Exam Date & Time: 02-May-2019 (02:00 PM - 05:00 PM)



#### MANIPAL ACADEMY OF HIGHER EDUCATION

Advanced Instrumental Analysis [PCH-MPA201T] Duration: 180 mins. Marks: 75 **SECTION - A** Answer all the questions. Answer the following (10 marks x = 50 marks) (10)Explain immobilised polysaccharides used in Enantiomeric separation (10)Write the principle, instrumentation and specific applications of 2) LC-MS. Explain the different steps involved in HPTLC (10)3) (10)4) What is chemical shift? Explain the parameters the chemical shift depends on and also the factors that influence the chemical shift values. (10)5) Expand HETCOR. Give its general feature. List out and explain about the various techniques included in it along with spectra. **SECTION - B** Answer all the questions. Answer the following (5 marks  $\times$  5 = 25 marks) (5)Assign the PMR chemical shift values for the following compounds: a) Anisole b) Ethyl benzene c) Ethyl alcohol d) Diethyl ether 7) Explain the various steps involved in the 2D NMR spectroscopy (5)with a diagram. (5) 8) Discuss the various types of fragmentation ions produced in Mass spectroscopy and their importance (5)9) Discuss the methods for the separation of Chiral compounds Discuss the methods involved in GC derivatisation (5) 10)

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10)

Exam Date & Time: 04-May-2019 (02:00 PM - 05:00 PM)



#### MANIPAL ACADEMY OF HIGHER EDUCATION

Manipal Academy of Higher Education, Manipal MPharm Theory End-Semester Examinations.

MPharm - Pharmaceutical Analysis Specialization

MPharm Semester II - End Semester Examination, May 2019

Date: 04/05/2019

Modern Bioanalytical Techniques [PCH-MPA202T] Marks: 75 Duration: 180 mins. SECTION - A Answer all the questions. Answer the following (10 marks  $\times$  5 = 50 marks) 1) Explain solid phase extraction and protein precipitation as sample (10) preparation methods in biopharmaceutical analysis 2) Discuss the various alternative methods in dissolution testing and (10) their applications 3) Write a note on different types and levels of bioanalytical method (10)validation and explain how stability is established during bioanalytical method validation as per FDA 4) What are the factors that can be determined with the help of (10)toxicokinetic principles? 5) Write in detail about safe laboratory practices to be followed in a (10)cell culture lab. **SECTION - B** Answer all the questions. Answer the following (5 marks  $\times$  5 = 25 marks) 6) What are biosimilars? Write a note on FDA recommendation on (5)factors that must be considered in assessing biosimilarity 7) Explain the pH-solubility curve for a weak acid and a weak base. (5) Add a note on BCS classification of drugs 8) What are the implications of Pharmacokinetic interactions? Explain (5) with suitable examples. 9) Explain the different Biosafety levels. (5)

(5)

Write about different types of Cell Culture Hoods

Exam Date & Time: 06-May-2019 (02:00 PM - 05:00 PM)



# MANIPAL ACADEMY OF HIGHER EDUCATION

Manipal Academy of Higher Education, Manipal MPharm Theory End-Semester Examinations.

MPharm - Pharmaceutical Analysis

MPharm Second Semester- End-Semester Examination- May-2019

Date: 06/05/2019

Quality Control and Quality Assurance [PCH-MPA203T]

Marks: 75

| Answe                                 | r all the questions.                                                                                          | SECTION - A                          | Duration: 180 mins.   |
|---------------------------------------|---------------------------------------------------------------------------------------------------------------|--------------------------------------|-----------------------|
| Answer<br>1)                          | the following (10 marks $x = 50$ marks                                                                        | s)                                   |                       |
| 2                                     | Explain in detail about packaging of sterile products. How will you (10) calculate percentage yield? Explain. |                                      |                       |
| 2)                                    | What is SOP? What are it preparation and mainten                                                              | ts benefits? Explain the met         |                       |
| 3)                                    | Explain specifications, ve finished products as per N                                                         |                                      | nd shelf life of (10) |
| 4)                                    | What is pharmaceutical q<br>GMP. How do you prepare                                                           |                                      | objectives of (10)    |
| 5)                                    | Mention the contents and material,                                                                            | specifications for the purch         | ase of raw (10)       |
| Answer all the questions. SECTION - B |                                                                                                               |                                      |                       |
| Answer the                            | following (5 marks x 5 = 25 marks)                                                                            | ry date? Discuss accelerate          | d stability (5)       |
| )                                     | Explain the In process quali                                                                                  | 4                                    | a stability (3)       |
|                                       | Explain the In process quali<br>Discuss packaging operation                                                   | ty control tests for supposit<br>ns. | ories (5)             |
|                                       | What is the difference betwee finished products? Explain.                                                     | een routine and periodic tes         | (5)<br>st for (5)     |
| )                                     | What are the secondary pact<br>transport packaging tests                                                      |                                      |                       |
|                                       |                                                                                                               |                                      |                       |

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Exam Date & Time: 08-May-2019 (02:00 PM - 05:00 PM)



## MANIPAL ACADEMY OF HIGHER EDUCATION

Manipal Academy of Higher Education, Manipal MPharm Theory End-Semester Examinations.

MPharm - Pharmaceutical Analysis

MPharm Semester I - End Semester Examination, May 2019

MPharm Semester I - End Semester Examination, May, 2019 Date: 08/05/2019 Herbal and Cosmetic Analysis [PCH-MPA204T] Marks: 75 Duration: 180 mins. SECTION - A Answer all the questions. Answer the following (10 marks  $\times 5 = 50$  marks) 1) How do you evaluate cosmetic products for moisture, ash and (10)volatile matter? 2) Explain WHO guidelines in quality assessment of herbal drugs. (10)3) Discuss the Indian and international patent law with its protocol. (10)4) What are the different methods used for screening of adulterant. (10)5) What are the general methods of analysis of raw materials used in (10) cosmetic products as per BIS? SECTION - B Answer all the questions. Answer the following (5 marks  $\times$  5 = 25 marks) 6) Discuss about various pharmacopoeias. (5)7) Briefly discuss about toxicity studies of herbal drugs. (5)8) What are AYUSH guidelines? Give the principle involved in homeopathy medicine. (5) 9) Define saponification value and acid value. Mention their (5) significance. 10) Give the specifications for sampling and testing of baby care (5) products.

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