



# BRAINWARE UNIVERSITY

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Barasat, Kolkata - 700125

**Term End Examination 2021 - 22**  
**Programme – Bachelor of Pharmacy**  
**Course Name – Medicinal Chemistry III**  
**Course Code - BP601T**  
**( Semester VI )**

**Time : 1 Hr.30 Min.**

**Full Marks : 75**

[The figure in the margin indicates full marks.]

### Group-A

(Multiple Choice Type Question)

1 x 75=75

*Choose the correct alternative from the following :*

- (1) Alkali hydrolysis of Penicillin produces
 

a) Penicillenic acid	b) Penillic acid
c) Penicilloic acid	d) Penilloic acid
- (2) Clavulanic acid has a beta lactum ring fused with
 

a) Clavulanic acid has a beta lactum ring fused with	b) Thiadiazole system
c) Oxazolidine system	d) d.Thiazolidine
- (3) Substitution of bulky groups on the alpha carbon of side chain of Penicillin provides
 

a) beta lactamase resistance	b) . acid resistance
c) Penicillinase resistance	d) both (a) and (b)
- (4) An electron withdrawing substituent on the alpha carbon of side chain of Penicillin provides
 

a) beta lactamase resistance	b) acid resistance
c) Penicillinase resistance	d) both (a) and (b)
- (5) The cephalosporin with the highest activity against gram positive cocci is
 

a) Cefaclor	b) Cephalothin
c) Cefuroxime	d) Cefepime
- (6) Which of the following is considered to be bacteriostatic?
 

a) Penicillin	b) Chloramphenicol
c) Ciprofloxacin	d) Cefoxitin
- (7) Flucloxacillin
 

a) Is ineffective against streptococci	b) Is active against enterococci and anaerobes
c) Blocks transpeptidation and inhibits peptido	d) Is poorly absorbed orally

glycan synthesis

- (8) Aminoglycosides
- a) Have a  $\beta$  lactam ring
  - b) Can produce neuromuscular blockade
  - c) Are DNA gyrase inhibitors
  - d) Normally reach high CSF concentrations
- (9) Benzyl penicillin is also known as
- a) Penicillin - G
  - b) Penicillin - V
  - c) Penicillin - F
  - d) Penicillin - K.
- (10) Which semi synthetic penicillin is bactericidal in nature?
- a) oxacilline
  - b) cloxacillin
  - c) amoxicillin
  - d) penicillin V
- (11) Which semi synthetic penicillin is used in pseudomonous infection?
- a) oxacilline
  - b) cloxacillin
  - c) amoxicillin
  - d) carbenicillin
- (12) Which antibiotic is used against septicaemia?
- a) oxacilline
  - b) gentamycin
  - c) amoxicillin
  - d) carbenicillin
- (13) What is the full form of 6-APA?
- a) 6-amino penicillanic-acid
  - b) 6-amino penicilloic-acid
  - c) 6-amino penicillanic-anhydride
  - d) 6-aceto penicillanic-acid
- (14) Chloroquine act by inhibiting following enzyme
- a) DNA and RNA polymerase
  - b) DNA gyrase
  - c) Dihydro folate reductase
  - d) DNA
- (15) Following drugs have a asymmetric centre except
- a) Primaquin
  - b) Pamaquin
  - c) Quinocide
  - d) Pentaquin
- (16) Clindamycin
- a) Inhibits bacterial cell wall synthesis
  - b) Is often used for prophylaxis of endocarditis in patients with Valvular disease who are un dergoing dental procedures
  - c) Penetrates through BBB into CSF well
  - d) Works well against enterococci and gram ne gative aerobic organisms
- (17) Ribosomal resistance occurs with
- a) Sulphonamides
  - b) Penicillin
  - c) Fluoroquinolones
  - d) Macrolides
- (18) Which of the following is an ester?
- a) aspirin
  - b) methadone
  - c) methoxyflurane
  - d) adrenaline
- (19) Esterification of  $-OH$  group in a drug may lead to
- a) Destabilization
  - b) Degradation
  - c) Prodrug formation
  - d) Epimerization
- (20) Starting material for the synthesis of chloroquine is
- a) p-chloro aniline
  - b) m-chloro aniline
  - c) o-chloro aniline
  - d) aniline.
- (21) The structure of biguanides contain

- a) 6 nitrogen atoms  
c) 3 nitrogen atoms
- (22) Quinine structurally  
a) (+) 8S : 9R isomer  
c) (-) 7S : 8R isomer
- (23) Ethambutol is used to treat  
a) hypertension  
c) poisoning
- (24) PAS is  
a) Para-amino-sulfuric acid  
c) Para-amino-salicylic acid
- (25) Mode of action of Biguanides  
a) Cross linking inhibition  
c) Protein synthesis inhibition
- (26) Regarding fluoroquinolones  
a) Ciprofloxacin is ineffective in the treatment of gonococcus  
c) Norfloxacin and Ciprofloxacin have long half lives (12 hours)
- (27) Regarding resistance to antibiotics  
a) Penicillinases cannot inactivate cephalosporins  
c) Mutation of aminoglycoside binding site is its main mechanism of resistance
- (28) Concerning toxicity of antibiotics  
a) Enamel dysplasia is common with aminoglycosides  
c) A disulfiram like reaction can occur with macrolides
- (29) Influenza causing virus is  
a) Herpes virus  
c) Retrovirus
- (30) Which of the following moieties are present in the structure of acyclovir ?  
a) Adenine  
c) Guanine
- (31) Amantadine is used as  
a) anti-viral drug  
c) antiprotozoal drug
- (32) Stavudine is a / an  
a) Antimetabolite  
c) Reverse transcriptase inhibitor
- (33) The antiviral drug which is a thiazole analogue is  
a) Nelfinavir  
c) Saquinavir
- b) 5 nitrogen atoms  
d) 7 nitrogen atoms
- b) (-) 8R : 9S isomer  
d) (+) 7S : 8R isomer
- b) tuberculosis  
d) HIV
- b) Para-amino-sorbitol  
d) Para-amino-sulfonamide
- b) dihydrofolate reductase inhibition  
d) Microtubule damaging
- b) Norfloxacin and Ciprofloxacin are predominantly faecally excreted  
d) May damage growing cartilage in children less than 18 years of age
- b) Macrolides can be inactivated by transferases  
d) Tetracycline resistance is a marker for multi drug resistance
- b) Grey Baby Syndrome occurs with rifampicin use  
d) Haemolytic
- b) Orthomyxovirus  
d) Adenovirus
- b) Cytosine  
d) Thymine
- b) anthelmintic drug  
d) antibacterial drug.
- b) HIV protease inhibitor  
d) DNA polymerase inhibitor
- b) Ritonovir  
d) Loviride





- a) Oxazole  
c) Thiazole
- (49) Which is the basic ring present in sulfadiazine?  
a) Pyridine  
c) Pyridazine
- (50) Thyroid function is regulated by  
a) TSH  
c) GTH
- (51) Cyclopropane having the formula  
a) C<sub>4</sub>H<sub>4</sub>  
c) C<sub>3</sub>H<sub>6</sub>
- (52) All of the following are true regarding metronidazole EXCEPT  
a) It is used to treat giardia  
c) It inhibits alcohol dehydrogenase
- (53) Diethylcarbamazine Citrate is which class of drug  
a) Antiviral  
c) Anthelmintic
- (54) Albendazole contains which of the following  
a) Imidazole  
c) Thazolidine
- (55) Natural anthelmintic agent is  
a) Avermectin  
c) Levamisole
- (56) Pyrethrum and Pyrethroids comes under  
a) Anthelmintic  
c) Antibacterial
- (57) In vivo, prontosil is converted to  
a) Sulphanilamide  
c) Sulphadiazine
- (58) Glibenclamide belongs to the class  
a) Thiazolidinediones  
c) Benzoic acid derivatives
- (59) The chemical name of sulphadiazine is  
a) N 1 -2-pyrimidinylsulphanilamide  
c) N 1 -2-pyridylsulphanilamide
- (60) An azole most commonly used for topical treatment of candidiasis  
a) amphotericin B  
c) griseofulvin
- (61) The process by which DNA is copied to produce two daughter DNA molecules is...  
a) translation  
c) reproduction
- (62) Which of the following is one of the rules in Lipinski's rule of five?  
a) A molecular weight equal to 500
- b) Isoxazole  
d) None of the above
- b) Pyrimidine  
d) Piperidine
- b) TSH  
d) ACTH
- b) C<sub>2</sub>H<sub>4</sub>  
d) C<sub>4</sub>H<sub>8</sub>
- b) It causes a metallic taste in the mouth  
d) It is used to treat gardnerella
- b) Antifungal  
d) Sulfonamide
- b) Fural  
d) Benzimidazol
- b) Metrifonate  
d) Niridazole
- b) Antiviral  
d) Cardiovascular agents
- b) Sulphacetamide  
d) Sulphathiazole
- b) Sulphonyl ureas  
d) Biguanides
- b) N 1 -5-methyl-3-isooxazolylsulphanilamide  
d) N 1 -acetylsulphanilamide.
- b) clotrimazole  
d) flucytosine
- b) transcription  
d) replication
- b) No more than five hydrogen bond acceptor g



- roups
- c) No more than 10 hydrogen bond donor groups
- d) A calculated logP value less than +5
- (63) Which of the following descriptions most accurately describes binding sites and binding regions?
- a) a binding site is part of a binding region
- b) a binding region is part of a binding site
- c) a binding region is the same as a binding site
- d) a binding region is on a drug whereas a binding site is on a macromolecular target
- (64) Identify the kind of interactions that are typically involved in binding a drug to the binding site of a protein.
- a) predominantly van der Waals interactions
- b) predominantly ionic bonds
- c) predominantly hydrogen bonds
- d) a combination of all of the above
- (65) Which of the following statements best describes a lead compound?
- a) A compound that contains the element lead
- b) A compound that contains the element lead
- c) A molecule that shows some activity or property of interest and serves as the starting point for the development of a drug.
- d) The first compound of a structural class of compounds to reach the market.
- (66) What does the symbol P represent in a QSAR equation?
- a) pH
- b) plasma concentration
- c) partition coefficient
- d) prodrug
- (67) What is the symbol  $\pi$  in a QSAR equation?
- a) The hydrophobicity of the molecule
- b) The electronic effect of a substituent
- c) The substituent hydrophobicity constant
- d) A measure of the steric properties for a substituent
- (68) A measure of the steric properties for a substituent
- a) Molar refractivity is a steric factor
- b) Molar refractivity is an electronic factor
- c) Molar refractivity is a hydrophobic factor
- d) Molar refractivity is a stereoelectronic factor
- (69) What software programme is used to determine the Verloop steric parameter?
- a) Alchemy
- b) Chem3D
- c) Sterimol
- d) ChemDraw
- (70) What does a negative value of  $\sigma$  signify for a substituent?
- a) It is electron donating
- b) It is electron withdrawing
- c) It is neutral
- d) It is hydrophobic
- (71) Which of the following statements is untrue when comparing 3D QSAR with conventional QSAR?
- a) Only drugs of the same structural class should be studied by 3D QSAR or QSAR
- b) 3D QSAR has a predictive quality unlike QSAR.
- c) Experimental parameters are not required by 3D QSAR, but are for QSAR.
- d) Results can be shown graphically in 3D QSAR, but not with QSAR.
- (72) Full form of QSAR
- a) Qualitative structure–activity retention
- b) Quantitative structure–activity retention
- c) Qualitative structure–activity relationship
- d) Quantitative structure–activity relationship
- (73) Which of the following is NOT a stage of Combinatorial chemistry?
- a) finding a lead compound
- b) structure–activity relationships of the lead compound
- c) optimising a lead compound
- d) structure determination of the lead compound

d

(74) Combinatorial chemistry can be useful at various stages of the drug design / development process. Which of the following is such a stage?

- a) Purifying a lead compound
- b) Optimising a lead compound
- c) Structure determination
- d) Pharmacological testing

(75) Chloroquine act by inhibiting following enzyme

- a) DNA and RNA polymerase
- b) DNA gyrase
- c) Dihydro folate reductase
- d) DNA