

Protein Synthesis and Cellular Control

- 1) Describe the process of transcription? (4)
- 2) Describe translation? (4)
- 3) Using a labelled diagram, describe the structure of tRNA? (3)
- 4) Describe the role of the following in protein synthesis (8);
 - a. RNA polymerase
 - b. Anticodon-codon complex
 - c. Stop codon
 - d. tRNA
 - e. mRNA
 - f. Start codon
 - g. Ribosome
 - h. Nucleus
- 5) Which type of bond joins the amino acids together and what is this reaction called? (2)
- 6) What happens to the polypeptide chain after it has been formed? (2)
- 7) A toxin works by inhibiting RNA polymerase; what effects is it likely to have and why? (2)
- 8) There are 64 codons and only 20 amino acids. Explain why? (2)
- 9) A transcription error results in premature insertion of a stop codon. What effect is this likely to have and why is it significant? (3)
- 10) Draw a simple labelled diagram to illustrate translation? (4)
- 11) How is protein synthesis controlled? (3)
- 12) How is protein activation controlled? (3)
- 13) What are (3);
 - a. Promoters

- b. Operators
- c. Transcription factors

- 14) Describe the 3 components of operons and give the functions of each? (6)
- 15) What are the two different parts of the control elements and why are they each important? (4)
- 16) How does the lac operon in E.coli work? (6)
- 17) What is apoptosis and how does it take place? (3)
- 18) Why is apoptosis important? (2)
- 19) What are homeotic genes, homeobox sequences and homeodomains and what are their significance? (3)
- 20) What is a frameshift mutation? (2)
- 21) Why is it that a frameshift mutation which appears earlier in the base sequence has a greater effect on the protein than one which appears later? (2)
- 22) Give 3 reasons why a mutation may have neutral effects? (3)
- 23) What are the different types of mutations? Give a brief description of each. (5)
- 24) Bacterial enzymes break down particular antibiotics and a mutation has resulted in the genes coding for these enzymes giving them the ability to break down a larger range of antibiotics. How has this helped them survive? (2)
- 25) Give one example of a harmful mutation and the results from that? (2)
- 26) If protein synthesis and/or activation are not controlled, cancer can result. Explain why? (3)

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