



PAPER ID-420748

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Subject Code: BP806ET

Roll No:

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BPHARMA
(SEM VIII) THEORY EXAMINATION 2021-22
QUALITY CONTROL AND STANDARDIZATION OF HERBAL

Time: 3 Hours

<https://pharmacyindia.co.in/>

Total Marks: 75

Note: 1. Attempt all Sections. If require any missing data; then choose suitably.

SECTION: A

1. Attempt all questions.

10 x 2 = 20

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| a. | Define processed and finished herb as per WHO. https://pharmacyindia.co.in/ |
| b. | What is chromatographic fingerprinting? |
| c. | Write a note on chemical markers with examples. |
| d. | Write briefly about authentication of medicinal plant. |
| e. | Define new drug application. |
| f. | Write the importance of GACP. https://pharmacyindia.co.in/ |
| g. | Define bitter value? |
| h. | Write any two adsorbents used in TLC. |
| i. | Define quality control and give its significance. |
| j. | Define export registration. |

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SECTION: B

2. Attempt any two parts of the following:

2 x 10 = 20

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| a. | Write a comparative note on various herbal pharmacopoeia. |
| b. | Describe the evaluation of commercial crude drugs intended for use. |
| c. | Explain briefly the current good manufacturing practices (cGMP) for herbal drugs |

SECTION: C

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3. Attempt any five parts of the following:

7 x 5 = 35

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|----|--|
| a. | Discuss stability testing methods and protocols for herbal medicines. |
| b. | Write a note on ICH guideline for quality control of herbal medicinal products |
| c. | What is standardization? Explain the application of HPTLC as a method for standardization. https://pharmacyindia.co.in/ |
| d. | Write a note on preparation of documents for new drug application |
| e. | What are the regulatory requirements of herbal medicines in India |
| f. | Explain briefly the good agriculture practices of herbal drugs |
| g. | Explain the WHO guidelines on safety monitoring of herbal medicines in pharmacovigilance systems https://pharmacyindia.co.in/ |

