

Question Paper

Exam Date & Time: 20-May-2024 (10:00 AM - 01:00 PM)



MANIPAL ACADEMY OF HIGHER EDUCATION

Quality Control and Standardization of Herbals [PCO-BP806ET-S1]

Marks: 75

Duration: 180 mins.

I Multiple Choice Questions (MCQs)

Answer all the questions.

Section Duration: 30 mins

- 1) 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
- 2) Fresh plant materials should be ideally stored as per GMP at (1)
- 2 - 8 °C
10 - 15
°C
20 - 30
°C
35 - 40
°C
- 3) Fertilizers should be applied sparingly to minimize (1)
- Erosion
Leaching
Tillage
Weeds
- 4) The domesticated, locally adapted, distinct identity of a traditional variety of species is known as (1)
- Ecotype
Chemotype
Phenotype
Landraces
- 5) National GLP compliance monitoring authority was established by DST India in the year (1)
- 1972
1970
2002
2010
- 6) One of the following is NOT useful for the Soil conservation (1)
- Crop rotation
Inter cropping

- Contour farming
Tillage
- 7) Shinoda test is performed to detect (1)
Saponins
Alkaloids
Carbohydrates
Flavonoids
- 8) Hemolytic activity is positive for which of the following (1)
Tannins
Cellulose
Flavonoids
Saponins
- 9) Rancidity of fixed oil is determined by (1)
High Ash value
High Acid value
High Extractive value
Low Ash value
- 10) Presence of Steroids in the extract is identified by (1)
Hager's test
Liebermann-Burchard test
Dragendorff test
Halphen's test
- 11) "Isoquinoline" containing chemical constituents (1)
Ephedra
Opium
Nux-vomica
Datura
- 12) Schedule T in ASU system refers to (1)
Good Manufacturing Practice
Good Agricultural Practice
Good Clinical Practice
Good storage practice
- 13) Limit of Heavy Metal Arsenic for exports as per ASU standard is (1)
0.3 ppm
3.0 ppm
0.5 ppm
1 ppm
- 14) Acceptance criteria indicates _____ limits in quality control of herbal drugs (1)
Alphabetical
Numerical
Therapeutic
Chemotaxonomy

15) Which of the following is NOT included in Herbal preparations? (1)

- [Tinctures](#)
- [Extracts](#)
- [Isolated compounds](#)
- [Expressed juices](#)

16) Tests, procedures, acceptance criteria, and limits are the part of (1)

- [Specifications](#)
- [Standardization](#)
- [Markers](#)
- [Drug extract ratio](#)

17) Control of herbal substances and preparations are described in (1)

- [CIMAP](#)
- [CHMP](#)
- [NBRI](#)
- [CFTRI](#)

18) Which of the following is NOT a healthcare professions? (1)

- [Pharmacy](#)
- [Medical](#)
- [Nursing](#)
- [Engineering](#)

19) Which Schedule deals with the manufacture of the herbal medicines? (1)

- [Schedule H](#)
- [Schedule T](#)
- [Schedule P](#)
- [Schedule O](#)

20) Drugs and cosmetics act was framed in which year? (1)

- [1937](#)
- [1938](#)
- [1939](#)
- [1940](#)

II Long Answers

Answer all the questions.

- 1) Describe the following (10)
 - a) Determination of Bitterness value. (5 Marks)
 - b) Foaming Index and Swelling Index. (5 Marks)
- 2) Explain various stages for obtaining Licensing Procedure for the manufacturing of Herbal medicines (10) in India.

III Short Answers

Answer all the questions.

- 1) Describe the SOPs for the controlled cultivation of medicinal plants. (5)
- 2) Describe the GLP (NGCMA) formation, scope and certification in India. (5)
- 3) Discuss on General considerations for herbal medicine research. (5)

- 4) Briefly discuss various steps involved in Research studies. (5)
- 5) Explain HPTLC as a tool for standardization and quality control of herbal drugs. (5)
- 6) List out various challenges in stability testing of herbal drugs. (5)
- 7) Give the components of The Ayurvedic Pharmacopoeia of India and The Unani Pharmacopoeia of India. (5)

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