



MANIPAL UNIVERSITY

FIRST YEAR M. PHARM. DEGREE EXAMINATION - MAY 2017
SUBJECT: ADVANCED PHARMACOGNOSY AND PHYTOCHEMISTRY (PCO 601T)
(SPECIALIZATION: PHARMACOGNOSY)
(2014 REGULATION)
Thursday, May 18, 2017 (10.00 - 13.00 Hrs.)

Marks: 100

Duration: 180 mins.

Answer ALL the questions.

- 1) Give a detailed account of various diseases affecting plants. (10)
- 2) Discuss the role of Adsorption Chromatography techniques used in separation of Phyto-constituents. (10)
- 3) What are Cardiac Glycosides? Describe their occurrence, chemistry, method of extraction and test for identification. (10)
- 4) Give an overview of *intrinsic* factors that control cultivation of crude drugs. (10)
- 5) Describe the source, constituents and uses of any four phyto-constituents used as Hepato-protective drugs. (10)
- 6) Elucidate the structure of Citral and Atropine. (10)
- 7) Describe the biogenesis of Tropane alkaloids. (10)
- 8) What are Terpenes? Classify them with examples? Add a note on the isolation of Terpenoids. (10)

Write short notes:

- 9A) Marine anti-inflammatory agents. (5)
- 9B) DNA Hybridization technique in Chemotaxonomy. (5)

Write briefly on the following:

- 10A) IUCN classification of endangered species with examples. (5)
- 10B) Mineral anti-oxidants. (5)



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FIRST YEAR M. PHARM. DEGREE EXAMINATION - MAY 2017
SUBJECT: HERBAL PRODUCT DEVELOPMENT AND FORMULATION (PCO 602T)
(SPECIALIZATION: PHARMACOGNOSY)
(2014 REGULATION)
Saturday, May 20, 2017 (10.00 - 13.00 Hrs.)

Marks: 100

Duration: 180 mins.

Answer ALL the questions.

- 1) Discuss pharmacovigilance for herbal drugs. (10)
- 2) Describe the effective application of spectroscopy in standardization of plant extracts. (10)
- 3) Discuss in detail the various excipients with their role used in herbal formulations. (10)
- 4) Give a detailed account of pre-processing of herbal raw materials. (10)
- 5) Enumerate with a detailed note on types of packaging and labeling of finished herbal preparations. (10)
- 6) Evaluation of herbal extracts with a special emphasis for their physical and chemical standardization parameters with suitable examples. (10)
- 7) Discuss the development of herbal formulations used in various skin diseases with their merits and demerits. (10)
- 8) Explain cGMP for the manufacture of herbal liquid dosage forms. (10)

Write short notes:

- 9A) Role of Phyto and Pharma equivalence studies. (5)
- 9B) Different types of Maceration techniques. (5)

Write briefly on the following:

- 10A) Commonly used herbs in Nutraceuticals. (5)
- 10B) Dissolution test with its significance. (5)



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FIRST YEAR M. PHARM. DEGREE EXAMINATION - MAY 2017
SUBJECT: MEDICINAL PLANT BIOTECHNOLOGY (PCO 603T)
(SPECIALIZATION: PHARMACOGNOSY)
(2014 REGULATION)
Tuesday, May 23, 2017 (10.00 - 13.00 Hrs.)

Duration: 180 mins.

Marks: 100

Answer ALL the questions.

- 1) What are protoplast cultures and give a detailed account of isolation and purification methods. (10)
- 2) Define and give an account of the factors affecting hairy root culture. Add a note on its applications. (10)
- 3) Explain cryopreservation and its impact on biomedicinals. (10)
- 4) Discuss on spontaneous genetic variation in detail. (10)
- 5) Give an account of screening methods and selection of high yielding cell lines. (10)
- 6) Give a detailed account of biotransformation in pharmacy with special reference to culture methods and precursors. (10)
- 7) Give the types, techniques and nutritional requirements for plant tissue culture. (10)
- 8) Write historical perspectives and prospects of medicinal plant biotechnology. (10)

Write short notes:

- 9A) Give the applications of cellular totipotency. (5)
- 9B) Describe the methods of gene identification. (5)

Write briefly on the following:

- 10A) Write a note on transgenic plants. (5)
- 10B) What are the general requirements for embryogenesis? (5)



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FIRST YEAR M. PHARM. DEGREE EXAMINATION - MAY 2017
SUBJECT: BIOLOGICAL SCREENING OF HERBAL DRUGS (PCO 604T)
(SPECIALIZATION: PHARMACOGNOSY)
(2014 REGULATION)

Thursday, May 25, 2017 (10.00 - 13.00 Hrs.)

Marks: 100

Duration: 180 mins.

Answer ALL the questions.

- 1) Give a detailed account of OECD guidelines for toxicity studies using staircase method. (10)
- 2) Discuss various in vitro models for evaluation of anticancer activity. (10)
- 3) What are Transgenic animals? Give the methods for their production and maintenance. (10)
- 4) Suggest screening protocol for Anti-Alzheimer's drug using appropriate in vitro and in vivo models. (10)
- 5) What do you understand by parametric data? Suggest and justify suitable statistical tests for a group of parametric data. (10)
- 6) Give an overview of Phase II clinical trial. (10)
- 7) Explain different in vivo methods applied for screening of drugs for nephroprotective activity. (10)
- 8) Give an overview of natural drug screening, explain the application of High Throughput screening in natural drug screening. Discuss its importance. (10)

Write short notes:

- 9A) Discuss the *in vitro* anti diabetic assays in brief. (5)
- 9B) DPPH anti-oxidant assay. (5)

Write briefly on the following:

- 10A) Brief outlook of CPCSEA guidelines. (5)
- 10B) One way ANNOVA. (5)