

**PARUL UNIVERSITY**  
**FACULTY OF PHARMACY**  
**D. Pharm., April-2018-19 Examination**

Year: 2

Subject Code: 08600204

Subject Name: Pharmaceutical Jurisprudence

Date: 22/04/2019

Time: 10:00 am to 01:00pm

Total Marks: 80

**Instructions:**

1. Figures to the right indicate maximum marks.
2. Make suitable assumptions wherever necessary.

**Q.1 Multiple Choice Questions (MCQs) (1 Mark Each)****(20)**

1. PCI is
 

a) Pharmacy committee of India	c) Central council of India
b) Pharmacy council of India	d) Both b and c
2. Standards for disinfectants are covered under,
 

a) Schedule O	c) Schedule Q
b) Schedule P	d) Schedule R
3. In DTAB, Director of Central Research Institute is,
 

a) Elected member	c) Ex-officio member
b) Nominated member	d) None of the above
4. One of the following is NOT a type of sales license.
 

a) Wholesale	c) Vendor
b) Retail	d) central drugs laboratory
5. Manufacture outside the bond is also called as,
 

a) Manufacture in bond	c) Bonded laboratories
b) Manufacture without the bond	d) Non-bonded laboratories
6. Opium can be manufacture only by,
 

a) Lambardar	c) Both a and b
b) Government opium factory	d) None of the above
7. What does the abbreviation OTC mean?
 

a) Off the counter	c) Over the cash
b) Over the counter	d) Outside the counter
8. License of import of drug is obtained on application to,
 

a) State government	c) Drug controller of India
b) Central government	d) Drug inspector
9. Which are the powers of Drug Inspectors?
 

a) To inspect the manufacture premises	c) Can stop vehicles and search for any wrong doing
b) To take samples of any drug or cosmetics	d) All of the above
10. \_\_\_\_\_ is given to applicant who does not have factory for manufacturing but wishes to avail the facilities owned by another license.
 

a) Loan license	c) Sales license
b) Repackaging license	d) Retail sale license
11. "Not for Human use – for treatment of animal only" is the label for \_\_\_\_\_ products.
 

a) Veterinary	c) Biological
b) Surgical dressings	d) Ophthalmic
12. For a scheduled formulation, the manufacturer selling price is,
 

a) R.P-20%	c) R.P-12%
b) R.P-16%	d) R.P-10%
13. The price of opium fixed by,
 

a) executive committee	c) state government
b) central government	d) none of above
14. What is the main objective of Drug and Magic Remedies Act 1954?

- a) To control marketing of the drugs  
b) Both a and b
- c) To control the advertisement of the drugs  
d) None of the above
15. M.A.P.E stands for  
a) Minimum allowable packing expenses  
b) Minimum allowable purchasing expenses  
c) Maximum allowable post-manufacturing expenses  
d) Maximum allowable pre- manufacturing expenses
16. DCC  
a) Drugs control committee.  
b) Drugs commercial committee.  
c) Drugs consultative committee.  
d) None of the above.
17. “Warning: To be sold by retail on the prescription of a Registered Medical Practitioner only” is for \_\_\_\_\_ Schedule.  
a) Y  
b) H  
c) X  
d) G
18. What are the functions of State Pharmacy council?  
a) To register pharmacists in the state  
b) To regulate pharmacy practice in India  
c) Inspection  
d) Both a and c
19. The Drug and Cosmetics Rules was passed in year,  
a) 1940  
b) 1945  
c) 1970  
d) 1947
20. Admission to the first year D.pharm program in any government college is based on \_\_\_\_\_ exam.  
a) Secondary  
b) Higher secondary  
c) Primary  
d) Graduation

**Q.2 Long Answers (any 8 out of 10) (05 Mark Each)**

**(40)**

1. What is AICTE? Write its role in education.
2. Discuss the constitution and function of drug technical advisory board (DTAB).
3. What are the qualifications for appointment of government analyst? Explain the duties of government analyst.
4. Write a note on Schedule X and M.
5. Explain the power, duties and responsibility of excise officer.
6. Enlist the types of sales license. Describe in detail about the loan license.
7. What are the requirements for bonded laboratories?
8. Which classes of drugs are prohibited to be imported in India?
9. Write note on pregnancy act.
10. Write a note on various offences and penalties under the Narcotic drugs and Psychotropic substance Act, 1985.

**Q.3 Short Answers (2 Mark Each){ Answer any 10}**

**(20)**

1. Explain the following terms:  
a) Adulterated drugs    b) Misbranded drugs
2. Write the functions of PCI.
3. Write a note on poison act.
4. Write a note on Drug price control order.
5. What are the objectives of Drugs and Magic remedies act?
6. Define the following terms:  
a) Drugs    b) Cosmetics
7. Explain the different labeling pattern of medicines.
8. What types of drugs without license are imported under the D&C Act 1940?
9. Enumerate various offences and penalties under the pharmacy Act 1948.
10. A brief account on the principles of pharmacist as mentioned in the Code of pharmaceutical Ethics.
- 11 Write a note on Pharmacy as a health care system.
- 12 Write a note on Central Drug laboratory.