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**TECHNICAL UNIVERSITY OF MOMBASA**

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FACULTY OF APPLIED AND HEALTH SCIENCES

DEPARTMENT OF PURE & APPLIED SCIENCES

**UNIVERSITY EXAMINATION FOR:**

MSC IN BIOTECHNOLOGY

ABT 5106: ADVANCED BIOINFORMATICS

SPECIAL/ SUPPLEMENTARY EXAMINATIONS

**SERIES: SEPTEMBER 2018**

**TIME: 3 HOURS**

**DATE: SEP 2018**

**Instructions to Candidates**

You should have the following for this examination

*-Answer Booklet, examination pass and student ID*

This paper consists of **SIX** questions, each **25 marks**. Attempt any **FOUR** questions Choose instruction.

**Do not write on the question paper.**

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**Question ONE (25 marks)**

- a. Outline the limitations of bioinformatics. 8 marks
  
- b. Explain the application of bioinformatics in the following fields;
  - i. Microbial genome applications 5 marks
  
  - ii. Comparative Studies 5 marks
  
  - iii. Health and Drug discovery 7 marks

**Question TWO (25 marks)**

- a. Outline the factor to consider when designing primers. 8 marks
  
- b. Describe any **FIVE** bioinformatic tools you would use to design primers. 10 marks

c. Distinguish between unigene and locus link.

7 marks

**Question THREE (25 marks)**

Explain the three primary nucleotide sequence databases.

25 marks

**Question FOUR (25 marks)**

a. Use the following substitution matrices to solve the following questions

**BLOSUM 62**

	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
A	4	-1	-2	-2	0	-1	-1	0	-2	-1	-1	-1	-2	-1	1	0	-3	-2	0	
R	-1	5	0	-2	-3	1	0	-2	0	-3	-2	2	-1	-3	-2	-1	-1	-3	-2	-3
N	-2	0	6	1	-3	0	0	0	1	-3	-3	0	-2	-3	-2	1	0	-4	-2	-3
D	-2	-2	1	6	-3	0	2	-1	-1	-3	-4	-1	-3	-3	-1	0	-1	-4	-3	-3
C	0	-3	-3	-3	9	-3	-4	-3	-3	-1	-1	-3	-1	-2	-3	-1	-1	-2	-2	-1
Q	-1	1	0	0	-3	5	2	-2	0	-3	-2	1	0	-3	-1	0	-1	-2	-1	-2
E	-1	0	0	2	-4	2	5	-2	0	-3	-3	1	-2	-3	-1	0	-1	-3	-2	-2
G	0	-2	0	-1	-3	-2	-2	6	-2	-4	-4	-2	-3	-3	-2	0	-2	-2	-3	-3
H	-2	0	1	-1	-3	0	0	-2	8	-3	-3	-1	-2	-1	-2	-1	-2	-2	2	-3
I	-1	-3	-3	-3	-1	-3	-3	-4	-3	4	2	-3	1	0	-3	-2	-1	-3	-1	3
L	-1	-2	-3	-4	-1	-2	-3	-4	-3	2	4	-2	2	0	-3	-2	-1	-2	-1	1
K	-1	2	0	-1	-3	1	1	-2	-1	-3	-2	5	-1	-3	-1	0	-1	-3	-2	-2
M	-1	-1	-2	-3	-1	0	-2	-3	-2	1	2	-1	5	0	-2	-1	-1	-1	-1	1
F	-2	-3	-3	-3	-2	-3	-3	-3	-1	0	0	3	0	6	-4	-2	-2	1	3	-1
P	-1	-2	-2	-1	-3	-1	-1	-2	-2	-3	-3	-1	-2	-4	7	-1	-1	-4	-3	-2
S	1	-1	1	0	-1	0	0	0	-1	-2	-2	0	-1	-2	-1	4	1	-3	-2	-2
T	0	-1	0	-1	-1	-1	-1	-2	-2	-1	-1	-1	-1	-2	-1	1	5	-2	-2	0
W	-3	-3	-4	-4	-2	-2	-3	-2	-2	-3	-2	-3	-1	1	-4	-3	-2	11	2	-3
Y	-2	-2	-2	-3	-2	-1	-2	-3	2	-1	-1	-2	-1	3	-3	-2	-2	2	7	-1
V	0	-3	-3	-3	-1	-2	-2	-3	-3	3	1	-2	1	-1	-2	-2	0	-3	-1	4

**PAM250**

	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V	
A	2	-2	0	0	-2	0	0	1	-1	-1	-2	-1	-1	-3	1	1	1	-6	-3	0	
R	-2	6	0	-1	-4	1	-1	-3	2	-2	-3	3	0	-4	0	0	-1	2	-4	-2	
N	0	0	2	2	-4	1	1	0	2	-2	-3	1	-2	-3	0	1	0	-4	-2	-2	
D	0	-1	2	4	-5	2	3	1	-2	-4	0	-3	-6	-1	0	0	-7	-4	-2		
C	-2	-4	-4	-5	12	-5	-5	-3	-3	-2	-6	-5	-5	-4	-3	0	-2	-8	0	-2	
Q	0	1	1	2	-5	4	2	-1	3	-2	-2	1	-1	-5	0	-1	-1	-5	-4	-2	
E	0	-1	1	3	-5	2	4	0	1	-2	-3	0	-2	-5	-1	0	0	-7	-4	-2	
G	1	-3	0	1	-3	-1	0	5	-2	-3	-4	-2	-3	-5	0	1	0	-7	-5	-1	
H	-1	2	2	1	-3	3	1	-2	6	-2	-2	0	-2	-2	0	-1	-1	-3	0	-2	
I	-1	-2	-2	-2	-2	-2	-2	-2	-3	-2	5	2	-2	2	1	-2	-1	0	-5	-1	4
L	-2	-3	-3	-4	-6	-2	-3	-4	-2	2	6	-3	4	2	-3	-3	-2	-2	-1	2	
K	-1	3	1	0	-5	1	0	-2	0	-2	-3	5	0	-5	-1	0	0	-3	-4	-2	
M	-1	0	-2	-3	-5	-1	-2	-3	-2	2	4	0	6	0	-2	-2	-1	-4	-2	2	
F	-3	-4	-3	-6	-4	-5	-5	-5	-2	1	2	-5	0	9	-5	-3	-3	0	7	-1	
P	1	0	0	-1	-3	0	-1	0	0	-2	-3	-1	-2	-5	6	1	0	-6	-5	-1	
S	1	0	1	0	0	-1	0	1	-1	-1	-3	0	-2	-3	1	2	1	-2	-3	-1	
T	-1	-1	0	0	-2	-1	0	0	-1	0	-2	0	-1	-3	0	1	3	-5	-3	0	
W	-6	2	-4	-7	-8	-5	-7	-7	-3	-5	-2	-3	-4	0	-6	-2	-5	17	0	-6	
Y	-3	-4	-2	-4	0	-4	-4	-5	0	-1	-1	-4	-2	7	-5	-3	-3	0	10	-2	
V	0	-2	-2	-2	-2	-2	-2	-1	-2	4	2	-2	2	-1	-1	-1	0	-6	-2	4	

Given the following pairs of sequences below;

- i. KYWSAVYA
- ii. KKAWVDVA

- i. DWSAEVKK
- ii. AWSADVKY

Calculate similarity scores between these sequences using:

- i. Identity matrix (assume match = 2, and mismatch = 1) 4 marks
- ii. PAM250 matrix 4 marks

- iii. BLOSUM 62 matrix 4 marks

b. Describe the needleman-Wunsch algorithm 10 marks

c. Distinguish between an identifier and accession code 3 marks

**Question FIVE (25 marks)**

a. Use the Standard Code for Information Interchange table below to solve the following.

	30	40	50	60	70	80	90	100	110	120
0		(	2	<	F	P	Z	d	n	x
1		)	3	=	G	Q	[	e	o	y
2		*	4	>	H	R	\	f	p	z
3	!	+	5	?	I	S	]	g	q	{
4	"	,	6	@	J	T	^	h	r	
5	#	-	7	A	K	U	_	i	s	}
6	\$	.	8	B	L	V		j	t	~
7	%	/	9	C	M	W	a	k	u	DEL
8	&	0	:	D	N	X	b	l	v	
9	'	1	;	E	O	Y	c	m	w	

Given characters 'k' and 'Y'

Calculate in each case the corresponding Phred quality score based on

- i. The fastq-sanger format. 6 marks
  - ii. The Solexa/Illumina read format. 6 marks
- b. Describe the tBLASTn bioinformatic tool 7 marks
- c. Using an illustration, describe the position specific iterative-BLAST (PSI-BLAST) approach. 6 marks

**Question SIX (25 marks)**

- a. Distinguish between MEGABLAST and BLASTn bioinformatic tools. 6 marks
- b. Contrast Parsimony and Maximum likelihood tree building algorithms. 5 marks
- c. Distinguish between ENZYME and REBASEA databases. 14 marks