



Name :

Roll No. :

Invigilator's Signature :

CS/B.Pharm/SEPARATE SUPPLE/SEM-8/PT-813/2011

2011

PHARMACEUTICAL JURISPRUDENCE AND ETHICS

Time Allotted : 3 Hours

Full Marks : 70

The figures in the margin indicate full marks.

*Candidates are required to give their answers in their own words
as far as practicable.*

GROUP – A

(Multiple Choice Type Questions)

1. Choose the correct alternatives for any *ten* of the following :

10 × 1 = 10

- i) The AICTE Act came into force w.e.f.
- a) 26th March, 1988 b) 28th March, 1989
- c) 28th March, 1988 d) 26th March, 1979.
- ii) The headquarters of AICTE is in
- a) Delhi b) Mumbai
- c) Kolkata d) None of these.
- iii) The medicinal and toilet preparation Rules were passed
in
- a) 1955 b) 1956
- c) 1957 d) 1958.

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- iv) According to MTP Act Alcohol means
- a) 50% methyl alcohol b) methyl alcohol
c) 50% ethyl alcohol d) ethyl alcohol.
- v) The pharmacy bill was introduced in
- a) 1948 b) 1949
c) 1945 d) 1946.
- vi) Antiserums and toxoids are tested at
- a) Chennai b) Lucknow
c) Izatnagar d) Delhi.
- vii) In case of schedule X drug, the prescription should be in
- a) duplicate b) triplicate
c) only one copy d) five copies.
- viii) Licence to sell drug specified C & C1 is given in form no.
- a) 19 b) 18
c) 21 d) 24.
- ix) The chairman of DTAB is
- a) President, PCI
b) Drug controller of India
c) Union Health Minister
d) Director General Health Service.
- x) In which year the Central Register of Pharmacist is maintained by the PCI as per as the provision of Pharmacy Act ?
- a) 1940 b) 1945
c) 1976 d) 1984.

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xi) The requirements and guidelines for clinical trials comes under which of the following schedules ?

- a) Schedule K b) Schedule V
c) Schedule Y d) Schedule U.

GROUP – B

(Short Answer Type Questions)

Answer any *three* of the following. 3 × 5 = 15

2. Write a short note on "Prevention of Cruelty to Animals Act".
3. Write about the ethics & code of conduct which a Pharmacist should follow according to the norms of Pharmacy Act.
4. What do you mean by the term "Misbranded" & "Spurious" drugs ?
5. Describe briefly the constitution of State Pharmacy Council.
6. Write briefly on Insecticides Act.

GROUP – C

(Long Answer Type Questions)

Answer any *three* of the following. 3 × 15 = 45

7. Give the objectives behind the Medical Termination of Pregnancy Act, 1971. Describe the circumstances under which the pregnancies may be terminated according to this act.

Give the offences and penalties of MTP Act, 1971.

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8. Give some basic differences between manufacture of alcoholic preparations in bond and outside bond.

Write the design and construction of bonded laboratory and non-bonded laboratory.

9. Define the following :

- a) Adulterated drugs
- b) Misbranded drugs
- c) Spurious drugs
- d) GMP
- e) Drug consultative committee.

10. Write a brief note about the ethics and code of conduct a pharmacist should follow according to the norms of the Pharmacy Act.

Define the term Advertisement and magic remedy.

Write in details about import, export and transshipment of narcotic drugs and Psychotropic Substances Act, 1985.

11. a) Give an account on "Education Regulation".
b) Mention the ex-officio members of DTAB.
c) Discuss the constitution of PCI. 3 + 4 + 8
12. a) What are qualification of Government Analyst ?
b) Discuss the procedure of inspection of Drug Inspector.
c) Discuss the manner of labelling with a special reference to schedule-H & schedule-X drugs. 4 + 6 + 5