

JAMES COOK UNIVERSITY

P O Box 6811 CAIRNS Qld 4870 Australia Tel: (07) 4042.1111 Fax: (07) 4042 1300

SCHOOL OF PHARMACY AND MOLECULAR SCIENCES Chemistry Department

FIRST SEMESTER EXAMINATIONS 2005

Cairns Campus

This paper must be handed in at the end of the Examination:	Yes
Release to Library:	No

PHONE NO:

TWO (2) HOURS

FIFTEEN (15) MINUTES

(07) 4042 1275

STUDENT NAME: (block letters)

STUDENT NUMBER:

SUBJECT CODE: CH1010:03

SUBJECT NAME: BIOLOGICAL CHEMISTRY

EXAMINER: Dr M. Liddell

DURATION OF EXAMINATION (hours):

PERUSAL TIME (minutes):

TOTAL NUMBER OF QUESTIONS: 27

INSTRUCTIONS TO STUDENTS:

 The exam is composed of two sections:
 Section A - Multiple choice 22 questions - 33%

 Section B - Short answer
 5 questions - 67%

 Total marks for paper = 100
 Answer ALL questions.
 All questions are not of equal value.

 Timings are indicated to allow 15 minutes of check-over time.

MATERIALS TO BE SUPPLIED BY EXAMINATION SECTION:

Examination Booklets required:	Yes
Multiple choice scanner sheets Scanner A- E:	Yes

MATERIALS STUDENTS MAY USE:	
Scientific calculator with no text storage facilities.	
Access to an English Dictionary:	Yes

Copyright Reserved

SECTION A

MULTIPLE CHOICE QUESTIONS (EACH QUESTION IS WORTH 1.5 MARKS). ANSWER ALL QUESTIONS – SHADE WITH A PENCIL THE MOST CORRECT ANSWER ON THE MULTICHOICE SCANNER SHEET.

Timing: you should complete the multi-choice section in 30 minutes (≈ 1.5 minutes per question).

This section has been deleted it is just multi-choice of the same calibre as the modules.

SECTION B

SHORT ANSWER QUESTIONS. (MARKS FOR EACH QUESTION ARE AS INDICATED) ANSWER EACH OF THE FIVE (5) QUESTIONS.

Question 1

Timing: you should complete this question in 15 minutes.

- (a) Hydrogen bonding, although weak, is one of the most important types of bonding in biology.
 - Define **hydrogen bonding**.
 - Provide an example of **intermolecular** hydrogen bonding.
 - Are **hydrophilic** or **hydrophobic** regions of a protein more likely to participate in hydrogen bonding with the surrounding solvent explain your answer.

(4 marks)

- (b) There are a variety of acids and bases found in cellular systems.
 - Provide an example of a common **weak acid** and its conjugate base.
 - Provide an example of a common **weak base** and its conjugate acid.
 - Explain how you could prepare a **buffer** from the acid and base components you have just written down.

(4 marks)

(6 marks)

(c) A colourless organic oil gave the following microanalytical results: C: 54.55 % H: 9.09 %

Infrared spectroscopy of a sample of the oil gave several strong bands at 3200 and 1721 cm⁻¹. In a mass spectrum of the compound a molecular ion was found at m/z 88.

- Provide a **molecular formula** and a **line-angle structure** for this compound. Explain clearly how you arrived at the formulae.
- Would the molecule you have drawn be likely show a UV absorption explain why or why not?
- How could you **verify the purity** of the sample?

Question 2

Timing: you should complete this question in 15 minutes.

- (a) Proteins have complex structures that relate to their function
 - What are the two major **structural types** of protein. Provide an example of a protein in each structural type and give the function each protein.
 - The primary structure of a protein is at the outset responsible for dictating which structural type a protein will be in. Draw a short stretch of a peptide chain (3 different amino acids) indicating what the **primary structure** is and how the amino acids are joined together.
 - Why is it important to know the **isolelectric point** important when purifying proteins?

(5 marks)

CH1010:03

- (b) Enzymes are cellular catalysts which control almost all reactions in a cell.
 - What is the definition of a catalyst.
 - Describe the following : **coenzyme**, **proenzyme**, **isoenzyme**, **active site**. Provide examples to illustrate your answer.

(4 marks)

- (c) There are two major classes of **vitamins: lipid soluble** and **water soluble**, give one example of each type of vitamin and explain the function of each.
 - What role does the protein component of the diet have?

(4 marks)

Question 3

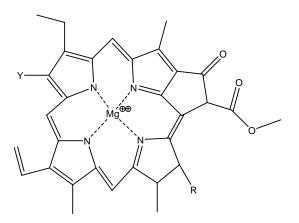
Timing: you should complete this question in 15 minutes.

- (a) Lipids play a variety of roles in cells and as such come in many different types.
 Provide an example of each of the following types of lipid (draw a structure) and indicate the function the lipid has.
 - Steroid.
 - Glycerophospholipid.

Describe the structure of a **cellular membrane**.

(4 marks)

(b) The **chlorophyll** molecule has the following coordinated metallic centre.



- What is the **coordination geometry** at the metal?
- What is the name of this type of ligand?
- Is **chelation** present in this molecule? Explain why or why not.
- What is the function of the R group which is referred to as a **phytol side chain**?.
- There are two different chlorophylls A and B which differ only in the group Y. What is the Y group in ChlA and the Y group in ChlB?

(4 marks)

- (c) Photosynthesis is divided into two major parts the light reactions and the dark reactions.
 - What are the **light reactions of photosynthesis** and where do they occur?
 - What are the **dark reactions of photosynthesis** and where do they occur?

(5 marks)

Question 4

Timing: you should complete this question in 15 minutes.

- (a) **Pyruvate** plays a major role in both aerobic and anaerobic respiration.
 - Define aerobic respiration.
 - Draw the structure of pyruvate.
 - Why is pyruvate important in both aerobic and anaerobic respiration?

- (b) **Fatty acid synthesis** involves the so called 'merry-go-round' synthesis.
 - Why is this called a 'merry-go-round' synthesis, indicate the starting substrate and the normal product from mammalian fatty acid synthesis and the enzyme system that is involved. (4 marks)
- (c) **Oxidative phosphorylation** when coupled to the citric acid cycle produces the bulk of the energy in oxidative respiration.
 - What is **oxidative phosphorylation**?
 - How is the citric acid connected to oxidative phosphorylation?

(4 marks)

Question 5

(a)

Timing: you should complete this question in 15 minutes.

- Chromosomes are the physical structures that DNA is found in inside the nucleus of the cell.
 - What are the different levels of structural organization inside a chromosome.
 - Why is this complex structural organisation necessary?
 - Define a gene.

(b) The copying of the genome of an organism requires great accuracy and reasonable speed.

- Describe briefly the 6 major steps in DNA replication
- How is the **error count** kept very low in this process?
- How is a **reasonable speed** maintained given that in many organisms there are millions of kb pairs to process in the genome?

(4 marks)

(4 marks)

- (b) The ribosomes are the site of **protein synthesis** in the cell.
 - What is the structure of the **ribosome**?
 - Describe the various steps in **DNA transcription**
 - What are **transcription factors** and why are they required in transcription.
 - List the various steps in **DNA translation**.

(6 marks)

(5 marks)

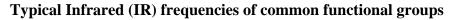
Equation List

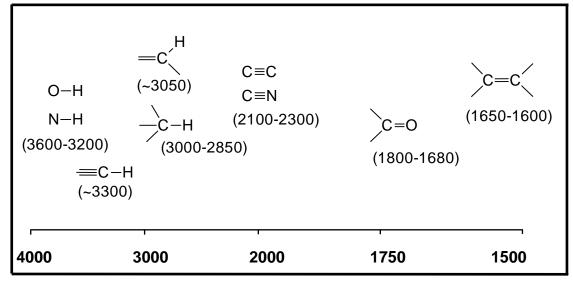
$$E = \frac{h c}{\lambda} = h \nu \qquad h = 6.63 \text{ x } 10^{-34} \text{ J s}^{-1}$$

$$A = \varepsilon 1 [i] \qquad c = 3.00 \text{ x } 10^8 \text{ m s}^{-1}$$

$$T(K) = T(^{\circ}C) + 273.15 \qquad R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$$

Spectroscopy Table





Wavenumber (cm⁻¹)

TABLE 1

Physical Quantity	Name of Unit	Symbol for Unit			
Length	metre	m			
Mass	kilogramme	kg			
Time	second	S			
Electric Current	ampere	а			
Thermodynamic Temperature	kelvin	K			
Amount of Substance	mole	mol			

TABLE 2

Physical Quantity	Name of S.I. Unit	Symbol for S.I. Unit
Volume	cubic metre	m ³
Frequency	hertz	Hz
Velocity	metre per second	ms ⁻¹
Acceleration	metre per second squared	ms ⁻²
Density	kilogramme per cubic metre	kg m ⁻³
Molar Mass	kilogramme per mole	kg mol ⁻¹
Concentration	mole per cubic metre	mol m ⁻³
Molality	mole per kilogramme	mol kg ⁻¹
Force	newton	Ν
Pressure	pascal	Pa
Energy	joule	J
Electric Charge	coulomb	С
Electron Potential Difference	volt	V

Page 10 of 10

PERIODIC TABLE CH1010:03

								-	1									18/VIII	l
								1										2	
	1	2						H 1.008					13/III	14/IV	15/V	16/VI	17/VII	He 4.003	
	3	4											5	6	7	8	9	10	
2	Li	Be											В	С	Ν	0	F	Ne	
	6.941	9.012											10.81	12.01	14.01	16.00	19.00	20.18	
	11	12											13	14	15	16	17	18	
3	Na	Mg											AI	Si	Р	S	CI	Ar	
	22.99	24.30	3	4	5	6	7	8	9	10	11	12	26.98	28.09	30.97	32.07	35.45	39.95	
	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	
р 4	K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr	
0	39.10	40.08	44.96	47.87	50.94	52.00	54.94	55.85	58.93	58.69	63.55	65.39	69.72	72.61	74.92	78.96	79.90	83.80	
<u> </u>	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	
O 5	Rb	Sr	Y	Zr	Nb	Мо	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Те		Xe	
ፈ	85.47	87.62	88.91	91.22	92.91	95.94	98.91	101.1	102.9	106.4	107.9	112.4	114.8	118.7	121.8	127.6	126.9	131.3	
	55	56		72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	
6	Cs	Ba	La-	Hf	Та	W	Re	Os	lr	Pt	Au	Hg	TI	Pb	Bi	Po	At	Rn	
	132.9	137.3	Lu	178.5	180.9	183.8	186.2	190.2	192.2	195.1	197.0	200.6	204.4	207.2	209.0	210.0	210.0	222.0	
	87	88	Ac-	104	105	106	107	108	109										
7	Fr	Ra	Lr	Unq	Unp	Unh	Uns	Uno	Une										
	223.0	226.0	Lſ																
			\	\backslash															
	s bloc	k	d blog										p block						
	3 0100	'n		JN									p DIOCK						
			\setminus	```	\backslash														
		L	anthani	des	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71
Landhadoo					La	Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu
				\backslash	138.9	140.1	140.9	146.2	144.9	150.4	152.0	157.2	158.9	162.5	164.9	167.3	168.9	173.0	175.0
\backslash				89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	
		A	Actinide	\setminus	Ac	Th	Ра	U	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No	Lr
				\setminus	227.0	232.0	231.0	238.0	237.0	239.1	241.1	244.1	249.1	252.1	252.1	257.1	258.1	259.1	262.1

fblock