19. bacterial form that is in a state of slow metabolism and that does not reproduce
- 20. process by which bacteria reproduce asexually
- 21. simple form of sexual reproduction
- 22. virus that infects bacteria
- 23. process by which some bacteria convert N_2 gas into ammonia
- 24. nonliving particle that can reproduce when in a living cell
- 25. poison produced by some bacteria

Short Answer

26. Compare and contrast provirus and retrovirus.

- 27. Compare and contrast lytic cycle and lysogenic cycle.
- 28. Describe the adaptations bacteria might have to live in an extreme habitat.
- 29. Discuss how bacteria have adapted to their diverse habitats.
- 30. Why aren't viruses named according to the rules of binomial nomenclature?
- 31. How are viruses classified according to nucleic acid? Give an example of a virus from each type.
- 32. Why is penicillin ineffective in destroying viruses or animal cells?
- 33. How does a virus recognize its host?
- 34. Why are viruses not considered to be living things?
- 35. Explain in your own words what happens in parts 1, 2, and 3 of the lysogenic cycle shown in the diagram in Figure 18-2.



Figure 18-2

- 36. For a period of time, bacteria were classified as plants. Why do you think bacteria were classified this way? Give at least two reasons why bacteria should not be classified as plants.
- 37. Why do physicians sometimes advise patients who are taking antibiotics to eat yogurt?
- 38. What conditions in developed countries may check the spread of bacteria that cause disease?
- 39. According to Table 18-1, what is the chief cause of death in developing countries? In developed countries? How does Table 18-1 reflect the fact that the availability of antibiotics affects the number of deaths due to infectious diseases?

Percentage of Deaths						
	Developed	countries		Developin	g countries	
Causes of death	Americas	Europe	Americas	Southeast Asia	Africa	Eastern Mediterranean
Infectious disease	3.6	8.6	31.1	43.9	49.8	44.5
Cancer	21.5	18.1	9.0	4.4	2.9	4.2
Circulatory diseases	54.5	53.8	24.5	15.6	11.7	14.1
Accidents	8.4	5.6	6.3	4.3	3.8	4.1

Table 18-1

In 1957, Heinz Fraenkel-Conrat and his coworkers were studying two viruses that infect tobacco plants. One of the disease-causing viruses was called TMV and the other, HRV. Both viruses were similar in structure. (See the diagrams in Figure 18-3.) It was easy to tell which virus had caused a disease because the lesions on the tobacco leaf differed, according to which virus was present. Fraenkel-Conrat knew that TMV and HRV are RNA viruses. He wanted to find out which part of the virus—the protein coat or the RNA—was carrying the genetic information needed to specify the reproduction of these viruses. He decided that he would find the answer by producing hybrids of the viruses. A hybrid has the RNA of one virus and the protein coat of another virus. In this case, the two hybrids are denoted H-T and T-H, where the first letter indicates the virus from which RNA was used.



Figure 18-3

- 40. What evidence in this experiment would show which part of the virus carried the genetic information? Refer to Figure 18-3.
- 41. What were the variables in the experiment? Refer to Figure 18-3.
- 42. What control would be used in this experiment? Refer to Figure 18-3.
- 43. Hypothesize what kind of lesions the T-H hybrid will cause. Explain. Refer to Figure 18-3.
- 44. In lesions caused by the H-T hybrid on tobacco leaves, the new viruses produced were not hybrids. They were all HRV. Why would this be so? Refer to Figure 18-3.
- 45. How might he determine whether the RNA or the protein coat of the H-T hybrid carried the genetic information of the virus? Refer to Figure 18-3.
- 46. Suggest how Fraenkel-Conrat might produce two hybrids. Refer to Figure 18-3.

Bio.10-Q2W4-Qs.Bank-Bacteria and Viruses Answer Section

MULTIPLE CHOICE

1.	ANS: D NAT: $C1 C4 C5$	PTS:	1	DIF:	В	OBJ:	18-3
2.	ANS: D NAT: $C1 C4 C5$	PTS:	1	DIF:	В	OBJ:	18-5
3.	ANS: B	PTS:	1	DIF:	В	OBJ:	18-4
4.	ANS: C	PTS:	1	DIF:	В	OBJ:	18-4
5.	NAT: C1 C4 C5 ANS: C	PTS:	1	DIF:	В	OBJ:	18-4
6.	NAT: C1 C4 C5 ANS: D	PTS:	1	DIF:	В	OBJ:	18-1
7.	NAT: A1 C3 C5 ANS: D	PTS:	1	DIF:	В	OBJ:	18-1
8.	NAT: A1 C3 C5 ANS: C	PTS:	1	DIF:	В	OBJ:	18-1
9.	NAT: A1 C3 C5 ANS: B	PTS:	1	DIF:	В	OBJ:	18-4
10.	NAT: C1 C4 C5 ANS: B	PTS:	1	DIF:	В	OBJ:	18-4
11.	NAT: C1 C4 C5 ANS: B	PTS:	1	DIF:	В	OBJ:	18-3
	NAT: C1 C4 C5						
COMPLETION							
12.	ANS: A						

13.	PTS: ANS:	1 D	DIF:	В	OBJ:	18-2	NAT:	A1 C3 C5
14.	PTS: ANS:	1 B	DIF:	В	OBJ:	18-2	NAT:	A1 C3 C5
15.	PTS: ANS:	1 E	DIF:	В	OBJ:	18-2	NAT:	A1 C3 C5
16.	PTS: ANS:	1 C	DIF:	В	OBJ:	18-2	NAT:	A1 C3 C5
	PTS:	1	DIF:	В	OBJ:	18-2	NAT:	A1 C3 C5

MATCHING

17.	ANS: B	PTS:	1	DIF:	В	OBJ:	18-2
	NAT: A1 C3 C5						
18.	ANS: I	PTS:	1	DIF:	В	OBJ:	18-2
	NAT: A1 C3 C5						
19.	ANS: E	PTS:	1	DIF:	В	OBJ:	18-4
	NAT: C1 C4 C5				_		
20.	ANS: F	PTS:	1	DIF:	В	OBJ:	18-4
	NAT: C1 C4 C5				_		
21.	ANS: A	PTS:	1	DIF:	В	OBJ:	18-4
22	NAT: C1 C4 C5	DTG	1	DIE	D	ODI	10.1
22.	ANS: C	PTS:	1	DIF:	В	OBI:	18-1
22	NAI: AI $ C_3 C_5$	DTC.	1	DIE	D	ODL	10 5
23.	AINS: D	P15:	1	DIF:	В	OBI:	18-5
24	$\begin{array}{c} \text{NAL: } CI \mid C4 \mid C3 \\ \text{ANG: } II \end{array}$	DTC.	1	DIE.	р	ODI	10 7
24.	ANS: Π NAT: $\Lambda 1 \mid C 2 \mid C 5$	P15:	1	DIF:	D	OP1:	10-2
25	NAL ALCOLO	DTC	1	DIE	P	OBI	18 /
<i>23</i> .	NAT: $C1 C4 C5$	115.	1	DIF.	D	ODJ.	10-4
	10111. CI CT CJ						

SHORT ANSWER

26. ANS:

A provirus is a DNA virus that has been inserted into a host cell chromosome. A retrovirus is an RNA virus that contains the enzyme reverse transcriptase, which copies viral RNA into DNA.

PTS:	1	DIF:	В	OBJ:	18-1	NAT: A1	C3	C5

27. ANS:

Both are viral reproductive cycles. In a lytic cycle, the virus causes the destruction of the host cell. In the lysogenic cycle, the viral DNA becomes integrated into the host cell's chromosome, after which it is passed on to future generations of the host cell.

	PTS: 1	DIF: B	OBJ: 18-2	NAT: A1 C3 C5
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28. ANS:

Answers may include: The bacteria should be able to reproduce quickly and live in the environment. The bacteria may also utilize substances that other organisms cannot use. They may have protective structures, such as capsules or endospores.

PTS: 1 DIF: A OBJ: 18-4 NAT: C1 | C4 | C5

29. ANS:

Rapid reproduction and simplicity of the cell have allowed bacteria to adapt to extreme environmental conditions. They may utilize poisonous substances, be anaerobic, and live in low pH environments.

	PTS: 1	DIF: A	OBJ: 18-4	NAT: C1 C4 C5
30.	ANS:			
	Viruses are not living	g things.		
	DTC. 1		ODI: 10.1	NAT. A1 C2 C5
	P15: 1	DIF: A	OBJ: 18-1	NAT: AT $ C_3 C_5$

31.	ANS: DNA viruses—chicken	pox, warts; RNA vi	ruses—HIV					
32	PTS: 1 D	DIF: A	OBJ: 18-1	NAT: A1 C3 C5				
52.	Penicillin interferes with viruses nor animal cells	h the enzyme that lir have cell walls, pen	iks the sugar chains in icillin has no effect or	the bacterial cell wall. Because neither them.				
33.	PTS: 1 E ANS:	DIF: A	OBJ: 18-4	NAT: C1 C4 C5				
	A protein on the surface allowing the virus to loc	e of a virus has a sha ck onto the host cell.	pe that matches a mol-	ecule in the plasma membrane of its host,				
34.	PTS: 1 D ANS:	DIF: B	OBJ: 18-2	NAT: A1 C3 C5				
	Viruses do not carry ou living cell.	t respiration, grow, c	or move. They can rep	roduce, but only when they are inside a				
35.	PTS: 1 D ANS:	DIF: B	OBJ: 18-1	NAT: A1 C3 C5				
	 A virus injects its DNA into a bacterium. The viral DNA becomes part of the host chromosome as a provirus. The provirus is inactive but is replicated when the host cell chromosome replicates. 							
36	PTS: 1 D	DIF: B	OBJ: 18-2	NAT: A1 C3 C5				
50.	Like plants, bacteria have cell walls and lack typical animal characteristics. Unlike plants, bacteria are prokaryotic and their cell walls lack cellulose.							
37	PTS: 1 D	DIF: A	OBJ: 18-4	NAT: C1 C4 C5				
57.	The antibiotics kill beneficial, as well as harmful, bacteria in the body. Because yogurt contains some of these beneficial bacteria, eating yogurt can help to replace them in the patient's body.							
38	PTS: 1 D	DIF: A	OBJ: 18-5	NAT: C1 C4 C5				
50.	Water purification and sanitary waste disposal help prevent the spread of disease-causing bacteria in developed countries.							
39.	PTS: 1 D	DIF: A	OBJ: 18-5	NAT: C1 C4 C5				
	In developing countries diseases. Developed cou do developing countries	, the chief cause of c untries have better ac s. That's why fewer c	leath is infectious dise ccess to doctors and an leaths from infectious	ases; in developed countries, it is circulatory ntibiotics for curing infectious diseases than diseases occur in developed countries.				
40	PTS: 1 D	DIF: A	OBJ: 18-5	NAT: C1 C4 C5				
40.	The H-T virus produces controlling the reproduce	s only HRV viruses t ction of the viruses.	hat would convince re	searchers that the RNA of the HRV was				

41.	PTS: 1 ANS:	DIF: A	OBJ: 18-1	NAT: A1 C3 C5
	The two kinds of hy	brids were the variab	les.	
42.	PTS: 1 ANS:	DIF: A	OBJ: 18-1	NAT: A1 C3 C5
	infecting tobacco pl	ants with the pure TM	IV and HRV; using pla	nts with no viruses
13	PTS: 1	DIF: A	OBJ: 18-1	NAT: A1 C3 C5
чэ.	The T-H hybrid will	cause the same kind	of lesions as the TMV	because it has the RNA of the TMV.
44	PTS: 1	DIF: A	OBJ: 18-1	NAT: A1 C3 C5
44.	Because the RNA of	f the HRV was provid	ling the genetic messag	e, only HRV copies could be made.
15	PTS: 1	DIF: A	OBJ: 18-1	NAT: A1 C3 C5
4J.	Answers will vary b hybrid causes the sa genetic information	ut may include that h me kind of lesions that because the hybrid ha	e could inject the H-T l at HRV typically cause as the RNA of HRV, ar	hybrid into a healthy tobacco plant. If the s, he would know that RNA provides the ind that is the information that was specified.
16	PTS: 1	DIF: A	OBJ: 18-1	NAT: A1 C3 C5
40.	Anse: Answers may vary b virus. Then he woul would mix the TMV	out may include that h d mix the RNA of the 7 RNA with the HRV	he would have to separa HRV with the TMV p protein coat to produce	ate the RNA from the protein coat of each protein coat to produce the H-T hybrid. He e the T-H hybrid.

PTS: 1 DIF: A OBJ: 18-1 NAT: A1 | C3 | C5