



Course Detailed Description – Procedures of the Course Plan Committee /Faculty of Pharmacy QF02/0408-2.1E

I	Department	Pharmacy
	F	

Course Name	Pharmaceutical Organic Chemistry (2)	Course No.	0201216
Prerequisite	Pharmaceutical Organic Chemistry (1)	Credit Hours	3
Number & date of course plan approval		Brief Description	See form QF02/0409

Course Objective	This course aims to teach the undergraduate-students the organic functional groups, in a continuation to those mentioned in Pharmaceutical Organic Chemistry (1), in terms of nomenclature, synthesis and reactions. In addition, applying these methodologies in synthesis of new derivatives of known active pharmaceutical ingredients.				
	1) To recognize most modern synthetic reagents and illustrate their common use with sample reactions of their devising.				
Intended	2) To understand and illustrate the mechanistic for all reactions to which they are exposed.				
Learning Outcomes	3) To locate accurate literature precedent for any transformation they wish to suggest, and to cite it the correct format.				
	4) To design an original synthesis for any molecule presented to them, complete with accurate, appropriate literature references for all non-trivial steps.				
	5) To communicate mechanistic logic with electron pushing arrows.				
Course Topics	Pharmaceutical organic chemistry is primarily a lecture and problem-solving course, which builds upon the first course of organic chemistry to prepare the student for other courses in pharmacy, biochemistry medicinal chemistry and phytochemistry. The curriculum is divided between advance topics in three areas of organic chemistry: (1) structures, properties, and nomenclatures of organic compounds, (2) mechanistic theory, and (3) synthesis and reactions. The course is composed of series of lectures, guided problem sets, and exams.				
Text Books	Organic Chemistry, T.W.G. Solomons & C.B.Fryhle, John Wiley & Sons, 10th edition.				
References	<ol> <li>Organic Chemistry By Morrison &amp; Boyd, 6th Edition.</li> <li>Organic Chemistry By McMurry, 7th Edition.</li> </ol>				
Grade	$1^{st} Exam = 25\%$ $2^{nd} Exam = 25\%$ $Practical Course Work = 50\%$ $(Reports, Term Papers, Quizzes)$				





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Determination	Fi	nal Exam = 50%	Grade Determination	Final Exam = 50	)%
		C011#50	Outline		
		Course	Outime	Chapters in	
Week	Hours	Sul	bjects	Textbook	Notes
1	1 1 1	Ethers and epoxides  *Ethers a- Structure b- Nomenclature c- Physical properties  Preparations of Ethers Williamson synthesis  *Reaction of Ethers Cleavage by strong acids  Epoxides a- Structure b- Nomenclature of epoxides  *Preparation of Epoxides: Epoxidation Reactions of Epoxides: a-Acid-catalyzed ring opening reactions b-Base -catalyzed ring opening reactions		11	
2	1 1 1	Aromatic Compound Reactions of Aromat *Introduction, nomer derivatives *Kekule's Structure of of benzene Aromaticity and Huck *Other polycyclic aro Reactions of Aromat *Electrophilic aromat	ds, Aromaticity and tic Compounds iclature of benzene f benzene and stability sel's rule, matic compounds	14	
3	1 1 1		ions on reactivity and bstitution, theory of	15	
4	1 1 1	*Structure and no derivatives, physical passing synthesis of phenols *Reactions of phenols		21	Ref. 1 & 2
5	1 1 1	*Nomenclature Physical properties	nes	16	Ref. 1 & 2





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	1	*Synthesis of aldehydes and ketones  *Reactions of Aldehydes and Ketones  Mechanism of nucleophilic addition to C=O.  The Addition of HCN  *The Addition of ylides (Wittig reaction) The Addition of organometallic reagents  * The Addition of alcohols: hemiacetals and acetals		Ref. 1
6	1 1	*The Addition of primary and secondary amines Oxidation & reduction of aldehydes and ketones	16	& 2
7	1 1 1	Reactions at the α-H of carbonyl compounds, Condensation and Conjugate addition reactions of carbonyl compounds.  *The acidity of the α-H of carbonyl compounds Reactions <i>via</i> enols and enolate anions  *Aldol reactions (without crossed Aldol reactions)  *Cyclizations <i>via</i> Aldol	18 + 19	Ref. 1 & 2
8	1 1 1	*Addition to α, β-unsaturated aldehydes and ketones *Carboxylic Acids and Their Derivatives, Nucleophilic Addition-Elimination at the Acyl Carbon	18 17	Ref. 1 & 2
9	1 1 1	*Nomenclature  *Physical properties  *Preparation of carboxylic acids  *Nucleophilic addition – elimination at the acyl carbon	17 17	Ref. 1 & 2
10	1 1 1	*Relative reactivity of acyl compounds *Acyl chlorides *carboxylic acid anhydrides	17	
Week	Hours	Subjects	Chapters in Textbook	Notes
11	1 1 1	*esters *amides *nitriles	17	
12	1 1 1	*Amines  *Nomenclature (one system) and physical properties Basicity of amines  *Synthesis of amines	20	Ref. 1 & 2





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13	1 1 1	*Reactions of amines, *reactions with nitrous acid. *Replacement reactions of arenediazonium salts.	20	Ref. 1 & 2
14	1 1 1	*Coupling reactions of arenediazonium salts *Reaction of amines with sulfonyl chloride. *Sulfa drugs-synthesis	20	Ref. 1 & 2
15	1 1 1	Heterocycilic compounds  *Five-membered ring systems and Six- membered ring systems-nomenclature  *Aromaticity and structure  *Simple examples of electrophilic substitution reactions and definition of the Fused rings: Indole, Benzofuran, Benzothiopheme, Quinoline and Isoquinoline	Based on the instructor's selection	
		Final Examination		

Approved by Dept. Chair	Date of Approval	

## **Extra Information**: (Updated every semester and filled by course instructor)

Course Instructor	
Office No.	
Extension	
Email	
Office hours	