

(3 Hours)

Total Marks: 75

Q. 1 Attempt all multiple-choice questions (MCQ)

20M

Sr No	Questions	Options
1	As per WHO guidelines Category 2 herbal medicines are _____	a Safe under specific conditions of use b Herbal medicines of uncertain safety c Safety established by use over long time d Not safe at all
2	Fluoride ion content in Non-fluoridate toothpaste should be maximum _____ parts per million as per BIS	a 50 b 200 c 800 d 500
3	Chemical method used for analysis of raw materials in cosmetics is _____	a Saponification Value b Relative density c Sieve Analysis d pH
4	Stability testing of herbal drugs and preparations is carried out at -----	a 40°C/75 per cent RH b 30°C/65 per cent RH c 25°C/60 per cent RH d 45°C/85 per cent RH
5	1gm of crude drug was heated at 110 C for 2 Hours and the weight after heating was 0.82 gm. The moisture content of crude drug is _____ % w/w	a 82 b 18 c 0.18 d 0.82
6	RAPD in DNA fingerprinting refers to ---	a Randomly Amplified Polymorphic DNA b Randomly Associated Polymorphic DNA c Regularly Amplified Polymorphic DNA d Randomly Amplified Polymeric DNA
7	Softening point of lipstick should be minimum _____ as per BIS standards	a 65°C b 75°C c 85°C d 55°C

- 8 which of the following modern analytical instruments is used for Adulterant Screening
- a Flame photometer
 - b UPLC-TOFMS
 - c Colorimeter
 - d AAS
- 9 Iodine value of fatty material used in cosmetics indicate
- a Molecular weight
 - b Unsaturation
 - c Free acid content
 - d Hydroxyl group content
- 10 Chemical method used for analysis of raw materials in cosmetics is _____
- a Saponification Value
 - b Relative density
 - c Sieve Analysis
 - d pH
- 11 Herbal Toxicity refers to
- a the relative ability of a substance to cause adverse effects in living organisms
 - b the amount of active constituents present in the drug
 - c the amount of substance which drug yields to the solvent
 - d The amount of starch present in the drug
- 12 The temperature at which the product starts oozing out the oil and becomes flattened out is -

- a breaking load
 - b melting point
 - c drooping point
 - d breaking point
- 13 In India, most of the traditional herbs are available as
- a Dispensing medicine
 - b OTC medicines
 - c Prescription medicine
 - d Schedule T drugs

- 14 Heavy metals content in Skin cream should be maximum _____ parts per million as per BIS
- a 500
b 200
c 1000
d 20
- 15 Authentication of herbal crude drug by size & appearance is called
- a Drug Fingerprinting
b Microscopic evaluation
c Macroscopic evaluation
d Bio-prospecting
- 16 Physical method used for analysis of raw materials in cosmetics is _____
- a Ash value
b Iodine value
c Microscopy
d Animal studies
- 17 Content of lead in crude drug should not be more than ---- ppm_____
- a 2
b 8
c 5
d 10
- 18 Majority of adverse events related to the use of herbal products are due to
- a unskilled labour used in collection
b poor quality control tests done on herbals
c High artificial drying of drugs
d bad storage conditions
- 19 Safety assessment of herbal drug is carried out by
- a Toxicological studies
b Physicochemical evaluation ash values and extractive values
c Authentication
d Organoleptic evaluation
- 20 Volatile oil content in crude drug is best determined by _____
- a Counter current Apparatus
b Soxhlet Apparatus
c Clavenger Apparatus
d titrimetry

Q2. Answer any TWO of the following

20M

- i) Briefly explain the Indian standard specification laid down for testing of Skin care & personal hygiene products
- ii) Discuss the spontaneous reporting schemes & challenges for the safety monitoring of herbal medicine.
- iii) Explain different aspects of WHO and AYUSH guidelines for Herbal drug standardization

Q3. Answer any SEVEN of the following

35M

- i) Justify with suitable example how is DNA fingerprinting technique useful in quality control of crude drugs.
 - ii) Define Herbal drug-drug interaction with suitable examples. Discuss Challenges in monitoring the safety of herbal medicine.
 - iii) Justify the importance of determination of viscosity & fineness of powders in evaluation of cosmetic products.
 - iv) Enlist different methods of Analysis for raw material used in cosmetic manufacture as per BIS.
 - v) Describe in detail about stability testing protocol and techniques applicable for natural product
 - vi) Discuss regulatory guidelines for setting up Herbal drug Industry.
 - vii) Explain different parts of herbal drug monograph of USP
 - viii) Explain determination of ash value & heavy metals for evaluation of cosmetic products.
 - ix) Discuss Indian and International patent law for herbal drug & mention their protocol.
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