[Total No. of Questions - 13] [Total No. of Printed Pages - 2] DEC-23-0099

BP-806 ET (Quality Control and Standardization of Herbals)

B.Pharm-8th (PCI)

Time: 3 Hours

Max. Marks: 75

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book (40 pages) issued to them and no supplementary/continuation sheet will be issued. The candidates shall limit their answers precisely within the answer-

Note: The question paper contains three sections. Section A is compulsory. Attempt any 2 questions from Section B and any 7 questions from Section C.

SECTION-A

(10×2=20)

(COMPULSORY SECTION)

Give short answers:

- What is a crude extract and a standardized extract?
- Mention 2 methods that are used to identify crude drugs
- What is the difference between GACP and GAP?
- Give 2 differences between HPLC and HPTLC.
- What are markers in plant extracts?
- Name the plant drug regulatory bodies in India
- φ Why do we need pharmacovigilance of herbal drugs?
- during cultivation. Mention 4 factors that effect quality of medicinal plants
- Define plant drug standardization.
- Give the conditions for the long term stability studies of a herbal product.

 $(2 \times 10 = 20)$

SECTION-B

BP-806 ET

Long Answer (Any Two)

- Ņ standardization of herbal drugs. Describe the need and processes involved in the
- traditional medicines. Describe evaluation of efficacy and safety of herbal drugs and
- What are different Chromatographic techniques? What are the various applications of these techniques in pharmacognosy?

SECTION-C

Short Note Answer (Any Seven)

- Ç Describe the regulation of herbal drugs in India
- for plant drugs. Compare the WHO, IP and API pharmacopoeial specifications
- application. Describe the requirements and steps involved for new drug
- œ Describe how will you determine the stability of a plant product?
- 9 What is the role of markers in plant drug standardization?
- Describe process for clinical trials of traditional plant drugs.
- Describe the role of microscopy in plant drug standardization.
- 12 You are given a traditional drug. How will you study its efficacy?
- 13. What are the essential parts of a GACP MONOGRAPH?