

M.Sc. CHEMISTRY
FOURTH SEMESTER
MEDICINAL CHEMISTRY
MSC - 402A

(Use Separate Answer Scripts for Objective & Descriptive)

Duration: 3 hrs.

Full Marks: 70

Time: 30 min.

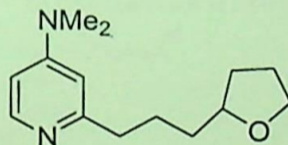
(PART-A: Objective)

Marks: 20

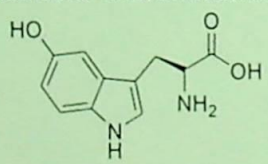
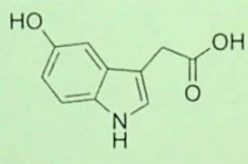
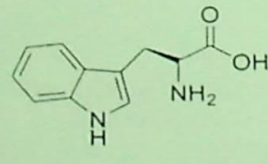
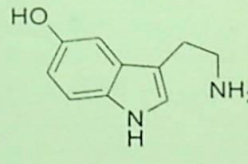
Choose the correct answer from the following:

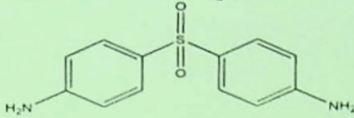
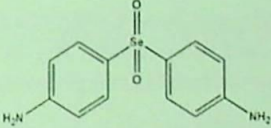
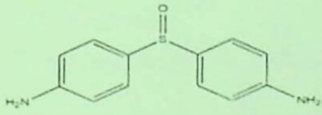
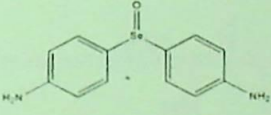
1X20=20

- Bacterial cell wall is a polymer of hetero-polysachharide crosslinked with peptide chain. The suger units of hetero-polysachharide are:
 - N-acetyl glucoseamine and N-acetyl muramic acid.
 - N-acetyl galactoseamine and N-acetyl muramic acid
 - N-acetyl glucuronic acid and N-acetyl muramic acid
 - N-acetyl glucoseamine and N-acetyl cialic acid.
- Any given drug will express its activity through
 - Enzyme inhibition
 - binding with specified cell receptors
 - Binding with DNA
 - any of these.
- The following compound has potent antifungal activity in a cell free system but poor activity in mice.



- Why is it not effective in mice? The reason could be.
- The compound is sterically crowded.
 - The compound is basic and cannot cross the membrane barrier.
 - The compound is highly basic and in physiological condition get protonated in GI track, becomes highly polar and cannot cross non-polar cell membrane.
 - All statements are wrong.
- Potency of a drug is found to increase when -Cl group is introduced in p-position in phenyl moiety of the lead compound of a drug. Which of the following group is likely to increase the potency further when the -Cl group is displaced with?
 - Cl in 3- and 4- position
 - NMe₂
 - OH
 - OMe
 - A highly polar drug:
 - Will be easily absorbed in the GI track.
 - Will be held up inside the lipid bi-layer of cell membrane.
 - Need to administered by injection for proper absorption.
 - None of the above is correct.

6. Chloramphenicol is -
- Antiseptic and disinfectant
 - Broad spectrum antibiotic
 - Antifertility drug
 - Antihistamine
7. β -Lactam compounds are effective inhibitors of cell wall synthesis. this is because, their geometry is similar to
- Bactoprene
 - N-acetyl-glucosamine
 - Alanyl-alanine residue
 - None of them
8. Sulbactam functions as an inhibition for the enzyme
- β -Lactamase
 - Ligase
 - Transpeptidase
 - Racemase
9. Crosslinking of peptidoglycan chain is catalyzed by
- Oxidase
 - Amidases
 - Transpeptidases
 - Hydrolases
10. 50S subunit contains
- 23S & 5S rRNA complexed with 34 polypeptides
 - 23S & 5S rRNA complexed with 38 polypeptides
 - 23S & 5S rRNA complexed with 43 polypeptides
 - 23S & 5S rRNA complexed with 21 polypeptides
11. Serotonin is a monoamine neuro-transmitter, its biosynthesis occurs from
- L-Tyrosine
 - L-Glycine
 - L-Tryptophan
 - L-Leucine
12. Monoamine neurotransmitter, serotonin is metabolized to
- 
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13. Drugs acting as monoamine oxidase inhibitors are
- Anaesthetics
 - Cardiovascular
 - Anti-depressants
 - Anti-neoplastic
14. Nephrotoxicity refers to the damages in
- Kidney
 - Liver
 - Heart
 - Eye
15. An example of non-halogenated anaesthetic is -
- Enflurane
 - Halothane
 - Isoflurane
 - Cyclopropane

16. NSAID refers to
- Steroidal anti-inflammatory drug
 - Non-steroidal Anti-inflammatory drug
 - Steroidal inflammatory narcotic drug
 - Non-steroidal inflammatory drug
17. Ibuprofen is a derivative of
- Butanoic acid
 - Propionic acid
 - Hexanoic acid
 - Ethanoic acid
18. Ethambutol is a
- Anti-cancer agent
 - Anti-pyretic agent
 - Antituberculosis agent
 - Cardiovascular drug
19. Correct structure of Dapsone is
- 
 - 
 - 
 - 
20. An important side effect of melphalan is
- sores or ulcers in the mouth
 - Ovarian cancer
 - Bone marrow depression
 - All of the above

[3]

(PART-B : Descriptive)

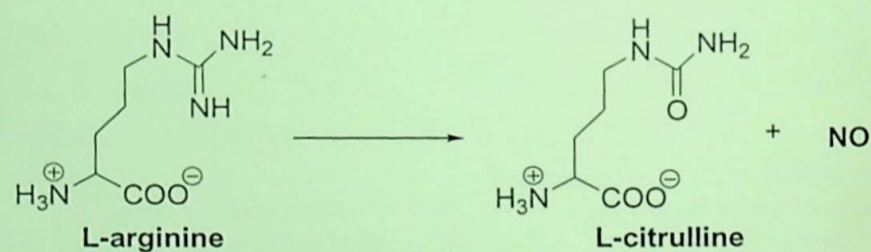
Time : 2 hrs. 40 min.

Marks : 50

[Answer question no.1 & any four (4) from the rest]

1. a. What do you mean by pinocytosis? 2
b. Discuss the Structure Activity Relationship (SAR) of penicillin. 3
c. Write the structure of enflurane. Give its industrial synthetic route. 2
d. Draw the structure of Melphalan. Describe its activity in treatment of cancer cells. 1+2 = 3

2. a. Nitric oxide synthase Catalyses the the conversion of L-arginine to L-citrulline and nitric oxide. If you want to interfere with the production of NO, design at least three potential leads. 3



- b. Write the linear Hansch equation correlating drug potency with lipophilicity. 2
- c. What do you mean by lead compound in drug discovery? Illustrate with example lead discovery and lead modification of any specific drug molecule. 5

3. a. Explain the mode of action of streptomycin. 2
b. Define translocation and transpeptidation? Explain their role in the synthesis of protein. 3
c. What are the inhibitors of protein synthesis? Discuss the mechanism of inhibition of protein synthesis by such inhibitors. 5

4. a. The penicillin on being hydrolyzed with hot dilute inorganic acids is known to give an amine known as D-penicillamine. Elucidate the structure of D-penicillamine. 5
b. Write the complete the synthesis of Penicillin-V 5

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|----|---|--------------|
| 5. | a. What is hepatotoxicity? Give the example of a drug having this side effect? Explain with example how it happens? | 2+1+2
=5 |
| | b. What is isosterism? Give short account of classical and nonclassical bio-isosterism. | 5 |
| 6. | a. What is the usage of an antidepressant drug? Give an example of monoamine transmitter and discuss its biosynthesis and metabolization. | 1+4=5 |
| | b. Write the structure of one anesthetic drug. Mention the type of anesthetic it belongs to. Discuss its synthetic route. | 1+1+3
=5 |
| 7. | a. Write down the structure of methyldopa. Mention one of its uses. Describe the synthesis of methyldopa. | 1+1+3
= 5 |
| | b. What is Tranquillizer? Draw the structure of Diazepam. Describe the synthesis of Diazepam. | 1+1+3
= 5 |
| 8. | a. Write down the structure of Dapsone. Mention one of its uses. Describe the synthesis. | 1+1+3
=5 |
| | b. Write the structure of Diclofenac and describe its synthetic procedure. | 1+4= 5 |

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