



UNIVERSITY OF
PATRAS
ΠΑΝΕΠΙΣΤΗΜΙΟ ΠΑΤΡΩΝ

DEPARTMENT OF PHARMACY

SCHOOL OF HEALTH SCIENCES

UNIVERSITY OF PATRAS
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DEPARTMENT OF PHARMACY
POSTGRADUATE PROGRAM: **DRUG DESIGN AND DEVELOPMENT**

**COURSE TITLE: APPLIED PHARMACEUTICAL ANALYSIS
AND CHARACTERISATION TECHNIQUES OF FINAL PRODUCTS
CODE: DPHA_B03**

**STATISTICS AND QUALITY MANAGEMENT IN PHARMACY
COURSE OUTLINE**

1. GENERAL

SCHOOL	HEALTH SCIENCES		
ACADEMIC UNIT	DEPARTMENT OF PHARMACY		
PARTICIPATING INSTITUTIONS	-		
TITLE of POSTGRADUATE PROGRAM	DRUG DESIGN AND DEVELOPMENT		
LEVEL	POSTGRADUATE		
COURSE CODE	DPHA_B03	SEMESTER	B'
COURSE TITLE	APPLIED PHARMACEUTICAL ANALYSIS AND CHARACTERISATION TECHNIQUES OF FINAL PRODUCTS		
INDEPENDENT TEACHING ACTIVITIES	WEEKLY TEACHING HOURS	CREDITS	
Courses	3	5	
COURSE TYPE	Skills Development		
PREREQUISITE COURSES	-		
LANGUAGE of INSTRUCTION and EXAMINATIONS	Greek		
COURSE OFFERED to ERASMUS STUDENTS	No		
COURSE (URL)	http://www.pharmacy.upatras.gr/images/DS/DPHA_B03_EN.pdf		

2. LEARNING OUTCOMES

Learning Outcomes
<p>The student is introduced to the Spectroscopic techniques for characterization of pharmaceutical formulation: (NIR, IR-ATR, Raman, X-ray diffraction, microscopy (optical and scanning electron), Elemental Analysis Techniques (XRF, AAS, AES, ICP-MS, ICP-OES), Diffraction, Polarimetry, Particle Size Characterization Techniques, Thermal Analysis Techniques (TGA, DTA, DSC). Porosity measurement (BET).</p> <p>Specifically, upon successful completion of the course, the graduate student is expected to have developed level 7 skills in the following topics:</p> <p>Ability to select and use the Spectroscopic technique for the identification and quantification of the individual components of a liquid or solid sample of a pharmaceutical formulation.</p>
General Competences
<ul style="list-style-type: none"> • Working independently • Team work • Search for, analysis and synthesis of data and information, with the use of the necessary technology

3. SYLLABUS

<ol style="list-style-type: none"> 1. Validation of analytical methods. The concept of traceability. Good practice rules (GLP, GMP) and quality procedures in the pharmaceutical industry. Stability control of active substances and excipients. 2. Techniques for the determination of physical characteristics of substances: 3. Diffractometry- Principles, instrumentation, applications in Pharmaceutical Analysis, 4. Polarimetry- Principles, instrumentation, applications in Pharmaceutical Analysis, 5. Particle size characterization Principles, instrumentation, applications in Pharmaceutical Analysis. 6. Methods of thermal analysis (TGA, DTA, DSC). 7. Measurement of porosity (BET). 8. Microscopy (Optical and scanning electron). 9. Polymorphism of active substances in formulations: NIR, IR-ATR, Raman, X-ray diffraction, Microscopy (optical and electron). Examples. 10. Elemental analysis (XRF, AAS, AES, ICP-MS, ICP-OES)
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4. TEACHING and LEARNING METHODS - EVALUATION

DELIVERY	Physical presence of students/teachers in a lecture hall (face-to-face)												
USE of INFORMATION and COMMUNICATIONS TECHNOLOGY	Learning process support through the e-class platform												
TEACHING METHODS	<table style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="text-align: left;"><i>Activity</i></th> <th style="text-align: right;"><i>Semester Workload</i></th> </tr> </thead> <tbody> <tr> <td>Interactive teaching</td> <td style="text-align: right;">39</td> </tr> <tr> <td>Study and analysis of bibliography</td> <td style="text-align: right;">47</td> </tr> <tr> <td>Project</td> <td style="text-align: right;">39</td> </tr> <tr> <td colspan="2">Course Total</td> </tr> <tr> <td>(25 hours of work-load per ECTS credit)</td> <td style="text-align: right;">125</td> </tr> </tbody> </table>	<i>Activity</i>	<i>Semester Workload</i>	Interactive teaching	39	Study and analysis of bibliography	47	Project	39	Course Total		(25 hours of work-load per ECTS credit)	125
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Project	39												
Course Total													
(25 hours of work-load per ECTS credit)	125												
STUDENT PERFORMANCE EVALUATION	<ol style="list-style-type: none"> 1. Written final examination (80%) including <ul style="list-style-type: none"> - Short development questions - Questions of a critical nature - Problem solving 2. Assignment - Presentation of an analytical problem from the international literature (20%) 												

5. RECOMMENDED BIBLIOGRAPHY

Related Bibliography

1. ΕΝΟΡΓΑΝΗ ΑΝΑΛΥΣΗ, ΘΕΜΙΣΤΟΚΛΗΣ Π. ΧΑΤΖΗΪΩΑΝΝΟΥ, ΜΙΧΑΗΛ Α. ΚΟΥΠΠΑΡΗΣ , 2014
2. ΕΝΟΡΓΑΝΗ ΧΗΜΙΚΗ ΑΝΑΛΥΣΗ, Ι. ΠΑΠΑΔΟΓΙΑΝΝΗΣ-Β. ΣΑΜΑΝΙΔΟΥ, 2η Έκδοση, Θεσσαλονίκη, 2011.
3. Φαρμακευτική ανάλυση, D.G. WATSON, , Επιμέλεια Ελληνικής Έκδοσης: Μ. Κουππάρης, Εκδόσεις Παρισιάνου, 2011.
4. ΘΕΜΕΛΙΩΔΕΙΣ ΑΡΧΕΣ ΑΝΑΛΥΤΙΚΗΣ ΧΗΜΕΙΑΣ, SKOOG, D. A. Skoog, D. M. West, F. James Holler, S. R. Crouch, Επιμέλεια Ελληνικής Έκδοσης: Μ. Ι. Καραγιάννης, Κ. Η. Ευσταθίου, Εκδόσεις Κωσταράκη, 2016