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MODULE DESCRIPTOR

Module Title

Advanced Pharmaceutical Analysis and Quality Control

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Reference	ASM040	Version	1
Created	January 2019	SCQF Level	SCQF 11
Approved	May 2019	SCQF Points	15
Amended		ECTS Points	7.5

Aims of Module

To enable students to evaluate, problem solve and develop skills in a range of advanced analytical techniques for the analysis of drugs and medicines. To enable students to understand and evaluate quality assurance and control systems for medicine manufacture.

Learning Outcomes for Module

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

- 1 Critically understand the principles and applications of advanced analytical techniques for the determination of a drug or medicine.
- 2 Critically evaluate the advantages and limitations of advanced analytical techniques for the determination of a drug or medicine.
- 3 Validate an appropriate quality assurance and control system for medicine manufacture.

Indicative Module Content

Pharmaceutical drug/medicine analysis, use of pharmaceutical monographs for the analysis of formulated preparations, extraction methods. Advanced analytical techniques include: liquid chromatography, gas chromatography, mass spectrometry, capillary electrophoresis, fluorescence spectroscopy, (derivative) ultra-violet spectroscopy, Raman spectroscopy, infrared spectroscopy. Quality assurance: Principles, procedures, test methods, records, reporting, data management, auditing and sampling; laboratory accreditation and accreditation regimes; standards GLP and GMP.

Module Delivery

The module will be delivered by lectures, tutorials and laboratory workshops.

Module Ref: ASM040 v1

Indicative Student Workload	Full Time	Part Time
Contact Hours	30	N/A
Non-Contact Hours	120	N/A
Placement/Work-Based Learning Experience [Notional] Hours		N/A
TOTAL	150	N/A
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: Coursework Weighting: 70% Outcomes Assessed: 1, 3

Description: Technical report critically evaluating analytical and quality control methods for a given medicine.

Component 2

Type: Coursework Weighting: 30% Outcomes Assessed: 2

Description: Critical appraisal of a contemporary analytical technique

MODULE PERFORMANCE DESCRIPTOR

Explanatory Text

To pass this module the student must achieve a grade D or better. The grading criteria are:

Module Grade	Minimum Requirements to achieve Module Grade:
Α	All components must have a minimum of 50% and the overall total (by weighting) must be equal to or greater than 70%.
В	All components must have a minimum of 40% and the overall total (by weighting) must be between 60-69%.
С	All components must have a minimum of 35% and the overall total (by weighting) must be between 50-59%.
D	All components must have a minimum of 35% and the overall total (by weighting) must be between 40-49%.
E	All components must have a minimum of 35% and the overall total (by weighting) must be between 35-39%.
F	Any component is less than or equal to 34%

Non-submission of work by published deadline or non-attendance for examination

Module Requirements

NS

Prerequisites for Module None, in addition to course entry requirements.

Corequisites for module None.

Precluded Modules None.

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INDICATIVE BIBLIOGRAPHY

- Aulton, M.E. (2017). 'Pharmaceutics the design and manufacture of medicines', 5th edition, Churchill Livingstone, Elsevier.
- 2 Khar, R.K. (2017). 'Lachman's/Lieberman's: the theory and practice of industrial pharmacy', 4th edition, CBS Publishers.
- 3 Prichard, E. (2007). 'Quality assurance in analytical chemistry', Wiley.
- Skoog, D.A., Holler, F.J., Crouch, S.R. (2018). 'Principles of instrumental analysis', 7th edition, Brookes Cole.
- Watson, D.G. (2016) 'Pharmaceutical analysis a textbook for pharmacy students and pharmaceutical chemists', 4th edition, Churchill Livingstone, Elsevier.