

Question Paper

Exam Date & Time: 08-Jul-2023 (10:00 AM - 01:00 PM)



MANIPAL ACADEMY OF HIGHER EDUCATION

Manipal College of Pharmaceutical Sciences

MAHE, Manipal

B.Pharm Semester VIII , End Sem Examination 2023

Quality Control and Standardization of Herbals [PCO-BP806ET-S3]

Marks: 75

Duration: 180 mins.

I Multiple Choice Questions (MCQs)

Answer all the questions.

Section Duration: 30 mins

- 1) Crude drug containing abundant mucilage and answers Ruthenium red test (1)
- [Clove](#)
[Linseed](#)
[Lemon](#)
[grass](#)
[Liquorice](#)
- 2) Extractive value is used (1)
- [To identify the presence of inorganic matter](#)
[To identify exhausted drugs](#)
[To identify impurities](#)
[To identify pesticides](#)
- 3) Bitter drugs used to (1)
- [Thirst](#)
[Urination](#)
[Induce immunity](#)
[Increase appetite](#)
- 4) Foreign matter in whole or cut medicinal plant materials should be (1)
- [NMT 2%](#)
[NMT 8%](#)
[NMT 10%](#)
[NMT 15%](#)
- 5) Which of the following method is used for determination of tannins? (1)
- [Lycopodium spore method](#)
[Hide powder method](#)
[LOD](#)
[None of the above](#)
- 6) Standard of self-generated alcohol for Asavas & Arishtas prescribed in (1)

[Rule 155](#)
[Rule 156](#)
[Rule 157](#)
[Rule 158](#)

7) Column Chromatography is used for (1)

[Identification of phytoconstituents](#)
[Quantification of phytoconstituents](#)
[Purification of phytoconstituents](#)
[All of the above](#)

8) Indian Herbal Pharmacopoeia was published in the year (1)

[1992](#)
[2002](#)
[2010](#)
[1975](#)

9) Spurious drugs has been defined and is under the section (1)

[33EEA](#)
[33EEB](#)
[33EEC](#)
[33EED](#)

10) Which of the following is "NOT" a part of protocol preparation? (1)

[Literature survey](#)
[Technology transfer](#)
[Working hypothesis](#)
[Ethical aspects](#)

11) Database related to herbal medicine and natural products are "NOT" established in which of the following countries? (1)

[China](#)
[Japan](#)
[United States](#)
[Bangladesh](#)

12) Which of the following professions come under healthcare? (1)

[Pharmacy](#)
[Judicial](#)
[Railway](#)
[Postal](#)

13) Ratio of herbal substance to genuine preparation is called (1)

[DAR](#)
[DUM](#)
[DER](#)
[DEP](#)

14) Active markers are responsible for which of the following? (1)

[Therapeutic](#)
[Analytical](#)

[Aeronautical](#)

[Biomedical](#)

- 15) Propagation materials in GACP should be free from contamination and tolerant to the following abiotic factor (1)

[Bacteria](#)

[Fungus](#)

[Animals](#)

-

[Minerals](#)

- 16) The ultimate goal of GMP is (1)

[The manufacturing process is as has been prescribed to maintain the standards](#)

[Safeguarding the health of the patient as well as producing good quality medicine](#)

[The manufactured drug which is released for sale is acceptable quality](#)

[All of the above](#)

- 17) Scientific name of the plant is recorded in the following order (1)

[Genus, species, author, family](#)

[Species, genus, author,](#)

[family](#)

[Author, genus, species, family](#)

[Family, genus, species, author](#)

- 18) National GLP compliance monitoring authority was established by DST India in the year (1)

[1972](#)

[1970](#)

[2002](#)

[2010](#)

- 19) The finished goods transferred from the production area after proper packing shall be stored in the finished goods stores, marked as (1)

[Finished goods stock](#)

[Quarantine](#)

[Finished product](#)

[None](#)

- 20) Fertilizers should be applied sparingly to minimize (1)

[Erosion](#)

[Leaching](#)

[Tillage](#)

[Weeds](#)

II Long Answers

Answer all the questions.

- 1) Explain the following; (10)
Bitterness value (3M)
Foaming Index (3M)
Determination of pesticide residue (4M)
- 2) Describe the comparison of various Herbal Pharmacopoeia (10)

III Short Answers

Answer all the questions.

- 1) Explain the Issues in Stability Testing of Herbal Products (5)
- 2) With the help of neat labelled diagram explain Gas liquid chromatography as tool for standardization of herbal drugs. (5)
- 3) What are the functions of National Pharmacovigilance centre and Uppsala monitoring centre (5)
- 4) Discuss briefly the control tests for herbal raw materials, preparations and medicinal products (5)
- 5) Briefly explain about the objectives for non-clinical studies and general considerations of pharmacological investigations (5)
- 6) Explain the SOPs for the careful cultivation of medicinal plants. (5)
- 7) What are Good Manufacturing Practices? State its objectives and significance (5)

-----End-----