Chapter 1—Sex and Development

MULTIPLE CHOICE

1.	What is the normal r a. 44 b. 42 c. 40 d. 46 e. 45	number of chro	omosomes for a	human?		
	ANS: D	PTS: 1	DIF:	Easy	REF:	p. 14
2.	 Which of the following a. Carries XX, XY b. Cell division occord c. Carries 1 pair of d. Carries an X or Ye. Undergoes multiplication 	, or YY chrom curs before fer sex chromoso Y chromosome	tilization. mes and 22 pair e and a single se	rs of autosomal t of 22 autosom	nal chro	
	ANS: C	PTS: 1	DIF:	Easy	REF:	p. 14
3.		ction of sperm, an males are c ge, the ratio of number of mal	we would expe onceived. F males to femal les and females	ct the sex ratio es is 1:1. are born.	to be 2	males for every 1
	ANS: C	PTS: 1	DIF:	Moderate	REF:	p. 14
4.	Whether a fetus devea complex interab. hormones produc. how many spernd. the number of aue. only on which se	iction of genes ced by the mot n enter the egg atosomes prese	and environme ther during preg ent.	nt.		
	ANS: A	PTS: 1	DIF:	Easy	REF:	p. 21
5.	What is the normal r a. 21 b. 22 c. 23 d. 24 e. 42 ANS: C	number of pairs		es for humans? Easy	REF:	p. 14

6. What is the sex chromosome make-up of a normal male?

a. XXY b. XY c. XX d. XXX e. XO ANS: B PTS: 1 DIF: Easy REF: p. 14 7. Autosomes are a. non-sex chromosomes. b. another term for the sex chromosomes. c. a pair of chromosomes. d. chromosomes with mutated genes. e. capable of forming Barr bodies. ANS: A PTS: 1 DIF: Easy REF: p. 14 8. Sperm sorting relies on ______ between sperm carrying the X chromosome and sperm carrying the Y chromosome. a. differences in electrical charges b. direct microscopic examination and recognition of visual differences c. differences in the swimming speed d. magnetic differences e. differences in the amounts of fluorescent dye binding. ANS: E PTS: 1 DIF: Moderate REF: p. 15 9. The process of sex selection a. is only possible after the egg has been fertilized. b. is only used to avoid having children with genetic disorders. c. carries a small risk of miscarriage for the mother. d. allows a couple to choose the sex of their child. allows a couple to alter the sex of their unborn child. e. ANS: D PTS: 1 REF: p. 15 DIF: Moderate 10. What is involved in *in vitro* fertilization? a. The woman receives hormone treatments. b. Eggs are surgically removed from the woman's ovaries. c. Both sperm and eggs are placed in a dish to allow fertilization. d. A sperm sample must be provided. e. All of these are steps in *in vitro* fertilization. ANS: E PTS: 1 DIF: Difficult REF: p. 16 11. In humans, fertilization usually occurs a. in the ovary. b. in the fallopian tube. c. in the vagina. d. in the uterus. e. right at the cervix. ANS: B PTS: 1 DIF: Easy REF: p. 17

12.	How many sperm us a. 1 b. 2 c. 3 d. 4 e. 5	sually en	ter an egg?				
	ANS: A	PTS:	1	DIF:	Easy	REF:	p. 17
13.	What are the differea. Inner cell massb. Inner cavityc. Middle layer ofd. Inner cell masse. All of these .	cells	-	2			
	ANS: D	PTS:	1	DIF:	Easy	REF:	p. 18
14.	The chorion a. contains human b. is a membrane i c. protects the emb d. forms in the last e. is an extension of	nside the oryo. trimeste	e embryo. er of pregnancy				
	ANS: C	PTS:	1	DIF:	Moderate	REF:	p. 18
15.	Human chorionic go a. nourishes the en b. stimulates the fo c. is present in uno d. prevents the exp e. is transported to	nbryo. ormation letectable oulsion o	of the placenta e amounts thro f the embryo.	ughout			
	ANS: D	PTS:	1	DIF:	Difficult	REF:	p. 18
16.	The villi a. extend into the s b. are fingerlike pr c. will eventually f d. come in contact e. All of these are	ojection form par with the	s. t of the placent	a.			
17.	 a. 1-2 weeks after b. 2-3 weeks c. 5 weeks d. 8 weeks e. 16 weeks 	fertilizat	bes the fetus ha				ves that can open?
	ANS: E	PTS:	1	DIF:	Easy	REF:	p. 20

- 18. During which trimester(s) does the most rapid fetal growth take place?
 - a. first trimester
 - b. second trimester
 - c. third trimester
 - d. There is equal fetal growth in all trimesters.
 - e. It varies depending on each individual pregnancy.

ANS: C PTS: 1 DIF: Easy REF: p. 18-19

19. At what point in development is the embryo considered a fetus?

- a. After the first month.
- b. After eight weeks.
- c. After the fourth month.
- d. After the sixth month.
- e. Just before birth.

ANS: B PTS: 1 DIF: Moderate REF: p. 20

- 20. When can the mother usually feel movements of the fetus' arms and legs?
 - a. within days after fertilization
 - b. during the second month
 - c. around the fourth month
 - d. not until the third trimester
 - e. only in the last two months of pregnancy

ANS: C PTS: 1 DIF: Moderate REF: p. 20

21. When can ultrasound be used to determine the sex of a fetus?

- a. within days after implantation
- b. 1-2 weeks after fertilization
- c. at the end of the first trimester
- d. at the beginning of the second trimester
- e. not until the third trimester

ANS: D PTS: 1 DIF: Difficult REF: p. 20

22. Which of the following is **FALSE**?

- a. Testes secrete the hormone testosterone.
- b. Testosterone promotes the development of male reproductive organs.
- c. Testosterone promotes the development of the ovary.
- d. Testosterone promotes the development of secondary male sex characteristics.
- e. The Y chromosome must be present for the secretion of testosterone.

ANS: C PTS: 1 DIF: Easy REF: p. 21

- 23. Which of the following is **FALSE** about androgen insensitivity?
 - a. Complete androgen insensitivity is caused by a mutation in a gene on the X chromosome.
 - b. Individuals have a mutation in the androgen receptor gene.
 - c. Development of individuals continues as if testosterone were absent.
 - d. Individuals with this condition have an XX sex chromosome combination.
 - e. Individuals with complete androgen insensitivity have the chromosome combination of a female but appear male.

ANS: D PTS: 1 DIF: Difficult REF: p. 23-24

- 24. Individuals with complete androgen insensitivity
 - a. menstruate after reaching puberty.
 - b. are phenotypically males.
 - c. often have testes present in their abdomens.
 - d. usually have no problems reproducing.
 - e. have a set of XX sex chromosomes.

ANS: C PTS: 1 DIF: Difficult REF: p. 23-24

- 25. Gonadal sex is determined
 - a. at fertilization.
 - b. at birth.
 - c. upon the formation of external genitalia.
 - d. when the presence or absence of the *SRY* gene determines the formation of testes or ovaries.
 - e. when either testosterone or estrogen is produced.

ANS: D PTS: 1 DIF: Difficult REF: p. 21

- 26. The _____ will eventually form the fetus.
 - a. inner cell mass of the blastocyst
 - b. outer layer of cells of the blastocyst
 - c. internal cavity of the blastocyst
 - d. outer layer of chorion
 - e. inner layer of placenta

ANS: A PTS: 1 DIF: Easy REF: p. 18

27. The _____ is the source for embryonic stem cells.

- a. inner cell mass of the blastocyst
- b. outer layer of cells of the blastocyst
- c. internal cavity of the blastocyst
- d. outer layer of chorion
- e. inner layer of placenta

ANS: A PTS: 1 DIF: Easy REF: p. 18

- 28. Sex testing in the Olympics prior to 2000 was based on
 - a. testosterone levels.
 - b. the presence or absence of the Y chromosome.
 - c. ultrasound scanning.
 - d. the presence or absence of Barr bodies.
 - e. estrogen levels.

ANS: D PTS: 1 DIF: Moderate REF: p. 25

29. Intersexuality

- a. is determined only by the number of sex chromosomes an individual possesses.
- b. is a condition in which the chromosomal and phenotypic sex of an individual match.
- c. describes a condition in which an individual's phenotype cannot be classified as either male or female.
- d. is determined only by the autosomes an individual possesses.
- e. is determined at fertilization.

	ANS: C	PTS: 1	DIF: Moderate	REF: p. 25
30.	a. 0 b. 1 c. 2 d. 3	ies would be found in Ild vary depending on		al with Turner syndrome? 1al.
	ANS: A	PTS: 1	DIF: Moderate	REF: p. 25
TRUI	E/FALSE			
1.	In an ultrasound scan	n, the sex organs can b	e seen as early as 7 we	eeks after implantation.
	ANS: F	PTS: 1	DIF: Moderate	REF: p. 20
2.	Development of the s	sex organs is influence	ed by the presence or a	bsence of the hormone testosterone.
	ANS: T	PTS: 1	DIF: Easy	REF: p. 21
3.	Chromosomal sex is	determined at fertiliza	tion.	
	ANS: T	PTS: 1	DIF: Easy	REF: p. 21
4.	Complete androgen i	nsensitivity is genetica	ally controlled.	
	ANS: T	PTS: 1	DIF: Moderate	REF: p. 23
5.		e X chromosome have ter in the sperm sorting		n containing the Y chromosome and
	ANS: T	PTS: 1	DIF: Easy	REF: p. 15
6.	Sex selection can use	e sperm sorting or prei	mplantation genetic di	iagnosis.
	ANS: T	PTS: 1	DIF: Easy	REF: p. 15
7.	In males, the develop located on the Y chro		external sex organs dep	pends on the actions of the SRY gene
	ANS: T	PTS: 1	DIF: Moderate	REF: p. 21
8.	In females, one X ch	romosome out of the t	wo becomes a Barr bo	ody.
	ANS: T	PTS: 1	DIF: Easy	REF: p. 24
9.	An individual who is	(45, X) would be phe	notypically female.	
	ANS: T	PTS: 1	DIF: Difficult	REF: p. 25

10. An individual who is (47, XXY) would be phenotypically female.

ANS: F PTS: 1 DIF: Difficult REF: p. 21

MATCHING

Match the appropriate chromosomal composition to the sex of the individual.

- a. Female
- b. Male
- 1. XX
- 2. XY
- 3. XO
- 4. XXY

1. ANS: A	PTS: 1	DIF: Easy	REF: p. 14
2. ANS: B	PTS: 1	DIF: Easy	REF: p. 14
3. ANS: A	PTS: 1	DIF: Moderate	REF: p. 14
4. ANS: B	PTS: 1	DIF: Moderate	REF: p. 14

Match the appropriate term with the description.

- a. finger-like projections
- b. large hollow ball of cells
- c. attachment of embryo to uterus
- d. source of embryonic stem cells
- e. hCG is produced by this membrane
- 5. Blastocyst
- 6. Implantation
- 7. Villi
- 8. Inner cell mass
- 9. Chorion

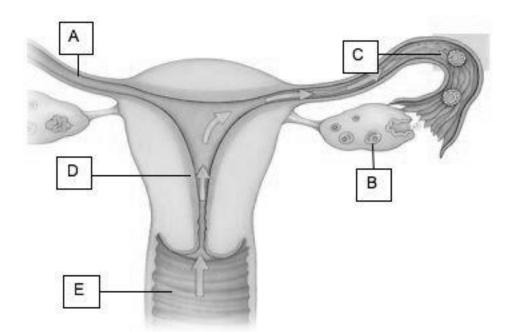
5. ANS: B	PTS: 1	DIF: Easy	REF: p. 18
6. ANS: C	PTS: 1	DIF: Easy	REF: p. 18
7. ANS: A	PTS: 1	DIF: Moderate	REF: p. 18
8. ANS: D	PTS: 1	DIF: Moderate	REF: p. 18
9. ANS: E	PTS: 1	DIF: Moderate	REF: p. 18

Match the description to the number of chromosomes.

- a. normal number of chromosomes in humans
- b. number of autosomal chromosomal pairs
- c. number of chromosomes present in an individual with Kleinfelter syndrome
- d. number of chromosomes present in an individual with Turner syndrome
- 10. 22
- 11. 46
- 12. 47
- 13. 45

10.	ANS: B	PTS: 1	DIF:	Easy	REF: p. 14
11.	ANS: A	PTS: 1	DIF:	Easy	REF: p. 14
12.	ANS: C	PTS: 1	DIF:	Moderate	REF: p. 25
13.	ANS: D	PTS: 1	DIF:	Moderate	REF: p. 25

Match the following descriptions with the appropriate letter.



- 14. sperm enter the female reproductive tract here
- 15. implantation will occur here
- 16. fertilization occurring
- 17. fallopian tube
- 18. the nonspecific gonad develops into this structure in human females

14. ANS: E	PTS: 1	DIF: Moderate	REF: p. 17-18
15. ANS: D	PTS: 1	DIF: Moderate	REF: p. 17-18
16. ANS: C	PTS: 1	DIF: Moderate	REF: p. 17-18
17. ANS: A	PTS: 1	DIF: Moderate	REF: p. 17-18
18. ANS: B	PTS: 1	DIF: Moderate	REF: p. 17-18

ESSAY

1. In countries like China and India, sex selection using ultrasound, amniocentesis, preimplantation genetic diagnosis, and sperm sorting has resulted in an imbalance of male and female offspring in the population. The trend has led to a gradual increase in the number of males born. How would you anticipate this sex selection could influence society in these countries? Do you think the same possibility exists in this country?

ANS:

Answers will vary but should discuss how the modified sex ratio could affect schooling, work, marriage, etc. Answers should also point out that this is less likely to occur in the United States because there is less of a demand for male children.

PTS: 1 DIF: Moderate REF: Entire chapter

2. One technology that has been valuable for couples who have a known increased risk of passing on a genetic disease (biochemical or chromosomal) is preimplantation genetic diagnosis (PGD). Ethically, some individuals feel this is "playing God" and should not be done; others consider the method a "scientific miracle" and are grateful for the opportunity to virtually insure that they will not have a disease-stricken child. Where would you fall on this spectrum of varying opinions? What factors have you considered in forming your opinion? Now suppose that you and your spouse are interested in starting a family but find out that the female (you or your spouse) is a carrier for Hunter syndrome, a genetic disorder in which the affected infants become blind, deaf, mentally retarded, and seldom live past age 5. Does this change your opinion of using PGD to select for a healthy child? Explain.

ANS:

Answers will vary but could include discussion about religious beliefs, societal costs, personal costs, mental, physical, and emotional stress, and how having a child with a severe genetic disorder can affect all the individuals involved. Answers may also delve into usage of PGD in cases of genetic disorders of varying severity.

PTS: 1 DIF: Moderate REF: Entire chapter

SHORT ANSWER

1. The X and Y chromosomes of the human genome are termed the sex chromosomes. Explain how the sex chromosome complement for a normal female and a normal male is determined at fertilization.

ANS:

Since females only have two X chromosomes, all of their eggs will have an X chromosome. Since males have an X and a Y chromosome, half of the males' sperm will contain an X and half a Y chromosome. If the egg is fertilized with an X chromosome-containing sperm, the resulting zygote will be XX, female. If the egg is fertilized with a Y chromosome-containing sperm, the resulting zygote will be XY, male.

PTS: 1 DIF: Easy REF: p. 14

2. Distinguish between chromosomal, gonadal, and phenotypic stages of sex development for a male. When does each occur and what is the key feature of each?

ANS:

Chromosomal sex is determined at fertilization and depends on whether the zygote carries two X chromosomes or an X and a Y. At about the seventh or eighth week of pregnancy, the presence of the Y chromosome causes the nonspecific gonads to become testes. In the phenotypic stage, sometime before the 12th week, production of masculinizing hormones such as testosterone lead to the formation of male external genitals.

PTS: 1 DIF: Difficult REF: p. 21

3. Describe the genetic factors required for the development of the male internal and external sex organs. Describe the genetic factors required for the development of the female internal and external sex organs.

ANS:

In males, the pathway begins with the action of a gene (*SRY*) on the Y chromosome, the presence of at least one X chromosome, and expression of genes carried on the other 22 chromosomes. In females, the pathway begins with the presence of two X chromosomes, the absence of Y chromosome genes, and expression of a female-specific set of genes on the X chromosome and the other 22 chromosomes.

PTS: 1 DIF: Moderate REF: p. 21

4. Explain why the sex ratio for humans changes over time.

ANS:

Although we cannot be completely certain about the numbers or the reasons, estimates indicate that more males are conceived than females. This does not mean that the sex ratio at birth reflects this imbalance. At birth the ratio is about 1:1.05 (100 females for every 105 males). As a generation ages, these numbers change. More males than females die in childhood, and when this generation reaches the age of 20, the ratio of males to females moves closer to 1:1. Beyond this age, females begin to outnumber males because men have a shorter life span than women.

PTS: 1 DIF: Moderate REF: p. 14

PROBLEM

1. During early embryonic development, one of the X chromosomes is randomly inactivated in each cell to form a Barr body, a condensed structure near the nuclear envelope that can be used to identify the sex of the individual and/or the presence of a sex chromosome abnormality. In individuals with abnormal numbers of X chromosomes, all but one of the X chromosomes will be inactivated. What would the number of Barr bodies, sex, and syndrome (if any) be for each of the following situations?

Sex chromosomes	Number of Barr bodies	Sex	Syndrome (if any)
XX			
XY			
XXX			
XO			
XYY			
XXY			
XXXY			
ANS:			
Sex chromosomes	Number of Barr bodies	Sex	Syndrome (if any)
XX	1	F	Normal female
XY	0	Μ	Normal male
XXX	2	F	Not listed in chapter

Human Genetics and Society 2nd Edition Yashon Test Bank

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XO	0	F	Turner syndrome
XYY	0	M	Not listed in chapter
XXY	1	M	Kleinfelter syndrome
XXXY	2	M	Kleinfelter syndrome
PTS: 1	DIF: Difficult	REF: p. 2	4-25