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國立臺灣大學 109 學年度碩士班招生考試試題

科目:遺傳學(C)

節次:

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一、單選題 (每題 3 分, 60%) ※ 注意:請於試卷內之「選擇題作答區」依序作答。 1. Which of the following is *not* one of the assumptions of the Hardy-Weinberg law? (A) The population is very large. (B) Non-random mating occurs within the population. (C) Mutations in the alleles do not occur. (D) No migration occurs into or out of the population. (E) The ability of all genotypes for survival and reproduction is the same. 2. Colorblindness is a recessive sex-linked disease that is caused by a defective gene on the X-chromosome. If a mother who is a carrier for the trait mates with a normal sighted male, what percentage of their daughters will have the disease? (A)0%(B) 25% (C) 50% (D) 100% (E) cannot be determined 3. Inversions can have an effect on the phenotype of the organism when (A) the transcribed region of a gene spans one or more of the inversion's breakpoints. (B) the inversion's breakpoints do not alter any of the inverted segments gene sequences. (C) the inversion is pericentric. (D) the inversion is paracentric. (E) Inversions never affect phenotypes. 4. Nondisjunction can occur at either the first or second division of meiosis. XYY individuals would most likely arise from nondisjunction at the \_\_\_\_\_ meiotic division in the (B) second, mother (A) first, mother (C) first, father (D) second, father (E) More than one of the choices could give rise to XYY individuals. 5. Which of the following statements is correct with respect to mitochondrial genes? (A) They are transmitted, largely intact, from parent to offspring. (B) Their inheritance shows Mendel's principle of segregation. (C) Their inheritance shows Mendel's principle of independent assortment. (D) A and B are correct. (E) A, B, and C are correct 6. In a series of two-factor crosses, 30% recombination occurs between gene A and gene B, and 26% recombination between gene B and gene C. If the order of genes on the chromosome is A-B-C, how much recombination will be seen between A and C in a two-factor cross? (E) 56% (C) 28%(D) 50% (A) 4%(B) 8% 7. A genetic analysis of an unknown infectious agent reveals that it contains only the nucleotides G, A, U, and C in the proportion 30%, 35%, 15%, and 20%, respectively. Based on this information, this infectious agent is most likely a

(B) double-stranded RNA virus

(E) single-stranded RNA virus

(C) bacterium

(A) double-stranded DNA virus

(D) single-stranded DNA virus

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<ol> <li>What is the molecular mechanism of the control of the</li></ol>		• •		rotain nhaanhamilation
• •				otem phosphorylation
(D) RNA methylation		(E) DNA cross-linking		
9. The nontemplate	strand of a portion of	fa gene reads: 5'-T	CACTGGTTCA. V	What is the sequence of the
resulting transcri	pt (RNA) for this por	tion?		
(A) 5'-AAGUGACCAAGU		(B) 5'-UGAACCAGUGAA (C) 5'-UUCACUGGUUCA		-UUCACUGGUUCA
(D) 5'-ACUUGO	GUCACUU (I	E) 5'-TGAACCAG	TGAA	
10. The nucleotide se	equence of a DNA rea	iding frame is 5'GT	A3'. A messenger R	NA molecule with a
	•	• •	_	synthesis, a transfer RNA
	codon. What is the nu			
(A) 5'CAT3'	(B) 5'CAU3'	(C) 5'GUA3'	(D) 5'UAC3'	(E) 5'AUG3'
11 77 1	MT4	. ,	. ,	. ,
11. The polarity of E	•	(6) 51 - 61	<i>(</i> () () ()	
(A) 5'>5'.	(B) 3'>5'.	(C) 5'>2'.	(D) 2'>5'.	(E) 2'>3'
12. Which of the following	lowing phenotypic rat	ios show incomplet	e dominance?	
(A) 2:1	(B) 3:1	(C) 1:2:1	(D) 1:1	(E) 4:1
13 Suppose the Lar	nd M genes are on the	came chromosome	hut congreted by 10	0 map units. What fraction
	n the cross LM/lm x li			o map ums. what naction
(A) 10%	(B) 25%	(C) 50%	(D) 75%	(E) 100%
,	• ,	. ,	• •	,
	molecule is amplified	l by PCR, how man	y DNA molecules w	ill there be after six
cycles?				
(A) 6	(B) 12	(C) 24	(D) 64	(E) 86
15. Robertsonian tr	anslocations result fro	om which of the foll	lowing?	
	r at or near the centron broken parts.	meres of two acroce	entric chromosomes	followed by the reciprocal
_	chromosome become	es attached to a non-	-homologous chrom	osome.
	ssing over occurs duri		_	
(D) There is fusi	on of two small chror	nosomes end-to-end	l such that a double	centromere occurs.
(E) None of abo	ve			
16 II da		1	DATA	. 6
	e regulators such as th	e lac repressor prev	ent KNA polymeras	e from initiating
transcription?				
(A) by blocking	passage of the polyme	erase tnrougn tne or	perator	
			1 1	
(B) by forming a	loop in the operator by blocking the DNA b	that restricts the pas		ase

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(E) by slowing down the RNA polymerase.

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17. Mutation of homeotic cluster genes often results in which of the following developmental defects in *Drosophila*?

- (A) Absence of a group of contiguous segments
- (B) Transformation of one segment into another
- (C) Polarity defects in every segment along the anterior-posterior axis
- (D) Tumor formation in imaginal discs
- (E) Absence of every other segment along the anterior-posterior axial
- 18. When bacteria produce mammalian proteins, cDNA is used rather than genomic DNA. Which of the following is the best explanation?
  - (A) It is easier to clone cDNA than genomic DNA of comparable size.
  - (B) It is easier to clone RNA than DNA.
  - (C) It is not possible to clone the entire coding region of the gene.
  - (D) Most eukaryotic genes have introns that cannot be removed in bacteria.
  - (E) Most eukaryotic gene promoters do not function in bacteria.
- 19. Consider the average in vivo turnover rates for proteins, DNA, and mRNA. Which of the following orders best describes the turnover rate from fastest (shortest average lifetime) to slowest (longest average lifetime)?
  - (A) mRNA > DNA > proteins
  - (B) mRNA > proteins > DNA
  - (C) Proteins > mRNA > DNA
  - (D) Proteins > DNA > mRNA
  - (E) DNA > mRNA > proteins
- 20. Which of the following most accurately describes a retrotransposon?
  - (A) A DNA sequence that can move from one site in the genome to another without replicating
  - (B) A DNA sequence that can be deleted from the genome without consequence
  - (C) A DNA sequence that replicates via an RNA intermediate
  - (D) A DNA sequence that replicates via a protein intermediate
  - (E) A DNA sequence that causes base-substitution mutations at some other site on the chromosome

## 二、解釋名詞 (每題 4 分, 40%) ※ 注意:請於試卷內之「非選擇題作答區」標明題號依序作答。

- 1. Allele
- 2. Homologous recombination
- 3. SNP (single nucleotide polymorphism)
- 4. 3'UTR
- 5. Oncogene
- 6. Syntenic block
- 7. Genome
- 8. Aneuploidy
- 9. Telomerase
- 10. Alternative splicing

試題隨卷繳回