

## CORPORATE AND ACADEMIC SERVICES

## MODULE SPECIFICATION

		Part 1: Basi	c Data			
Module Title	Pathophysiology	of Disease (Pre	emedical Sciences	s)		
Module Code	USSK63-30-1		Level	1	Version	1
Owning Faculty	Health and Appl	ied Sciences	Field	Applied Sciences		
Contributes towards	Premedical Scie	nces Cert HE				
UWE Credit Rating	30	ECTS Credit Rating		30	ECTS Cr	redit Rating
Pre-requisites	None		Co- requisites	None		
Excluded Combinations	None         Module Entry requirements         N/A					
Valid From	September 2014         Valid to         September 2020					

CAP Approval Date 28/03/2014

Part 2: Learning and Teaching				
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Learning Outcomes	<ul> <li>On successful completion of this module students will be able to:</li> <li>describe current understanding of topical issues in microbiology and understand how the discipline of Medical Microbiology has evolved (A1, B2, 3)</li> <li>discuss the diversity of microorganisms and their ubiquity and discuss the interactions of microorganisms with each other and with animals (A1, B2, 3)</li> <li>explain the importance of pathogenic bacteria, viruses, fungi and parasites and in particular with regard to the medically important diseases they may cause (A1, B3)</li> <li>describe some of the other major causes of human disease and explain their biological basis (A1, B3)</li> <li>explain the basis of disease response mechanisms such as inflammation, necrosis and cell death (A1, B3)</li> <li>discuss approaches to the investigation and diagnosis of selected disease processes (A1, B2, 3)</li> <li>demonstrate basic skills of observation, measurement and data analysis and interpretation in experiments concerning human diseases and basic skills in the safe handling and containment of microorganisms (in particular in a simulated clinical diagnostic work that allows appreciation of the interface with patients) (B1)</li> <li>All learning outcomes will be assessed under the module components and elements therein as indicated.</li> </ul>			
Syllabus Outline	<ul> <li>Introductory microbiology: range of size, nutrition and taxonomy of</li> </ul>			
	microorganisms. The Bacteria - main groups based on primary characteristics.			

	<ul> <li>Archaea. Fungi - main groups based on sexual reproduction.</li> <li>Cultivation and control of microorganisms: Aseptic technique, microbiological culture media, selective and differential media, microbial growth. Laboratory safety; physical and chemical methods of control. Hazard groupings of microorganisms, containment categories for laboratories.</li> <li>Food and industrial microbiology: microbial food spoilage, food poisoning and food-borne infections. Microorganisms used by the food industry, microbial production of antibiotics and complex organic molecules.</li> <li>Microbial interactions: intermicrobial relationships; animal-microbe interactions, including an introduction to the human microbiota and to pathogenicity.</li> <li>Medical microbiology - Development of the discipline: The history of medical microbiology: a review of the "golden age" of microbiology and its leading figures; the role of the medical microbiologist today, including developments which aid in the understanding of pathogens and diagnostics.</li> <li>Medical microbiology - Diseases: Coverage of a range of medically important bacteria, viruses, fungi and parasites: an overview of the range of diseases that microbes cause, from the trivial to the life-threatening.</li> <li>Current issues in Medical Microbiology - Emerging and re-emerging pathogens: an evaluation of the re-emergence of illnesses (e.g. tuberculosis) to attempt to identify reasons for their return; consideration of the emergence of new diseases (e.g. SARS, haemorrhagic viruses).</li> <li>General concepts and introduction to human disease.</li> <li>Overview of haemopoeisis, normal blood parameters and haemostasis and those relating to infection scenarios</li> <li>Disorders of the liver and heart (atherosclerosis) will be described, also diabetes</li> <li>Clinical cytogenetics, karyotype analysis and phenotypic expression of genetic abnormality</li> <li>Cellular injury and death, The cell as the basis of life and disease, Cell death, necrosis and apoptosis</li> <li></li></ul>
Contact Hours	foreign material, Cells and tissues of the immune system. Students undertaking this 30 credit module can expect 78h of scheduled learning contact time with teaching staff, spread over the academic year. This contact time will occur during lectures (36h), practical sessions (24h), tutorials (12h) and during timetabled in class assessments (6h) in the form of MCQ tests.
Teaching and Learning Methods	Theoretical material within the module will be presented to the students in the form of weekly lectures throughout each of the semesters in the academic year. The learning of lecture content will be reinforced through time spent in independent learning by the directed reading of recommended texts and through the use of technology enhanced learning resources that will be provided online. A number of relevant practical sessions will be incorporated during each of the semesters and will be used to highlight important aspects of both biochemistry and genetics within an integrated biomedical and medical context. Practical sessions will both drive hands on learning and the acquisition of technical skills at both an individual and group working level. Online MCQ assessments will be used to further engage students in the development of their continual learning skills.
	Students undertaking this module can expect to receive 2h of lectures per week of the teaching period and would be expected to spend another 3h in independent learning while undertaking directed reading in relation to each of the lecture sessions. In addition to the lectures the students will undertake 8x2h practical classes across both semesters. For each of the practical classes the students should again expect to spend the same time in reading around the subject before and after each of these sessions. Each practical class will be followed by a 1h tutorial session. The students will also receive fortnightly 30min in class assessments that comprise online MCQs

	and v	vhich will te	est their knowle	edge gained d	uring both lec	ture and prac	tical sessions	3.
		neduled le essment pe	arning includ eriods.	les lectures, <sub>l</sub>	practical clas	ses, tutorials	and in clas	S
			learning inclu		gaged with es	sential readir	ıg, assignmer	nt
		ow. Schedu	ns constitute Iled sessions r					
Key Information Sets Information	this m comp prosp	nodule cont arable sets	Sets (KIS) are tributes to, wh s of standardis dents to compa olying for.	ich is a require ed information	ement set by H about under	HESA/HEFCE graduate cou	E. KIS are rses allowing	
		Key Inform	nation Set - Mo	odule data				
		Numbero	f credits for this	s module		30		
		Hours to be allocated	Scheduled learning and teaching study hours	Independent study hours	Placement study hours	Allocated Hours		
		300	78	222	0	300		
	Pleas nece	r <b>sework</b> : W se note tha	Unseen writte Vritten assignn t this is the tot ect the compor lescription:	nent or essay, al of various ty	/pes of asses	sment and wi		n
		т	otal assessm	ent of the mod				
		·						
			Vritten exam as	•	-	40%		
		C	Coursework as	sessment per	centage	60% 100%		
Reading Strategy	•	resource include through pages p library c presente	ents will be e es available t a range of ele web sites ar provide access atalogue. Mar ed with opport and evaluatio	o them throu ctronic journal nd information s to subject re ny resources c unities within	gh membersh s and a wide gateways. T levant resour an be access the curriculun	hip of the U variety of res The Universit ces and serv ed remotely. In to develop	niversity. The ources availal y Library's w ices, and to t Students will their informat	ese ble veb the be
	•		sential readin					

	sold a print study pack or be referred to texts that are available electronically, <i>etc.</i> 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 <b>further reading</b> is expected, this will be indicated clearly. If specific texts are listed, a clear indication will be given regarding how to access them and, if appropriate, students will be given guidance on how to identify relevant sources for themselves, e.g. through use of bibliographical databases.
Indicative	Latest versions of the following:
Reading List	
	Core texts
	<ul> <li>An introduction to human disease, pathology and pathophysiology correlations Crowley, Publisher: Jones and Bartlett</li> <li>Currell G and Downman AA, Mathematics and Statistics for Science, Wiley- Disease</li> </ul>
	<ul> <li>Blackwell.</li> <li>Willey, J. M., Sherwood, L. M. &amp; Woolverton, C. J. (2011). <i>Prescott's Microbiology 8<sup>th</sup> edition</i>. New York: McGraw Hill.</li> </ul>
	<ul> <li>Brooks, G. F., Carroll, K. C., Butel, J. S., Morse, S. A. &amp; Mietzner, T. (2010). Jawetz, Melnick &amp; Adelberg's Medical Microbiology 25<sup>th</sup> edition. New York: McGraw Hill.</li> </ul>
	Other useful texts include:
	Haematology:
	<ul> <li>A Beginner's Guide to Blood Cells. Bain. Publisher: Blackwell Publishers</li> <li>Lecture Notes on Haematology. Hugh-Jones, Wickramasinghe and Hatton. Publisher: Blackwell Publishers</li> <li>Case-Based Haematology. McCann, Foa, Smith and Conneally. Publisher: Blackwell Publishers</li> </ul>
	<ul> <li>Clinical Biochemistry:</li> <li>Clinical Chemistry, Luxton. Publisher: Butterworth-Heinemann</li> <li>Clinical Chemistry 5th edition, Marshall, Bangert, Publisher: C.V. Mosby</li> <li>Clinical Biochemistry 3rd edition, An illustrated colour text, Gaw, Murphy, Cowan, Denis O'Relly, Stewart, Shepherd, Publisher: Churchill Livingstone</li> </ul>
	Immunology:
	<ul> <li>How the Immune system works. Sompayrac, Publisher: Blackwell Publishers</li> <li>Immunology (5th edition). Goldsby, Kindt, Kuby, Osborne, Publisher: WH Freeman and Co.</li> </ul>
	Pathology:
	<ul> <li>Pathology. Steven and Lowe. Publisher: C.V. Mosby</li> <li>Basic Pathology: An introduction to the Mechanisms of Disease Lakhani, Dilly, Finlayson, Dogan, Publisher: Arnold</li> <li>The Biology of Disease, Phillips, Murray, Kirk. Publisher: Blackwell Publishers</li> </ul>
	Microbiology
	<ul> <li>Madigan, M., Martinko, J., Stahl, D. &amp; Clark, D. (2012). Brock Biology of Microorganisms 13<sup>th</sup> edition. San Francisco: Pearson.</li> <li>Strelkauskas, A., Strelkauskas, J. &amp; Moszyk-Strelkauskas, D. (2010). Microbiology: a clinical approach. New York: Garland Science.</li> </ul>

Assessment Strategy	<ul> <li>Summative assessment for this module will be provided usi number of approaches. The nature of the premedical scie programme to which this module contributes requires continuous final assessment of student learning and a measure of acquisition of both oral and written presentation skills of anal data.</li> </ul>	nces and their
	<ul> <li>Continuous assessment within component B will be provided by use of frequent multiple choice question tests throughout the mo- and following blocks of learning provided in the form of lect These tests will be provided online, marked automatically and results provided to the module leader. Feedback at this level will be provided online and will be by review of the tests after they been completed and will include the correct answers and rationale behind these.</li> </ul>	odule ures. 1 the also have
	<ul> <li>The ability of the students to write scientifically, analyse data present their work will be assessed under component B in the for an oral presentation of a scientific poster and also a w assignment, either an essay based or practical report write up. T will be marked and feedback provided in the form of w comments.</li> </ul>	rm of ritten hese
	<ul> <li>Final assessments under component A will take the form of examination that comprises short answer and multiple cl questions.</li> </ul>	

Identify final assessment component and element			
% weighting between components A and B (Standard modules only)		B: 60	
First Sit Component A (controlled conditions)	Element	veighting	
Description of each element		(as % of component)	
1. EX1 Examination Exam Period 2 (3h) FINAL ASSESSMENT		100%	
Component B Description of each element	Element v (as % of co		

Description of each element	(as % of component)
1. CW1 Poster Presentation	25%
2. CW2 Written assignment	25%
3. CW3 MCQ Tests	50%

Resit (further attendance at taught classes is not required)				
Component A (controlled conditions) Description of each element	Element weighting (as % of component)			
1. EX2 Examination Exam Period 3 (3h) FINAL ASSESSMENT 100%				
Component B Description of each element	Element weighting (as % of component)			
1. CW1 Poster Presentation	50%			

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.