

ZSCI H4102: Pharmaceutical Science

Module Title:			Pharmaceutical Science			
Language of Instruction:		n:	English			
Credite:		5				
oreans.		19				
NFQ Level:		8				
Module Deliv	vered In		1 programme(s)			
module Dell						
Teaching & Learning Strategies:			This module will be delivered via a one-hour lecture and a three-hour practical 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 understand the procedures in drug production from synthesis to finished product manufacture. To develop analytical procedures for routine pharmaceutical analysis. To understand drug registration requirements.			
Learning Ou	tcomes					
On successfu	Il completio	n of th	nis module the learner should be able to:			
LO1	Understan	nd the	procedures in drug production from synthesis to finished product manufacture.			
LO2	Develop analytical procedures for routine pharmaceutical analysis.					
LO3	Understand drug registration requirements.Validate existing drug manufacturing plant and instrumentation.					
Pro-requisit	learning					
	e leanning					
Module Recommendations This is prior learning (or a practical skill) that is recommended before enrolment in this module.						
No recomme	ndations lis	ted				
<i>Incompatible Modules</i> 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						
No Co-requisite modules listed						
Requirements This is prior learning (or a practical skill) that is mandatory before enrolment in this module is allowed.						
No requirements listed						



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Module Content & Assessment

Indicative Content

Pharmaceutical preparation

1. Introduction requirements for product registration 2. Preformulation: solubility, solubilisation. Polymorphism, eutectic points, degradation profiles. 3. Formulation theory and practice: simple drug delivery vehicles. 4. Accelerated stability and shelf life. Modified release medications. 5. Nanotechnological innovation. 6. Emulsion preparation: surface tension and surface energy. Experimental techniques. 7. Surfactants, critical micelle concentration, spreading and contact angle. Lapace excess pressure. Microemulsions and liquid crystal phases. Aerosols 8. Stability evaluation and chemical assay. Product registration 9. Regulatory bodies, IMB, FDA, etc. 10. Good manufacturing processes: quality management, production activities, internal audits, calibration 11. Validation: policy, documentation, qualification of equipment and ancillaries, process and analytical validation. Cleaning validation and change control. 12. Rejection and reuse of materials. Complaints and recalls. Contract manufacture 13. Materials management. Production & in-process controls.

Assessment Breakdown	%
Continuous Assessment	10.00%
Practical	20.00%
End of Module Formal Examination	70.00%

Special Regulation

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

Continuous Assessment					
Assessment Type	Assessment Description	Outcome addressed	% of total	Assessment Date	
Short Answer Questions	Regular written examinations to evaluate student understanding of course content	1,2,3	10.00	n/a	

No Project

Practical				
Assessment Type	Assessment Description	Outcome addressed	% of total	Assessment Date
Practical/Skills Evaluation	Dissolution studies of rapid release and controlled-release medications. Formulation of simple emulsion and tablets Assessment of physical stability. Assay development. 16. Use of conductivity to determine solubility products of a number of sparingly-soluble drugs. 17. Extraction by Soxhlet apparatus of podophyllin from podophyllum rhizomes.	1	20.00	Sem 1 End

End of Module Formal Examination					
Assessment Type	Assessment Description	Outcome addressed	% of total	Assessment Date	
Formal Exam	Final year evaluation	1,2,3	70.00	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	2.00	
Practicals	12 Weeks per Stage	2.50	
Independent Learning Time	15 Weeks per Stage	5.00	
	Total Hours	129.00	

Module Delivered In				
Programme Code	Programme	Semester	Delivery	
CW_SABTP_B	Bachelor of Science (Honours) in Biosciences with Biopharmaceuticals	7	Mandatory	