UNIVERSITY OF ESWATINI

FACULTY OF SCIENCE AND ENGINEERING

DEPARTMENT OF BIOLOGICAL SCIENCES

MAIN EXAMINATION PAPER 2020/2021

COURSE CODE:

BIO451

TITLE OF PAPER:

BIOINFORMATICS

TIME ALLOWED:

THREE (3) HOURS

INSTRUCTIONS:

NUMBERS IN BRACKETS DENOTE THE NUMBER OF MARKS

THIS PAPER COMPRISES OF TWO SECTIONS and ONE ANNEX.

SECTION A COMPRISES OF EIGHT (8) QUESTIONS.

SECTION B COMPRISES OF SEVEN (7) QUESTIONS.

ANSWER ALL QUESTIONS IN SECTIONS A AND B.

TOTAL MARKS AVAILABLE: 70

NO ADDITIONAL MATERIAL (E.G. NOTES, BOOKS, PHONES, SMART WATCHES ETC) MAY BE TAKEN INTO THE EXAMINATION.

THE USE OF A SCIENTIFIC CALCULATOR IS PERMITTED.

DO NOT OPEN THIS PAPER UNTIL PERMISSION HAS BEEN GRANTED BY THE CHIEF INVIGILATOR

Section A

Total marks available: 35

A poacher has been arrested by the police in possession of several processed animal parts. Unfortunately, it is not possible to identify the animal(s) just by looking at them. The police have sent samples of the processed animal parts to you for forensic analysis. You have extracted and sequenced the samples using your next-generation sequencer.

AQ1: State what **type of molecule** and **file format** of the data you expect to receive from the sequencer. [2]

AQ2 Describe what you would do with the sequence data to help identify the organism(s). In your answer, include what databases you would use. [4]

AQ3: The results have come back for one of the samples. Explain what the four columns inside the black oval (Query cover, E value, Per. Ident and Accession) tells you about your search.

| Descriptions Graphic Summary Alignments Taxonomy | | _ | | | | | on the second | · · · · · · · · · · · · · · · · · · · | portunine de l'acceptant de l'accept | DW Specific (MACCONTRACTOR) |
|--|--------------------|--------------------|---------------|--------------|-------|----------------|---|---------------------------------------|--|-----------------------------|
| Sequences producing significant alignments | Dow | nload Y | | Mana | ge Co | olumn | s Y (| Show [| 100 | V . |
| select all 0 sequences selected | | | | | igla. | | | D _E C | | er i i i i i i i |
| Description | Scientific Name | Common | Taxid | Max Score | Score | Query Cover | E value | Per. Ident | Acc. Len | Accessio |
| Tragelaphus strepsiceros isolate PHC11 mitochondrion, complete genome | Tragelag | greater | 9946 | 3195 | 3195 | 100% | 0.0 | 99.77% | 16382 | JN 1632708. |
| Tragelachus spakii mitochondrion complete genome | Tragelao | . <u>Sitatunga</u> | 69298 | 2809 | 2809 | 100% | 0.0 | 95.75% | 16408 | EF536357 |
| Tragelachus oryx isolate PHC13 mitochondrion_conrolete genome | Tragelao | . <u>eland</u> | 9945 | 2776 | 2776 | 100% | 0.0 | 95.42% | 16397 | JN632704 |
| Taurotragus derbianus mitochondrion, complete genome | <u>Taurotra</u> | giant el | <u>303930</u> | 2776 | 2776 | 100% | 0.0 | 95,46% | 16399 | EF536354 |
| Tragelachus eurycerus isotate SU/18 mitochandrion, complete genome | Tragelap., | bongo | 69297 | 2771 | 2771 | 100% | 0.0 | 95,36% | | JM632703 |
| Tragelachus scriptus isolate MBP11 mitochondrion, comclete ganoma | Tragelan | . <u>bushbuck</u> | 66440 | 2748 | 2748 | 100% | 0.0 | 95.18% | | JN632706 |
| Tragelanhus scrintus isolate PhC17 mitochondrion, complete genome | Tragelap | <u>bushbuck</u> | 66440 | 2732 | 2732 | 100% | 0.0 | | | <u>JN632707</u> |
| Tragelaghus buxtoni mitochondrion, complete genome | Tragelao. | <u>mountai</u> | <u> 69296</u> | 2723 | 2723 | 100% | 0.0 | 94.80% | | HC 0380 |
| Tragelachus scriptus isolate CAR31 mitochondrion, comclete genome | Tranelao | <u>bushbuck</u> | 66440 | 2553 | 2553 | 99% | 0.0 | 93.26% | | |
| Tragelachus angasii isulate MBP15 mitochondrion. complete genome | Tragelap. | <u>nyala</u> | <u>66437</u> | 2514 | 2514 | 100% | 0.0 | 92.77% | | |
| Tragelaphus imberbis mitochondrial complete genome | Tragelao. | <u>lesser k</u> | 9947 | 2507 | 2507 | 100% | 0.0 | 92.62% | | |
| Tragelaphus imbarbis mitochondrion, complete genome | <u>Tragelan</u> . | <u>. lesser k</u> | 9947 | 2507 | 2507 | 100% | 0.0 | 92.62% | | EF53635 |
| Tragelactius imberbis mitochondrial 12S and 16S ribosomal RHAs and Phe- Val-, and Leu-IRHA cenes | Tragelap. | . lesser k | 9917 | 2507 | 2507 | 100% | 0.0 | 92.62% | | |
| Boselaphus tragocamelus mitochondrion, controlete genome | <u>Boselap</u> . | . <u>nilgai</u> | <u>9917</u> | 2318 | 2318 | 100% | 0.0 | 90.73% | 16322 | EF63635 |
| | | | | | | | | | [2] | |

AQ4: Based on the results above, what accession number would you pursue? [1]

AQ5: After further investigation, using GenBank, you come across this result (see Annex 1) for a second sample. Identify the following:

| ۵1 | The size and type of molecule | [2] |
|----|--|-----|
| , | • • | [2] |
| b) | The full name of the GenBank Division | [1] |
| c) | The protein name | |
| d) | Organism (common and scientific) | [3] |
| e) | Name of the gene encoding for this protein | [2] |

f) The accession number of the gene encoding for this protein

[2]

AQ6: You have sent your results to the legal teams handling the case. Unfortunately, there is some dispute over the results from the second sample (AQ5). Both the prosecution and defense teams have requested some further analysis. Specifically, they request you to analyse the sample that they gave you against the very same organism identified in from Annex 1. Explain how you could go about getting a tissue sample from the organism (in Annex 1) to run your investigation.

[5]

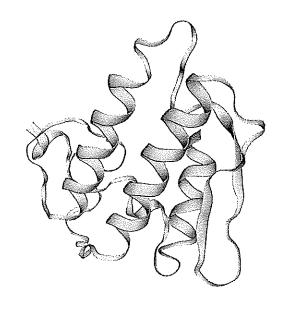
AQ6: Describe **how** you would use **bioinformatics** to determine the structure and function of the protein (from AQ5), including the database(s) you would use. [5]

AQ7: List two reasons for determining the structure of proteins.

[2]

AQ8: Describe the key feature(s) of the protein below.

[2]



Section B

Total marks available: 35

BQ1: Which BLAST application would you use to identify the following sequences:

- a) AGAAGTTCCAGATCATTCCAAGA [1] b) AUCCCAGGGUUCACCAGAGCCAGGAGA [1]
- c) KAGTAKCATVADDCHUABCCAGAAGNMWSS [1]
- d) VLPWGQMSFWGATVITNLLSAIPYIGNTLV [1]

BQ2: Identify which of the following sequence(s) would you enter into the tBLASTn database: [1]

- a) MGGNTAKCRRADDCHUABCCAGAAGNMWSS
- b) CGAGAUUAUCCCAGGGUUCACCAGAGA
- c) TAGGTTCCCAGATCATTCCAAGA
- d) ATPYISSAFVGYVLPWGPTVQMSFWGATVITNLLS

BQ3: What is the default scoring matrix for BLASTp?

[1]

BQ4: Assuming a scoring method: match = +1, mismatch = -2, and gap = -1, using the Needleman-Wunsch algorithm:

a) Initialize the scoring matrix below.

| | Т | С | Т | G | Α | Т | Т | С | Т | С |
|---|-------|---|---|---|---|---|---|---|---|---|
| | | | | | | | | | | |
| T | | | | | | | | | | |
| С | | | | | | | | | | |
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| С | | | | | | | | | | |
| С | | | | | | İ | | | | |

- b) Calculate the score of the matrix and fill the traceback matrix. Use arrows to indicate how you determined the path forward. Use circles and arrows to determine your trace back. [5]
- c) Deduce the best alignment based on the traceback matrix. [3]

BQ5: Assuming a scoring method: match = +1, mismatch = -2, and gap = -2, using the Smith and Waterman algorithm:

a) Initialize the scoring matrix below.

| Initialize | e the scor | ing ma | trix bei | ow. | | | | | | | |
|------------|------------|--------|----------|-----|---|----------|---|---|---|---|---|
| | Т | С | Т | G | Α | T | T | С | T | C | С |
| | | | _ | | | | | | | | |
| Т | | | | | | | | | | | |
| Α | | | · | | | | | | | | |
| G | | | 1 | | | | | | | | |
| С | | | | | | | | | | | |
| Α | | | | | | <u> </u> | | | | | |
| Т | | | | | | | | | | | |
| Т | | | | | | | | | | | |
| Α | | | | | | | | | | | |
| G | | | | | | | | | | | |
| С | | | | | | | | | | | |

- b) Calculate the score of the matrix and fill the traceback matrix. Use arrows to indicate how you determined the path forward. Use circles and arrows to determine your trace back. [5]
- c) Deduce the best alignment based on the traceback matrix.

[3]

[2]

[2]

d) Were there any sub-optimal alignments? Explain your answer.

[2]

BQ6: Align the following two sequences using a dot plot

[3]

- TCTGATTCTC
- TCGCATCAC

BQ7: You have four sets of nucleic acid sequences of very different lengths to align. Which type of algorithm(s) would be most appropriate? Explain your answer and give ONE advantage for your algorithm(s). [4]