## **MODIFIED TRUE/FALSE**

	ANS: T REF: p. 250 STA: F.12.10 ANS: F, oxygen		9.1.1 Explain Foundation Ec				L2 gy they need for life processes. analysis
	PTS: 1 respiration.	DIF:			p. 251		9.1.2 Define cellular
3.	STA: F.12.1 ANS: F, oxygen	TOP:	Foundation Ec	lition		BLM:	knowledge
	PTS: 1 respiration.	DIF:	L2	REF:	p. 252	OBJ:	9.1.2 Define cellular
4.	STA: F.12.1 ANS: T	TOP:	Foundation Ec	lition PTS:	1	BLM: DIF:	comprehension L2
	REF: p. 253		9.1.3 Compare		synthesis and co		respiration.
	STA: F.12.1   F.12.9	9		TOP:	Foundation Ed	lition	
F	BLM: application ANS: F, All						
5.	ANS: F, All						
	PTS: 1	DIF:	L3	REF:	p. 253		
	OBJ: 9.1.3 Compare	e photo	synthesis and c	ellular	respiration.	STA:	F.12.1   F.12.9
	BLM: application						
6.	ANS: T	0.5.4		PTS:		DIF:	
	REF: p. 255				happens during		
7	STA: F.12.1 ANS: T	TOP:	Foundation Ec	PTS:	1	BLM: DIF:	comprehension
7.	REF: p. 256	OBI-	922 Describe		appens during		
	STA: F.12.1		Foundation Ed				knowledge
8.	ANS: T			PTS:	1	DIF:	÷
	REF: p. 256   p. 258						
	OBJ: 9.2.3 Explain			rons ar	e used by the el	ectron	transport chain.
0	STA: F.12.1		knowledge				
9.	ANS: F, cell membra	ane					
	PTS: 1	DIF:	L2	REF∙	p. 258		
	OBJ: 9.2.3 Explain				•	ectron t	transport chain.
	STA: F.12.1		comprehensio		2		L L
10.	ANS: T			PTS:	1	DIF:	L2
	REF: p. 256   p. 258						
	OBJ: 9.2.4 Identify				ration generates		
11	STA: F.12.1		Foundation Ed	11t10n		BUM:	analysis
11.	ANS: F, lactic acid f	erment	auon				
	PTS: 1	DIF:	L2	REF:	p. 263		
	OBJ: 9.3.1 Explain	how or	ganisms get ene	ergy in	▲ ·	oxygen	
	STA: F.12.1	TOP:	Foundation Ed	lition		BLM:	application

12. ANS: F, oxygen PTS: 1 DIF: L3 REF: p. 252 OBJ: 9.3.1 Explain how organisms get energy in the absence of oxygen. BLM: synthesis STA: F.12.1 13. ANS: T PTS: 1 DIF: L3 REF: p. 265 OBJ: 9.3.2 Identify the pathways the body uses to release energy during exercise. STA: F.12.10 **BLM**: application 14. ANS: F, ATP PTS: 1 DIF: L2 REF: p. 264 | p. 265 OBJ: 9.3.2 Identify the pathways the body uses to release energy during exercise. TOP: Foundation Edition STA: F.12.10 BLM: comprehension 15. ANS: T PTS: 1 DIF: L1 REF: p. 264 | p. 265 OBJ: 9.3.2 Identify the pathways the body uses to release energy during exercise. TOP: Foundation Edition STA: F.12.10 BLM: knowledge 16. ANS: F, decreases PTS: 1 DIF: L2 REF: p. 275 OBJ: 10.1.1 Explain the problems that growth causes for cells. STA: F.12.1 BLM: comprehension TOP: Foundation Edition 17. ANS: F, larger PTS: 1 DIF: L2 REF: p. 274 | p. 275 OBJ: 10.1.1 Explain the problems that growth causes for cells. STA: F.12.1 TOP: Foundation Edition BLM: comprehension 18. ANS: T PTS: 1 DIF: L3 REF: p. 275 OBJ: 10.1.1 Explain the problems that growth causes for cells. STA: F.12.1 BLM: analysis 19. ANS: F, asexual PTS: 1 DIF: L3 REF: p. 277 | p. 284 OBJ: 10.1.2 Compare asexual and sexual reproduction. STA: F.12.1 **TOP:** Foundation Edition BLM: synthesis 20. ANS: T PTS: 1 DIF: L2 OBJ: 10.2.1 Describe the role of chromosomes in cell division. REF: p. 280 STA: F.12.1 TOP: Foundation Edition BLM: application 21. ANS: F, G<sub>1</sub> DIF: L2 REF: p. 281 PTS: 1 OBJ: 10.2.2 Name the main events of the cell cycle. STA: F.12.1 BLM: application TOP: Foundation Edition 22. ANS: F, nuclear envelope REF: p. 282 PTS: 1 DIF: L3 OBJ: 10.2.3 Describe what happens during the four phases of mitosis. STA: F.12.1 TOP: Foundation Edition BLM: analysis 23. ANS: F, prophase

PTS: 1 DIF: L2 REF: p. 282 OBJ: 10.2.3 Describe what happens during the four phases of mitosis. STA: F.12.1 BLM: comprehension 24. ANS: T PTS: 1 DIF: L2 REF: p. 283 OBJ: 10.2.3 Describe what happens during the four phases of mitosis. STA: F.12.1 TOP: Foundation Edition BLM: comprehension 25. ANS: T DIF: L1 PTS: 1 OBJ: 10.2.4 Describe the process of cytokinesis. REF: p. 284 TOP: Foundation Edition BLM: knowledge STA: F.12.1 26. ANS: T PTS: 1 DIF: L2 OBJ: 10.3.1 Describe how the cell cycle is regulated. REF: p. 286 STA: F.12.2 TOP: Foundation Edition BLM: application PTS: 1 27. ANS: T DIF: L1 REF: p. 289 OBJ: 10.3.2 Explain how cancer cells are different from other cells. **TOP:** Foundation Edition BLM: knowledge PTS: 1 28. ANS: T DIF: L2 REF: p. 293 OBJ: 10.4.1 Describe the process of differentiation. STA: F.12.2 TOP: Foundation Edition BLM: evaluation 29. ANS: F, pluripotent PTS: 1 REF: p. 295 DIF: L1 OBJ: 10.4.2 Define stem cells and explain their importance. STA: F.12.2 BLM: knowledge 30. ANS: F destroys kills PTS: 1 DIF: L2 REF: p. 297 OBJ: 10.4.3 Identify the possible benefits and issues relating to stem cell research. STA: G.12.3 | H.12.3 **TOP:** Foundation Edition BLM: analysis 31. ANS: T PTS: 1 DIF: L1 REF: p. 309 OBJ: 11.1.1 Describe Mendel's studies and conclusions about inheritance. STA: F.12.3 TOP: Foundation Edition BLM: comprehension 32. ANS: F, dominant PTS: 1 DIF: L3 REF: p. 310 OBJ: 11.1.1 Describe Mendel's studies and conclusions about inheritance. STA: F.12.3 BLM: application 33. ANS: F, separate PTS: 1 DIF: L1 REF: p. 325 OBJ: 11.1.2 Describe what happens during segregation. STA: F.12.3 TOP: Foundation Edition BLM: knowledge 34. ANS: T PTS: 1 DIF: L3 REF: p. 324 | p. 325 OBJ: 11.1.2 Describe what happens during segregation. STA: F.12.3 BLM: evaluation PTS: 1 35. ANS: T DIF: L2

REF: p. 313 OBJ: 11.2.1 Explain how geneticists use the principles of probability to make Punnett squares. STA: F.12.3 TOP: Foundation Edition BLM: comprehension 36. ANS: F 50% PTS: 1 DIF: L2 REF: p. 313 | p. 314 OBJ: 11.2.1 Explain how geneticists use the principles of probability to make Punnett squares. **TOP:** Foundation Edition STA: F.12.3 **BLM**: application 37. ANS: F, contradicted PTS: 1 DIF: L3 REF: p. 317 OBJ: 11.2.2 Explain the principle of independent assortment. STA: F.12.3 BLM: evaluation 38. ANS: T PTS: 1 DIF: L2 REF: p. 316 | p. 318 OBJ: 11.2.3 Explain how Mendel's principles apply to all organisms. STA: F.12.3 TOP: Foundation Edition BLM: analysis 39. ANS: F, 50% PTS: 1 REF: p. 319 DIF: L2 OBJ: 11.3.1 Describe the other inheritance patterns. STA: F.12.3 BLM: analysis 40. ANS: F, trait controlled by multiple alleles PTS: 1 DIF: L2 REF: p. 320 STA: F.12.3 OBJ: 11.3.1 Describe the other inheritance patterns. **TOP:** Foundation Edition BLM: application PTS: 1 41. ANS: T DIF: L2 REF: p. 323 OBJ: 11.4.1 Contrast the number of chromosomes in body cells and in gametes. TOP: Foundation Edition BLM: analysis STA: F.12.4 42. ANS: F, anaphase I PTS: 1 DIF: L3 REF: p. 324 | p. 325 OBJ: 11.4.2 Summarize the events of meiosis. STA: F.12.4 BLM: analysis 43. ANS: F, four cells PTS: 1 REF: p. 326 | p. 327 DIF: L1 OBJ: 11.4.3 Contrast meiosis and mitosis. STA: F.12.4 **TOP:** Foundation Edition BLM: knowledge 44. ANS: F, four REF: p. 328 | p. 329 PTS: 1 DIF: L2 OBJ: 11.4.4 Describe how alleles from different genes can be inherited together. TOP: Foundation Edition STA: F.12.4 BLM: analysis 45. ANS: F, together

PTS: 1 DIF: L2 REF: p. 328 | p. 329 OBJ: 11.4.4 Describe how alleles from different genes can be inherited together. STA: F.12.4 TOP: Foundation Edition BLM: application 46. ANS: T DIF: L1 PTS: 1 REF: p. 338 | p. 339 OBJ: 12.1.1 Summarize the process of bacterial transformation. STA: F.12.3 TOP: Foundation Edition BLM: knowledge 47. ANS: F, smaller PTS: 1 DIF: L1 REF: p. 340 OBJ: 12.1.2 Describe the role of bacteriophages in identifying genetic material. TOP: Foundation Edition STA: F.12.3 BLM: evaluation 48. ANS: F, DNA, genetic material PTS: 1 DIF: L1 REF: p. 340 | p. 341 OBJ: 12.1.2 Describe the role of bacteriophages in identifying genetic material. TOP: Foundation Edition STA: F.12.3 BLM: comprehension 49. ANS: T DIF: L2 PTS: 1 OBJ: 12.1.3 Identify the role of DNA in heredity. REF: p. 342 | p. 343 STA: F.12.3 TOP: Foundation Edition BLM: comprehension 50. ANS: F, nucleotides PTS: 1 DIF: L1 REF: p. 333 OBJ: 12.2.1 Identify the chemical components of DNA. STA: F.12.3 TOP: Foundation Edition BLM: knowledge 51. ANS: T PTS: 1 DIF: L2 REF: p. 344 | p. 345 OBJ: 12.2.1 Identify the chemical components of DNA. STA: F.12.3 TOP: Foundation Edition BLM: comprehension 52. ANS: T PTS: 1 DIF: L2 REF: p. 345 OBJ: 12.2.2 Discuss the experiments leading to the identification of DNA as the molecule that carries STA: F.12.3 the genetic code. TOP: Foundation Edition BLM: comprehension 53. ANS: T PTS: 1 DIF: L2 REF: p. 346 | p. 347 OBJ: 12.2.2 Discuss the experiments leading to the identification of DNA as the molecule that carries TOP: Foundation Edition the genetic code. STA: F.12.3 BLM: comprehension 54. ANS: T PTS: 1 DIF: L3 REF: p. 341 OBJ: 12.1.2 Describe the role of bacteriophages in identifying genetic material. STA: F.12.3 BLM: analysis 55. ANS: F, hydrogen PTS: 1 DIF: L1 REF: p. 348 OBJ: 12.2.3 Describe the steps leading to the development of the double-helix model of DNA. TOP: Foundation Edition STA: C.12.5 | F.12.3 BLM: knowledge 56. ANS: F, 100%

57.	the gen	12.2.2 Discuss netic code. analysis		periments leadi	REF: p. 345 ng to the identification TOP: Foundation Ed		A as the molecule that carries
58.	TOP:			L2 events of DNA	REF: p. 350 A replication. BLM: application	STA:	F.12.3
59.	STA:				REF: p. 352 prokaryotes with that of	of euka	ryotes.
60.	TOP:		lition	L3 events of DNA	REF: p. 350 A replication. BLM: application	STA:	F.12.3
					REF: p. 352   p. 353 prokaryotes with that o lition	of euka	ryotes. comprehension
COMPLE	ΓION						
1.	ANS:	the sun					
2	PTS: OBJ: STA:	9.1.1 Explain			REF: p. 250 he energy they need for	r life pr	ocesses.
2.	ANS:	F.12.10 mitochondria	101.	Foundation EC	lition		application
	PTS: respira STA:	mitochondria 1 tion. F.12.1	DIF: TOP:		REF: p. 252	BLM: OBJ:	application 9.1.2 Define cellular application
3.	PTS: respira STA: ANS: PTS: OBJ: BLM:	mitochondria 1 tion. F.12.1 photosynthesis 1	DIF: TOP: DIF:	L2 Foundation Ed	REF: p. 252	BLM: OBJ: BLM:	9.1.2 Define cellular

DIF: L1 REF: p. 254 PTS: 1 OBJ: 9.2.1 Describe what happens during glycolysis. STA: F.12.1 TOP: Foundation Edition BLM: knowledge 6. ANS: 2 PTS: 1 DIF: L3 REF: p. 256 | p. 257 OBJ: 9.2.2 Describe what happens during the Krebs cycle. STA: F.12.1 BLM: synthesis 7. ANS:  $H^+$  ions PTS: 1 DIF: L3 REF: p. 258 OBJ: 9.2.3 Explain how high-energy electrons are used by the electron transport chain. STA: F.12.1 BLM: synthesis 8. ANS: electron transport chain PTS: 1 DIF: L2 REF: p. 258 OBJ: 9.2.3 Explain how high-energy electrons are used by the electron transport chain. TOP: Foundation Edition BLM: comprehension STA: F.12.1 9. ANS: 2 PTS: 1 REF: p. 254 DIF: L2 OBJ: 9.2.4 Identify how much ATP cellular respiration generates. TOP: Foundation Edition STA: F.12.1 BLM: comprehension 10. ANS: alcoholic PTS: 1 REF: p. 263 DIF: L2 OBJ: 9.3.1 Explain how organisms get energy in the absence of oxygen. TOP: Foundation Edition STA: F.12.1 BLM: analysis 11. ANS: 2 PTS: 1 DIF: L1 REF: p. 254 | p. 263 OBJ: 9.3.1 Explain how organisms get energy in the absence of oxygen. TOP: Foundation Edition STA: F.12.1 BLM: knowledge 12. ANS: C PTS: 1 DIF: L2 REF: p. 252 | p. 263 OBJ: 9.1.2 Define cellular respiration. STA: F.12.1 **TOP:** Foundation Edition BLM: analysis 13. ANS: oxygen PTS: 1 REF: p. 265 DIF: L1 OBJ: 9.3.2 Identify the pathways the body uses to release energy during exercise. STA: F.12.10 BLM: knowledge 14. ANS: lactic acid PTS: 1 REF: p. 265 DIF: L2 OBJ: 9.3.2 Identify the pathways the body uses to release energy during exercise. STA: F.12.10 TOP: Foundation Edition **BLM**: application 15. ANS: more

PTS: 1 DIF: L3 REF: p. 252 OBJ: 9.3.2 Identify the pathways the body uses to release energy during exercise. STA: F.12.10 BLM: synthesis 16. ANS: less DIF: L1 PTS: 1 REF: p. 274 OBJ: 10.1.1 Explain the problems that growth causes for cells. STA: F.12.1 BLM: knowledge 17. ANS: sexual, asexual PTS: 1 REF: p. 277 DIF: L2 OBJ: 10.1.2 Compare asexual and sexual reproduction. STA: F.12.1 TOP: Foundation Edition BLM: comprehension 18. ANS: 92 PTS: 1 DIF: L3 REF: p. 280 | p. 282 OBJ: 10.2.1 Describe the role of chromosomes in cell division. **TOP:** Foundation Edition STA: F.12.1 BLM: synthesis 19. ANS: mitosis REF: p. 282 PTS: 1 DIF: L1 OBJ: 10.2.2 Name the main events of the cell cycle. STA: F.12.1 **TOP:** Foundation Edition BLM: knowledge 20. ANS: binary fission PTS: 1 DIF: L2 REF: p. 281 OBJ: 10.2.2 Name the main events of the cell cycle. STA: F.12.1 **TOP:** Foundation Edition BLM: comprehension 21. ANS: 10 PTS: 1 DIF: L3 REF: p. 281 | p. 282 OBJ: 10.2.2 Name the main events of the cell cycle. STA: F.12.1 **TOP:** Foundation Edition BLM: synthesis 22. ANS: anaphase PTS: 1 DIF: L2 REF: p. 282 OBJ: 10.2.3 Describe what happens during the four phases of mitosis. **TOP:** Foundation Edition STA: F.12.1 BLM: comprehension 23. ANS: G<sub>1</sub> phase, Interphase PTS: 1 DIF: L3 REF: p. 284 OBJ: 10.2.2 Name the main events of the cell cycle. STA: F.12.1 **TOP:** Foundation Edition BLM: synthesis 24. ANS: cell plate REF: p. 284 PTS: 1 DIF: L2 OBJ: 10.2.4 Describe the process of cytokinesis. STA: F.12.1 **TOP:** Foundation Edition BLM: application

25. ANS: plant PTS: 1 DIF: L2 REF: p. 284 STA: F.12.1 OBJ: 10.2.4 Describe the process of cytokinesis. **TOP:** Foundation Edition BLM: analysis 26. ANS: apoptosis PTS: 1 DIF: L1 REF: p. 288 OBJ: 10.3.1 Describe how the cell cycle is regulated. STA: F.12.2 **TOP:** Foundation Edition BLM: knowledge 27. ANS: chemotherapy REF: p. 290 PTS: 1 DIF: L2 OBJ: 10.3.2 Explain how cancer cells are different from other cells. **TOP:** Foundation Edition BLM: comprehension 28. ANS: differentiation PTS: 1 DIF: L2 REF: p. 293 OBJ: 10.4.1 Describe the process of differentiation. STA: F.12.2 **TOP:** Foundation Edition BLM: comprehension 29. ANS: differentiation PTS: 1 DIF: L2 REF: p. 294 | p. 295 OBJ: 10.4.2 Define stem cells and explain their importance. STA: F.12.2 **TOP:** Foundation Edition BLM: comprehension 30. ANS: adult REF: p. 297 PTS: 1 DIF: L3 OBJ: 10.4.3 Identify the possible benefits and issues relating to stem cell research. STA: G.12.3 | H.12.3 BLM: evaluation 31. ANS: P REF: p. 310 PTS: 1 DIF: L2 OBJ: 11.1.1 Describe Mendel's studies and conclusions about inheritance. STA: F.12.3 TOP: Foundation Edition BLM: application 32. ANS: gametes, sex cells REF: p. 314 PTS: 1 DIF: L2 OBJ: 11.1.2 Describe what happens during segregation. STA: F.12.3 **TOP:** Foundation Edition BLM: comprehension 33. ANS: 19 PTS: 1 DIF: L3 REF: p. 324 OBJ: 11.2.1 Explain how geneticists use the principles of probability to make Punnett squares. **TOP:** Foundation Edition STA: F.12.3 BLM: analysis 34. ANS:  $1/2 \ge 1/2 \ge 1/2 \ge 1/2 \ge 1/32$ DIF: L3 REF: p. 313 PTS: 1 OBJ: 11.2.1 Explain how geneticists use the principles of probability to make Punnett squares.

STA: F.12.3 BLM: analysis 35. ANS: TT and Tt PTS: 1 DIF: L2 REF: p. 314 OBJ: 11.2.1 Explain how geneticists use the principles of probability to make Punnett squares. TOP: Foundation Edition BLM: application STA: F.12.3 36. ANS: genes, chromosomes PTS: 1 REF: p. 317 | p. 329 DIF: L1 OBJ: 11.2.2 Explain the principle of independent assortment. STA: F.12.3 TOP: Foundation Edition BLM: knowledge 37. ANS: round yellow seeds only PTS: 1 DIF: L2 REF: p. 317 OBJ: 11.2.2 Explain the principle of independent assortment. STA: F.12.3 BLM: analysis 38. ANS: Mendel PTS: 1 DIF: L1 REF: p. 318 OBJ: 11.2.3 Explain how Mendel's principles apply to all organisms. TOP: Foundation Edition BLM: application STA: F.12.3 39. ANS: polygenic trait REF: p. 320 PTS: 1 DIF: L2 OBJ: 11.3.1 Describe the other inheritance patterns. STA: F.12.3 BLM: comprehension 40. ANS: light energy PTS: 1 DIF: L3 REF: p. 321 OBJ: 11.3.2 Explain the relationship between genes and the environment. STA: F.12.3 BLM: evaluation 41. ANS: genes and environmental conditions PTS: 1 DIF: L1 REF: p. 321 OBJ: 11.3.2 Explain the relationship between genes and the environment. TOP: Foundation Edition BLM: knowledge STA: F.12.3 42. ANS: incomplete dominance DIF: L1 PTS: 1 REF: p. 319 | p. 320 OBJ: 11.3.1 Describe the other inheritance patterns. STA: F.12.3 TOP: Foundation Edition BLM: analysis 43. ANS: half PTS: 1 REF: p. 323 DIF: L2 OBJ: 11.4.1 Contrast the number of chromosomes in body cells and in gametes. STA: F.12.4 TOP: Foundation Edition BLM: comprehension 44. ANS: prophase I PTS: 1 DIF: L2 REF: p. 324 | p. 325

45.		11.4.2 Summa analysis gene	rize the	events of meio	osis.		STA:	F.12.4
46.	OBJ: STA:	1 11.4.4 Describ F.12.4 developed pne	TOP:	alleles from dif Foundation Ec	ferent g	p. 328   p. 329 genes can be inh		together. knowledge
47.	STA:	1 12.1.1 Summa F.12.3 protein coat			terial tr	ansformation.	BLM:	comprehension
48.	STA:			le of bacteriop	hages in	p. 340 n identifying ge		naterial. knowledge
49.	OBJ: BLM:	1 12.1.3 Identify analysis nucleotide		L2 e of DNA in he		▲	STA:	F.12.3
50.	OBJ: TOP:	1 12.2.1 Identify Foundation Ec AGCT			ents of	p. 345 DNA. comprehension		F.12.3
51.	TOP:				ents of	p. 345 DNA. application	STA:	F.12.3
52.	the ger BLM:		DIF: s the exp STA:	periments leadi	-	▲		A as the molecule that carries
53.	PTS: OBJ: STA:	-	TOP:		terial tr		BLM:	analysis
	STA:				the deve	p. 348 elopment of the Foundation Ed		e-helix model of DNA.

54. ANS: replicate PTS: 1 DIF: L3 REF: p. 348 OBJ: 12.2.3 Describe the steps leading to the development of the double-helix model of DNA. **TOP:** Foundation Edition STA: C.12.5 | F.12.3 BLM: evaluation 55. ANS: 20% PTS: 1 DIF: L3 REF: p. 345 OBJ: 12.2.2 Discuss the experiments leading to the identification of DNA as the molecule that carries **TOP:** Foundation Edition the genetic code. STA: F.12.3 BLM: analysis 56. ANS: histones PTS: 1 REF: p. 352 DIF: L1 OBJ: 12.3.2 Compare DNA replication in prokaryotes with that of eukaryotes. STA: F.12.4 BLM: knowledge 57. ANS: enzymes PTS: 1 DIF: L2 REF: p. 351 OBJ: 12.3.1 Summarize the events of DNA replication. STA: F.12.3 **TOP:** Foundation Edition BLM: analysis 58. ANS: telomeres PTS: 1 DIF: L1 REF: p. 352 OBJ: 12.3.1 Summarize the events of DNA replication. STA: F.12.3 **TOP:** Foundation Edition BLM: knowledge 59. ANS: DNA, chromosome PTS: 1 DIF: L2 REF: p. 353 OBJ: 12.3.2 Compare DNA replication in prokaryotes with that of eukaryotes. STA: F.12.4 TOP: Foundation Edition BLM: synthesis 60. ANS: origins of replication PTS: 1 DIF: L3 REF: p. 353 OBJ: 12.3.2 Compare DNA replication in prokaryotes with that of eukaryotes. BLM: synthesis STA: F.12.4 **SHORT ANSWER** 1. ANS: There are 1000 calories in 1 Calorie. Eating 2000 calories is the same as eating 2 Calories. PTS: 1 DIF: L3 REF: p. 250 OBJ: 9.1.1 Explain where organisms get the energy they need for life processes.

STA: F.12.10 BLM: synthesis

2. ANS:

Cellular respiration is the process that releases energy by breaking down food molecules in the presence of oxygen.

	PTS: 1	DIF:	L1	REF:	p. 251	OBJ:	9.1.2 Define cellular
3	respiration. STA: F.12.1 ANS:	TOP:	Foundation E	dition		BLM:	knowledge
5.	The three stages are						n), the Krebs cycle (which occurs in the inner mitochondrial
	PTS: 1	DIF:	L2	REF:	p. 252	OBJ:	9.1.2 Define cellular
4	respiration. STA: F.12.1 ANS:	TOP:	Foundation E	dition		BLM:	synthesis
							rgy from plants to animals. on.
	PTS: 1 OBJ: 9.1.3 Compar BLM: analysis	DIF: e photo			p. 253 respiration.	STA:	F.12.1   F.12.9
5.	ANS:	•	•	nospher	e as a product,	wherea	s cellular respiration uses oxygen
				DEE	252		
	PTS: 1 OBJ: 9.1.3 Compar TOP: Foundation E			ellular	p. 253 respiration. knowledge	STA:	F.12.1   F.12.9
6.	ANS: During glycolysis, th acid.	e bond	s of glucose are	broker	and rearranged	l to pro	duce 2 molecules of pyruvic
	PTS: 1	DIF:			p. 254		
7	OBJ: 9.2.1 Describ TOP: Foundation E		happens during		/sis. knowledge	STA:	F.12.1
7.	ANS: Glycolysis requires a net gain of 2 ATP me			P moleo	cules and produ	ces an	output of 4 ATP molecules, for a
	PTS: 1 OBJ: 9.2.1 Describ	DIF: e what			p. 255 /sis.	STA:	F.12.1
8.	BLM: analysis ANS:						
	Citric acid is the first	t compo	ound formed in	the pro	cess.		
	PTS: 1 OBJ: 9.2.2 Describ TOP: Foundation E		L1 happens during	the Kre	p. 256 ebs cycle. knowledge	STA:	F.12.1
9.	ANS: The movement of H <sup>+</sup> ADP into ATP.	ions b	ack across the i	nner mi	tochondrial me	mbrane	e through ATP synthase converts

PTS: 1 DIF: L1 REF: p. 258 OBJ: 9.2.3 Explain how high-energy electrons are used by the electron transport chain. STA: F.12.1 **TOP:** Foundation Edition BLM: knowledge 10. ANS: Oxygen is the final electron acceptor in the electron transport chain, which means that it is needed to get rid of low-energy electrons and H<sup>+</sup> ions. PTS: 1 DIF: L2 REF: p. 258 OBJ: 9.2.3 Explain how high-energy electrons are used by the electron transport chain. TOP: Foundation Edition STA: F.12.1 BLM: comprehension 11. ANS: Sample answer: Lactic acid fermentation occurs in the muscles, and alcoholic fermentation occurs in rising bread dough. DIF: L2 REF: p. 263 PTS: 1 OBJ: 9.3.1 Explain how organisms get energy in the absence of oxygen. **TOP:** Foundation Edition STA: F.12.1 **BLM**: application 12. ANS: Alcoholic fermentation produces carbon dioxide, alcohol, and NAD<sup>+</sup>, whereas lactic acid fermentation produces lactic acid and NAD<sup>+</sup>. PTS: 1 DIF: L2 REF: p. 263 OBJ: 9.3.1 Explain how organisms get energy in the absence of oxygen. TOP: Foundation Edition STA: F.12.1 BLM: analysis 13. ANS: Pathway A and pathway B can both take place when there is no oxygen. When cells run out of oxygen, they can still produce some energy, even though they do so inefficiently. PTS: 1 DIF: L3 REF: p. 262 | p. 263 OBJ: 9.3.1 Explain how organisms get energy in the absence of oxygen. STA: F.12.1 BLM: synthesis 14. ANS: Sample answer: Cellular respiration, shown in pathway C, is most efficient, because it produces the most ATP using the same amount of glucose as the other two pathways. PTS: 1 DIF: L3 REF: p. 252 OBJ: 9.2.4 Identify how much ATP cellular respiration generates. STA: F.12.1 BLM: analysis 15. ANS: The body uses ATP that is already present in the muscle cells, ATP released from lactic acid fermentation, and ATP released from cellular respiration. PTS: 1 DIF: L2 REF: p. 264 | p. 265 OBJ: 9.3.2 Identify the pathways the body uses to release energy during exercise. **TOP:** Foundation Edition BLM: comprehension STA: F.12.10 16. ANS: As a cell grows larger, more demands are placed on its DNA, and the cell has more trouble moving enough nutrients and wastes across the cell membrane.

17.	PTS:1DIF:L2REF:p. 274   p. 276OBJ:10.1.1 Explain the problems that growth causes for cells.STA:F.12.1TOP:Foundation EditionBLM:comprehensionANS:A large cell carries out its activities less efficiently than a small one does.
18.	PTS:1DIF:L2REF:p. 274OBJ:10.1.1 Explain the problems that growth causes for cells.STA:F.12.1BLM:applicationANS:Because the offspring of asexual reproduction are genetically identical to parents, they have the characteristics that help them survive in the conditions in which the parent cells survived. They might not have characteristics to survive should the conditions change.
19.	PTS:1DIF:L3REF:p. 278OBJ:10.1.2 Compare asexual and sexual reproduction.STA:F.12.1TOP:Foundation EditionBLM:evaluationANS:Packaging genetic material into chromosomes helps the cell separate the DNA precisely during celldivision. If the genetic material was spread out into smaller pieces, some of the material might getlost more easily when the cell divided into two cells.
20.	PTS:1DIF:L2REF:p. 280OBJ:10.2.1 Describe the role of chromosomes in cell division.STA:F.12.1TOP:Foundation EditionBLM:analysisANS:Chromatids are two identical DNA strands joined by a centromere, and chromatin is the material (DNA and proteins) that makes up chromosomes.
21.	PTS:1DIF:L3REF:p. 280   p. 282OBJ:10.2.1 Describe the role of chromosomes in cell division.STA:F.12.1TOP:Foundation EditionBLM:synthesisANS:A:G1 phase, cell growth;B:S phase, DNA replication;C:G2 phase, preparation for mitosis;D:M phase,cell division (mitosis and cytokinesis).
22.	PTS:1DIF:L2REF:p. 281   p. 282OBJ:10.2.2 Name the main events of the cell cycle.STA:F.12.1TOP:Foundation EditionBLM:analysisANS:I is anaphase. 2 is prophase. 3 is interphase (or $G_2$ phase). 4 is telophase. 5 is metaphase.They occur in the following order:3, 2, 5, 1, and 4 (or: 2, 5, 1, 4, 3).
23.	PTS:1DIF:L3REF:p. 280   p. 282OBJ:10.2.3 Describe what happens during the four phases of mitosis.STA:F.12.1TOP:Foundation EditionBLM: analysisANS:

the cell. PTS: 1 REF: p. 280 | p. 282 DIF: L2 OBJ: 10.2.3 Describe what happens during the four phases of mitosis. STA: F.12.1 TOP: Foundation Edition BLM: analysis 24. ANS: In plant cells, a cell plate forms in the cytoplasm midway between each new nucleus. The cell plate gradually develops into a separating membrane, and a cell wall begins to appear in the cell plate. In animal cells, there is no cell plate. The cell membrane is drawn inward until the cytoplasm is pinched into two nearly equal parts. PTS: 1 DIF: L3 REF: p. 284 OBJ: 10.2.4 Describe the process of cytokinesis. STA: F.12.1 TOP: Foundation Edition **BLM:** synthesis 25. ANS: A cell that lacked cyclins would probably not undergo mitotic division, and then it would continue to grow, have DNA overload, and exchange materials inefficiently until it dies. PTS: 1 DIF: L3 REF: p. 286 OBJ: 10.3.1 Describe how the cell cycle is regulated. STA: F.12.2 BLM: evaluation 26. ANS: Cancer cells do not respond to the signals that control the growth of normal cells. As a result, cancer cells form tumors and can spread throughout the body. PTS: 1 DIF: L2 REF: p. 289 OBJ: 10.3.2 Explain how cancer cells are different from other cells. TOP: Foundation Edition BLM: comprehension 27. ANS: Cancer cells are not constrained by crowding and would probably continue to grow after forming a thin layer covering the bottom of the petri dish. PTS: 1 DIF: L3 REF: p. 289 OBJ: 10.3.2 Explain how cancer cells are different from other cells. BLM: synthesis 28. ANS: Differentiation is the process by which cells become specialized. PTS: 1 DIF: L1 REF: p. 293 OBJ: 10.4.1 Describe the process of differentiation. STA: F.12.2 TOP: Foundation Edition BLM: knowledge 29. ANS: Embryonic stem cells come from embryos and are pluripotent, whereas adult stem cells come from adults and are only multipotent. REF: p. 295 PTS: 1 DIF: L2 OBJ: 10.4.2 Define stem cells and explain their importance. STA: F.12.2 TOP: Foundation Edition BLM: analysis

In metaphase the sister chromatids are still attached to one another and are found in the middle of the cell, whereas in anaphase the sister chromatids have separated and are beginning to move to opposite sides of

#### 30. ANS:

Harvesting adult stem cells do not generally harm the donor, whereas harvesting embryonic stem cells usually destroys the embryo.

PTS: 1 DIF: L2 REF: p. 297 OBJ: 10.4.3 Identify the possible benefits and issues relating to stem cell research.

STA: G.12.3 | H.12.3 BLM: evaluation

31. ANS:

Garden pea plants produce many offspring, they have traits that come in only two forms, and crosses between the plants can be controlled easily.

PTS: 1	DIF: L3	REF: p. 308   p. 309
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OBJ: 11.1.1 Describe Mendel's studies and conclusions about inheritance.

STA: F.12.3 TOP: Foundation Edition BLM: synthesis

### 32. ANS:

Allowing the  $F_1$  pea plants to self-pollinate caused the recessive phenotype to reappear in the  $F_2$  generation. Self-pollination of the  $F_1$  plants also allowed the 3:1 phenotype ratios to occur, supporting Mendel's theory. Self-pollination showed that traits controlled by recessive alleles could reappear in the  $F_2$  generation.

PTS: 1 DIF: L3 REF: p. 311 | p. 312 OBJ: 11.1.2 Describe what happens during segregation. STA: F.12.3 BLM: synthesis

33. ANS:

If all of the individuals in a generation receive one dominant allele and one recessive allele, then they would all show the dominant trait. If they are bred, then they will pass on the dominant allele to some of their offspring and the recessive allele to others. Offspring receiving two recessive alleles will show the recessive trait, so it will reappear.

OBJ: 11.1.2 Describe what happens during segregation. STA: F.12.3

TOP: Foundation Edition BLM: analysis

34. ANS:

The phenotype ratio is 9 round, yellow seeds : 3 round, green seeds : 3 wrinkled, yellow seeds : 1 wrinkled, green seed.

PTS: 1 DIF: L2 REF: p. 317

OBJ: 11.2.2 Explain the principle of independent assortment. STA: F.12.3

TOP: Foundation EditionBLM: application

35. ANS:

Thirty of the offspring are expected to be tall and have yellow seeds.

PTS: 1 DIF: L3 REF: p. 316 | p. 317

OBJ: 11.2.2 Explain the principle of independent assortment. STA: F.12.3

BLM: analysis

36. ANS:

Both pea plants and fruit flies are small organisms and can be easily manipulated. Both can produce large numbers of offspring in a relatively short period of time.

PTS: 1 DIF: L2 REF: p. 318 OBJ: 11.2.3 Explain how Mendel's principles apply to all organisms. STA: F.12.3 TOP: Foundation Edition BLM: synthesis 37. ANS: 100% PTS: 1 DIF: L2 REF: p. 314 | p. 315 | p. 316 OBJ: 11.2.3 Explain how Mendel's principles apply to all organisms. STA: F.12.3 TOP: Foundation Edition BLM: analysis 38. ANS: Keep an arctic fox warm when its native environment would be cool. PTS: 1 DIF: L3 REF: p. 321 OBJ: 11.3.2 Explain the relationship between genes and the environment. **BLM:** evaluation STA: F.12.3 39. ANS: A diploid cell has two sets of chromosomes. DIF: L1 PTS: 1 REF: p. 323 OBJ: 11.4.1 Contrast the number of chromosomes in body cells and in gametes. STA: F.12.4 TOP: Foundation Edition BLM: knowledge 40. ANS: Homologous chromosomes are the two sets of chromosomes found in a body cell-one set inherited from the male parent and the other inherited from the female parent. PTS: 1 DIF: L2 REF: p. 323 OBJ: 11.4.1 Contrast the number of chromosomes in body cells and in gametes. **TOP:** Foundation Edition STA: F.12.4 BLM: comprehension 41. ANS: The number of chromosomes is cut in half. PTS: 1 DIF: L1 REF: p. 324 OBJ: 11.4.2 Summarize the events of meiosis. STA: F.12.4 TOP: Foundation Edition BLM: knowledge 42. ANS: Mitosis produces diploid body cells, whereas meiosis produces haploid gametes. PTS: 1 REF: p. 327 DIF: L2 OBJ: 11.4.3 Contrast meiosis and mitosis. STA: F.12.4 **TOP:** Foundation Edition BLM: analysis 43. ANS: sex cells, gametes PTS: 1 DIF: L1 REF: p. 323 | p. 325 OBJ: 11.4.3 Contrast meiosis and mitosis. STA: F.12.4 TOP: Foundation Edition BLM: knowledge 44. ANS: The genes that Mendel studied were located on different chromosomes or were located far apart on the same chromosome.

45.	PTS:1DIF:L2REF:p. 328   p. 329OBJ:11.4.4 Describe how alleles from different genes can be inherited together.STA:F.12.4BLM:evaluationANS:Crossing-over occurs most frequently between the star eye gene and the black body gene.
46.	PTS:1DIF:L3REF:p. 328   p. 329OBJ:11.4.4 Describe how alleles from different genes can be inherited together.STA:F.12.4BLM: synthesisANS:The harmless living bacteria took in pneumonia-causing DNA (genes) from the heat-killed, pneumonia-causing bacteria, as a result of which the harmless bacteria changed into bacteria that cause pneumonia.
47.	PTS:1DIF:L3REF:p. 350   p. 351OBJ:12.1.1 Summarize the process of bacterial transformation.STA:F.12.3BLM:synthesisANS:He concluded that a chemical factor, a gene, had transformed the bacteria.
48.	PTS:1DIF:L1REF:p. 339OBJ:12.1.1 Summarize the process of bacterial transformation.STA:F.12.3TOP:Foundation EditionBLM: comprehensionANS:A bacteriophage is a kind of virus that infects and kills bacteria.
49.	PTS:1DIF:L1REF:p. 340OBJ:12.1.2 Describe the role of bacteriophages in identifying genetic material.STA:F.12.3TOP:Foundation EditionBLM: comprehensionANS:DNA stores, copies, and transmits information.
50.	PTS:1DIF:L1REF:p. 342   p. 343OBJ:12.1.3 Identify the role of DNA in heredity.STA:F.12.3TOP:Foundation EditionBLM:knowledgeANS:It is most important during the formation of reproductive cells, because the loss of any genetic material then means the loss of valuable information for offspring.
51.	PTS: 1 DIF: L2 REF: p. 343 OBJ: 12.1.3 Identify the role of DNA in heredity. STA: F.12.3 BLM: synthesis ANS: The circles are the phosphate group, the pentagons are deoxyribose, and the A and T (adenosine and thymine) are the bases.
	PTS: 1 DIF: L2 REF: p. 345 OBJ: 12.2.1 Identify the chemical components of DNA. STA: F.12.3

OBJ:12.2.1 Identify the chemical components of DNA.TOP:Foundation EditionBLM:application

### 52. ANS:

The nucleotides in a strand of DNA are joined by covalent bonds between their sugar and phosphate groups, and by hydrogen bonds between the complimentary bases.

PTS: 1 DIF: L3 REF: p. 344 OBJ: 12.2.1 Identify the chemical components of DNA. STA: F.12.3 TOP: Foundation Edition BLM: synthesis 53. ANS: Avery repeated Griffith's experiment, and identified the component of the cell that caused transformation. PTS: 1 DIF: L3 REF: p. 338 | p. 339 OBJ: 12.1.1 Summarize the process of bacterial transformation. TOP: Foundation Edition BLM: synthesis STA: F.12.3 54. ANS: Hershey and Chase labeled the DNA of a bacteriophage with <sup>32</sup>P, and found that after the bacteria were infected with the bacteriophage, the <sup>32</sup>P was in the bacteria. PTS: 1 REF: p. 341 DIF: L1 OBJ: 12.1.2 Describe the role of bacteriophages in identifying genetic material. TOP: Foundation Edition STA: F.12.3 BLM: comprehension 55. ANS: He systematically destroyed all the other kinds of molecules besides DNA in the dead-cell mixture before using the mixture to successfully transform harmless bacteria into helpful bacteria. PTS: 1 DIF: L2 REF: p. 340 OBJ: 12.1.1 Summarize the process of bacterial transformation. **TOP:** Foundation Edition STA: F.12.3 BLM: synthesis 56. ANS: Rosalind Franklin used powerful X-ray beams to make diffraction photographs that gave Watson and Crick the clues they needed to determine DNA's structure. REF: p. 346 | p. 347 PTS: 1 DIF: L2 OBJ: 12.2.2 Discuss the experiments leading to the identification of DNA as the molecule that carries STA: F.12.3 **TOP:** Foundation Edition the genetic code. BLM: synthesis 57. ANS: The percentage of adenine would have increased by about 5 percent. PTS: 1 DIF: L3 REF: p. 345 OBJ: 12.2.2 Discuss the experiments leading to the identification of DNA as the molecule that carries the genetic code. STA: F.12.3 BLM: analysis 58. ANS: The hydrogen bonds between the base pairs must be broken, and the molecule must unwind. DIF: L2 REF: p. 350 p. 351 PTS: 1 OBJ: 12.3.1 Summarize the events of DNA replication. STA: F.12.3 TOP: Foundation Edition BLM: analysis 59. ANS: The molecule is DNA polymerase, an enzyme that joins individual nucleotides to make a strand of DNA.

DIF: L1

PTS: 1

60.	OBJ: 12.3.1 Summ TOP: Foundation E ANS: In prokaryotes, DNA places.	dition	BLM: knowledge	STA: F.12.3 ryotes DNA replication starts in many
	PTS: 1 OBJ: 12.3.2 Compa STA: F.12.4	DIF: L1 are DNA replication TOP: Foundation	REF: p. 353 in prokaryotes with tha Edition	t of eukaryotes. BLM: knowledge
SCIENCE	SKILLS			
1.	ANS: Sample answer: The mouse should give o		respiration is $6O_2 + C_6$	$_{5}H_{12}O_{6} \rightarrow 6CO_{2} + 6H_{2}O + Energy.$ The
	PTS: 1 respiration.	DIF: L2	REF: p. 251	OBJ: 9.1.2 Define cellular
	respiration. STA: F.12.1	DIF: L2 TOP: Foundation	I.	BLM: application
2.	respiration. STA: F.12.1 ANS: Sample answer: The respiration. Fresh air C, and D. Air mixed	TOP: Foundation mouse requires oxys containing oxygen f with whatever the m	Edition gen and sugar from food flows in through the tub nouse gives off flows fro	
2.	respiration. STA: F.12.1 ANS: Sample answer: The respiration. Fresh air C, and D. Air mixed	TOP: Foundation mouse requires oxys containing oxygen f with whatever the m	Edition gen and sugar from food flows in through the tub nouse gives off flows fro	BLM: application d (glucose) to carry out cellular bes from outside the flasks into flasks B, om flask B into flask A. The mouse

REF: p. 351

3. ANS:

Sample answer: If the mouse is carrying out cellular respiration, it will give off CO<sub>2</sub>. The CO<sub>2</sub> will flow into flask A, and the phenolphthalein in flask A will change from pink to clear.

PTS:	1	DIF:	L2	REF:	p. 251	OBJ:	9.1.2 Define cellular
respira	ation.						
STA:	F.12.1	TOP:	Foundation Ed	lition		BLM:	application
DICA							

4. ANS:

Sample answer: The cricket, like all living organisms, is carrying out cellular respiration. However, the mouse is larger than the cricket and gives off more  $CO_2$  than the cricket. After one hour, the cricket probably has not given off enough  $CO_2$  to measure. If the scientist allows the experiment to continue for several hours, she will see that more  $CO_2$  is given off by the cricket over time.

PTS: 1	DIF: L3	REF: p. 251	OBJ: 9.1.2 Define cellular
respiration.			
STA: F.12.1	BLM: synthesis		
ANS:			

5. NS:

Sample answer: The mouse that had been exercising should give off more  $CO_2$  because this mouse will be breathing more heavily. This mouse might even have an oxygen debt to repay, which means it is making up for the oxygen and energy it used up during the exercise.

PTS:1DIF:L2REF:p. 265OBJ:9.3.2 Identify the pathways the body uses to release energy during exercise.STA:F.12.10BLM: analysis

## ESSAY

1. ANS:

If the energy in glucose were released in just one step, most of the energy would be lost as heat. The gradual process of cellular respiration allows the cell to control the release of energy into packages of ATP that can be used more efficiently for cell activities.

PTS: 1	DIF: L3	REF: p. 250	OBJ: 9.1.2 Define cellular	BJ: 9.1.2 Define cellular	
respiration.					
STA: F.12.1	BLM: synthesis				

2. ANS:

Sample answer: Glycolysis is the breakdown of glucose into 2 molecules of pyruvic acid, producing 4 ATP molecules. An initial input of 2 ATP molecules is required to start glycolysis; thus, there is a net gain of 2 ATP molecules. This process produces 2 high-energy electrons, which are passed to NAD<sup>+</sup> to form NADH. If oxygen is present, glycolysis leads to the Krebs cycle and the electron transport chain. If oxygen is not present, glycolysis is followed by the rest of fermentation.

PTS: 1 DIF: L2 REF: p. 254 | p. 262

OBJ: 9.2.1 Describe what happens during glycolysis. STA: F.12.1

TOP: Foundation Edition BLM: synthesis

3. ANS:

Sample answer: During the Krebs cycle, pyruvic acid is broken down into carbon dioxide in a series of energy-extracting reactions. Coenzyme A forms acetyl-CoA, which later becomes citric acid. Citric acid is then broken down, CO<sub>2</sub> is released, and electrons are transferred to energy carriers. One molecule of pyruvic acid gives 4 molecules of NADH, 1 molecule of FADH<sub>2</sub>, and 1 molecule of ATP.

PTS: 1 DIF: L2 REF: p. 256 | p. 257

OBJ: 9.2.2 Describe what happens during the Krebs cycle. STA: F.12.1

TOP: Foundation Edition BLM: synthesis

4. ANS:

During brief periods of intense activity, muscle cells may use oxygen faster than it can be supplied by the body. When the oxygen supply gets very low, the electron transport chain cannot function because oxygen serves as its final electron acceptor. This forces the Krebs cycle to stop. In this anaerobic situation, the muscle cells can produce ATP only by means of lactic acid fermentation.

PTS:1DIF:L2REF:p. 262 | p. 265OBJ:9.3.2 Identify the pathways the body uses to release energy during exercise.STA:F.12.10TOP:Foundation EditionBLM: analysis