Section 2 - Module Specifica	ation					
Module Title Clinical Pharmacology and the Assessment of Drug Safety Module Code BMD273						
Credit Value 15 Level	5 Mode of Delivery		Semester	Semester B		
Pre-requisite modules	Co-requisite modules	Overlapping mod	ules			
1) Content Description Provide a description of the mod System (approx. 70-80 words).	dule, as it will appear in the N	lodule Directory and o	on the Studen	t Information		
This module will introduce to the stuthe progression of the diseases they enhancers in sport. Introductory lectures will be followe formal lectures and interactive semily papers. We will offer practical works	are used to treat. It will include and the second of the s	consideration of drugs of of the subject given by e rials with opportunities t	of abuse and dru experts in their f	ugs as performance		
2) Module Aims Specify the aims of the module,	i.e. the broad educational pu	rposes for offering th	is module.			
A critical understanding of the the di	rugs that are used to treat comm	on diseases and their me	chanisms of act	tion		
An awareness of benefits, side effects, risks, contra-indications and interactions of drugs.						
A critical understanding for evidence	-based prescribing in clinical pra	rtice				

3) Learning Outcomes

Identify the learning outcomes for this module, i.e. knowledge, skills and attributes to be developed through completion of this module. Outcomes should be referenced to the relevant QAA benchmark statements and the <a href="Framework for Higher Education Qualifications in England, Wales and Northern Ireland (2008). The SEEC Credit Level Descriptors for Further and Higher Education 2003 and Queen Mary Statement of Graduate Attributes should also be used as a guiding framework for curriculum design.

Acad	demic Content:
A 1	Critique of the pathophysiology of common disorders in the neurological, endocrine, immune and cardiovascular disorders
A 2	An analysis of the mechanisms of drug actions in the treatment of disease.

Disci	Disciplinary Skills - able to:			
B 1	Critically evaluate published research studies and clinical audits			
B 2	Conduct laboratory experiments safely with care and precision			
В3	Write scientific reports and present scientific data			
B 4	Recognise safe and unsafe prescribing activities			

Attrik	Attributes:			
C 1	Have the intellectual curiosity to learn continuously from diverse sources of information			
C 2	Be able to critique complex scientific concepts clearly and logically			
C 3	Make judgments based on evidence			
C 4	Effective time management and independent learning			

4) Reading List

Provide an indicative reading list for the module. This should include key texts and/or journals but should not be an exhaustive list of materials.

- * Rang & Dale's Pharmacology: with STUDENT CONSULT Online Access by Humphrey P. Rang, Maureen M. Dale, James M. Ritter and R. J. Flower, Publisher: Churchill Livingstone; 7th Revised edition edition (25 Mar 2011), ISBN-10: 0702034711
- * Oxford Textbook of Clinical Pharmacology and Drug Therapy by David Grahame-Smith and Jeffrey Aronson ISBN-10: 0192632345

Topical research papers in relevant journals, for example: British Journal of Clinical Pharmacology

Lancet New England Journal of Medicine Current opinion in Pharmacology

5) Teaching and Learning Profile

Provide details of the method of delivery (lectures, seminars, fieldwork, practical classes, etc.) used to enable the achievement of learning outcomes and an indicative number of hours for each activity to give an overall picture of the workload a student taking the module would be expected to undertake. This information will form the Key Information Set for each undergraduate programme and will be used to populate the KIS widget found on the QMUL programme information pages. More information can be found online about KIS. You may also wish to refer to the QAA guidance on contact hours when completing this section.

Activity Type	KIS Category	Time Spent (in hours)
Lecture	Scheduled	20
Seminar	Scheduled	4
Practical Classes and workshops	Scheduled	8
Guided independent study	Independent	118
	Total	150

Specify the total module notional study hours. This should be a total of the hours given for each activity. The notional study hours for each academic credit point is 10. A 15 credit point module therefore represents 150 notional study hours.

Activity Type	Total Time Spent (in hours)	Percentage of Time Spent
Scheduled learning and teaching	32	21
Placement		
Independent Study	118	79
Total	150	100

Use the information provided in the box above to specify the total time spent and the percentage time spent in each category of teaching and learning activity.

6) Assessment Profile

Provide details of the assessment methods used to assess the achievement of learning outcomes.

Description o Assessment	I Assessment Lyne	KIS Category	Duration / Length	% Weighting	Final element of assessment?	Qualifying Mark
Report	Report	Written	2h	50%	No	

Essay	Study Design	Coursework	1500 words	25%	yes	
Essay	Drug-Drug interactions	Coursework	1500 words	25%	No	

Qualifying mark: A specified minimum mark that must be obtained in one or more elements of assessment in order to pass a module. This is in addition to, and distinct from, the requirement to achieve a pass in the module mark to pass the module.

Reassessment

Provide details of the reassessment methods used, specifying whether reassessment is either standard reassessment or synoptic reassessment.

Standard Reassessment	 Synoptic Reassessment

Synoptic reassessment details (if you have indicated synoptic reassessment above, please give details)					
Brief Description of Assessment Assessment Type Duration / Length of Examination / Coursework					
	Written Exam	2 hours			
	Practical Report	15000			