

CORPORATE AND ACADEMIC SERVICES

MODULE SPECIFICATION

Part 1: Basic Data					
Module Title	Medicinal Chemistry				
Module Code	USSKB5-15-2	Level	2	Version 1	
Owning Faculty	Health and Applied	Field	Biological, Biomedical and Analytical Sciences		
Contributes towards	BSc Forensic Science, BSc Forensic Science (Chemistry), BSc Biomedical Science, BSc Biomedical Science (Clinical)			Biomedical Science,	
UWE Credit Rating	15	ECTS Credit Rating	7.5	Module Type	Standard
Pre-requisites	USSJRT-30-1 Chemistry in Context or USSKA4-30-1 Cell Biochemistry and Genetics		Co- requisites	None	
Excluded Combinations	USSKB7-15-2 Molecular Genetics, USSKB8-15-2 Forensic Biology for the Forensic Science (Biology) pathway only.		Module Entry requirements	N/A	
Valid From	September 2014		Valid to	current	

CAP Approval Date	

Part 2: Learning and Teaching				
Learning Outcomes	On successful completion of this module students will be able to:			
	 apply chemical knowledge to rationalise drug design, structural modification and synthesis (components A1, B1); 			
	 comment on chemical strategies used in drug development to control pharmacokinetics, formulation and delivery (component A1). 			
	 carry out experimental techniques commonly used in drug synthesis (component B1) 			
	 recognise and explain how physicochemical properties of drugs can be measured, and used to study and predict structure-activity relationships (components A1, B1) 			
	 describe and explain the action and development of selected examples of medicines and drugs (component A1) 			
Syllabus Outline	The general principles of medicinal chemistry:			
	General classes of drugs and their medicinal activity. Types of drug target and the origins of drug-target interactions. Identification of the essential structural features (pharmacophore) for bioactivity.			

	Examples of synthetic methods in medicinal chemistry to develop leads, to optimise drug-target binding interactions, to alter drug solubility/stability, to improve drug pharmacokinetics, to enhance drug delivery/formulation, to mass produce drugs and to control drug stereochemistry. Relationships between chemical structure and physicochemical properties of a drug (structure-activity relationships, SARs), and their quantitative measurement (QSAR). Hydrophobicity and Hammett constants, Hansch analysis, Craig plots and Topliss decision trees. Types of prodrug and the use of classical and non-classical bioisosteres. Applications to drug design and development in medicinal chemistry. The use of X-ray crystal structure determination and spectroscopy to identify drug intermediates and to inform drug development. <u>Case Studies in Medicinal Chemistry</u> : To illustrate major classes of medicines and their chemical development and action, a selection of the following will be discussed; The design and development of different types of antidepressant; Anti-cancer drugs acting as alkylating and intercalating agents; Antiulcer treatment - QSAR in the development of omeprazole or cimetidine; Captopril, an antihypertensive agent – the development of a lead compound;
	Development of the morphinans and methadone from morphine;
	Structure and activity of the antibacterial penicillins.
Contact Hours	The contact hours (36) are distributed as follows:
	21 hours of lectures,
	3 hours of workshops,
Teaching and Learning Methods	 The material will be delivered using a combination of lectures, workshops and laboratory work. Lectures will be augmented by directed reading in the recommended text and in selected publications from the scientific literature, e.g. Drug Discovery Today, Journal of Medicinal Chemistry. The topics selected for delivery by workshops and practical work will be designed to enhance problem solving skills and to provide experience of relevant laboratory techniques. Technology enhanced learning will be embedded within teaching materials via links to supplementary electronic online resources of the textbook and other relevant information portals, e.g. <u>http://www.chemspider.com</u> Use will also be made of various in-house electronic resources and flash videos in chemistry for biologists available at <u>http://calcscience.uwe.ac.uk</u>. Student learning will be further supported through a variety of materials posted on the University's E-Learning Environment, Blackboard. Independent learning will take the following forms with an approximate indication of time required for each: Essential reading to support acquisition of knowledge and completion of problem solving chills avariate relating to loctures.

Key Information Sets Information	Key Information Sets (KIS) are produced at programme level for all programmes that this module contributes to, which is a requirement set by HESA/HEFCE. KIS are comparable sets of standardised information about undergraduate courses allowing prospective students to compare and contrast between programmes they are						
	Key Inform	ation Set - Mo	odule data				
	Number of a	credits for this	module		15		
	Hours to be Schoduled Independent DI			Placement	Allocated		
	allocated	learning and teaching study hours	study hours	study hours	Hours		
	150	36	114	0	150	\bigcirc	
	The table below indicates as a percentage the total assessment of the module which constitutes a - Written Exam : One unseen written exam Coursework : One portfolio of workshop and laboratory worksheets Total assessment of the module:						
	Written exam assessment percentage			50)%		
	Coursework assessment percentage			50)%		
	Practical exam assessment percentage		0	%			
Deading	All students will be appoured to make full use of the print and electronic resources						
Strategy	An students will be encouraged to make full use of the print and electronic resources available to them through membership of the University. These include a range of electronic journals and a wide variety of resources available through web sites and information gateways. The University Library's web pages provide access to subject relevant resources and services, and to the library catalogue. Many resources can be accessed remotely. Students will be presented with opportunities within the curriculum to develop their information retrieval and evaluation skills in order to identify such resources effectively. Any essential reading will be indicated clearly, along with the method for accessing it,						
	pack or be referred to texts that are available electronically, etc. This guidance will be available either in the module handbook, via the module information on Blackboard or through any other vehicle deemed appropriate by the module/programme leaders.						
If further reading is expected, this wil a clear indication will be given regardir students will be given guidance on how e.g. through use of bibliographical data				l be indicated ng how to acco v to identify re abases.	clearly. If spe ess them and levant source	ecific texts are listed, d, if appropriate, es for themselves,	
Indicative Reading List	The followin	ng book is rec of the course.	commended fo	or purchase by	/ students as	it covers the majority	
	Patrick, G. University F	L. (2013) <i>An</i> Press.	Introduction to	o Medicinal Cl	nemistry 5 th e	d. Oxford: Oxford	

Students are also advised to consult related texts on the topic, of which the following are representative:
Thomas, G. (2007) <i>Medicinal Chemistry – an introduction</i> ,2 nd ed. Chichester: John Wiley.
Silverman, R. B. (2004) <i>The Organic Chemistry of Drug Design and Drug Action</i> ,2 nd ed. Oxford: Academic Press.
Students who wish to access chemistry texts to underpin their chemical knowledge are advised to consult some of the following in Frenchay library:
Volhardt, P, Schore, N., (2009) <i>Organic Chemistry - structure and function</i> 6 th ed. New York: W.H. Freeman
Crow, J. Bradshaw, T. and Monk, P. (2006) <i>Chemistry for the Biosciences</i> . Oxford: Oxford University Press.
Lewis, R. and Evans, W. (2011) Chemistry, 4 th ed. Basingstoke: Palgrave Macmillan,
Additional useful texts can be accessed at shelf marks 615.1 and 541.22

Part 3: Assessment			
Assessment Strategy	Students will undertake practical work and workshops based on synthesis of selected drugs and the principles of drug development.		
	The assessed worksheets will contain questions and responses for students to complete during these timetabled sessions and further questions for students to research in their own time.		
	The examination will assess the students' knowledge acquired during lectures and workshops, and from their own directed, independent learning.		

Identify final assessment component and element		
	A:	B :
% weighting between components A and B (Standard modules only)	50	50
First Sit		
Component A (controlled conditions)	Element weighting	
Description of each element	(as % of co	omponent)
1. 3 hour written examination)%
Component B	Element v	veiahtina
Description of each element	(as % of co	omponent)
		20/
1. Portfolio of worksheets		J%

Resit (further attendance at taught classes is not required)	
Component A (controlled conditions)	Element weighting
Description of each element	(as % of component)

1. 3 hour written examination	100%	
2.		
Component B Description of each element	Element weighting (as % of component)	
1. Portfolio of worksheets	100%	
2.		
If a student is permitted an EXCEPTIONAL RETAKE of the module the assessment will be that indicated by the Module Description at the time that retake commences.		