

Genetics 301 Sample Final Examination  
Spring 2003

Multiple Choice Questions (Choose the best answer) (The actual test will have 30 questions worth 4 points each. Coverage will be 25 questions over new material, 5 questions over old material. This exam has 50 questions and a somewhat higher proportion of questions from the earlier portions of the course).

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

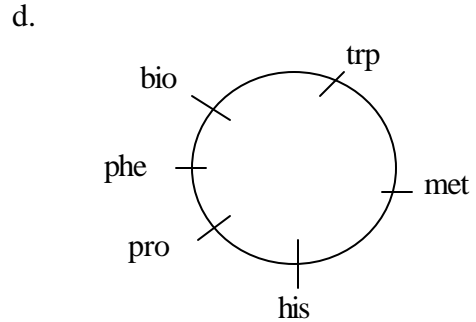
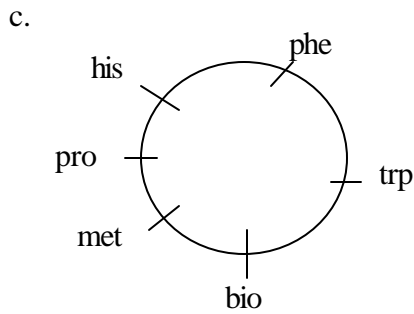
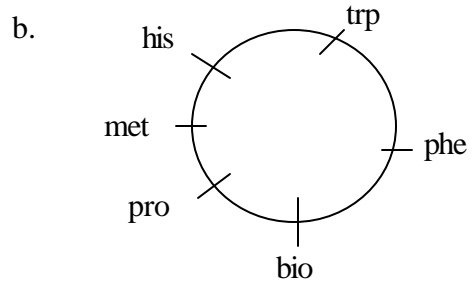
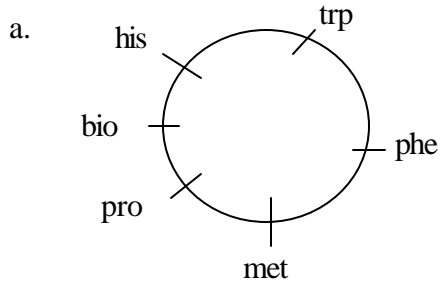
5. Polyploidy refers to:
- extra copies of a gene adjacent to each other on a chromosome
  - an individual with complete extra sets of chromosomes
  - a chromosome which has replicated but not divided
  - multiple ribosomes present on a single mRNA
  - an inversion which does not include the centromere
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:

- a. takes place in a “conservative” manner
- b. takes place in a “dispersive” manner
- c. takes place in a “semi-conservative” manner
- d. usually involves one origin of replication per chromosome in eukaryotes
- e. takes place only in the 3' to 5' direction

17. A duplication is:

- a. an exchange between non-homologous chromosomes, resulting in chromosomes with new genes adjacent to each other.
- b. loss of genes in part of a chromosome
- c. an extra copy of the genes on part of a chromosome
- d. a reversal of order of genes on a chromosome
- e. 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

- a. .10
- b. .20
- c. .30
- d. .40
- e. .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?

- a. nonsense mutation
- b. missense mutation
- c. frameshift mutation
- d. promoter mutation
- e. 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
- can generally accommodate larger inserts than phage vectors can
  - grow within bacteria, and are present in bacterial colonies on an agar plate
  - can accommodate inserts of over 100 kilobases
  - include centromeres to allow propagation in yeast
  - burst bacteria and form plaques on a “lawn” of bacteria
32. Simple tandem repeat polymorphisms in humans are most useful for:
- solving criminal and paternity cases
  - reconstructing the relationships of humans and chimps.
  - estimating relationships of humans and Neanderthals
  - transferring disease resistance factors into bone marrow cells
  - estimating matches for blood transfusions
33. The polymerase chain reaction or PCR is a technique that
- was used to demonstrate DNA as the genetic material
  - is used to determine the content of minerals in a soil sample
  - uses short DNA primers and a thermostable DNA polymerase to replicate specific DNA sequences in vitro.
  - measures the ribosome transfer rate during translation
  - detects the level of polymerases involved in replication
34. Positional cloning refers to:
- using a selection procedure to clone a cDNA
  - cloning a portion of a gene using PCR
  - isolating a gene by PCR using primers from another species
  - isolating a gene from a specific tissue in which it is being expressed
  - 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:
- bacteria containing recombinant plasmids
  - yeast carrying foreign genes
  - transgenic plants
  - mammals producing substances in their milk
  - 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. 0.001
- c. 0.02
- d. 0.01
- e. 0.05

50. The following genotypes are found in a population:

$\frac{AA}{70}$	$\frac{Aa}{50}$	$\frac{aa}{20}$
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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  $a = 0.63$