GPAT QUESTION PAPER 1989 WITH ANSWER KEY

PY- PHARMACEUTICAL SCIENCES

Time: 3 hours Maximum Marks: 200

- N. B. 1. This question paper contains two parts A and B.
 - 2. Answer all the question from part A.
 - 3. Answer Any 20 Question from part B.

PART - A

- N. B. 1. There are 2 sections in this part
 - 2. Answer all the question in both sections 1 and 2.
 - 3. Answer should be given serial order in the answer book.
 - 4. Do not skip question while writing the answers.
 - Write the question number and show your answer by writing the alphabet (against the No.) in Capital letters.
 - 6. In section 1 each question carriers 1-Marks.
 - 7. In section 2 each question carries 2-marks.
 - 8. A model is shown at the beginning of each section in part A.
 - 9. Answer to the question in this part must be Witten in the first three pages only.

CHOOSE THE CORRECT ANSWER

Model Question

- 1. Repeated administration of Tyramine results in its decreasing effectiveness:
 - (a) Gets detoxicated easily
 - (b) Displaces nor-adrenaline from nerve ending binding site
 - (c) Displaces adrenaline from nerve ending binding site
 - (d) None of the above
- 2. Atropine on hydrolysis with Barium hydroxide gives:
 - (a) Tropanol and Tropic acid

(b) Scopine and Tropic acid

(c) Ecgonine and Benzoic acid

- (d) Benzyl Ecgonine and Methanol
- 3. The concentration of sucrose in simple Syrup BP is:
 - (a) 85% w/w

(b) 60.70% w/w

(c) 66.70% w/w

- (d) 40.74% w/w
- 4. Stratified cork and forked are the characteristic diagnostic features of:
 - (a) Apocynaceae

(b) Scrophularaceae

(c) Gentianaceae

(d) Polygonaceae

| 5. | Most accepted mechanism for developing bacterial resistance to sulphonamides is: | | | | | | | | |
|-----|--|---------------------------|----------------------|--|--|--|--|--|--|
| | (a) An increasing capacity to inactivate or destroy the drug | | | | | | | | |
| | (b) An alternative metabolic pathway for synthesis of an essential metabolite | | | | | | | | |
| | (c) An increasing product of drug antagonist | | | | | | | | |
| | (d) An alternation in enzyme that utilize PABA | | | | | | | | |
| 6. | C 17 α - β unsaturated lactone ring is a common t | feature in: | | | | | | | |
| | (a) Digitalis and squill glycosides | (b) Digitalis and strop | hantus glycosides | | | | | | |
| | (c) Digitalis and Senna glycosides | (d) Digitalis and Amys | gdalin | | | | | | |
| 7. | For drying blood plasma the following techinique | e is used: | | | | | | | |
| | (a) Spray drying | (b) Freeze drying | | | | | | | |
| | (c) Vacuum drying | (d) Fluid bed drying | | | | | | | |
| 8. | C3 O-glycoside digitoxin is used for: | | | | | | | | |
| | (a) Cardiac action | (b) Hypotensive actio | n | | | | | | |
| | (c) Precipitating steroids from solution | (d) Precipitating Anth | raquinone glycosides | | | | | | |
| 9. | Chemical name of amoxicillin is: | | | | | | | | |
| | (a) 6 - [D-(-) α – amino p-hydroxyacetamido] pe | enicillanic acid | | | | | | | |
| | (b) 4 - [D-(-) α - amino p-hydroxyacetamido] pe | enicillanic acid | | | | | | | |
| | (c) β - [Hydroxy analogue of Benzyl penicillin | | | | | | | | |
| | (d) α - Carboxy benzyl penicillin | | | | | | | | |
| 10. | The HLB value of sodium lauryl sulphate is: | | | | | | | | |
| | (a) 6.5 (b) 13.8 | (c) 25.0 | (d) 40.0 | | | | | | |
| 11. | Claviceps purpurea yields after infecting ovaries | of Graminaceous plants: | | | | | | | |
| | (a) Digitoxin | (b) Lysergic acid deri | vatives | | | | | | |
| | (c) Reserpine | (d) Polypeptides | | | | | | | |
| 12. | In the official bioassay of Erythromycin strain us | sed is: | | | | | | | |
| | (a) Bacillus subtilis | (b) Micrococcus luteu | S | | | | | | |
| | (c) Salmonella typii | (d) Escherichia coil | | | | | | | |
| 13. | The disintegration time for sugar coated tablet is | | | | | | | | |
| | (a) 30 minutes (b) 45 minutes | (c) 60 minutes | (d) 75 minutes | | | | | | |
| 14. | Idioblasts of crystal layer of calcium oxalate is a di | agnostic feature of | | | | | | | |
| | (a) Hyoscyamus Niger leaves | (b) Deadly nightshade | leaves | | | | | | |
| | (c) Cinchona bark | (d) Senna leaves | | | | | | | |
| 15. | Antibiotic which interacts with calcium ion is: | | | | | | | | |
| | (a) Erythromycin (b) Streptomycin | (c) Tetracycline | (d) Ampicillin | | | | | | |
| 16. | Flow rate of granules from the hopper can be imp | proved by adding; | | | | | | | |
| | (a) Disintegrant (b) Glidant | (c) Binder | (d) Lubricant | | | | | | |
| 17. | Silicon carbide rod heated to a high temperature i | | | | | | | | |
| | (a) Detector in infra red spectroscope | (b) Source of light in ir | | | | | | | |
| | (c) Source of light fluorimetery | (d) Detector in gas chr | omatography | | | | | | |

| 18. | Anomocytic type stomata are found in the leaves of: | | |
|-----|--|--------|--|
| | (a) Fox glove | (b) | Urginea maritime |
| | (c) Cassia acutifolia | (d) | Atropa belladonna |
| 19. | Liver microsomal enzymes are stimulated (enzymic | indu | ction) by: |
| | (a) Cimetidine (b) Phenobarbitone | (c) | Procaine (d) Adrenaline |
| 20. | Enteric coating is achieved by using: | | |
| | (a) Hydroxy propyl methyl cellulose | (b) | Carboxy methyl cellulose |
| | (c) Cellulose acetate Phthalate | (d) | Povidone |
| 21. | Car price reaction is applied for the photometric ev | aluat | ion of: |
| | (a) Vitamin A | (b) | Tocopherol |
| | (c) Nandrolone Phenyl Propionate | (d) | Benzodiazepine |
| 22. | Peroxide enzyme present in acacia is identified by: | | |
| | (a) Borntragers test | (b) | Molisch's test |
| | (c) Oxidation and extraction in Benzene | (d) | Oxidation and treatment with Benzididine |
| 23. | Prostaglandins are a group of related: | | |
| | (a) Alcohols (b) Aldehydes | (c) | Fatty acid (d) Alkaloids |
| 24. | Licence to sell drug specified in schedule C and C1 is | s give | en from number |
| | (a) -19 (b) -18 | (c) | -21 (d) -24 |
| 25. | Liqiude paraffin exhibits: | | |
| | (a) Plastic flow | (b) | Newtonian flow |
| | (c) Pseudoplastic flow | (d) | Dilatant flow |
| 26. | Estrogenic and Progestrogenic combination mainly: | | |
| 20. | (a) Inhibits the ovulation | | Inhibits the implantation of the fertilized ovum |
| | (c) Inhibits the fertilization of ovum | | Inhibits development of endometrium |
| 27. | More of earthy matter in a Rhizome is determine by | | |
| | (a) Total ash value | | |
| | (b) The earthy material is separated and then weight | ghed | |
| | (c) The Rhizome is washed in water and the in hyd | lroch | loric acid finally it is weighed |
| | (d) Acid insoluble ash value | | |
| 28. | Lidocaine is synthesized from: | | |
| | (a) 2 : 6-dimethyl-5-amino methyl benzene | (b) | 2: 6-dimethyl-5-nitro methyl benzene |
| | (c) 2:6-xylidene | (d) | 2-methyl-6-ethyl-5-amino methyl benzene |
| 29. | Sterilization temperature for aqueous solution in au | itocla | ve (Moist heat) is: |
| | (a) 72°C (b) 121°C | (c) | 147 °C (d) 160 °C |
| 30. | Following combination is suggested in the treatmen | t of I | Leprosy: |
| | (a) Dapsone + Ampicinllin + Clofazimine | (b) | Dapsone + Clofazimine + Rifampin |
| | (c) Dapsone + Erythromycin + Rifampin | (d) | Dapsone + Teracycline + Streptomycine |

| 31. | The gummy nature Astragalus gummifer is depend of | n: | | | |
|------|---|---------------|------------------------|-------|----------------------|
| | (a) More of Methoxly group of Bassorin | (b) | The carbohydrate co | ntent | t |
| | (c) More of hydroxyl groups of the sugar moiety | (d) | More of protein cont | ain o | f the drug |
| 32. | The vitamin administered with isoniazid to minimize | e its a | adverse reaction is | | |
| | (a) Vitamin A (b) Pyridoxine | (c) | Biotin | (d) | Pantothenic acid |
| 33. | For the synthesis of Nitrofurantoin which one of the | follo | owing combination of | chen | nicals are used: |
| | (a) 5-Nitro 2-furaldehyde and 2-amino hydantoin | (b) | 5-Nitro 2-furaldehyde | and | hydantoin |
| | (c) 5-amino 2-furaldehyde and 2-amino hydantoin | (d) | 5-Nitro 2-furaldehyde | and | barbituric acid |
| 34. | To get the optimum optical density of the solution fo | r 1 c | m thick layer the conc | entra | tion should be about |
| | (a) 10 ⁻⁴ mole/lit (b) 10 ⁻⁷ mole/lit | (c) | 0.1 gm/lit | (d) | 0.5 gm/lit |
| 35. | The sugar moiety of Digitails purpurea is: | | | | |
| | (a) 2:6-deoxy allose | (b) | 2 : 6-dedoxy glucose | | |
| | (c) 2:6-deoxy Rhamnose | (d) | 2 : 6-deoxy galactose | | |
| 36. | Additional of sodium chloride to sodium Oleate emuk | sion | will: | | |
| | (a) Stabilize emulsion | (b) | Destabilize emulsion | | |
| | (c) Decrease the globule size of the emulsion | (d) | None of the above | | |
| 37. | Anti hypertensive drug inhibits the rennin angiotens | sin sy | stem is: | | |
| | (a) Reserpine (b) Captopril | (c) | Methyl dopa | (d) | Propranalol |
| 38. | Acidity of Ascorbic acid is due to the presence of: | | | | |
| | (a) Free carboxylic acid | (b) | A number of hydrox | yl gr | oup |
| | (c) Enolic groups | (d) | None of the above | | |
| 39. | Progesterone injection BP is a sterile solution in: | | | | |
| | (a) Water (b) Ethyl oleate | (c) | Propylene glycol | (d) | Glycerol |
| 40. | Thiamine on treatment with sodium sulfite solution | and | sulfur dioxide yields: | | |
| | (a) Pyrimidine and a thiazole derivative http://w | ww.x | camstudy.com | | |
| | (b) Pyridine and thiazole derivative | | | | |
| | (c) 2:3:4-Thihydropyridine and Thiophene deriva | tives | 5 | | |
| | (d) Pyrimidine and Thiophene derivatives. | | | | |
| | | | | | |
| | SECTION | V - II | | | |
| | MATCH THE FO | OLL | OWING | | |
| | | | | | |
| 2.1. | Identify the correct skeleton ring present in the follo | wing | g compounds from the | ring | system listed from A |

(A) Perhydro cyclopentanophenanthrene

(B) 1:8 Naphthyridine

to E.

Riboflavin

Estrone

1.

2.

| | 3. | Indomethacin | (C) | Indole | | |
|------|------|--|-------|------------|--------|---|
| | 4. | Nalidixic acid | (D) | Quinolin | n | |
| | | | (E) | Iso allo | xagin | ne |
| | (a) | 1-E, 2-A, 3-C, 4-D | (b) | 1-D, 2-0 | c, 3-B | , 4-A |
| | (c) | 1-B, 2-C, 3-A, 4-D | (d) | 1-D, 2-A | A, 3-C | C, 4-B |
| 2.2. | Cho | sse the instrument or apparatus listed f | rom | A to E st | udy | the following: |
| | 1. | Rheology of semi solids | (A) | Andreas | sen P | Pipette |
| | 2. | Hardness of tablets | (B) | Monasa | nto t | tester |
| | 3. | Particle size in suspension | (C) | Ultrasor | nifier | • |
| | 4. | Homogenization of emulsion | (D) | Viscom | eter | |
| | | | (E) | Zeta me | ter | |
| | (a) | 1-D, 2-B, 3-C, 4-A | (b) | 1-E, 2-B | , 3-A | , 4-C |
| | (c) | 1-D, 2-C, 3-A, 4-B | (d) | 1-C, 2-B | 3, 3-D | , 4-A |
| 2.3. | Give | en below are some microscopical diagno | stic | of the dr | ug li | sted in A to E. Chosse the appropriate one. |
| | 1. | Unlignified septate fiber | | | (A) | Rhubarb |
| | 2. | Raphides of calcium oxalate embedded i | n mı | ıcilage | (B) | Solanaceous plant |
| | 3. | Anisocytic type of stomata | | | (C) | Ginger |
| | 4. | Star spots | | | (D) | Squill |
| | | | | | (E) | Solanaceous plants |
| | (a) | 1-A, 2-B, 3-C, 4-D | | | (b) | 1-D, 2-C, 3-B, 4-A |
| | (c) | 1-B, 2-C, 3-A, 4-D | | | (d) | 1-A, 2-D, 3-E, 4-A |
| 2.4. | Cho | sse the most appropriate drug for the f | ollov | ving | | |
| | 1. | Potassium-sparing diuretic | | | (A) | Spiranolactone |
| | 2. | Loop diuretic | | | (B) | Mannitol |
| | 3. | Osmotic diuretic | | | (C) | Furosemide |
| | 4. | Carbonic anhydrase inhibitor | | | (D) | Acetazolamide |
| | | | | | (E) | Aldosterone |
| | ` ' | 1-A, 2-C, 3-E, 4-D | | | , , | 1-A, 2-B, 3-C, 4-D |
| | . , | 1-A, 2-C, 3-D, 4-B | | | ` ' | 1-C, 2-B, 3-D, 4-A |
| 2.5. | | • | wav | e length i | range | es as listed under A to E. Choose the correct |
| | wav | ve length for the colour. | | | | |
| | 1. | Green | | | ` ' | 635-700 |
| | 2. | Orange | | | | 520-560 |
| | 3. | Yellow | | | | 560-590 |
| | 4. | Red | | | | 590-635 |
| | | | | | | 650-780 |
| | ` ' | 1-A, 2-B, 3-C, 4-D | | | | 1-B, 2-C, 3-A, 4-D |
| | (c) | 1-B. 2-D. 3-C. 4-E | | | (d) | 1-B. 2-C. 3-D. 4-A |

| 2.6. | Give | en below equipment used in the manufac | cture | of the follow | Wi | ing product A | to E | . Match them correctly. |
|------|---|---|------------|---------------|-----|-------------------|--------------------|---------------------------|
| | 1. | Zanasi | | (A) |) | Tablet granule: | S | |
| | 2. | HEPA Filter | | (B) |) | Tablet coating | | |
| | 3. | Chilsonator | | (C) | | Emulsion | | |
| | 4. | Accela cota | | (D) |) | Injectable | | |
| | | | | (E) | | Capsules | | |
| | (a) | 1-D, 2-A, 3-C, 4-B | | (b) | | 1-E, 2-D, 3-A, | 4-B | |
| | (c) | 1-B, 2-C, 3-A, 4-D | | (d) | | 1-C, 2-B, 3-D, 4 | 4-A | |
| 2.7. | Ma | tch the following with the schedules listed | d in A | A to E correc | ct | ly. | | |
| | 1. | Requirements of factory premises | | | | (| (A) | P |
| | 2. | Standards for disinfectant fluids | | | | (| (B) | V |
| | 3. | Life period of drugs | | | | | (C) | N |
| | 4. | List of minimum equipment for the effi | cient | t running of | P | harmacy (| (D) | 0 |
| | | | | | | | (E) | M |
| | (a) | 1-E, 2-D, 3-A, 4-C | | | | (| (b) | 1-B, 2-C, 3-D, 4-A |
| | (c) | 1-B, 2-C, 3-A, 4-D | | | | (| (d) | 1-C, 2-B, 3-D, 4-A |
| 2.8. | 2.8. Following are the reaction/tests observed in case of drugs listed in A to E. Match | | | | | | th them correctly. | |
| | 1. | When fixed oil is exposed to U.V. rays, I | olue | (A) |) | Digoxin | | |
| | 2. | On oxaidation with $KMnO_4$, Benzaldehyo | l e | (B) |) | Benzoin | | |
| | 3. | With ammoniacal Quaxom characterist | ic ba | llooned (C) |) | Cinchona | | |
| | 4. | Bark powder exhibits fluorescence with | 1 | (D) |) | Palmolein | | |
| | | sulphuric acid | | (E) |) | Gossypium ba | arba | adance |
| | (a) | 1-A, 2-B, 3-C, 4-D | | (b) |) | 1-D, 2-B, 3-E, | 4-C | |
| | (c) | 1-B, 2-C, 3-A, 4-D | | (d) |) | 1-C, 2-B, 3-D, | 4-A | |
| 2.9. | Me | chanism of Antitubercular action of the drug. | ug lis | ted are indic | ca | te are in A to E. | . Ch | oose the most appropriate |
| | 1. | Ethambutol | (A) | Prevents the | e | synthesis of p | rot | ein and DNA and reduces |
| | | | | R.N.A. synth | ıe | esis. | | |
| | 2. | P. A. S. | (B) | Interferes w | νi | th several of p | orot | ein synthesis |
| | 3. | Cycloserine | (C) | Competitive | e i | inhibiton | | |
| | 4. | Ethionamide | (D) | Inhibits pep | ot | ide synthesis i | n M | ycobacteria |
| | | | (E) | Inhibits DNA | Α | directed RNA | Syr | nthesis |
| | (a) | 1-A, 2-B, 3-C, 4-D | (b) | 1-D, 2-A, 3-0 | C, | . 4-B | | |
| | (c) | 1-B, 2-C, 3-A, 4-D | (d) | 1-D, 2-C, 3-H | Β, | . 4-A | | |
| | | | | | | | | |
| | | | | | | | | |

| 2.10.Given below are the receptor and their antagonist (A to E). Match them correctly. 1. Histamine H₂ Receptor (A) Atropine 2. Muscarinic Receptor (B) Ranitidine 3. Adrenaline α receptor (C) Pentolamine 4. Adrenaline α receptor (D) Metaraminol (E) Metoprolol (a) 1-B, 2-A, 3-C, 4-E (b) 1-D, 2-C, 3-B, 4-A |
|---|
| Muscarinic Receptor Adrenaline α receptor Adrenaline α receptor Adrenaline α receptor Metaraminol Metoprolol |
| 3. Adrenaline α receptor 4. Adrenaline α recptor (C) Pentolamine (D) Metaraminol (E) Metoprolol |
| (E) Metoprolol |
| |
| (a) 1-B, 2-A, 3-C, 4-E (b) 1-D, 2-C, 3-B, 4-A |
| |
| (c) 1-B, 2-C, 3-A, 4-D (d) 1-C, 2-B, 3-D, 4-A |
| 2.11. Match the following regions in GIT with the pH levels indicated from A to E. |
| 1. Mouth $(A) = 5.0 - 6.0$ |
| 2. Stomach (B) = $6.8 - 7.5$ |
| 3. Deodenum (C) = $6.8 - 7.0$ |
| 4. Large intestine $(D) = 3.0 - 5.0$ |
| (E) $= 1.5 - 3.0$ |
| (a) 1-A, 2-D, 3-B, 4-C (b) 1-A, 2-D, 3-B, 4-A |
| (c) 1-B, 2-E, 3-D, 4-C (D) 1-C, 2-B, 3-D, 4-A |
| 2.12.Listed in A to E are some of the analytical constants. Match them correctly with the drugs given below. |
| 1. A Leafy drug (A) Total ash value |
| 2. A Bark (B) Cineole content |
| 3. Eucalyptus oil (C) Fibre length |
| 4. A fixed oil having more of unsaturated (D) Iodine value |
| fatty acid glycerides (E) Stomatal index |
| (a) 1-A, 2-B, 3-C, 4-D (b) 1-D, 2-C, 3-B, 4-A |
| (c) 1-E, 2-C, 3-B, 4-D (d) 1-C, 2-B, 3-D, 4-A |
| 2.13. Match the ingredients listed A to E with the purpose for which they are used in the formulations. |
| 1. Film coating (A) Sodium benzoate |
| 2. Syrups (B) Ethyl cellulose |
| 3. Emulsification (C) Eudragit |
| 4. Enteric coating (D) Sucrose |
| (E) Sodium oleate |
| (a) 1-B, 2-D, 3-A, 4-C (b) 1-C, 2-D, 3-E, 4-B |
| (c) 1-B, 2-C, 3-A, 4-D (d) 1-C, 2-B, 3-D, 4-A |
| 2.14. Match the biological listed under A to E for the following compounds: |
| 1. 1: 3-Propanediol, 2-methyl 2-propyl Carbamate (A) Antimalarial |
| 2. 2 Chloro-10[3-(dimethylamino) propyl] Phenothiazine (B) Bactericidal to anaerobic an Microerophilic organisms |
| 3. 5 Nitro-2-furaldeyde semicarbazone (C) Antibacterial |
| 4. 2 Methyl-5-Nitro Imidazole –ethanol (D) Relief of anxiety and tension |
| (E) Tranquilizer |

| 1 | (a) | 1-A, 2-B, 3-C, 4-D | (b) 1-D, 2-A, 3-C, 4-B |
|--------|------|--|--|
| | (c) | 1-B, 2-C, 3-A, 4-D | (d) 1-E, 2-E, 3-D, 4-C |
| 2.15.0 | Give | en below are the drug A to E and the | nilments for which they are recommended. Match them correct |
| | l. | Parkinsonism | (A) Methyl dopa |
| 2 | 2. | Hypertension | (B) Levodopa with decarboxylase inhibitor |
| 3 | 3. | Nasal congestion | (C) Neostigmine |
| 4 | 1. | Myasthenia gravis | (D) Phenyl Propanolmine |
| | | , | (E) Ibuprofen |
| (| a) | 1-A, 2-B, 3-C, 4-D | (b) 1-B, 2-A, 3-D, 4-C |
| (| (c) | 1-B, 2-C, 3-A, 4-D | (d) 1-C, 2-B, 3-D, 4-A |
| 2.16.0 | Give | en below are some of the drugs and t | neir mode action in A to E. Match them correctly. |
| 1 | l. | Hydralazine | (A) Vasodilator by direct action |
| 2 | 2. | Phenothiazine | (B) Inhibits the Vasoconstrictor and presor effect of 5 HT |
| 3 | 3. | Methylsergide | (C) Antagonist D2 receptor of Dopamine |
| 4 | 1. | Tolazmide | (D) Stimulate the islet tissue to secrete insulin |
| | | | (E) Inhibiting the enzyme carbonic anhydrase |
| (| (a) | 1-A, 2-B, 3-C, 4-D | (b) 1-D, 2-C, 3-B, 4-A |
| (| (c) | 1-B, 2-C, 3-A, 4-D | (d) 1-A, 2-C, 3-B, 4-D |
| 2.17.0 | Give | en below in A to E are the names of | f drugs,. Appropriate tests are given below for drugs. Mat |
| t | her | n correctly. | |
| 1 | l. | Alcoholic solution of $\alpha\text{-naphthol}$ and | sulphuric acid E (A) Atropine |
| 2 | 2. | Murexide test | (B) Resepine |
| 3 | 3. | Para-dimethylamino Benzaldehyde | (C) Caffeine |
| 4 | 1. | Ninhydrine | (D) Gelatin |
| | | | (E) Triticum sativum powder |
| (| (a) | 1-A, 2-B, 3-C, 4-D | (b) 1-D, 2-C, 3-B, 4-A |
| (| c) | 1-E, 2-C, 3-A, 4-D | (D) 1-C, 2-B, 3-D, 4-A |
| 2.18.0 | Give | en below in A to E are the names of | instruments used for the determination of the following. Mat |
| t | her | n correctly | |
| 1 | L. | Particle volume | (A) Clarity apparatus |
| 2 | 2. | Presence of Foreign particle | (B) Du Nouy ring |
| 3 | 3. | Surface tension | (C) Coulter counter |
| 4 | 1. | Presence of polymorph | (D) Compactor |
| | | | (E) Differential thermal calorimeter |
| | | 1-C, 2-A, 3-B, 4-E | (b) 1-D, 2-C, 3-B, 4-A |
| (| (c) | 1-B, 2-C, 3-A, 4-D | (d) 1-C, 2-B, 3-D, 4-A |
| | | | |

2.19. Chosse the correct starting material listed from A to E for the synthesis of the following drugs.

1. Cortisone

(A) Diosgenin

2. Progesterone

(B) β-ionone

3. Testosterone

(C) Spirostanol

4. Vitamin A

- (D) Sarmentogenin
- (E) Anthracin
- (a) 1-A, 2-B, 3-C, 4-D

(b) 1-D, 2-C, 3-B, 4-A

(c) 1-B, 2-C, 3-A, 4-D

(d) 1-C, 2-A, 3-A, 4-B

2.20. Given below are the types of ointment bases. Match them with the correct ointments in A to E.

1. Absorption base

(A) Emulsifying ointmen

2. Oleogenous base

(B) Hydrophilic ointment

3. Emulsion base

(C) Oily cream

4. Water soluble base

(D) Kaolin poultice

(E) Simple ointment

(a) 1-A, 2-B, 3-C, 4-D

(b) 1-D, 2-A, 3-C, 4-B

(c) 1-B, 2-E, 3-C, 4-A

(d) 1-C, 2-B, 3-D, 4-A



N.B.: Answer any twenty questions

If more than 20 questions are attempted only the first 20 will be considered.

Answer should not exceed 15 lines

All Question carry equal marks.

- 3. Write the structure of the following indicating to what pharmacological category they belong
 - (a) [1-dimethylamino-3-(4-Chlorophenyl 3.2-Pyridyl) Propane]
 - (b) 2. Hydroxy methylene-17 β hydroxy -17-methyl 5 α-androstan 3-one
 - (c) 2, 4-diamino-5-(3, 4, 5-trimethoxy phenyl) methyl pyridine.
- 4. (a) What is a barrier-layer cell?
 - (b) What are the different ways by which a molecule can absorb energy
- 5. Explain briefly the improved artificial method for producing Sclerotium.
- 6. Give the characteristics of the ideal preservative for Pharmaceutical preparation.
- 7. Give the principal of official assay of INH. Given Equations for the reactions involved.
- 8. Give the exact mode action of the following drugs:
 - (a) Dicoumarol
 - (b) Vinblastin
 - (c) Valprolic acid

- 9. Give the mode of action of the following anti arrhythmic drugs: (a) Procainamide (b) Propranolol (c) Verapamil 10. Mention the various factors governing transdernal absorption of drugs. 11. (a) What is Hoffmann's exhaustive Methylaation? (b) Show the complete step of reactions when Isoquinoline is subjected to Hoffmann's exhaustive Methylaation. 12. How the solid samples are prepared for the measurement of IR Absorption spectra? Why such a process is adopted 13. Name the various Insulin injections which are official in IP. Mention time onset and duration of action. 14. Give the names of the drugs, their source. And one chemical test for identification of any one important constituent in each of the following. (a) Drug obtained as latex after incisions on capsule. (b) Dried juice obtained from the leaves of plant belonging to Liliaceaefamily. (c) A seed having action on heart 15. Enumerate the problems associated with use of plastic as a material for packaging Pharmaceuticals. 16. With the help of IR absorption readings how you can distinguish the following pairs of compounds. Predict the bands and interpret http://www.xamstudy.com 17. Define the following: (a) Liposome (b) Polymorphism (c) Prodrug 18. Name the various methods in the preparation of micro capsules and give only the process involved in the Cocaervation phase separation technique. 19. Give the mode of action of following antibiotics: (a) Ampicillin
- 20. What are the possible adulterants of fox glove leaves? How are they detected?
- 21. List the physic chemical factors affecting drug absorption.

(b) Tetracycline(c) Erythromycin

22. Write the equation for the following synthesis:
O-Chloro benzoic acid is condensed with 2, 3 Xylidine with the aid of Potassium carbonate and the resulting
Potassium salt is treats with mineral acid.

- 23. Give the possible Drug/Drug interaction of the following combination:
 - (a) Penicillin with probenecid
 - (b) Lithium carbonate with Chlorthiazide
 - (c) Levodopa with pyridoxine
- 24. (a) What concentration of Dextrose willbe used for the preparation of 100ml of Dextrose solution isotonic with blood serum. Molecular weight of Dextrose = 180
 - (b) In what proportion 80% and 30% alcohol mixed to obtain 50% alcohol.
- 25. Give the structure and specification relationship in the following compounds:
 - (a) Phenobarbital
 - (b) Amobarbital
 - (c) Cyclobarbital
 - (d) Pentobarbital
- 26. (a) An alkaloid gave E_1^1 at 310 nm = 180. The Extinction of 0.003% solution in water at 310 nm was found 0.500 (1 cm cell). Calculate the percentage of alkaloid.
 - (b) Find the HLB value of a center which has Saponificatio number 40.5 and acid number of the fatty acid 260.0
- 27. Name the endogenous neurohormones and give their structure.

End of paper

ANSWER KEY GATE 1989

Section - I

| 1 - c | 2 – b | 3 – c | 4 – a | 5 – d | 6 – b | 7 – b | 8 – a | 9 – b | 10 – d |
|--------|--------|--------|--------|--------|--------|--------|--------|--------|--------|
| 11 - b | 12 - b | 13 - с | 14 - с | 15 - b | 16 - b | 17 - b | 18 – a | 19 - b | 20 – c |
| 21 – a | 22 – d | 23 – c | 24 – c | 25 – c | 26 – a | 27 – d | 28 – c | 29 – b | 30 – b |
| 31 – a | 32 - b | 33 – a | 34 - c | 35 – b | 36 – b | 37 – b | 38 – a | 39 – b | 40 - a |

Section - II

| 2.1 - a | 2.2 - b | 2.3 - d | 2.4 - a | 2.5 - c | 2.6 - b | 2.7 - a | 2.8 - b | 2.9 - d |
|----------|----------|----------|----------|----------|----------|----------|----------|----------|
| 2.10 - a | 2.11 - c | 2.12 - с | 2.13 - b | 2.14 - d | 2.15 - b | 2.16 - d | 2.17 - с | 2.18 – a |
| 2.19 - d | 2.20 - c | | | | | | | |