PHARMACEUTICAL JURISPRUDENCE(BP505T)-17/07/2020

B.Pharm First Internal Examination 2020-21
Date:17/07/2020
Semester:5th
Subject:Pharmaceutical Jurisprudence(BP505T)
Instructions:
1.All questions are compulsory
2.Each questions carry 1 Mark

* Required

Email address *

Your email

ENROLLMENT NO. *

Your answer

1.Schedule F(ii)prescribes the standards for_____.

- ◯ a.Opthalmic preparations
- ◯ b.Cosmetics
- ◯ c.Umbilicaltaps
- ◯ d.None

2.Schedule H prescribes _____.

- a.the list of drugs to be sold by retail only on the prescription of RMP
- b.the list of drugs to be taken only under the supervision of medical practitioner
- c.the list of narcotic drugs and psychotropic substances
- d.the list of diseases which drugs may not claim to cure

3.Requirements and guidelines on the clinical trials for the import and manufacture of new drug is specified in_____.

- a.Schedule D
- b.Schedule X
- c.Schedule K
- d.Schedule Y

4.As per drugs and cosmetic act ,What is schedule S?

- a. Standards of poisons
- b. Standards for cosmetics
- c. Standards for medical devices
- d. Standards for ophthalmic preparation

5.Standards of patent and proprietary medicine is specified in_____.

- a.Schedule M
- b.Schedule Q
- c.Schedule V
- d.Schedule T

6.Schedule B is related with_____.

- a.Forms
- b.GMP
- c.Fees
- d. Equipment

7.List of permitted coaltar colors for use in cosmetics is specified in _____.

- a.Schedule X
- b.Schedule Q
- c.Schedule S
- d.Schedule T

8.Standards of quality in respect to drugs means the drug comply the standards as specified in_____.

- a. Second Schedule
- b. Schedule S
- c. First Schedule
- d. None

9. _____ drugs are manufactured in accordance with the formula described in the first schedule.

- a. Standard quality
- b. Ayurvedic, Siddha and Unani
- c. Homoeopathic
- d. Allopathic

10.Cosmetic includes the following articles except_____.

- a.Soap
- b.Facepowder
- c.Nail Polish
- d.Lipstick

11.Drug shall be deemed to be adulterated except if it_____.

- a.Contains different colors other than those prescribed.
- b. Is imported under a name that belong to another drug.
- c.Contains filthy,putrid or decomposed substances.
- d.All

12.Drug store means a licensed premises for the sale of drugs that_____.

- a.Requires the services of qualified person and where drugs are not compounded against the prescription
- b. Do not require the services of qualified person and where drugs are not compounded against the prescription
- c. Do not require the services of qualified person and where drugs are compounded against the prescription
- d.None

13.Drugs Technical Advisory Board is purely_____body.

- a.Administrative
- b.Analytical
- c.Both a and b
- d.None

14.Drugs Consultative Committee is constituted by_____.

- a.DTAB
- b.Central Council
- c.State Government
- d.Central Government

15.Government Analyst is appointed by Central Government or State Government under section _____in relation to Ayurvedic, Siddha and Unani systems of medicine.

- a.20
- b.21
- c.33F
- d.21A

16.Inspection book for the repacking of drugs shall be maintained in Form_____

- a.35
- b.24B
- c.25B
- d.25A

17.Loan licence can be granted for the manufacture of drugs specified in Schedule_____.

- a.CandC(i)
- b.X
- c. other than C, C(i) and X
- d.none

18.In pharmacy, a separate cupboard should be provided for storage of_____.

- a.Poisons
- b.Cosmetics
- c.Patent and proprietary medicine
- d.Ayurvedic, Siddha and Unani drugs

19. Prescriptions containing Schedule _____drugs should not be dispensed more than once unless prescriber given the directions to do so.

- a. G
- b. W
- c. H and X
- d. C and C(i).

20.No_____drugs should be supplied as physician sample.

- a. Schedule H
- b. Schedule G
- c. Schedule C and C(i)
- d. Schedule X.

21.The symbol on left top corner in red on the label is XRx denotes

- a. Schedule M
- b. Schedule X
- c. Schedule X1
- d. Schedule H

22.Which indication must be prescribed on the label of ophthalmic ointments.

- a. Use within 1 month of opening
- b. It is dangerous to take this preparation except under medical supervision
- c. Consult to Physician if irritation persist, discontinue the use
- d. Not for hurnan use

23.Schedule M(i)prescribes the requirements of factory premises, plant, equipment's, etc for the manufacture of_____.

- a.Homoeopathic medicine
- b.Cosmetics
- c.Drugs
- d.Medical devices

24. PCI is reconstituted at every

- a. 1 year
- b. 2 year
- c. 3 year
- d. 5 year

25.Example of Schedule C1 drug is

- a. Fish liver oil
- b.Hydroxyurea
- c. Pyrimidine
- d.Alprazolam

26.Central drug laboratory is located at_____.

- a. Mumbai
- b. Lacknow
- c. Calcutta
- d. Hyderabad

27.Specify the form number required for licence for retail sale of schedule C and C1 drug.

- a. 22
- b. 21
- c. 19
- d. 18

28.The First Schedule to the Drugs and Cosmetics Act, 1940 prescribes_____.

- a. standards for cosmetics
- b. standards for medical devices
- c. authoritative books of Ayurvedic, Sidhha and Unani system
- d. standards of the drugs to be complied with by imported drugs

29.Who among the following is the Chairman of Drug Technical Advisory Board?

- a. The Drugs Controller of India
- b. The President of Pharmacy Council of India
- c. The President of Medical Council of India
- d. The Director General of Health Services

30.A drug not labelled in the prescribed manner shall be treated as _____.

- a. adulterated drug
- b. spurious drug
- c. misbranded drug
- d. mischievous drug

31.Powers of the Inspectors appointed under the Drugs and Cosmetics Act, 1940 are mentioned under which of the following Sections of the Act?

- a. 20
- b. 21
- c. 22
- d. 23

32.Schedule F to the Drugs and Cosmetics Rules, 1945 prescribes the_____.

- a. requirements for the functioning and operation of a blood bank
- b. standards for surgical dressings
- c. list of drugs to be prescribed
- d. standards for disinfectant fluids

33.Schedule FF to the Drugs and Cosmetics Rules, 1945 contains the list of_____.

- a. drugs which can be marked under generic names only
- b. drugs which are habit forming
- c. standards for ophthalmic preparation
- d. drugs which are exempted from certain provisions applicable to manufacturing

34.Example of Schedule C drug is

- a. fish liver oil
- b. digitalis drugs
- c. ergot
- d. Amphetamines

35.Drug and cosmetic act was passed in _____.

- a. 1940
- b. 1942
- c. 1945
- d. 1840

36. Misbranded drug is_____.

- a. If it is so colored, coated, powdered or polished that damage is concealed, or if it is made to appear of better or greater therapeutic value than it really is
- b. if it is an imitation of, or a substitute for, another drug or resembles another drug in a manner likely to deceive or bears upon it or upon its label or container the name of another drug
- c. if it is imported under a name which belongs to another drug
- d. None

37. What is the meaning of DTAB?

- a. Drugs Technical Advisory British rule
- b. Drugs Technical Action Board
- c. Drugs Testing Advisory Board
- d. None

38.What is the meaning of DCC?

- a. Drugs Consultative Committee
- b. Drugs Consultative Control
- c. Drugs Center of Committee
- d. None

39. What is import?

- a. To bring something out of india
- b. To send something out of india
- c. To export cosmetics
- d. None

40.Schedule C?

- a. Deal with biological product
- b. Deal with ophthalmic product
- c. both of above
- d. None

41.Schedule B?

- a. List of forms for test or analysis by CDL or Govt. analysts
- b. Performa for forms (Application, issue, renewal, etc.)
- c. List of drugs that are exempted from provisions of import
- d. None

42.Schedule D?

- a. Rates of fee for test or analysis by CDL or Govt. analysts
- b. Performa for forms (Application, issue, renewal, etc.)
- c. List of drugs that are exempted from provisions of import
- d. None

43.Provisions applicable to blood bank is _____.

- a. Schedule E1
- b. Schedule H
- c. Schedule O
- d. None

44.Schedule G?

- a. Rates of fee for test or analysis by CDL or Govt. analysts
- b. Performa for forms (Application, issue, renewal, etc.)
- c. List of drugs that are exempted from provisions of import
- d. None

45.Classes of drugs prohibited to import?

- a. Misbranded drugs,Adulterated drugs,Substandard drugs
- b. Drugs of substandard quality,Standard drugs, Quality drugs
- c. Adulterated drugs,Spurious drugs, Standard drugs
- d. All

46.Licence for wholesaler if drugs specified in schedule C and C1 is issued in the form_____.

- a. 21B
- b. 21BB
- c. 20B
- d. 20A

47. List of minimum equipment to run a Pharmacy is_____.

- a. schedule N
- b. schedule F
- c. schedule J
- d. schedule R

48. Standards for disinfectant fluids is _____.

- a. Schedule P
- b. Schedule M
- c. Schedule y
- d. Schedule O

49.Life period of drug is _____.

- a. Schedule P1
- b. Schedule M
- c. Schedule Y
- d. None

50.List of substances required to be used under medical supervision and labelled accordingly is_____.

- a. Schedule X
- b. Schedule H
- c. Schedule G1
- d. Schedule G

51. Standard to be complied with by imported drugs and by drugs manufactured for sale, sold, stocked or exhibited for sale or distribution is _____.

- a. Schedule N
- b. Schedule S
- c. Schedule Q
- d. None

52. Standards for medical devices is_____.

- a. Schedule R2
- b. Schedule S1
- c. Schedule R
- d. Schedule R1

53. Type of sale of drugs?

- a. Manufacturer label
- b. Dispensing label
- c. Retail and Wholesale
- d. None

54. What is Adulterated drug?

- a. If it is not consisting, in whole or in part, of any filthy, putrid or decomposed substance
- b. If it has been prepared, packed or stored under sanitary conditions whereby it may have been contaminated with filth or whereby it may have been rendered injurious to health
- c. If its container is composed in whole or in part, of any poisonous or deleterious substance which may render the contents injurious to health
- d. All

55.List of drugs marked under generic name_____.

- a. Schedule X
- b. Schedule E
- c. Schedule Y
- d. Schedule W

56. What is Spurious drugs?

- a. if it is imported under a name which not belongs to another drug
- b. if it is an imitation of, or a substitute for, another drug or resembles another drug in a manner likely to deceive or bears upon it or upon its label or container the name of another drug
- c. Both a and b
- d. if it is exported under a name which belongs to another drug

57.Drug inspector is appointed under the act of?

- a. 19 of IPC
- b. 21 of IPC
- c. 42 of IPC
- d. 48 of IPC

58. GMP requirement of factory premises,plants and equipments_____,

- a. Schedule–M
- b. Schedule-E1
- c. Schedule-Y
- d. Schedule -D

59. The minimum qualification for registration as pharmacist is_____.

- a. BSc.
- b. Pharm D
- c. B.pharmacy
- d. D.pharmacy

60.Given the following are schedule X drugs except:

- a. Amobarbital
- b. Metformin
- c. Amphetamine
- d. Glutethimide

61.One of the forms mentioned below is used to issue licence for wholesale of drugs other than specified in schedule C,Cl and X.

- a. Form 20C
- b. Form21 B
- c. Form 20BB
- d. Form 20 A

62.The drug samples taken by drug inspector for analysis are sending to?

- a. Drug Controller
- b. Govt. Analyst
- c. Chemical Analyst
- d. Testing Laboratories

63. Standards for mechanical contraceptive are given in schedule?

- a. Schedule S
- b. Schedule Q
- c. Schedule R
- d. Schedule W

64. Insulin injection comes under the schedule_____.

- a. G
- b. C
- c. C1
- d. H1

65.For registration of pharmacist in the various states, the pharmacy act provides for the constitution of_____.

- a. Central pharmacy Council
- b. Registration Tribunals
- c. Corporative Societies
- d. State Pharmacy Council

66.Loan licenses are issued for:

- a. Drugs specified in Schedule-C/C1
- b. Drugs other than specified in C/C1 & X.
- c. Both a and b
- d. None

67.Repackaging of drugs is granted of_____.

- a. Drugs other than Schedule-C/C1 and X.
- b. Drugs specified in Schedule-C/C1
- c. Both a and b
- d. None

68.Administration of the D&C act 1940 and rules 1945 the executive body contain_____.

- a. Controlling authorities
- b. Government Analysts
- c. Both a and b
- d. None

69.Which form number should be used for the application to import drugs for personal use.

- a. 24C
- b. 8
- c. 19
- d. 12A

70.Narcotics drugs and psychotropic substances act was passed in_____.

- a. 1945
- b. 1955
- c. 1985
- d. 1995

Submit