

Genetics 301 Sample Final Examination
Spring 2003

50 Multiple Choice Questions-(Choose the best answer)

1. A cross between two true breeding lines one with dark blue flowers and one with bright white flowers produces F1 offspring that are light blue. When the F1 progeny are selfed a 1:2:1 ratio of dark blue to light blue to white flowers is observed. What genetic phenomenon is consistent with these results?
 - a. epistasis
 - b. incomplete dominance
 - c. codominance
 - d. inbreeding depression
 - e. random mating

2. Mutations which occur in body cells which do not go on to form gametes can be classified as:
 - a. auxotrophic mutations
 - b. somatic mutations
 - c. morphological mutations
 - d. oncogenes
 - e. temperature sensitive mutations

3. What would be the frequency of AABBCC individuals from a mating of two AaBbCc individuals?
 - a. 1/64
 - b. 1/32
 - c. 1/16
 - d. 1/8
 - e. 3/16
 - f. 1/4

4. The stage of meiosis in which chromosomes pair and cross over is:
 - a. prophase I
 - b. metaphase I
 - c. prophase II
 - d. metaphase II
 - e. anaphase II

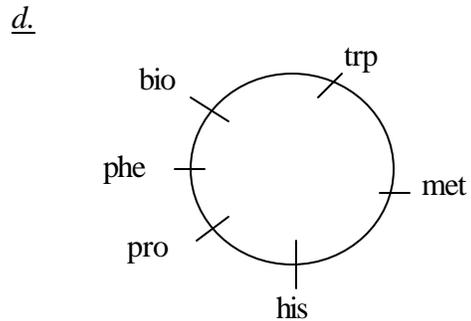
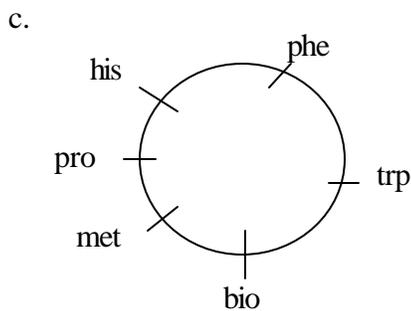
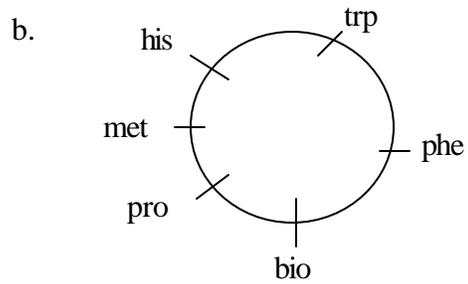
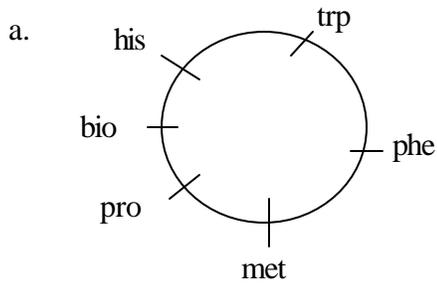
5. Polyploidy refers to:
- extra copies of a gene adjacent to each other on a chromosome
 - an individual with complete extra sets of chromosomes
 - a chromosome which has replicated but not divided
 - multiple ribosomes present on a single mRNA
 - an inversion which does not include the centromere
6. A gene showing codominance-
- has both alleles independently expressed in the heterozygote
 - has one allele dominant to the other
 - has alleles tightly linked on the same chromosome
 - has alleles expressed at the same time in development
 - has alleles that are recessive to each other
7. The phenomenon of “independent assortment” refers to:
- expression at the same stage of development
 - unlinked transmission of genes in crosses resulting from being located on different chromosomes, or far apart on the same chromosome.
 - association of an RNA and a protein implying related function
 - independent location of genes from each other in an interphase cell
 - association of a protein and a DNA sequence implying related function
8. Mendel’s law of segregation, as applied to the behavior of chromosomes in meiosis, means that:
- pairing of homologs will convert one allele into the other, leading to separation of the types.
 - alleles of a gene separate from each other when homologs separate in meiosis I, or in meiosis II if there is a single crossover between the gene and the centromere.
 - genes on the same chromosome will show 50% recombination
 - alleles of a gene will be linked and passed on together through meiosis
9. Which component of transcribed RNA in eukaryotes is present in the initial transcript but is removed before translation occurs:
- Intron
 - 3’ Poly A tail
 - Ribosome binding site
 - 5’ cap
 - codons coding for the protein to be produced

10. Choose the correct statement about the genetic code.
- includes 61 codons for amino acids and 3 stop codons
 - almost universal; exactly the same in most genetic systems
 - three bases per codon
 - some amino acids are coded by multiple codons
 - all of the above
11. X-chromosome inactivation
- normally takes place in males but not females
 - is the cause of the Y chromosome being genetically inactive
 - takes place in humans so that the same X chromosome is inactive in all of the cells of a female
 - occurs in fruit flies but not in mammals
 - results in genetically turning off one of the two X chromosomes in female mammals
12. DNA ligase is:
- an enzyme that joins fragments in normal DNA replication
 - an enzyme involved in protein synthesis
 - an enzyme of bacterial origin which cuts DNA at defined base sequences
 - an enzyme that facilitates transcription of specific genes
 - an enzyme which limits the level to which a particular nutrient reaches
13. An Hfr strain of E. coli contains:
- a vector of yeast or bacterial origin which is used to make many copies of a particular DNA sequence
 - a bacterial chromosome with a human gene inserted
 - a bacterial chromosome with the F factor inserted
 - a human chromosome with a transposable element inserted
 - a bacterial chromosome with a phage inserted

14. An experiment was conducted in *E. coli* to map the following genes (pro, his, bio, met, phe and trp) on a circular map using 3 different Hfr strains.

Strain 1 Order of transfer (early to late):	trp	met	his	pro
Strain 2 Order of transfer (early to late):	his	met	trp	bio
Strain 3 Order of transfer (early to late):	pro	phe	bio	trp

Based on the results what is the most likely map?



15. Generation of antibody diversity in vertebrate animals takes place through:

- the presence of as many genes in the germ line as there are types of antibodies possible.
- infection with bacteria carrying antibody genes
- infection with viruses carrying antibody genes
- polyploidy in antibody-forming cells
- rearrangement of DNA in tissues that go on to produce antibodies

16. Replication of DNA:

- takes place in a “conservative” manner
- takes place in a “dispersive” manner
- takes place in a “semi-conservative” manner
- usually involves one origin of replication per chromosome in eukaryotes
- takes place only in the 3’ to 5’ direction

17. A duplication is:
- an exchange between non-homologous chromosomes, resulting in chromosomes with new genes adjacent to each other.
 - loss of genes in part of a chromosome
 - an extra copy of the genes on part of a chromosome
 - a reversal of order of genes on a chromosome
 - an extra set of chromosomes in an organism
18. What is the co-transduction frequency for the A and B genes, from the following dataset? (Assume that there has been selection for the A⁺ form of the A gene).
- | Genotype | Number |
|--|--------|
| A ⁺ B ⁺ C ⁺ | 10 |
| A ⁺ B ⁺ C ⁻ | 30 |
| A ⁺ B ⁻ C ⁺ | 20 |
| A ⁺ B ⁻ C ⁻ | 40 |
- .10
 - .20
 - .30
 - .40
 - .50
19. A mutation in a codon leads to the substitution of one amino acid with another. What is the name for this type of mutation?
- nonsense mutation
 - missense mutation
 - frameshift mutation
 - promoter mutation
 - operator mutation
20. Mapping of human chromosomes:
- has been restricted to the sex chromosomes because of small family sizes
 - proceeded much more successfully as large numbers of DNA markers became available.
 - has determined that the number of linkage groups is about twice the number of chromosomes
 - has demonstrated that almost all of the DNA is involved in coding for genes
 - has shown that there are more genes on the Y than on the X chromosome

21. Homeobox sequences
- are present in the genome of many animal species
 - are found in prokaryotes but not in eukaryotes
 - were identified as the integration sites for bacterial viruses
 - represent integration sites for transposable elements
 - represent the termination signals for transcription
22. Tracing of a cell lineage during development means that:
- the cells giving rise to and derived from a specific cell are known
 - the sequence of the enhancers for developmental genes is known
 - the regulatory genes for the organism have been genetically mapped
 - cell components in the membrane involved in signaling have been isolated
 - cell components in the nucleus involved in signaling have been isolated
23. Zinc finger proteins and helix-turn-helix proteins are:
- types of DNA-binding proteins
 - involved in the control of translation
 - components of ribosomes
 - part of the hemoglobin in blood cells
 - bound to transfer RNA during replication
24. Transcriptional activator proteins:
- transcribe a messenger off a DNA template
 - bind to ribosomes to activate the production of specific proteins
 - are produced during an infection of bacteria by a phage
 - are essential to function of transfer RNAs during translation
 - bind regions near a eukaryotic gene and allow an RNA polymerase to transcribe a gene
25. Differential distribution of substances in the egg most typically results in:
- differences in gene expression which may establish a pattern in the embryo as the cells divide
 - amplification of specific genes during development
 - development of polyploid tissues
 - loss of specific genes during development
 - dominance of genes derived from the father

26. Arabidopsis is advantageous for plant genetic research because:
- it is commercially important as a food crop
 - it is an endangered species
 - it is the closest to humans of any existing plant
 - it is a small plant with a small genome size which can be raised inexpensively
 - it is a close relative of corn and results with this species can be applied to problems in corn
27. A homeotic mutation is one which:
- is present in only one form in an individual
 - substitutes one body part for another in development
 - results in development of a tumor
 - is wild type at one temperature and abnormal at another
 - leads to increased body size in an organism
28. Assuming that the level of glucose is low, a mutation in the repressor of the lac operon in E. coli, preventing binding of the repressor to the operator, should result in:
- constitutive expression of the lac operon genes
 - lack of expression or reduced expression of the lac operon genes under all circumstances
 - expression of the genes only when lactose is present
 - expression of the genes only when lactose is absent
29. Assuming that the level of glucose is low, a mutation in the repressor associated with the lac operon of E. coli which prevents binding of the repressor to lactose should result in:
- constitutive expression of the lac operon genes
 - lack of expression or reduced expression of the lac operon genes under all circumstances
 - expression of the genes only when lactose is present
 - expression of the genes only when lactose is absent
30. RFLP analysis is a technique that
- uses hybridization to detect specific DNA restriction fragments in genomic DNA
 - is used to determine whether a gene is transcribed in specific cells
 - measures the transfer frequency of genes during conjugation
 - is used to detect genetic variation at the protein level.
 - is used to amplify genes for producing useful products

31. Plasmid vectors for cloning

- a. can generally accommodate larger inserts than phage vectors can
- b. grow within bacteria, and are present in bacterial colonies on an agar plate
- c. can accommodate inserts of over 100 kilobases
- d. include centromeres to allow propagation in yeast
- e. burst bacteria and form plaques on a “lawn” of bacteria

32. Simple tandem repeat polymorphisms in humans are most useful for:

- a. solving criminal and paternity cases
- b. reconstructing the relationships of humans and chimps.
- c. estimating relationships of humans and Neanderthals
- d. transferring disease resistance factors into bone marrow cells
- e. estimating matches for blood transfusions

33. The polymerase chain reaction or PCR is a technique that

- a. was used to demonstrate DNA as the genetic material
- b. is used to determine the content of minerals in a soil sample
- c. uses short DNA primers and a thermostable DNA polymerase to replicate specific DNA sequences in vitro.
- d. measures the ribosome transfer rate during translation
- e. detects the level of polymerases involved in replication

34. Positional cloning refers to:

- a. using a selection procedure to clone a cDNA
- b. cloning a portion of a gene using PCR
- c. isolating a gene by PCR using primers from another species
- d. isolating a gene from a specific tissue in which it is being expressed
- e. mapping a gene to a chromosomal region and then identifying and cloning a genomic copy of the gene from the region

35. Large quantities of useful products can be produced through genetic engineering involving:

- a. bacteria containing recombinant plasmids
- b. yeast carrying foreign genes
- c. transgenic plants
- d. mammals producing substances in their milk
- e. all of the above

36. On average, how many fragments would a restriction enzyme which recognizes a specific 4 base sequence in DNA be expected to cleave a double-stranded bacteriophage with a genome size of 5,000 bp into?
- about 2
 - about 4
 - about 20
 - about 50
 - about 1250
37. The “sticky ends” generated by restriction enzymes allow:
- selection for plasmids lacking antibiotic resistance
 - easy identification of plasmids which carry an insert
 - replication of transfer RNA within the bacterial cell
 - insertion of centromeres into ribosomes lacking them
 - pieces of DNA from different sources to hybridize to each other and to be joined together
38. QTL analysis is used to:
- identify RNA polymerase binding sites
 - map genes in bacterial viruses
 - determine which genes are expressed at a developmental stage
 - identify chromosome regions associated with a complex trait in a genetic cross
 - determine the most rapidly-evolving parts of genes
39. Assuming Hardy-Weinberg equilibrium, the genotype frequency of heterozygotes, if the frequency of the two alleles at the gene being studied are 0.6 and 0.4, will be:
- 0.80
 - 0.64
 - 0.48
 - 0.32
 - 0.16
40. The likelihood of an individual in a population carrying two specific alleles of a human DNA marker, each of which has a frequency of 0.2, will be:
- 0.4
 - 0.32
 - 0.16
 - 0.08
 - 0.02

41. A threshold trait is one which:
- is expressed on a continuous scale (such as blood pressure)
 - is present in a few discrete classes, but is influenced by both genetics and the environment (such as diabetes or schizophrenia)
 - is caused by only a single gene, with no environmental influence
 - is present in a very low frequency in the population
 - is associated with superior survival of the heterozygote
42. Mitochondrial DNA is advantageous for evolutionary studies because:
- it is inherited only through the female parent and thus evolves in a way that allows trees of relationship to be easily constructed
 - it is inserted into the X chromosome
 - it first appeared in humans and is not found in other animals
 - it evolves more slowly than the genes in the nucleus
 - it was derived from the globin genes as an extra copy
43. What are the assumptions of Hardy Weinberg equilibrium?
- Small population size, random mating, no selection, no migration, no mutation
 - large population size, random mating, no selection, no migration, no mutation
 - large population size, random mating, heterozygotes survive the best, no migration, no mutation
 - large population size, like individuals mate, no selection, no migration, no mutation
 - large population size, random mating, no selection, migrants enter from other populations, no mutation
44. Twin studies in humans are useful because:
- they allow more refined estimates of chromosome location to be made
 - twins have a greater likelihood of being heterozygous
 - they allow improved expression of genes
 - cloning of genes is facilitated by the presence of extra copies.
 - they allow genetic as opposed to environmental influences on variation in a trait to be estimated
45. Which of the following statements about heritability are true?
- is a measure of level of gene linkage
 - is a measure of inbreeding
 - is a measure of proportion of repeated DNA in an organism
 - is a measure of the level of heterozygotes in a population
 - is a measure of the proportion of variation that is due to genetic causes

46. The allele associated with sickle cell anemia apparently reached a high frequency in some human populations due to:

- a. random mating
- b. superior fitness of heterozygotes in areas where malaria was present
- c. migration of individuals with the allele into other populations
- d. a high mutation rate at that specific gene
- e. genetic drift

47. An increase in the inbreeding coefficient, F , is likely to result in:

- a. reduced likelihood of heterozygotes being present in a population
- b. higher proportion of genes that show linkage
- c. higher proportion of genes with introns
- d. lower level of difference between proteins in two daughter cells
- e. higher level of difference between RNA molecules in two daughter cells

48. Most new mutations appear to be:

- a. beneficial
- b. neutral or deleterious
- c. present in homozygotes rather than heterozygotes
- d. detectable using allozyme studies (protein electrophoresis)
- e. present within pericentric inversions

49. If the frequency of males affected with an X-linked recessive condition in a human population is .10 (one in ten), what will be the expected frequency of affected females?

- a. 0.0001
- b. .001
- c. .02
- d. .01
- e. .05

50. The following genotypes are found in a population:

<u>AA</u>	<u>Aa</u>	<u>aa</u>
70	50	20

What are the allele frequencies of A and a?

- a. $A = 0.86$ and $a = 0.14$
- b. $A = 0.68$ and $a = 0.32$
- c. $A = 0.63$ and $a = 0.36$
- d. $A = 0.32$ and $a = 0.68$
- e. $A = 0.36$ and $a = 0.63$