## COMPUTER SCIENCE TRIPOS Part II – 2024 – Paper 8

## 2 Bioinformatics (pl219)

(a) Discuss how to use bioinformatics algorithms to detect the specific pathogenic sequences in the genome of a pathogenic species, by comparing its genome with the genome of an evolutionarily-close non-pathogenic species.

*Hint.* Most pathogenic bacteria have long DNA sequences containing diseasecausing genes that are not present in the genome of similar non-pathogenic species. Consider how to detect extra material, and perhaps inverted repeats (which are usually formed during the insertion of the disease-causing genes.)

[5 marks]

- (b) Compute the global alignment and the best score of the sequences {CGTGT, TGGCGCC} with the following parameters: match score = +2, mismatch score = -1, gap penalty = -2. Report the final score and alignment(s). [4 marks]
- (c) Dimerisation occurs when two similar proteins (P) join together to form a dimer
  (D), and dissociation reverses this process. The Gillespie algorithm may be applied to model dimerisation and dissociation of proteins, with species P and D, and the following rate constants:
  - Dimensition:  $2P \rightarrow D$  with rate  $c_1$
  - Dissociation:  $D \rightarrow 2P$  with rate  $c_2$

Dimerisation is rare and dimers are unstable, therefore  $c_2 \gg c_1$ . Explain how to use the Gillespie algorithm to model dimerisation and dissociation reactions.

[7 marks]

(d) Discuss advantages and disadvantages of using DNA to store information.

[4 marks]