Question Paper

Exam Date & Time: 15-May-2023 (10:00 AM - 01:00 PM)

MANIPAL ACADEMY OF HIGHER EDUCATION

PHARMACY LAW and ETHICS [PMA-ER20-26T-S2]

Marks: 80 **Duration: 180 mins.**

MCQs

Answer all the questions. Section Duration: 30 mins

20 Q x 1 mark = 20 marks

1) What are rules established by an agency to interpret Laws are referred to as? (1)

- Legislations
- Regulations
- Guidelines
- Processes

2) Under the provisions of Poisons Act, following are the functions of State Government except (1)

- Possession for Sale
- Sale of Poison
- Import of Poison
- Possession of Poison

3) According to Medical Termination of Pregnancy Act, the District level committee can have a maximum of how many members in the committee? (1)

- 5 members
- 7 members
- 9 members
- 11 members

4) What is the full form of RMP? (1)

- Right Medical Practitioner
- Registered Marketing Practitioner
- Right Marketing Practitioner
- Registered Medical Practitioner

5) What is "any living creature except human being" is referred to as what according to Prevention of Cruelty to Animals Act? (1)

- Animal
- Bird
- Insect
- Reptile

6) The Code of Pharmaceutical Ethics in India is given by whom? (1)

ICI
DCI
MCI
PCI

7) Drugs and Magic Remedies (Objectionable Advertisements) Act was passed in which year? (1)

1948
1919
1954
1940

8) Minimum haemoglobin value required for a donor to donate blood is ____ gm/dl (1)

12
12.5
12.25
11

9) Maximum age for first time blood donor is ____ years (1)

50 years
60 years
40 years
45 years

10) Which of these is not a biomedical waste? (1)

Fumes
Syringes
Sharp
Amputated body parts

11) Cytotoxic and expired drugs are disposed by (1)

Dumping
Autoclave
Incineration
Chemical disinfection

12) Which of the following item falls under non-hazardous waste? (1)

Chemotherapy waste
Chemicals
Dextrose solution
Gases

13) Autoclaving and microwaving are done for which of the following types of medical waste? (1)

Human anatomical waste
Recyclable contaminated waste
Cytotoxic drugs
Microbiological waste

14) Grant of license to manufacture a drug requires (1)

Form 24
Form 25

Form 26
Form 27

15) Biological and Biotechnological Products are included in schedule (1)

A
B
C and C1
X

16) Schedule C is related to (1)

Biological and Immunological Product
Homeopathy Product
Ayurvedic Product
Allopathic Product

17) Which of the following is prohibited to be imported (1)

Herbal drugs
Schedule G drugs
Ayurvedic drugs
Misbranded drugs

18) In 1985, one of the following Act was passed (1)

Narcotic and Psychotropic substances Act
Drugs and Magic Remedies Act
Medical Termination of Pregnancy Act
Poisonous Act

19) Government opium factory is situated at (1)

Ghazipur
Lucknow
Srinagar
Calcutta

20) Schedule X is related to (1)

List of equipments
Conduct of clinical trials
List of Generic drugs
None of the above

Long Answers

Answer all the questions.

6 Q x 5 marks = 30 marks

1) List out the provisions of Education Regulation and process for withdrawal of approval as specified by the PCI. (5)

2) Analyse the recommendations given by Chopra committee for the development of pharmaceutical sector in India. (5)

3) Explain the contents of New Drug Application (NDA). (5)

4) Explain the procedure for registration of clinical establishment under Clinical Establishment Act 2010. (5)

5)	Write a note on Central Drug Laboratory	(5)
6)	Explain in detail Schedule N as per D&C Act.	(5)

Short Answers

Answer all the questions.

10 Q x 3 mark = 30 marks

1)	What are the classes of Conditionally Exempted advertisement according to Drugs and Magic Remedies (Objectionable Advertisement) Act?	(3)
2)	Enlist any 6 functions of Animal Welfare Board of India.	(3)
3)	Provide formula for calculation of retail price based on DPCO 2013.	(3)
4)	What are the qualifications required to carry out Medical Termination of Pregnancy, according to Medical Termination of Pregnancy Act?	(3)
5)	Write a short note on pre- transfusion sample testing in a blood bank	(3)
6)	Explain the categories of colour code of collecting Bio-Medical Waste (BMW).	(3)
7)	Write a short note on documentation purchase record of drugs in community pharmacy.	(3)
8)	Give offenses and penalties under the import of drugs as per D&C Act	(3)
9)	Explain the qualifications and duties of Drug Inspector.	(3)
10)	Define Narcotic drugs and Psychotropic substances as Per NDPS Act.	(3)

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