# 考生注意事項:所有考題務必在答案卷上作答。凡在問題卷上作答者無效。

### 一、選擇題(均為單選,每題1分,答錯倒扣0.25分)

Select a correct name for each of the following compounds.

- 1. A. Cytosine
  - B. Uracil
  - C. Thymine
  - D. 5 Methylpyrimidine E. 5 Methylcytosine



- 2. A. Adenine
  - B. Xanthine

  - C. Guanine D. N<sup>7</sup>- methylguanine
  - E. Uracil

- 3. A. 3.7 dimethyl-xanthine

  - A. 3.7 dimethyl-xanthine
    B. 1.6 dimethyl-xanthine
    C. 3.7 dimethyl-hypoxanthine
    D. 1.6 dimethyl-hypoxanthine

  - E. Caffeine



- 4. At neutral pH, what is the least soluble of the bases ?
  - A. Adenine
  - B. Guanine
  - C. Xanthine
  - D. Thymine
  - E. Cytosine

# 二選擇題(均為單選,每題2分,答錯倒扣0.5分)

- 5. Inosinic acid is
  - A. a precursor of all purine ribonucleotides synthesized de novo. B. deficient in the condition known as gout.

  - C. a degradation product of cytidine.
    D. a competitive inhibitor of xanthine oxidase.
  - E. a end product of purine catabolism.
- 6. Which of the following statements about allopurinol ( 4 hydroxypyrazolopyrimide ) is <u>Incorrect</u> ?
  - Allopurinol is
  - A. an inhibitor of xanthine oxidase
    B. used for the treatment of hyperuricemia.

  - C. an inhibitor of de novo purine biosynthesis.
    D. effective in the local treatment of herpetic Keratitis
  - E. used to treat chronic gout.
- 7. What is the main product of purine in lower primates and other mammals ? A. Uric acid

  - B. Allantoin
  - C. Urea
  - D. Guanine
  - E. None of the above
- 8. The severe combined immunodeficiency disease ( SCID ) is an inherited autosomal recessive disorder in which both T cells and B cells are sparse and dysfunctional.

  - The defect is due to the deficiendy of A. hypoxanthine guanine phosphoribosyl transferase
  - B. adenosine deaminase
  - C. xanthine oxidase
  - D. purine nucleoside phosphorylase
  - E. hypoxanthine oxidase

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#### 國立成功大學七十九學年度 集物化學 考試( 試題) 中間代謝 茅 ţĪ

考生注意事项:所有考题務必在答案卷上作答。凡在問題卷上作答者無效。

三選擇題(毎題2分,答錯倒扣0.5分)

Answer question 9-12 according to the following key.

- A. If 1,2 and 3 are correct.
- B. If 1 and 3 are correct.
- C. If 2 and 4 are correct.
- D. If only 4 is correct
- E. If all are correct9. The uptake of glucose in which of the following cells is insulin independent.
  - 1. red blood cells
  - liver parenchymal cells
     brain cells
     heart cells
- 10. Fructose -2.6- bisphosphate
  - 1. is an activator of fructose -1.6- diphosphatase.
  - 2. is increased when glucagon/insulin ratio is high.
  - 3. is an extracellular messenger.
  - stimulates the conversion of fructose -6- phosphate to fructose-1.6diphosphate.
- 11. The pyruvate dehydrogenase complex reaction includes the
  - 1. reduction of FAD by oxidized lipoic acid. 2. oxidation of FADH $_2$  by NAD $^+$ . 3. fromation of ATP.

  - 4. formation of acetyl CoA
- 12. Cyclic AMP mediates the stimulation of glycogenolysis in liver by which of the following hormones or compounds.
  - 1. inositol triphosphate
  - 2. epinephrine
  - 3. insulin
  - 4. glucagon

#### 四簡答题

- 13. (3%) How the triacylglycerol is formed from glycerol and fatty acids?
- 14. (4%) Cholesterol plays an important role in cellular function. How the cholesterol content of cells is regulated?
- 15. (5%) Where the ketone bodies are formed? Why they are formed?
- 16. (8%) Triacylglycerol content in the blood of a diabetic patient is also higher than that of a normal person. Why?
- 17. (10%) Compare the following aspects of fatty acid oxidation and synthesis.
  - (a) Site of the process.

  - (b) Acyl carrier
    (c) Reductants and oxidants
    (d) Stereochemistry of the intermediates.
  - (e) Direction of synthesis or degradation.
- 18. (4%) Write the reactions and enzymes that catalyze substrate level phosphorylation during glycolysis.
- 19. (8%) Discuss the regulation of glycogen synthase by covalent modification.
- 20. (6%) Oxaloacetate-4-14C ( O=C-COOH  $CH_2-*COOH$  ) is incubated with aectyl CoA and all the mecessary enzymes and cofactors of the tricarboxylic acid cycle. Would the oxaloacetate be labeled after one turn of the cycle ? Give your explanation.

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國立	成功大學七十九學年度 集物化學	·考試(中間代謝 試題)共 3 月
考生注意事项:所有考题務必在答案卷上作答。凡在問題卷上作答者無效。		
21.	(4%) Which enzyme deficiency in eryth hemolytic anemia ? Why ?	rocyte would produce drug - induced
22.	(3%) 5 - Fluorodeoxyuridylate ( $FdUMP$ ) is a potent antitumor agent, how can this drug inhibit growth of cancer cell ?	
23.	(2%) Give the name of the coenzyme which is essential in most aminotrans- ferase catalytic activity.	
24.	(4%) Complete the following equation of the reaction catalyzed by glutamate dehydrogenase.	
	Glutamate + +	+ + + + + + + + + + + + + + + + + + +
25.	(4%) Complete the overall chemical balance of urea biosynthesis from ammonium.	
	2NH <sub>3</sub> + + Urea	+ + + + 2Pi
26.	. (3%) Name three amino acids which is involved in the biosynthesis of creatine.	
27.	(3%) Name the enzyme deficiency of the following genetic diseases.	
	Genetic disease	Name of the enzyme
	Maple syrup urine disease	1
	Phenylketonuria	2.
	Alkaptonuria	3
28.	(4%) Draw the structure of S-adenosylmethionine.	
29.	(5%) Describe the role of vitamine A in vision.	