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دور مايو ۲۰۱۶ الزمن: ساعتـــان التاريخ:۲۰۱۲/۲/۱۲



كلية العلوم - قسم الرياضيات

الفرقة: التانيـــــة الشعب: كيمياء كيمياء حيوية كيمياء نبات – كيمياء حيوية كيمياء نبات كيمياء حيوان علوم بيئة جيولوجيا المادة: ر ٢٠١ رياضيات بحتة

# أجب على الأسئلة الآتية:

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مع التمنيات بالتوفيق



Mansoura University Faculty of Science Zoology Department Date: 5<sup>th</sup> June 2014

Time: 2 hrs.



Program: Biophysics Z 225, Blood & Endocrine

Full Mark: 60 Marks

# Answer All the following questions:

# I. Answer All the following questions:

[20 Marks]

# (I.a.): True or False Questions ...... (10 Marks)

- 1. Mineralocorticoid hormones are produced in the zona glomerulosa.
- 2. Most hormones are regulated by a positive feedback mechanism.
- 3. One of the potential effects of hormones is to activate enzymes.
- 4. Hormones are secreted from endocrine glands into the bloodstream. Exocrine glands secrete hormones into ducts.
- 5. The sex hormones are produced in the male and female gonads.
- 6. Of the three categories of hormones (steroids, proteins, and amines), only the steroids and amines can be administered orally.
- 7. The presence of specific receptor sites on the cell membrane ensures the correct signaling of hormones to target cells.
- 8. Hyperglycemia results when alpha cells continuously secrete glucagon.
- 9. Insulin hormone stimulates the liver to release sugar into the blood.
- 10. Gastrin, Secretin, cholecystakinin, and entrokinin are hormones secreted by the small intestine.

# (I.b.): Choose the correct answer: .....(10 Marks)

## 1- Parafollicular cells are related to...

a. thyroid gland

b. parathyroid gland

c. pituitary gland

d. pineal gland.

#### 2- Which is TRUE statement regarding glucagon?

a. secreted by Alpha cells

b. reduce blood glucose rate

c. stimulate gluconeogenesis

d. promote protein production.

# 3- A meal rich in proteins but low in Carbohydrates does not cause hypoglycemia because:

- a. glucagon secretion is stimulated by hypoglycemic meals.
- b. the meal causes compensatory increase in T4 secretion.
- c. cortisol in circulation prevents glucose from entering the muscles.
- d. the amino acids in the meal are converted to glucose

## 4- Which of the following is not involved in regulation of plasma Ca++ levels?

- a. kidneys.
- b. skin.
- c. lungs.
- d. intestine.

# 5-) Hypopituitarism is characterized by: d. excessive growth of the b. intolerance to heat. c. weight gain. a. Infertility soft tissue. 6- Excessive growth hormone secretion in adults causes: b. acromegaly c. increased entry of glucose in muscles d. hypothyroidism a. gigantism. 7- Parathyroid hormone: a. decreases Ca++ mobilization of bone b. increases Ca++ mobilization from bone. c. decreases circulating d. increases urinary excretion of Ca++ levels of free Ca++ 8- Calcitonin a. increases Ca++ absorption by stomach. b. is secreted by hypothalamus. d. is secreted by thyroid gland. c. is secreted by parathyroid gland. 9- When would plasma insulin levels be expected to be higher? a- After intravenous administration of somatostatin. b- Following a carbohydrate-rich meal. d- Following vigorous exercise c- During a surgical procedure. 10- ADH travel in the blood to the main target: a- Hypothalamus b- pituitary c- Collecting ducts d- Renal cortex (II) Answer the following questions using labeled diagram. (25marks) (II.a.): Write on the following: ......(10 Marks) a-The Physiological importance of Testosterone & Oestrogen b- Diabetes mellitus. (II.b.): Answer the following questions: ......(15marks) a) Shortly illustrate the Blood clotting b) Define Diapedesis, Leucopenia, polycythemia (15 marks) (III) Answer the following items: a) Briefly illustrate the sites and essential factors of erythrocyte production in the adult human. b) Calculate the MCV and the MCHC for a subject with a red blood cell count of $4x10^6$ per cubic mm, a hematocrit of 40% and a hemoglobin concentration of 12 g/dl. Describe the subject's red cells: are they normocytic? Normochromic c) Tabulate a comparison between types of leucocytes with respect to percentages, location

Best wishes

and functions.

Prof. Dr. M. Amr El-Missiry & Prof. Dr. Maher Amer

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	Mansoura University Faculty of Science Chemistry Department
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Second Term	May 2013
Second Level Biochemistry	Date: 9
Course Title: Amino acids & Proteins	/ 6/ 2014
Metabolism	Time allowed: 2 hours
Code No.: Biochemistry 277	Full Mark: 80 Marks

Note: Express your answers by formulae, equations, pathways, figures and diagrams wherever possible,

## **Answer The Following Questions**

# Question I: (30 Marks)

# A- Give a scientific name of the following definitions: (12 Marks)

- 1- The proportion of essential amino acids in a food relative to their proportion in a protein that will be synthesized in the body.
- 2- A rare autosomal disorder resulted from a metabolic defect in reaction 5 of urea biosynthesis.
- 3- Nitrogen excreted nitrogen intake = positive value.
- 4- In contrast to transamination reactions, a reaction results in the liberation of the amino group as free ammonia, replacing the amino group with oxygen from O2.
- 5- An enzyme can release or incorporate ammonium to or from glutamate.
- 6- A disease results from a metabolic block at glutamate-γ- semialdehyde in proline catabolism.
- 7- A metabolic disorder produced from a defect at fumarylacetoacetate hydroxylase (reaction 4) in tyrosine metabolism, the untreated acute and chronic cases leads to death from liver failure.
- 8- Serves as a precursor for methyl transfer reactions e.g. the conversion of norepinephrin to epinephrine.
- 9- It is the most important route for disposing of nitrogen from the body.
- 10- An essential amino acid is only required during growth.
- 11- It is the source in metabolism of: Serine?
- 12- It is the source in metabolism of: Glycine.

#### B- Complete the following: (10 Marks)

a- Hydroxylation of peptide-bound prolyl and lysyl residues is catalyzed byand
of skin and skeletal muscles.
b- All amino acids except and undergo transaminations.
c- The urea cycle is Upregulated by and BUT it's Downregulated
by
<b>d-</b>
breakdown.
e- Folic acid deficiency can be detected by excretion of N-formiminoglutamate (Figlu) following
a dose of amino acid.
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#### C- Choose the correct answer: (8 Marks)

Don't give more than one answer to a question. – Copy the table in your answer sheet.

No.	1	2	3	4	5	6	7	8	Total Marks = 8
Answer									8

1- Nitric oxide and urea have in common	the fact that they both have as an intermediate
precursor the amino acid:	
a) Aspartate.	b) arginine.
c) glutamate.	d) phenylalanine.
2- The NH3 produced in muscle degradat	ion of nitrogenated compounds is transported
through blood to the liver using as carr	
a) alanine and glutamine.	b) urea and alanine.
c) NH4 and glutamate.	d) Glutamate and glutamine.
e) Alpha-ketoglutarate and urea.	
3- The necessary coenzyme for transamin	ation reactions is:
a) Coenzyme.	b) Tetrahydrobiopterin.
c) FMN.	d) Folate.
e) Pyridoxal Phosphate.	f) Thiamine Pyrophosphate.
4- A person with phenylketonuria cannot of	
a) phenylalanine to tyrosine.	b) phenylalanine to isoleucine.
c) phenol into ketones.	d) phenylalanine to lysine.
5- Lacked argininosuccinate synthase acti	ivity leads to <u>one</u> of the following metabolic disorder
of urea biosynthesis:	
a) Citrullinemia.	b) Hyperargininemia.
c) Hyperammonemia Type 1.	d) Hyperammonemia Type 2.
6- These are neurotransmitters derived from	om amino acids, <u>except</u> :
a) Histamine.	b) Epinephrine.
c) Dopamine.	d) γ-aminobutyrate.
7- The metabolic defect of tyrosine aminor	transferase <u>(reaction 1)</u> is detected in
a) Neonatal tyrosinemia.	b) Tyrosinemia Type 2
c) Tyrosinemia Type 1.	d) Alkaptonuria.
8- These are the characteristic features of	Maple Syrup Urine Disease, <u>except:</u>
a) Accumulation of branched-chain a as	
b) Accumulation of branched-chain α-k	
c) Deficiency of branched-chain α-keto	
d) Early diagnosis prior to birth is usele	
f) Brain damage, mental retardation and	l early mortality.
Question 2: (30 Marks)	
Give a brief account on:	
i- Hartnup disease.	
ii- Alkaptonuria.	
iii- β-alanyl dipeotides and related disorder	S.
iv-Biosynthesis of creatine and creatinine.	Caracia de la casa de la
v- Alternative pathways of phenylalanine c	atabolism in phenylketonuria.
Question 3: (20 Marks)	
Demonstrate the following by chemical e	quations
i- Biosynthesis of serotonin and melatonin	
ii- Biosynthesis of epinephrine and norepin	
iii- Two pathways of L-cysteine catabolism	
iv- Biosynthesis of $\gamma$ -aminobutyrate and re	
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With My Best Wishs

Examiner: Dr. Nivin A. Salah



# المستورالمان كيميا, جوس أعن الإعامل النوية وع ٧٦

**Mansoura University Faculty of Science Chemistry Department Subject: Chemistry** Course(s): Nucleic acids Metabolism **Biochem 276** 



2<sup>nd</sup> Level Biochemistry Students

Date : May 2014

**Time Allowed: 2 hours** Full Mark: 80 Marks

# ANSWER THE FOLLOWING QUESTIONS

Express your answer by formulae equations, pathways, diagrams or figures wherever possible.

I. a)	Ti	rue or false	[30 Marks] [16 Marks]	
	1.	Dietary purines do not participate in the salvage pathways and converte	ed to uric acid	( )
		The building block of nucleic acids is nucleotides, which includes in		. ,
		phosphate group		( )
	3.	Conversion of ribonucleotides to the deoxy forms is catalyzed by	ribonucleotide diphosp	hate
		kinase		( )
		If one purine and one pyrimidine formed a base pair their width will fit		
	5.	Purines bond to the first carbon of the sugar at their N9 atoms, while prat their N1 atoms.	yrimidines bond to the s	ugar ()
	6.	Cytarabine is a suicide inhibitor and works through irreversible inhibit	tion of thymidylate synt	hase
				( )
	7.	Allopurinol inhibits xanthine oxidase and reduces the levels of serum u	ric acid	( )
	8.	In marine invertebrates and crustaceans, allantoin is metabolized to am	monia	( )
b)	di	rimidine analogues include several drugs that are useful in seases. Comment and describe <u>THREE</u> examples.  escribe the different pathways for purine catabolism.	treatment of neopla [14 Marks]	nstic
III.	De	1) Gout. 2) Tautomerization of nitrogenous bases. 3) Folate analogues.	[24 Marks]	

Best wishes for our dear students, Dr. Amr Negm