

DO NOT WRITE ANYTHING ON YOUR QUESTION PAPER EXCEPT YOUR ROLL NO.
QUESTION PAPER CONTAINING ANYTHING WOULD BE TREATED AS MALPRACTICE
ANSWER THE QUESTIONS SERIALY AND CONTINUOUSLY

Subject: PHARMACY LAW & ETHICS (Theory)

Full Mark -80

Time -3 Hrs.

1. Answer any six (6x5)
- Write the objectives and salient features of Pharmacy Practice Regulation 2015.
 - Discuss briefly about offences & penalties of the NDPS Act 1985.
 - Write down the objectives and registration of medical devices.
 - Write down the principles of good regulatory practices.
 - Write about the registration of clinical establishments.
 - What are the requirements for the functioning & operation of a blood bank?
 - How does the state disaster management authority get established?
2. Answer any ten (3x10)
- Write a short note on DCC.
 - What are the objectives of the Pharmacy Act 1948?
 - Distinguish between community pharmacy license & hospital pharmacy license.
 - Write a short note on CDL.
 - Write down the principles of Bioethics.
 - What are the salient features of the Prevention of Cruelty to Animals Act 1960?
 - Write down the offenses & penalties for manufacturing of Spurious drugs.
 - Write down different steps involved in the manufacturing of dietary supplements.
 - What are the prerequisite conditions required for the grant of Repacking license?
 - What are the objectives of the D&C Act 1940?
 - Differentiate between trademark & patent.
3. Answer all questions: (1x20)
- | | | | |
|---------------------|-----------------|-----------------|---------------------------|
| i) Biomedical waste | ii) Novel foods | iii) IAEC | iv) Apprentice pharmacist |
| v) SUGAM | vi) ANDA | vii) Schedule-Y | viii) Contract research |
| ix) Bioethics | x) Schedule-F | xi) Lunatic | xii) Controlled substance |
| xiii) Patent | xiv) NDRF | xv) CDSCO | xvi) Cord blood centre |
| xvii) Phooka | xviii) CPCSEA | xix) FSMP | xx) Education regulation |

