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UNIVERSITY OF PETROLEUM AND ENERGY STUDIES
End Semester Examination, December 2021

Course: Pharmaceutical Jurisprudence **Semester :V**
Program: BPharm **Duration : 03 Hours**
Course Code: BP505T **Max. Marks: 75**

Instructions: Read each question carefully. Attempt all questions under Section A (20 x 1 marks). Attempt any two questions out of three under Section B (2 x 10 marks). Attempt any seven questions out of nine under Section B (7 x 5 marks).

SECTION A

Type the answers in test box

(20Qx1M=20 Marks)

S. No.		Marks	COs
Q.1	As per Drugs and Cosmetics Act and Rules, the Good Manufacturing Practice is included under Schedule (a) W (b) S (c) P (d) M		CO1
Q.2	Frusemide is included in (a) Schedule 2 (b) Schedule 3 (c) Schedule 4 (d) Schedule 1		CO2
Q.3	A medical practitioner must NOT prescribe the drug (a) temazepam (b) morphine (c) acitretin (d) diclofenac		CO1
Q.4	List of drugs exempted from the provisions of import of drugs (a) Schedule H (b) Schedule V (c) Schedule A (d) Schedule D		CO2
Q.5	In which one of the following schedules, drugs are marketed under generic name only? (a) Schedule W (b) Schedule X (c) Schedule Y (d) Schedule N		CO1
Q.6	Which of the following should appear on the manufacturer's package of thyroxine? (a) Pharmacy only Medicine (b) Pharmacist only Medicine (c) Prescription only Medicine (d) Controlled Medicine		CO2
Q.7	The incharge of state drug laboratory is (a) Drug Inspector (b) Government Analyst (c) Drug Registrar (d) Drug Controller of INDIA		CO1
Q.8	The standards for cosmetics are included in (a) Schedule C (b) Schedule S (c) Schedule X (d) Schedule C1		CO2
Q.9	Schedule Q describes (a) list of dyes, colours and pigments permitted to be used in cosmetics (b) standard for cosmetics (c) standard for disinfectant fluids (d) standard for surgical dressings.		CO1
Q.10	Class 100 in a parenteral production department indicates (a) area which contains not more than 100 particles/ft3 (b) area which is free from the particles of size more than 100 µ (c) area which is free from the particles of size more than 1000 µ (d) an area which is 100% sterile .		CO2
Q.11	As per drugs and cosmetics act, list of substances that should be sold only on prescription of registered medical practioner is given in (a) Schedule H (b) Schedule V (c) Schedule X (d) Schedule Q		CO1
Q.12	Schedule C drugs belongs to (a) standards for surgical dressings (b) biologicals to be administered parentrally (c) list of ayurvedic, siddha and unani drugs (d) requirement of factory premises		CO2
Q.13	Standards of ophthalmic preparations are included in Schedule (a) FF (b) U (c) W (d) C		CO1
Q.14	For manufacturing blood products or to operate blood bank, license is issued in the form number (a) 28 A (b) 28 (c) 28 B (d) 28 C		CO2
Q.15	If the drug is not labelled in prescribed manner then it is known as _____ drug. (a) spurious (b) misbranded (c) adulterated (d) toxic		CO1
Q.16	As per Schedule P of drugs and cosmetics act, diphtheria toxoid has expiry period of (a) 6 months (b) 12 months (c) 24 months (d) 60 months		CO2

Q.17	The total area required for the manufacture of cosmetics as per Schedule M is (a) 15 sq.m. (b) 25 sq.m. (c) 30 sq.m. (d) 35 sq.m		**CO1**
Q.18	All the following schedules are omitted EXCEPT (a) Schedule E (b) Schedule I (c) Schedule L (d) Schedule W		**CO2**
Q.19	Repacking of drugs means (a) formulation of drugs in bulk and packing in small units (b) breaking of drug from bulk container into small packs and labelling them for sale (c) packing, dispensing or formulation of drugs in retail sale (d) compounding of drugs in wholesale business		**CO1**
Q.20	VDRL antigen is to be tested and analysed by the (a) Drug Inspector (b) Excise Commissioner (c) Serologist and Chemical Examiner (d) Drug Controller of India .		**CO2**

SECTION B (20 Marks)

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(2Qx10M=20 Marks)

Attempt 2 Question out of 3

Q.1	Write the qualifications and duties of a Drug Inspector (Drugs Control Officer). Describe the procedures followed by the drug inspector in obtaining a sample. (5M+5M)		**CO4**
Q.2	What are the various types of patents and explain the procedure for obtaining a patent. (5M+5M)		**CO5**
Q.3	Describe the Drugs Prices Control Order and explain the salient features and the objectives of Drugs Prices Control Order.		**CO2**

SECTION-C (35 Marks)

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(7Qx5M=30 Marks)

Attempt 7 Question out of 9

Q.1	Write a note on offences and penalties prescribed under Pharmacy Act 1948. (2.5M+2.5M)		**CO1**
Q.2	Discuss as to how following GMP in the manufacture of drugs helps in maintaining the quality of drugs produced. .		**CO4**
Q.3	Discuss briefly the objectives of the Narcotic Drugs and Psychotropic Substances Act, 1985 and how it is achieved? (2.5M+2.5M)		**CO3**
Q.4	What are the objectives and salient features of Right to Information Act,2005? (2.5M+2.5M)		**CO1**
Q.5	Write notes on Schedules M, N, U of Drugs and Cosmetics Rules.		**CO4**
Q.6	Explain in brief the legal procedure for cultivation of poppy plant and extraction of opium & its salts.		**CO3**
Q.7	Write briefly on constitution and functions of "Committee for Control and Supervision of Experiments on Animals".		**CO1**
Q.8	Explain the circumstances under which a Registered Medical Practitioner can terminate the pregnancy.		**CO1**
Q.9	What are the advertisements prohibited under the Drugs and Magic. Remedies Act, 1954.		**CO1**