REV-01
DPH/11/16

2024/07

SET A

D.PHARM.
SECOND YEAR
PHARMACY LAW & ETHICS
ER 20-206T [SPECIAL REPEAT]

[USE OMR FOR OBJECTIVE PART]

Duration: 3 hrs. **Full Marks: 80**

[PART-A : Objective]

Choose the correct answer from the following: ***1×20=20***

1. Which drugs have low permeability, low solubility
   a. Class II
   b. Class III
   c. Class IV
   d. Class I

2. Which among the following is a drug regulatory authority
   a. MTP act
   b. Indian Pharmacopoeia
   c. CDSCO
   d. GMP

3. GRP means-
   a. Good Reasoning Practices
   b. Good regulatory process
   c. Good review Process
   d. Good regulatory practices.

4. When was MTP act enacted-
   a. 1917
   b. 1971
   c. 1947
   d. 2002

5. Which among the following blood group is the universal donor-
   a. O +ve
   b. AB -ve
   c. B +ve
   d. O -ve

6. How many elected members state pharmacy council have-
   a. 6
   b. 7
   c. 8
   d. 12

7. Insulin comes under-
   a. Schedule H
   b. Schedule J
   c. Schedule G
   d. Schedule O

8. ANDA means-
   a. Abbreviated new drug application
   b. Associated new design application
   c. Approved new data application
   d. Additional new drug application.

9. According to ER practical training for the apprentice pharmacist should be-
   a. 400 hours
   b. 500 hours
   c. 450 hours
   d. 250 hours

10. Chopra committee is also known as
    a. Drug legislative committee
    b. Drugs Enquiry committee
    c. Pharmacy enquiry committee
    d. Health survey committee

11. MTP act allows termination of pregnancy up to-
    a. 8 weeks
    b. 20 weeks
    c. 12 weeks
    d. 32 weeks

12. BCS means-
    a. Bio-pharmacological classification system
    b. Biological classification system
    c. Biopharmaceutical classification system
    d. Basic Control system.

13. The disaster management act was passed on-
    a. 1995
    b. 1955
    c. 2015
    d. 2005

14. Which drugs have high permeability, high solubility-
    a. Class III
    b. Class IV
    c. Class I
    d. Class II

15. Which regulatory body updates and reviews the Indian Pharmacopoeia
    a. ANDA
    b. CDSCO
    c. NIHFW
    d. DTAB

16. Drug and magic remedies act enacted in-
    a. 1985
    b. 1944
    c. 1945
    d. 1954

17. Animal welfare board is established by-
    a. Central govt.
    b. PCI
    c. State govt.
    d. Drug Inspector

18. In 1986, which among the following act was implemented-
    a. Drug and magic remedies act.
    b. Disaster management act
    c. Consumer protection act
    d. Biomedical waste management act.

19. The code of moral principles or the science of morals is termed as-
    a. Ethics
    b. Integrity
    c. Ideals
    d. Reliability

20. The licensing authority does not grant a license for import of drugs under-
    a. Schedule P
    b. Schedule X
    c. Schedule A
    d. Schedule C 1

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(PART-B : Short Answers)

[Answer any ten (10) from the following]

[3x10=30]

1. What are the objectives of GRP. 3
2. Write the objectives of Drugs and Cosmetics Act. 3
3. Write a short note on "biowaste management". 3
4. Write about the objectives of MTP act. 3
5. Define the following Schedule: X , C , H 3
6. What are the documents required for renewal of license. 3
7. What are medical devices. Classify them. 1+2=3
8. Write a short note on possession for sale of poison. 3
9. Write a short note on Spurious drugs. 3
10. What is the significance of the Disaster management act,2005. 3
11. Write a short note on community pharmacist. 3

(PART-C : Long Answers)

[Answer any six (6) from the following]

[5x6=30]

1. Explain NDA. 1+4=5
2. Classify drugs by BCS system of classification. 5
3. What is FSSAI. Mention its functions. 5
4. Discuss about the Committee members of DTAB. 5
5. Write about the roles of CDSCO. 5
6. Write about the functions of Blood bank 5
7. Explain about the members of PCI. 5

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