

MODULE DESCRIPTOR

Module Title

Drug Analysis And Toxicology

Reference	PLM306	Version	1
Created	February 2023	SCQF Level	SCQF 11
Approved	March 2023	SCQF Points	30
Amended	August 2021	ECTS Points	15

Aims of Module

To enable the students to critically evaluate the principles, applications and limitations of instrumental analysis in Drug Analysis and Toxicology and to select and implement a range of appropriate analytical techniques to solve a given analytical problem.

Learning Outcomes for Module

On completion of this module, students are expected to be able to:

- 1 Critically appraise key factors related to the quality/safety/authenticity of prescription drugs.
- 2 Critically evaluate the appropriateness of the different techniques available for sampling, preparation and analysis for an analytical problem related to the quality/safety/authenticity of prescription drugs.
- 3 Critically review the results and conclusions from an analytical group project related to quality/safety/authenticity of prescription drugs.

Indicative Module Content

Drug screening and confirmation. Analytical procedures for the determination of drugs of abuse in biological matrices e.g. blood, serum, plasma, saliva, urine etc. Structure elucidation, Pharmaceutical drug analysis, drug formulation, use of general pharmaceutical monographs for formulated preparations. Measuring toxicology: LD50, exposure limits, thresholds, reversibility, sensitivity. Toxicokinetics: absorption, distribution, metabolism and excretion. Solution of an analytical problem appropriate to Drug Analysis or Toxicology requiring a combination of Laboratory techniques.

Module Delivery

Full-time and Part-time; the module is delivered by formal lectures and some external speakers. Mandatory attendance for 5 days of laboratory work as part of a designated group solving a drug analysis or toxicology problem.

Indicative Student Workload

	Full Time	Part Time
Contact Hours	70	70
Non-Contact Hours	230	230
Placement/Work-Based Learning Experience [Notional] Hours	N/A	N/A
TOTAL	300	300
Actual Placement hours for professional, statutory or regulatory body		

ASSESSMENT PLAN

If a major/minor model is used and box is ticked, % weightings below are indicative only.

Component 1

Type:	Practical Exam	Weighting:	100%	Outcomes Assessed:	1, 2, 3
Description:	Demonstrate competent laboratory planning & analysis of an abused or common pharmaceutical drug and individual performance in a PowerPoint presentation.				

MODULE PERFORMANCE DESCRIPTOR**Explanatory Text**

The grade represents Component 1 (PE1). A minimum of Module Grade D is required to pass the module. Non-submission will result in an NS grade.

Module Grade	Minimum Requirements to achieve Module Grade:
A	A
B	B
C	C
D	D
E	E
F	F
NS	Non-submission of work by published deadline or non-attendance for examination

Module Requirements

Prerequisites for Module	None, in addition to the course entry requirements.
Corequisites for module	None.
Precluded Modules	None.

INDICATIVE BIBLIOGRAPHY

- 1 AULTON, M.E. 2021. *Pharmaceutics, The Design and Manufacture of Medicines*. Churchill Livingstone Elsevier.
- 2 WATSON, D.G. 2020. *Pharmaceutical Analysis*. Churchill Livingstone Elsevier.
- 3 ANSEL, H.C. 2014. Eight edition. *Pharmaceutical Dosage Forms and Drug Delivery Systems*. Wolters Kluwer Health.
- 4 CREAN, A. 2010. *The physiochemical basis of Pharmaceuticals*. Oxford University Press.
- 5 SKOOG, W. CROUCH, S. WEST, D and HOLLER, F. 2021. Tenth edition. *Skoog and West. Fundamentals of Analytical Chemistry*. Cengage Learning