

24/03/25

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10  
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Duration: 3 Hours

Total Marks: 75

- N.B. :** 1. All questions are compulsory  
2. Figures to right indicate full marks

**Q. 1** Choose the appropriate option for following multiple choice based questions. **20**  
(Write the correct option and the correct answer.)

- 1 Drug consisting of filthy and putrid substance are called \_\_\_\_\_ drug
- Spurious
  - Misbranded
  - Adulterated
  - Proprietary
- 2 Cosmetic Colors are included in the schedule.
- T
  - M
  - Q
  - X
- 3 The application to import drug as part of bonafide luggage is made to the licensing authority in....
- Form 12A
  - Form 122
  - Form 129B
  - Form 26
- 4 Schedule II of D and C Act states
- List of Ayurvedic Siddha and Unani books
  - Fees for test or analysis by CDL or Government Analyst
  - List of diseases and ailments which a drug may not claim to prevent or cure
  - Standards to be complied with by imported drugs and by drugs manufactured for sale, sold, stocked or exhibited for sale or distributed
- 5 Which authority would you apply for license for manufacture of large volume parenterals for manufacture or sale or distribution of drugs
- Government Analyst
  - Drugs controller of India
  - Drugs consultative committee
  - Drug Technical Advisory Board
- 6 Schedule B of the D and C rules stand for
- List of forms used for making applications for issuing licences, granting licences, sending memorandums.
  - Standards for ophthalmic preparations
  - Fees for test or analysis by CDL or Government analyst
  - Biological and special products

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- 7 Following labeling direction must be mentioned on toothpaste containing fluoride
- Fluoride content of Toothpaste is not more than 100 units
  - Fluoride content of Toothpaste is not more than 1000 ppm
  - Fluoride content of Toothpaste is not more than 1500 ppm
  - Fluoride content of Toothpaste is not more than 20 ppm
- 8 \_\_\_\_\_ is advisory to the central government in matters concerning administration of the D and C act.
- Drugs Consultative committee
  - Central Drugs Laboratory
  - Customs collectors
  - Government Analyst
- 9 \_\_\_\_\_ is Chairman of DTAB.
- Drug Controller of India
  - Director Central Drug Laboratory
  - Director of Central drug research institute
  - Director General of Health Services
- 10 The name of Registered Pharmacist can be removed from the register, if
- His name was entered by error in the register
  - His name is recommended to enter by another person
  - He was not aware about it
  - Anytime it can be removed without specifying the reason
- 11 Name the medicinal preparations which are considered as capable of being misused as ordinary alcoholic beverages.
- |                             |                           |
|-----------------------------|---------------------------|
| a. Toilet preparations      | b. Restricted preparation |
| c. Unrestricted preparation | d. Misbranded drugs       |
- 12 Cannabis means
- Charas
  - Opium
  - Medicinal Preparation
  - Manufactured drug
- 13 Find the odd one out:
- Advertisement published by Government
  - Displayed signboards by Registered Practitioners
  - Advertisement of drugs in medical, pharmaceutical scientific-technical journals
  - Advertisement of Magic remedy
- 14 Under the Prevention of Cruelty to Animals Act, which of the following is considered an act of cruelty?
- Providing food and water to animals
  - Inflicting unnecessary pain or suffering on an animal
  - Adopting an animal from a shelter
  - Training animals for competitions

- 15 \_\_\_\_\_ What is the % margin for retailers as per DPCO?  
 a. 06  
 b. 16  
 c. 26  
 d. 36
- 16 \_\_\_\_\_ was chairman of the Drugs Enquiry committee established in 1931.  
 a. Lt. col. Chopra  
 b. Jaisukhlal Haathi  
 c. Dr. B. Mukharjee  
 d. Mr. Bhor
- 17 What must be obtained from the woman seeking an abortion under the MTP Act?  
 a. Written consent  
 b. Verbal consent  
 c. Consent from her family  
 d. No consent is required
- 18 Who can file an RTI application?  
 a. Only Indian citizens  
 b. Foreign nationals  
 c. Corporate entities  
 d. NGOs only
- 19 Which of the following best describes a trademark?  
 a. A design or symbol representing a product or service  
 b. A legal document proving ownership of a property  
 c. A type of patent for inventions  
 d. A copyright for artistic works
- 20 What is the long form of TRIPS?  
 a. Trade-Related organization of Intellectual Property Rights  
 b. Trade-Related audits of Intellectual Property Rights  
 c. Trade-Related Acts of Intellectual Property Rights  
 d. Trade-Related Aspects of Intellectual Property Rights

Q. 2 Answer any two of the following

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- I. a) Define 'Government analyst' as per D and C Act. Give the functions and qualifications for appointment of a person as government analyst. Give the procedure followed by government analysts on receipt of samples for analysis. 6  
 b) Discuss composition and functions of PCI as per Pharmacy Act 4
- II a) Define 'Drug' as per D and C Act 1940. Enlist classes of drugs which are prohibited for manufacture and sale. 6  
 b) State the objectives of RTI. What are the obligations of public authorities towards the Right to information. 4

III	a) Define 'Coca Leaf' under NDPS Act. What are the operations controlled under the NDPS act? Enlist any two offenses under NDPS Act	6
	b) Elaborate on Labeling of ophthalmic solution, suspension and ointment preparations as per D and C act 1940	4
Q. 3	Answer any seven of the following	35
I	Discuss the ethics pharmacists should follow in relation to his/her trade and in relation to the medical profession	5
II	Differentiate between restricted and non-restricted preparation .Explain in detail about bonded manufactory	5
III	What are the provisions for import of drugs for personal use and give examples of any two drugs prohibited for import as per D and C Act.	5
IV	Define 'Scheduled formulation' and explain calculation of ceiling price of scheduled formulation as per DPCO 2013.	5
V	a) Describe the functions of Institutional animal ethics	2.5
	b) Under what circumstances pregnancy can be terminated medically as per MTP Act	2.5
VI	Write a note on Schedule N.	5
VII	Define 'Advertisement'. Elaborate on classes of exempted advertisements as stated by DMR(OA) Act 1954	5
VIII	Explain what you mean by patentable invention. Explain briefly the filling of a patent for an invention.	5
IX	Elaborate on powers and duties of Drug inspector	5