

SECOND-YEAR OF MASTER OF SCIENCE CHEMISTRY REVISED SYLLABUS ACCORDING TO CBCS NEP2020

COURSE TITLE: PHARMACEUTICAL ANALYSIS
SEMESTER-IV
W.E.F. 2024-2025

RECOMMENDED BY THE BOARD OF STUDIES IN CHEMISTRY AND

APPROVED BY THE ACADEMIC COUNCIL

Devrukh Shikshan Prasarak Mandal's
Nya. Tatyasaheb Athalye Arts, Ved. S. R. Sapre Commerce, and
Vid. Dadasaheb Pitre Science College (Autonomous), Devrukh.
Tal. Sangameshwar, Dist. Ratnagiri-415804, Maharashtra,
India

Academic Council Item No:

Nya. Tatyasaheb Athalye Arts, Ved. S. R. Sapre
Commerce, and Vid. Dadasaheb Pitre Science
College (Autonomous), Devrukh. Tal.
Sangameshwar, Dist. Ratnagiri-415804,
University of Mumbai
Master of Science
Chemistry
Second Year
Two
02
Pharmaceutical Analysis
S614CHT
Elective
Chemistry Graduate learner seeking Admission to
Post Graduate Programme in adherence to Rules
and Regulations of the University of Mumbai and
Government of Maharashtra
40%
Formative
PG
60:40
NEP-CBCS
2024-2025

Syllabus for Second Year of Master of Science in Chemistry (With effect from the academic year 2024-2025)

SEMESTER-IV Paper No.- V

Course Title: Pharmaceutical Analysis No. of Credits - 02

Type of Vertical: Elective Course Code: S614CHT

Learning Outcomes Based on BLOOM's Taxonomy:

After completing the course, the learner will be able to					
Course	Blooms				
Learning	Taxonomy	Course Learning Outcome			
Outcome No.					
CLO-01	Understand	Discuss sources, classification, standardization and test			
		involved in pharmaceutical analysis.			
CLO-02	Apply	Explain methods used in analysis of pharmaceutical			
		products.			

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SEMESTER-IV Paper No.- V

Course Title: Pharmaceutical Analysis

No. of Credits - 02

Type of Vertical: Elective Course Code: S614CHT

COURSE CONTENT						
Module No.	Content	Credits	No. of Hours			
1	 UNIT-I: Pharmaceutical Analysis General idea regarding the Pharmaceutical Industry, definition and classification of drugs, introduction to pharmaceutical formulations, classification of dosage forms. Role of FDA in pharmaceutical industries. Sources of impurities in pharmaceutical products and raw materials. Standardization of finished products and their characteristics, official methods of quality control. 	01	15			
2	 Analysis of compounds based on functional groups, instrumental methods for analysis of drugs, assays involving chromatographic separations, proximate assays, assays of enzyme containing substances, biological and microbiological assays and tests. Limit tests, solubility tests, disintegration tests, stability studies, impurity profile of drugs, bioequivalence and bioavailability studies. Polymers in pharmaceuticals and novel drug delivery systems. 	01	15			
	Total	02	30			

Access to the Course

The course is available for all the students admitted for Second year of Master of Science.

Methods of Assessment

The assessment pattern would be 60:40, 60% for Semester End Examination (SEE) and 40% for Continuous Internal Assessment (CIA). The structure of the SEE and CIA would be as recommended by the Board of Studies and approved by the Board of Examination and the Academic Council of the college.

References:

- 1. Analytical Biochemistry, David J Holmes and Hazel Peck, Longman, 1983.
- 2. Bioanalytical Chemistry, Susan R Mikkelesen and Eduardo Cotton, John Wiley and Sons, 2004
- 3. Encyclopedia of Industrial Chemical Analysis, Foster Dee Snell et al, Interscience Publishers, 1967.
- 4. Government of India Publications of Food, Drug and Cosmetic Act and Rules.
- 5. The Handbook of Drug Laws, M L Mehra, University Book Agency, Ahmedabad, 1997.
- 6. Chemical Analysis of Drugs, Takeru Higuchi, Interscience Publishers, 1995.
- 7. Text book of Pharmaceutical Analysis, Kenneth Antonio Connors, Wiley, 2001