

I. 解釋名詞 (18%) :

- |                         |                     |
|-------------------------|---------------------|
| (1) Epistasis           | (4) Retrotransposon |
| (2) Dosage compensation | (5) Homeobox        |
| (3) RNA interference    | (6) C-value paradox |

II. 單選題 (66%) : ※ 注意：請於試卷上「選擇題作答區」依序作答。

- A new phenotype not seen in the parents can arise by:
  - gene interaction
  - co-dominance
  - complementation
  - none of the above
  - all of the above
- What aspect of chromosome behavior most clearly accounts for Mendel's law of segregation?
  - Movement of sister chromatids to opposite poles at anaphase II of meiosis.
  - Movement of homologous chromosomes to opposite poles at anaphase I of meiosis.
  - Crossing over between homologous chromosomes during prophase I of meiosis.
  - Replication of chromosomes prior to meiosis.
  - Independent alignment of different homologous pairs on the metaphase I spindle.
- In chickens, it is the females that have two different sex chromosomes (Z and W) while the males have two Z chromosomes. A Z-linked gene controls the pattern of the feathers with the dominant B allele causing the barred pattern and the b allele causing non-barred feathers. You cross a barred female with a non-barred male. What do you expect for the phenotype of the progeny?
  - daughters all one type, sons all the other type
  - daughters and sons of both types
  - sons of one type, daughters of both types
  - daughters of one type, sons of both types
  - none of the above
- In tetrad analysis, second-division segregations result from:
  - single crossovers between linked genes
  - double crossovers between linked genes
  - single crossovers between a gene and a centromere
  - independent assortment of unlinked genes
  - nondisjunction of homologs
- A suppressor tRNA can suppress a:
  - nonsense mutation
  - missense mutation
  - deletion mutation
  - frameshift mutation
  - silent mutation

6. The hydrolysis of an -NH<sub>2</sub> group from a base is called \_\_\_\_\_, while intercalating agents such as proflavin function as mutagens by causing \_\_\_\_\_:

- A) deamination; transversions
- B) deamination; deletions or insertions
- C) excision repair; deletions or insertions
- D) excision repair; transversions
- E) deletion; transitions

7. The fact that the human proteome is larger than predicted by the genome is due to:

- A) alternative splicing
- B) chemical modifications of proteins
- C) large number of paralogs
- D) variations in domain architecture
- E) all of the above

8. For linkage analysis, what variable most determines a marker's ability to predict genotype?

- A) presence on the X chromosome
- B) map distance from allele
- C) size of restriction fragment
- D) size of disease gene
- E) recognition sequence

9. A normal female who is homozygous for a mutation in Xist which renders it inactive would have which of the following characteristics?

- A) no telomerase activity
- B) An inability to form Barr bodies
- C) Inability to form constitutive heterochromatin
- D) Reduced levels of DNA replication
- E) None of the above

10. Crossover suppression is a characteristic of \_\_\_\_\_:

- A) translocations
- B) transposition
- C) deletions
- D) aneuploidy
- E) inversions

11. Which of the following can contribute to evolution of a species?

- A) amphidiploidy
- B) transposable elements
- C) translocations
- D) all of the above
- E) none of the above

12. A transposon inserts into an operon. What are the likely consequences?

- A) all genes are knocked out
- B) downstream genes are knocked out
- C) flanking genes cotransduce less
- D) a & c
- E) b & c

13. You perform a generalized transduction mapping experiment by growing phage on a wild-type host and transducing a recipient that is mutant for three genes. The results are shown in the table below:

phenotype	number of transductants
F <sup>+</sup> M <sup>-</sup> R <sup>-</sup>	116
F <sup>+</sup> M <sup>+</sup> R <sup>-</sup>	447
F <sup>+</sup> M <sup>-</sup> R <sup>+</sup>	7
F <sup>+</sup> M <sup>+</sup> R <sup>+</sup>	490

What is the order of the genes?

- A) RFM
- B) MRF
- C) FMR
- D) a or b
- E) b or c



14. A cross is conducted between isogamous algae that show uniparental inheritance where mtDNA is inherited from one parent and cpDNA is inherited from the other. A mutation in cpDNA causes a yellow color, and a mutation in mtDNA confers chloramphenicol resistance. From the following mating:

yellow/chloramphenicol resistant × green/chloramphenicol sensitive

Which of the following ratios might be observed in the resulting haploid progeny?

- A) 2 yellow/resistant, 2 yellow/sensitive
- B) 2 green/resistant, 2 green/sensitive
- C) 4 green/sensitive
- D) 4 green/resistant
- E) 1 yellow/resistant, 1 yellow/sensitive, 1 green/resistant, 1 green/sensitive

15. Which of the following is a common consequence of mutations that eliminate cell-cycle checkpoints?

- A) increased DNA repair
- B) decreased frequency of cell division
- C) arrest
- D) aneuploidy
- E) decreased mutation

16. Eukaryotic cells are able to carefully regulate precise levels of transcription in specific genes encoding structural proteins through:

- A) complex enhancer elements that can associate with multiple activator and repressor proteins

- B) production of different types of sigma factors
- C) attenuation
- D) all of the above
- E) none of the above

17. If a yeast *cdc28* mutant arrests as unbudded cells and a *cdc7* mutant arrests as budded cells after a shift from permissive to restrictive temperature, what will the phenotype of the double mutant be at restrictive temperature if the *cdc28* gene product acts before the *cdc7* gene product in the cell cycle?

- A) arrests as budded cells
- B) arrests as unbudded cells
- C) arrests as a mixture of budded and unbudded cells
- D) lethal
- E) wild-type (cells in various stages of the cell cycle)

18. Which of the following statements is false?

- A) Haploinsufficiency describes a situation where one wild-type copy of a gene is not enough for normal development to occur.
- B) RNAi can be used to create a phenocopy that mimics a loss-of-function mutation
- C) Gain-of-function mutations produce either excess protein or a new form of a protein
- D) The ectopic phenotype is the one that is expressed in the double mutant
- E) *In situ* RNA hybridization can be used to determine where in a developing organism a gene is expressed

19. Which of the following is not a form of a loss-of-function mutation?

- A) ectopic gene expression
- B) hypomorphic mutation
- C) conditional mutation
- D) knockout mutation
- E) all of the above are loss-of-function mutations

20. In humans, *brachydactyly* is a dominant condition. 173 people in a population of 372 show the disease [50 are *BB*, 123 are *Bb*] and 199 are normal phenotypes (*bb*). The frequency of the *b* allele in this generation is:

- A) 0.58
- B) 0.30
- C) 0.70
- D) 0.53
- E) 0.13

21. The enzyme primase is required for DNA replication because:

- A) primase breaks the hydrogen bonds holding the two strands together
- B) DNA polymerase can only add bases to an existing nucleic acid strand
- C) the enzyme binds small pieces of DNA together
- D) the primase corrects any errors made by the DNA polymerase
- E) None of the above

22. A feature of chromatin remodeling is that:

- A) it can turn on transcription but cannot silence genes
- B) the modified state of chromatin can be passed on when DNA replicates
- C) it involves only the addition of methyl groups to DNA
- D) it requires RNAi
- E) None of the above

III. 問答題 (16%):

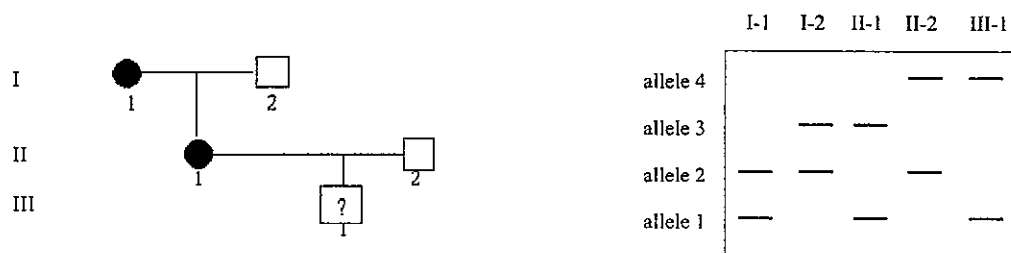
1. A trihybrid (3 genes heterozygous) produces the following gametes:

ABC	46
Abc	63
ABc	4
aBc	381
abc	38
aBC	71
abC	2
<u>AbC</u>	<u>395</u>
Total	1000

What is the order and map distance between the three genes? (8%)

2. In a certain breed of plants, thorns are determined by the dominant allele  $T$  and thornless is determined by the recessive allele  $t$ .  $T$  is 80% penetrant in the heterozygote; 20% of heterozygotes will appear thornless. If you make the parental cross  $TT \times tt$ ; what would be the expected number of phenotypes (thorny: thornless) observed in a population of 500  $F_2$  plants? (4%)

3. Microsatellites are very useful as DNA markers. The multiple alleles can be detected as PCR products of different sizes. One particular microsatellite has 4 alleles in the population and is closely linked to an autosomal dominant form of early Alzheimer's disease. A pedigree and microsatellite analysis of a family with a history of early Alzheimer's disease is indicated below:



- (1) Which allele is associated with early Alzheimer's disease in this family? (2%)
- (2) Will the male III-1 develop early Alzheimer's disease? (2%)