

Requirements
This is prior learning (or a practical skill) that is mandatory before enrolment in this module is allowed.

No Co-requisite modules listed

No requirements listed

ZSCI C4105: Assay Development

University					
Module Title:		Assay Development			
Language of Instruction:		English			
Credits: 10					
NFQ Level:	8				
Module Delivered In		1 programme(s)			
Teaching & Learning Strategies:		This module will be delivered via three one-hour lectures and two three-hour practicals per week Students may be required to carry out assignments, give presentations and take multiple choice questions. Independent learning will be facilitated during the preparation of assignments, presentations and practicals.			
Module Aim:		To develop separation and extraction strategies in the analysis of drugs in pharmaceutical preparations			
Learning Outcomes					
On successful comple	etion of t	his module the learner should be able to:			
LO1 Develo	LO1 Develop spectroscopic and other methods in the principles of validation of an analytical method.				
LO2 Validate	LO2 Validate the major extraction and separation techniques used in sample recovery				
LO3 Explore the ass		ay methods to determine contaminant degradation by-products using standard chromatographic techniques			
LO4 Evaluate the methods used in		ethods used in the analysis of various drug delivery vehicles			
Pre-requisite learning	g				
Module Recommendations This is prior learning (or a practical skill) that is recommended before enrolment in this module.					
No recommendations listed					
Incompatible Modules These are modules which have learning outcomes that are too similar to the learning outcomes of this module.					
No incompatible modules listed					
Co-requisite Modules					



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Module Content & Assessment

Indicative Content

Assay Development

• Discuss the principles of validation of an analytical method. • Evaluate the major extraction and separation techniques used in sample recovery • Develop assay methods separating out drug and primary metabolites using standard chromatographic methods • Explore the robustness of the methods in the analysis of sustained-release, multi-dose and other drug-delivery forms

Assessment Breakdown	%
Continuous Assessment	10.00%
Practical	30.00%
End of Module Formal Examination	60.00%

Special Regulation

Students must achieve a minimum grade (35%) in both practical/CA and final exam.

Continuous Asse	essment				
Assessment Type	Assessment Description	Outcome addressed	% of total	Assessment Date	
Case Studies	Regular written examinations to evaluate student understanding of course content	1,4	10.00	n/a	

No Project

Practical	actical			
Assessment Type	Assessment Description	Outcome addressed	% of total	Assessment Date
Practical/Skills Evaluation	Solvent extraction, flocculation, Sep-pack® and pre-column separation.	2,3	30.00	n/a

End of Module Formal Examination				
Assessment Type	Assessment Description	Outcome addressed	% of total	Assessment Date
Formal Exam	Final year evaluation	1,4	60.00	End-of-Semester

SETU Carlow Campus reserves the right to alter the nature and timings of assessment



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Module Workload

Workload: Full Time				
Workload Type	Frequency	Average Weekly Learner Workload		
Lecture	12 Weeks per Stage	3.00		
Practicals	12 Weeks per Stage	6.00		
Independent Learning	15 Weeks per Stage	9.47		
	Total Hours	250.00		

Module Delivered In

Programme Code	Programme	Semester	Delivery
CW_SAPHA_B	Bachelor of Science (Honours) in Pharmaceutics and Drug Formulation	7	Mandatory