



1	Course	N	ame:
	Course		anne

#### Organic Pharmaceutical Chemistry IV

2. Course Code:

#### 547 ChPOp4

3. Semester / Year:

1stSemester/5thYear

4. Description Preparation Date:

9-2025

5. Available Attendance Forms:

Students' signatures on attendance sheets

6. Number of Credit Hours (Total) / Number of Units (Total)

2 hours theory (30) / 2 units

7. Course administrator's name (mention all, if more than one name)

Name: Lecturer Dr. Marwan Imad

Email: marwan.imad.jihad@bcms.edu.iq

#### 8. Course Objectives

Course Objectives	<ul> <li>Introducing</li> </ul>	the	students	to	advanced	concepts
	pharmaceutica	al chem	istry such as	prodr	ugs, drug targ	eting
	combinatorial	chemis	stry			

#### 9. Teaching and Learning Strategies

Strategy

• Theory lectures with teaching aids such as videos and diagrams

#### 10. Course Structure

Week	Hours	Paguirad Lagraing	Unit or subject name	Lagraina	Evaluation
week	Hours	Required Learning	Unit or subject name	Learning	
		Outcomes		method	method
1-3	6	Understanding	Basic concept of prodrugs;	• Lectures	• Paper-
		the concept of	Covalent bonds (cleavable);		based exams
		prodrugs	Prodrugs of functional groups;		
			Types of prodrugs		
4-6	6	Understanding	• Chemical delivery systems;	• Lectures	• Paper-
		the role of	Polymeric prodrugs; Types and		based exams
		polymers as	structure of polymers; Cross-		
		delivery systems	linking reagents		
		for drugs			
7+8	4	Understanding	Drug targeting	• Lectures	• Paper-
		the concept of			based exams
		targeting drugs			
		to specific			
		tissues and			
		organs			





8-15	14	Understanding	Combinatorial chemistry;	• Lectures	• Paper-
		the concept of	Peptides and other linear		based Exams
		combinatorial	structures; Drug like		
		chemistry and	molecules; Support and		
		library design	linker; Solution- phase		
			combinatorial chemistry		
			Detection, purification and		
			analgesics; Encoding		
			combinatorial libraries;		
			High- throughput screening;		
			Virtual screening; Chemical		
			diversity and		
			library design		
44.6	- 1				

#### 11. Course Evaluation

- 30 M: Theoretical assessment (paper-based midterm exam, attendance)
- 70 M: paper-based theoretical final exam

100 Marks total

100 Marks total						
12. Learning and Teaching Resources	12. Learning and Teaching Resources					
Required textbooks (curricular books, if any)	Wilson and Gisvold Textbook of Orga medicinal and					
	Pharmaceutical chemis Delgado JN, Remers WA, (Eds);					
	12thediti 2010					
Main references (sources)	Wilson and Gisvold Textbook of Orga medicinal and					
	Pharmaceutical chemis Delgado JN, Remers WA, (Eds);					
	12thediti 2010					
Recommended books and references						
(scientific journals, reports)						
Electronic References, Websites						
Update percentage	1 % change in the theoretical lectures					





#### 1. Course Name:

#### Industrial Pharmacy I

2. Course Code:

#### 548 Phlp2

3. Semester / Year:

first Semester

4. Description Preparation Date:

9-2025

5. Available Attendance Forms:

On campus

- 6. Number of Credit Hours (Total) / Number of Units (Total):
- 2 hours/week (Theory), 2hours/week (Practical), Total units=4
- 7. Course administrator's name (mention all, if more than one name)

Theory:

Name: Prof. Dr.Alaa Abdul Hussein Email: alaa.abdulhussein@bcms.edu.iq

Practical:

Name: Assistant lecturer Rana Kadum Email: ranakadum@bcms.edu.iq

#### 8. Course Objectives

#### Course Objectives

The subject aims to teach pharmacy students the steps and lines

Upon which the Per formulation processing of pharmaceutical dosage forms. This

fundamental course provides the

required principles to integrate knowledge of

Pharmaceutical Technology in Per formulation of perfect dosage form. It includes milling, mixing, drying and filtration,

besides sterilization to achieve proper processing of dosage for

9. Teaching and Learning Strategies

#### Strategy

- 1-Lectures and Presentation
- 2-Discussions
- 3- Laboratory experiments
- 4- Inverted classrooms

#### 10. Course Structure

Week	Hours	Required Learning Outcomes	Unit or subject name	Learning method	Evaluation method
1	3	Understand the Principles of pharmaceutical processing, mixing	fluid mixing; Flow characteristics; mechanisms of mixing; mixing equipment; batch and continuous mixing	- Lectures -White board -Data show -Power point -Explanatory	-Written exams - Oral exams -Laboratory reports
2	3	Knowledge of the mixer and best selection of mixer	batch and continuous mixing; mixer selection.	diagrams -Scientific YouTube	
3	3	Describe the Milling	pharmaceutical application of milling; size distribution and	videos -laboratory experiments	





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			measurement; Theory of	
			comminution	
		Understand types of	types of mills; factors	
		mills	influencing milling;	
4	3		selection of mill	
			techniques and techniques	
			of milling	
		Understand the	Definition of drying;	
		Drying industrial	purpose; Psychrometry	
5	3	process	(humidity measurement);	
		,	theory of drying; drying of	
			solids,	
		Define drying	classification of dryer;	
6	3	equipment's	specialized drying	
			methods	
		Understand process	Theory; filter media; filter	
		of Clarification and	aids; selection of drying	
7	3	filtration	method; non- sterile and	
			sterile operations.	
			integrity testing	
		Understand the	equipment's and systems	
		equipment's and	(commercial and	
	_	systems	laboratory) of filtration	
8	3	(commercial and		
		laboratory) of		
		filtration.		
		Describe	Sterilization; validation of	
		Sterilization;	methods; microbial death	
9	3	validation of	kinetics	
		methods; microbial		
		death kinetics		
		To understand	Methods of sterilization	
		Methods of	(thermal and non-	
10	3	sterilization	thermal); mechanisms;	
		Stermzation	evaluation.	
11	3	Describe	development; formulation	
''		Pharmaceutical	acveropment, formulation	
		Dosage forms;		
		sterile products		
12	3	Learn production,	production; processing;	
12		processing of sterile	quality control.	
		product	quanty control.	
		product		





#### 11. Course Evaluation

Midterm examination 15 marks

 $Quiz\ and\ classroom\ activities\ 5\ marks$ 

Practical part 20 marks

Final examination 60 Marks arks

### 12. Learning and Teaching Resources

12. Learning and reacting resources	
Required textbooks (curricular books, if any)	The Theory and Practice of Industrial Pharmacy by
	Leon Lachman et al.
Main references (sources)	Pharmaceutics: The Science of Dosage Form Design,
, ,	by Michael E. Aulton
Recommended books and references (scientific	Ansel's Pharmaceutical Dosage Forms and Drug
journals, reports)	Deliv Systems by Loyd Allen
Electronic References, Websites	





#### 1. Course Name:

#### Applied Therapeutics I

2. Course Code:

#### 549 ACIAt1

3. Semester / Year:

First semester/Fifth Year

4. Description Preparation Date:

09-2025

5. Available Attendance Forms:

On campus

- 6. Number of Credit Hours (Total) / Number of Units (Total)
- 3 Hours /3 Units
- 7. Course administrator's name (mention all, if/more than one name)

Name: Prof. Dr. Hayder Al-Tukmagi

E-mail: Tukmagi@bcms.edu.iq

#### 8. Course Objectives

#### **Course Objectives**

- The course provides students with basic knowledge about pathophysiology, symptoms and aims of treatment.
- In addition to the basic knowledge on the drug's use, kinetics, drug interactions, dose calculations, side effects, treatment algorithms and patient awareness are provided.

#### 9. Teaching and Learning Strategies

Strategy	Lectures
	Seminars
	Simple
	quizzes
	Brainstorming questions

#### 10. Course Structure

Wee	Hour	Re	quired Learning Outcomes	Unitor	Learnin	Evaluatio
k	S			subject name	g method	n method
1	2	1. 2. 3. 4. 5.	Differentiate between Sensitivity and Specificity of lab tests. Identify reference ranges of lab tests. Identify normal and abnormal liver function tests. Identify normal and abnormal renal function tests. Interpretation of complete blood count test results.			Simple quizzes.
		<ul><li>6.</li><li>7.</li></ul>	Interpretation of urinalysis main findings. Interpretation of hematological lab			
			investigations			





LEGE OF N		1.	Identify the common types of lipid disorders.	Dyslipidem	Lectures.	Simple
				ia.		•
		2.	Identify the statin-benefit groups and		Simple	quizzes.
		2	intensity of statin therapy.		discussions.	
		3.	Recommend appropriate therapeutic lifestyle			
			changes (TLC) and pharmacotherapy			
			interventions for dyslipidemia.			
		4.	'			
			cardiovascular disease risk and corresponding			
2	2	_	treatment goals.			
		5.	Identify patients who are indicated for non-			
			statin therapy.			
		6.	Describe components of a monitoring plan to			
			assess effectiveness and adverse effects of			
			pharmacotherapy for dyslipidemias.			
		7.	Educate patients about the state of the			
			disease, appropriate TLC, and drug therapy			
			required for effective treatment.			
		1.	Differentiate types of cerebrovascular disease	Stroke.	Lectures.	Simple
			including transient ischemic attack (TIA),		Simple	quizzes.
	1		ischemic stroke (cerebral infarction), and		discussions.	
			hemorrhagic stroke.			
		2.	Identify modifiable and nonmodifiable risk			
			factors associated with ischemic stroke and			
			hemorrhagic stroke.			
		3.	Explain the pathophysiology of ischemic			
			stroke and hemorrhagic stroke.			
		4.	Describe the clinical presentation of TIA,			
			ischemic stroke, and hemorrhagic stroke.			
3		5.	Formulate strategies for primary prevention			
			of acute ischemic stroke.			
		6.	Evaluate treatment options for acute ischemic stroke.			
		7.	Determine whether fibrinolytic therapy is			
			indicated in a patient with acute ischemic			
			stroke.			
		8.	Evaluate the role of endovascular therapy in a			
			patient with acute ischemic stroke.			
		9.	•			
			prevention of acute ischemic strokes.			
		10	. Evaluate treatment options for acute			
			hemorrhagic stroke.			
	<u> </u>		<u> </u>		<u> </u>	





		1.	Assess a patient's kidney function based on	Acute	Lectures.	Simple
			clinical presentation, laboratory results, and	kidney	Simple	quizzes.
			urinary indices.	injury	discussions.	
		2.	Identify pharmacotherapeutic outcomes and			
			endpoints of therapy in patients with acute			
			kidney injury (AKI).			
4	1	3.	Apply knowledge of the pathophysiology of			
			AKI to develop a treatment plan.			
		4.	Develop strategies to minimize the			
			occurrence of drug-induced AKI.			
		5.	Monitor and evaluate the safety and			
			effectiveness of the treatment plan.			
		1.	List the risk factors that increase susceptibility	Chronic and	Lectures.	Simple
			for chronic kidney disease (CKD).	end-stage	Simple	quizzes.
		2.	Explain the mechanisms associated with the	kidney	discussions.	
			progression of CKD.	disease.		
		3.	Outline  the  desired  outcomes  for  treatment  of			
			CKD.			
5	2	4.	Develop a therapeutic approach to slow			
3	۷		progression of CKD including lifestyle			
			modifications and pharmacologic therapies.			
		5.	Identify specific consequences associated			
			with CKD.			
		6.	Design an appropriate therapeutic approach			
			for specific consequences associated with			
			CKD.			
		1.	Identify indications for dialysis.	Hemodialy	Lectures.	Simple
		2.	List advantages and disadvantages of	sis and	Simple	quizzes.
		_	hemodialysis and peritoneal dialysis.	peritoneal	discussions.	
6	1	3.	1 1 1	dialysis.		
			hemodialysis and peritoneal dialysis.			
		4.	Identify complications of hemodialysis and			
			peritoneal dialysis and their management.			
		1.	Definition of pharmacovigilance.	Pharmacovi	Lectures.	Simple
		2.	Recognize who should report the	gilance e.	Simple	quizzes.
			pharmacovigilance reports.		discussions.	
_	4	3.	Describe the importance of			
7	1		pharmacovigilance.			
		4.	Historical events reported ADRs.			
		5.	Describe Causality Assessment.			
		6.	Identify terms used in pharmacovigilance.			
		7.	Identify the importance of pharmacovigilance.			





GE OF W		1.	Explain the pathophysiology of cirrhosis and	Cirrhosis	Lectures.	Simple
		1.	portal hypertension.	and portal	Simple	quizzes.
		2.	Identify signs and symptoms of cirrhosis.	hypertensio	discussions.	quizzes.
		3.	Identify laboratory abnormalities that result	n.	discussions.	
		٥.	from liver disease and describe the associated	11,		
			pathophysiology.			
8	2	4.	Describe the consequences associated with			
			decreased hepatic function.			
		5.	Identify treatment goals for a patient with			
			complications of cirrhosis.			
		6.	Recommend a specific treatment regimen for			
			a patient with cirrhosis that includes lifestyle			
			changes, nonpharmacologic measures, and			
			pharmacologic therapy.			
		1.	Differentiate the five types of viral hepatitis by	Viral	Lectures.	Simple
			epidemiology, etiology, and clinical	hepatitis.	Simple	quizzes.
			presentation.		discussions.	
		2.	Identify modes of transmission and risk			
			factors among the major types of viral			
			hepatitis.			
		3.	Evaluate hepatic serologies to understand			
9	1		how the type of hepatitis is diagnosed.			
		4.	Create treatment goals for a patient infected			
			with viral hepatitis.			
		5.	Recommend appropriate pharmacotherapy			
			for prevention of viral hepatitis.			
		6.	Develop a care plan for treatment of			
			chronic viral hepatitis.			
		1.	Characterize the pathophysiologic	Inflam	Lectures.	Simple
			mechanisms underlying inflammatory bowel	matory	Simple	quizzes.
			disease (IBD).	bowel	discussions.	
		2.	Recognize the signs and symptoms of IBD,	disease		
			including major differences between			
			ulcerative colitis (UC) and Crohn disease	•		
			(CD).			
10	1	3.	Identify appropriate therapeutic outcomes for			
			patients with IBD.			
		4.	Describe pharmacological treatment options			
			for patients with acute or chronic symptoms			
			of UC and CD.			
		5.	Create a patient-specific drug treatment plan			
			based on symptoms, severity, and location of			
			UC or CD.			





		6.	Recommend appropriate monitoring			
			parameters for drug treatments for IBD			
		1.	List the types and etiologies of shock	Shock	Lectures.	Simple
			syndromes.	syndromes.	Simple	quizzes.
		2.	Describe the major hemodynamic		discussions.	
			abnormalities that occur in patients with			
			shock.			
		3.	Describe the clinical presentation including			
			signs, symptoms, and laboratory test			
11	1		measurements for the typical shock patient.			
		4.	Prepare a treatment plan with clearly defined			
			outcome criteria for a shock patient that			
			includes both fluid management and			
			pharmacologic therapy.			
		5.	$Compare\ and\ contrast\ the\ relative\ advantages$			
			and disadvantages of crystalloids, colloids,			
			and blood products in the treatment of shock.			
		1.	Estimate the volumes of various body fluid	Disor	Lectures.	Simple
			compartments.	ders	Simple	quizzes.
		2.	Identify the electrolytes primarily found in the	of	discussions.	
	2		extracellular and intracellular fluid	fluids		
			compartments.	and		
12		3.	Describe the unique relationship between	electr		
	_		serum sodium concentration and total body	olytes		
			water (TBW).			
		4.	Review the etiology, clinical presentation, and			
			management for disorders of sodium,			
			potassium, calcium, phosphorus, and			
			magnesium.	F 1		
		1.	Describe the epidemiology and social impact	Epilepsy.	Lectures.	Simple
		2	of epilepsy.		Simple	quizzes.
		2.	Define terminology related to epilepsy,		discussions.	
		3.	including seizure, convulsion, and epilepsy.			
		3.	Describe the basic pathophysiology of			
		4.	seizures and epilepsy.  Differentiate and classify seizure types given a			
13	1	٦.	description of the clinical presentation of the			
			seizure and electroencephalogram.			
		5.	Identify key therapeutic decision points and			
		].	therapeutic goals in the treatment of epilepsy.			
		6.	Discuss nonpharmacologic treatments for			
		.	epilepsy.			
		7.	Recommend an appropriate			
		, <b>.</b>	The second and appropriate			





		pharmacotherapeutic regimen with			
		monitoring parameters for the treatment of			
		epilepsy.			
		8. Devise a plan for switching a patient from one			
		antiepileptic regimen to a different regimen.			
		9. Manage potential drug interactions with			
		antiepileptic drugs (AEDs).			
		10. Determine when and how to discontinue AED therapy.			
		1. Identify risk factors for multiple sclerosis	Multiple	Lectures.	Simple
		(MS).	sclerosis.	Simple	quizzes.
		2. Distinguish between forms of MS based on		discussions.	
		patient presentation and disease course.			
		3. Compare and contrast MS			
14	1	disease-modifying treatment choices for a given			
		patient. 4. Determine appropriate symptomatic			
		treatment choices for a given patient.			
		5. Develop a monitoring plan for a patient			
		placed on specific medications.			
		1. Evaluate patient-specific parameters to	Enteral	Lectures.	Simple
		determine whether EN is appropriate.	nutrition.	Simple	quizzes.
		2. Compare clinical efficacy, complications, and		discussions.	
		costs of EN versus parenteral nutrition (PN).			
		3. Describe the components of EN and their role			
		in nutrition support therapy.			
		4. Develop a plan to design, initiate, and adjust			
15	1	an EN formulation for an adult patient based			
		on patient- specific factors.			
		5. Describe the etiology and risk factors for EN-			
		associated complications in adult patients			
		receiving EN.			
		6. Select appropriate medication administration			
		techniques for an EN patient.			
		1. List appropriate indications for parenteral	Paren	Lectures.	Simple
		nutrition (PN) in adult patients.	teral	Simple	quizzes.
		2. Describe the components of PN and their role	nutriti	discussions.	
		in nutrition support therapy.	on.		
10	4	3. Develop a plan to design, initiate, and adjust a			
16	1	PN formulation for an adult patient based on			
		patient- specific factors.			
		4. Describe the etiology and risk factors			
		for PN macronutrient-associated			
		complications in adult patients			





			receiving PN.			
		5.	Describe the etiology and risk factors			
			for refeeding syndrome, as well as measure to			
			prevent refeeding syndrome.			
		1.	Identify risk factors and signs and symptoms	Deep	Lectures.	Simple
			of deep vein thrombosis (DVT) and	veno	Simple	quizzes.
			pulmonary embolism (PE).	us	discussions.	
		2.	Describe the processes of hemostasis and	thro		
			thrombosis.	mbos		
		3.	Determine a patient's relative risk of	is.		
			developing venous thrombosis.			
		4.	Formulate an appropriate prevention strategy			
			for a patient at risk for DVT.			
		5.	Select and interpret laboratory test(s) to			
			monitor antithrombotic drugs.			
		6.	Identify factors that place a patient at high risk			
			of bleeding while receiving antithrombotic			
4=			drugs.			
17	1	7.	State at least two potential advantages of			
			newer anticoagulants (ie, low molecular			
			weight heparins [LMWHs], fondaparinux,			
			oral direct thrombin inhibitors [DTIs], and			
			$or al\ direct\ factor\ Xa\ in hibitors)\ over\ traditional$			
			anticoagulants (ie, unfractionated heparin			
			and warfarin).			
		8.	Manage a patient with toxicity secondary to			
			warfarin (elevated international normalized			
			ratio [INR] with or without bleeding).			
		9.	Identify anticoagulant drug-drug and drug-			
			food interactions.			
		10	. Formulate an appropriate treatment plan for a			
			patient who develops a DVT or PE.			
		1.	Describe the phases of cardiac action	Arrhythmias.	Lectures.	Simple
			potential.		Simple	quizzes.
		2.	Describe the modified Vaughan Williams		discussions.	
			classification of antiarrhythmic drugs.			
		3.	Compare and contrast risk factors for and			
18	2		features, mechanisms, etiologies, symptoms,			
			and goals of therapy of (a) sinus bradycardia,			
			(b) atrioventricular (AV) block, (c) atrial			
			fibrillation (AF), (d) paroxysmal			
			supraventricular tachycardia (PSVT), (e)			
			premature ventricular complexes (PVCs), (f)			





			ventricular tachycardia (VT, including			
			, and the second			
			torsades de pointes [TdP]), and (g) ventricular			
			fibrillation (VF).			
		4.	Compare and contrast appropriate treatment			
			options for sinus bradycardia and AV block.			
		5.	Compare and contrast mechanisms of action			
			of drugs used for ventricular rate control,			
			conversion to sinus rhythm and			
			maintenance of sinus rhythm in patients with			
			AF.			
		6.	Compare and contrast the advantages and			
			disadvantages of warfarin and the non-			
			vitamin K antagonist oral anticoagulants			
			(NOACs) for prevention of stroke and			
			systemic embolism in patients with AF.			
		7.	Discuss nonpharmacologic methods for			
			termination of PSVT, compare mechanisms			
			of action of drugs used for acute			
			termination of PSVT and compare			
			appropriate treatment options for long-term			
			prevention of PSVT recurrence.			
		8.	Compare and contrast mechanisms of action			
			of drugs used for treatment of acute episodes			
			of VT and describe options and indications for			
			nonpharmacologic treatment of VT and VF.			
		9.	Design individualized drug therapy treatment			
			plans for patients with (a) sinus bradycardia,			
			(b) AV block, (c) AF, (d) PSVT, (e) PVCs, (F)VT			
			(including TdP), and (g) VF.			
		1.		Pain	Lectures.	Simple
		••	nociceptive, inflammatory, neuropathic, and	mana	Simple	quizzes.
			functional.	geme	discussions.	70.2200.
		2.	Explain the mechanisms involved in pain	nt.	discussions.	
		2.	transmission.	116.		
		3	Select an appropriate method of pain			
		٥.	assessment.			
19	2	4.				
		4.	Recommend an appropriate choice of			
			analgesic, dose, and monitoring plan for a			
			patient based on type and severity of pain and			
		_	other patient- specific parameters.			
		5.	Perform calculations involving equianalgesic			
			doses, conversion of one opioid to another,			
			rescue doses, and conversion to a continuous			





			infusion.			
		6.	Educate patients and caregivers about			
			effective pain management, dealing with			
			chronic pain, and the use nonpharmacologic			
			measures.			
		1.	Differentiate types of headache syndromes	Headache.	Lectures.	Simple
			based on clinical features.		Simple	quizzes.
		2.	Recommend nonpharmacologic measures		discussions.	
			for headache treatment and prevention.			
		3.	Determine when the pharmacologic			
20	1		treatment of headache is indicated.			
		4.	Construct individualized treatment regimens			
			for the acute and chronic management of			
			headache syndromes.			
		5.	Monitor headache treatment to ensure its			
			safety, tolerability, and efficacy.			
		1.	Describe the pathophysiology of Parkinson	Parki	Lectures.	Simple
			disease (PD) related to neurotransmitter	nson'	Simple	quizzes.
			involvement and targets for drug therapy.	S	discussions.	
		2.	Recognize the  cardinal  motor  symptoms  of  PD	diseas		
			and determine a patient's clinical status and	e.		
			disease progression.			
		3.	For a patient initiating therapy for PD,			
			recommend appropriate drug therapy and			
			construct patient- specific treatment goals.			
		4.	Recognize and recommend appropriate			
21	2		treatment for nonmotor symptoms.			
		5.	Formulate a plan to minimize patient "off-			
			time" and maximize "on-time" including			
			timing, dosage, and frequency of medications.			
		6.	Recognize and treat various motor complications in PD.			
		7.	•			
			regarding medications and lifestyle			
			modifications for PD.			
		8.	Develop a monitoring plan to assess			
			effectiveness and adverse effects of			
			treatment.			
		1.	Explain the pathophysiology of benign	Benign	Lectures.	Simple
			prostatic hypertrophy (BPH).	prostatic	Simple	quizzes.
22	1	2.	Recognize the symptoms and signs of BPH.	hyperpla	discussions.	
		3.	List the desired treatment outcomes for BPH.	sia.		
		4.	Identify factors that guide selection of a			
<u> </u>	L	1	<u> </u>	<u> </u>	<u> </u>	





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			particular $\alpha$ 1- adrenergic antagonist for an individual patient.			
		5	Compare and contrast α1- adrenergic			
		٥.	antagonists versus $5\alpha$ -reductase inhibitors in			
			terms of mechanism of action, treatment			
		6	outcomes, adverse effects, and interactions.			
		0.	Describe the indications, advantages, and			
			disadvantages of various combination drug			
			regimens that include an $\alpha$ 1-adrenergic			
			antagonist, $5\alpha$ -reductase inhibitor,			
			anticholinergic agent, tadalafil, or mirabegron.			
		7.	Describe the indications for surgical			
			intervention.			
		8.	Apply the patient care process to develop			
			an individualized treatment plan.			
		1.	Identify risk factors for the development of	Glaucoma.	Lectures.	Simple
			primary open-angle glaucoma (POAG) and		Simple	quizzes.
			acute angle- closure glaucoma.		discussions.	
		2.	Recommend a frequency for glaucoma			
			screening based on patient-specific risk			
			factors.			
		3.	Compare and contrast the			
			pathophysiologic mechanisms responsible			
			for open-angle glaucoma and acute angle-			
			closure glaucoma.			
		4.	Outline the clinical presentation of chronic			
			open- angle glaucoma and acute angle-			
22	4		closure glaucoma.			
23	1	5.	List the goals of managing patients with			
			POAG suspect, POAG, and acute angle-			
			closure glaucoma.			
		6.	Choose the most appropriate therapy based			
			on patient- specific data for open-angle			
			glaucoma, glaucoma suspect, and acute angle-			
			closure glaucoma.			
		7.	Develop a monitoring plan for patients on			
			specific pharmacologic regimens.			
		8.	Counsel patients about glaucoma, drug			
			therapy options, ophthalmic administration			
			techniques, and the importance of adherence			
			to the prescribed regimen.			
11. (	Course	e Eva	aluation			

Midterm exam 25 marks, Quizzes and attendance 5 marks, Final exam 70 marks





12. Learning and Teaching Resources					
Required textbooks (curricular books,	Clinical pharmacy and therapeutics. Pharmacotherapy handbook.				
if any)	ACCP updates in therapeutics.				
	Pharmacotherapy: A pathophysiologic approach.				
	Pharmacotherapy: principles and practice. Applied therapeutics.				
Main references (sources)	Pharmacotherapy: Pathophysiologic approach.				
	Pharmacotherapy: principles and practice.				
	Applied therapeutics. ACCP updates in therapeutics.				
Recommended books and references	Pharmacotherapy: A pathophysiologic approach.				
(scientific journals, reports)	Pharmacotherapy: principles and practice				
Electronic References, Websites	Electronic books and review articles.				





1. Course Name:
Clinical Chemistry
2. Course Code:

#### 550 ACICc

3. Semester / Year:

First semester / Fifth

4. Description Preparation Date:

9-2025

5. Available Attendance Forms:

In-person attendance

6. Number of Credit Hours (Total) / Number of Units (Total)

3 hours theory+ 2 hours practical (75) / 4 units

7. Course administrator's name (mention all, if more than one name)

Theory:

Name: Assistant prof. Dr. Zainab Al-Shamaa

Email: z.alshamma@bcms.edu.iq

Practical:

Name: Assistant lecturer Yousef Alwan Email: yousefalwan@bcms.edu.iq

#### Course Objectives

## **Course Objectives** 1) Understanding of human body chemistry in both healthy and diseased states, enabling to diagnose, monitor, and manage disease through laboratory data analysis. 2) Interpreting the results of biochemistry analyses that augment the clinical examination to achieve definite diagnosis of the disease. 3) Evaluating data accuracy and applying this knowledge to therapeutic decisionmaking and patient care.

9. Teaching and Le	9. Teaching and Learning Strategies						
Strategy	Presentation and recitation						
	Reading & research						
	Interactive discussions						
	Brainstorming						

#### 10. Course Structure

Week	Hours	Required Learning Outcomes	Unit or subject name	Learning method	Evaluation method
1	4	Disorders of Carbohydrates metabolism, Hyperglycemia & Diabetes mellitus, Hypoglycemia	Carbohydrates disorders	Lectures, discussions, and reports	Theoretical exam, and classroom activities
2	3	Understanding the abnormalities in the metabolism of lipids and the laboratory assessment	Disorders of lipid metabolism	=	Ш





		Understanding of the	Liver function tests		
		metabolic, synthetic and			
	,	excretory functions of the		_	_
3	3	liver and the related		=	=
		disorders; and the laboratory			
		assessment of liver functions			
		Understanding of the excretory	Kidney function		
		functions of the kidney and its	tests		
		role in maintaining blood			
		hemostasis and elimination of			
		waste products			
4	3	Study of the acute and chronic		=	=
		kidney diseases and the			
		laboratory tests of kidney			
		functions; and types of kidney			
		stones			
		Study of different diseases			
5	3	associated with change in	Diagnostic	=	=
	5	Enzymatic Activity in blood	enzymology		
		Understand of hormone	Hypothalamus &		
		types, functions and	pituitary		
		regulation, with special	endocrinology,		
		emphasis on the	adrenal gland		
		hypothalamic hormones			
6-7	6	The pituitary gland hormones			_
0-7	0	actions and disorders; and the		=	=
		laboratory analyses of			
		pituitary gland disorders			
		The adrenal gland hormones			
		actions and disorders; and the			
		laboratory analyses of adrenal			
		gland disorders The male and female			
		reproductive glands hormones	Reproductive		
8-9	6	and the physiologic and	glands hormones		
		pathologic alterations in their	and diseases		
		levels			
		The thyroid gland hormones	Thomasdal		
	_	actions and disorders; and the	Thyroid gland		
10	3	laboratory analysis of thyroid	hormones and	=	=
		gland disorders	diseases		
11	3	Drug interaction with	Drug interaction	=	=
L	<u>I</u>	1			





		laboratory Tests	with laboratory	
			Tests	
		Disorders of calcium	Disorders of	
12	3	metabolism	calcium	
			metabolism	
	2	Study of different tumor	Tumor markers	
12		markers in blood that can be		
13		used for detection and		
		monitoring tumors		
14-15	6	Inborn errors of metabolism	Inborn errors of	
14-15		indom endrs of metabolism	metabolism	

#### 11. Course Evaluation

Midterm examination 15 marks

 $Quiz\ and\ classroom\ activities\ 5\ marks$ 

Practical part 20 marks

Final examination 60 Marks arks

42 Lauriana J.T. akina D						
12. Learning and Teaching Resources						
Required textbooks (curricular books,	1) Crook, Clinical Chemistry & Metabolic Medicine, 8th edition					
if any)	2) Tietz Clinical Chemistry& Molecular Diagnostics 6th edition;					
	2018					
	3) Kaplan,Clinical Chemistry, 5th edition					
Main references (sources)	Tietz Clinical chemistry& Molecular Diagnostics 7th edition;					
	2015.					
Recommended books and references	Clinical Chemistry, Kaplan 2012					
(scientific journals, reports)						
Electronic References, Websites						





1.	Course	Ν	ame:

#### Hospital training

2. Course Code:

#### 551 AClHt

3. Semester / Year:

Second semester/Fifth

4. Description Preparation Date:

9-2025

5. Available Attendance Forms:

Training in hospital

6. Number of Credit Hours (Total) / Number of Units (Total)

Four hours practical (60) / 2 Units

7. Course administrator's name (mention all, if more than one name)

Name: Kawther Faris

Email: kawther.kf@gmail.com

Name: Assistant lecturer Ibraheem Kais
Email: <u>ibraheem.kais0@bcms.edu.iq</u>
Name: Assistant lecturer Ameer Ali Kazal

Email: ameerali@bcms.edu.iq

#### 8. Course Objectives

#### Course Objectives

To teach students the application of pharmacy practice in differ hospital wards; it

- Training in case evaluation and follow up
- Evaluation of the rapeutic regimens and registration of err related to drug therapy and presenting ideas to solve problems

#### 9. Teaching and Learning Strategies

#### Strategy

Explaining cases of patients in different hospital wards

Discussions with board students in hospital Brainstorming

questions

#### 10. Course Structure

Wee	Hours	Required Learning	Unitor	Learning	Evaluation
k		Outcomes	subject	method	method
			name		
1-4	4	To provide the students the essential clinical pharmacy skills with emphasis on dealing with patients, medical charts laboratory information, and clinical monitoring. The following topics will be covered: (Cardiology, Nephrology, Gastroenterology, Pulmonology, and Endocrinology)	Internal medicine	Explanation of patients cases and discussion with Board students	Quizzes and case presentation





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		To provide the students the essential	pediatrics	Explanation of	Quizzes and case
		clinical pharmacy skills with emphasis on		patients	presentation
		dealing with patients, medical charts,		cases and	
		laboratory information, and clinical		discussion	
		monitoring. The following topics will be		with Board	
		covered:		students	
5-8	4	Pediatric Neonatology,			
5-8	4	Pediatric Nephrology,			
		Pediatric Infections,			
		Pediatric Neurology,			
		Pediatric Cardiology,			
		Pediatric Gastroenterology,			
		Pediatric Respiratory Disorders, and			
		pediatric endocrinology			
		To provide the students the	surgery	Explanation	Quizzes and case
		essential clinical pharmacy skills with		of patients	presentation
		emphasis on dealing with patients,		cases and	
		medical charts, laboratory information, and		discussion	
		clinical monitoring. The following topics		with Board	
		will be covered:		students	
		Surgical Prophylaxis, Types of Surgical			
		Operations, Preoperative bowel			
9-12	4	preparation, Intravenous fluid therapy,			
		Blood transfusion and blood products, Peri-			
		operative care and diabetes,			
		Perioperative medication			
		management, Acute appendicitis,			
		Gallstones, Common bile duct stones,			
		Thyroidectomy, Bowel Obstruction,			
ſ		Pancreatitis, Hernia, Guidelines on			
ſ		Parenteral Nutrition in Surgery.			





13-16	4	To provide the students the essential	Obstetrics	Explanation of	Quizzes and case	
		clinical pharmacy skills with emphasis on	gynecology	patients cases and	presentation	
		dealing with patients, medical charts,		discussion with		
		laboratory information, and clinical		Board students		
		monitoring. The following topic will be				
		covered: Abortion, Common				
		Complications of Pregnancy, Induction				
		and Augmentation of labor, Obstetric				
		hemorrhage, Caesarean section, Ectopic				
		Pregnancy, Heavy and irregular				
		Menstruation, Polycystic Ovarian				
		Syndrome, Molar Pregnancy, some drugs				
		that are used in obstetrics and				
		gynecology				
11	. Course E	valuation				
34 qu	izzes, 6 ca	se presentation, 60 final exams.				
12	. Learning	and Teaching Resources				
Requir	ed textboo	oks (curricular books, if any)		Clinical Training A	dopted by the	
Main ro	eferences	(sources)	Manuals for Clinical Training Adopted by Department			
Recom	mended	books and references	Pharmacy times (journal) Us pharmacist (journal)			
		ls, reports)		· ·	, 	
Electro	nic Refere	nces, Websites	UpToDate resource, Medscape			





Oral and

written exams

Lectures

Metformin

toxicity

1. C	1. Course Name							
Clin	Clinical Toxicology							
2.	2. Course Code							
552	ACICt							
3. S	emester/\	/ear						
1 <sup>st</sup> s	semester / I	ifth ye	ar					
4. C	escription	Prepara	ation Date:					
9-2	025							
5. A	wailable At	tendan	nce Forms					
on	campus							
6. N	Number of (	Credit H	Hours (Total) / Number of Un	its (Total)				
2 h	ours theore	etical + :	2 hours practical (60) / 3 unit	S				
7. C	Course adm	inistrat	tor's name (if more than one r	name)				
The	eory:							
Nar	ne: Assista	nt prof.	. Elham Mahmood					
Ema	ail: <u>elham.r</u>	<u>nahmo</u>	ood@bcms.edu.iq					
pra	ctical:							
Nar	ne: Assista	nt lectu	ırer Muhee Nimma					
Ema	ail: <u>muhee.</u>	nimma	salman@bcms.edu.iq					
8. Cou	rse Objecti	ves						
			Clinical toxicology aims to	study the harmful	effects of agents a	and drugs on		
Course (	rse Objectives humans,							
			Study how to diagnose, tre	eat and prevent the	e harmful effects o	ftoxic		
			substances.					
9. Tead	ching and L	earnin	g Strategies					
			Lecturing					
	strategy		Homework					
	Quiz							
10. Co	ourse Struc	ture						
The	The Hours Required Learning Outsomes			Unit or subject	Learning	Evaluation		
week	week Hours Required Learning Outcomes			name	method	method		
		Stude	nts will be able to study					
		how t	o assess the general					
1	3	physic	cal health of a person, give	Vital signs				

clues to possible diseases, and

show progress toward recovery.

Students will be able to study

interventions and outcomes of

how to analyses the

symptomology, clinical

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		patients presenting with severe		
		metformin toxicity.		
		Students will be able to study <i>the</i>		
		etiology of hydrocarbon toxicity	Hydrocarbon	
3	2	and the typical presentation of a	Toxicity	
		patient with hydrocarbon		
		toxicity.		
		Students will be able to study the		
4	2	signs and symptoms of toxicity	ACEI poisoning	
		and treatment of toxicity.		
		Students will be able to study the		
		objective of poison centers and		
5	2	how to reduce morbidity and	Poison centers	
		mortality associated with		
		poisonings.		
		Students will be able to study the		1
	-	caustics and corrosives and how	Caustics and	
6	2	cause tissue injury by a chemical	corrosives	
		reaction.		
		Students will be able to study the		1
7	2	sign and symptom of toxicity and	Opioid toxicity	
		antidote of toxicity.		
		Students will be able to study the		1
8		potential toxicity of NSAIDs	NSAID toxicity	
		Students will be able to study the		
9	2	sign and symptoms of toxicity	Adrenergic	
		and treatment of toxicity.	toxicity	
		Students will be able to study the		
		toxicokinetic of cocaine toxicity,		
		physical exam findings for a		
		patient with cocaine toxicity and	Cocaine	
10	2	the management options for	toxicity	
		cocaine toxicity.	toxicity	
		the management options for		
		cocaine toxicity		
		· · · · · · · · · · · · · · · · · · ·		1
		Students will be able to study the		
11	2	most common drugs and chemicals associated with	Dysrhythmia	
11	۷		toxicity	
		ventricular dysrhythmias and		
		their outcomes.	D: :	١
12	3	Students will be able to study the	Digoxin	
		etiology of digoxin toxicity, the	toxicity	



Web sites



OLLEGE OF MEDICA					1000
		pathophysiology of digoxin			
		toxicity. And the management of			
		digoxin toxicity.			
11. Co	ourse Evalu	ation	·		
15 mark	s: Theoreti	cal Midterm Exam:			
20 marl	ks: Practica	I			
5 marks	: Daily Exa	ms			
Marks a	rks: final e	xam 60			
12. Le	earning and	Teaching Resources			
Require	d textbool	ks (curricular books, if any)	Goldfrank's To	xicologic Emerger	ncies, Ninth
			Edition		
Main re	ferences (	sources)	Toxicology Re	call Toxicology Re	call 1st Edition
		oks and references			





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#### **Pharmacoeconomics**

2. Course Code:

#### 554 ACIPco

3. Semester / Year:

2<sup>nd</sup> semester/5<sup>th</sup> year students

4. Description Preparation Date:

2-9-2025

5. Available Attendance Forms:

Class attendance (on-campus)

6. Number of Credit Hours (Total) / Number of Units (Total)

Two hours theory (30) /2 Units

7. Course administrator's name (mention all, if more than one name)

Name: Lecturer Dr. Humam Tawfiq Hadi

Email: humam.hadi@bcms.edu.iq
Name: Lecturer Dr. Nawfel Ayad
Email: nawfel.ayad@bcms.edu.iq

#### 8. Course Objectives

# Course Objectives Understand the basic terms of Pharmacoeconomics, how to build the model for economic feasibility studies, and how to extract statistical data from clinical studies to include them in the model for the economic feasibility study.

#### 9. Teaching and Learning Strategies

Strategy Lecturing Quiz

#### 10. Course Structure

Week	Hours	Required Learning Outcomes	Unit or subject	Learning	Evaluation
			name	method	method
		1. Introducing Pharmacoeconomic	Basic principle of	Interactive	Simple
		principles.	Pharmacoeconomics	lectures and	quizzes
		2. Demonstrate types of healthcare		related	
		costs with examples		articles	
1	4	3. Learn about ECHO models for			
		the 3 patient outcome types.			
		4. Explain and differentiate among			
		the 4 methods of			
		Pharmacoeconomic analyses.			
		1. Identifying costs	Cost analysis	Interactive	Simple
		2. Types of costs (Direct Medical		lectures and	quizzes
2	4	Costs, Direct Nonmedical Costs,		related	
		Indirect costs, Intangible costs)		articles	
		3. Incremental costs and marginal			





		,			
		COSTS			
		4. Opportunity costs			
		5. How are costs valued? Timin			
		Adjustments for Costs			G. I
		1. Understand the Cost-	Cost-minimizing	Interactive	Simple
		effectiveness analysis	analysis and Cost	lectures and	quizzes
3	4	2. Outcome measures in cost-	effectiveness	related	
		effectiveness analysis	analyses (CEA).	articles	
		3. Knowing how to calculate Cost-			
		effectiveness Ratios			
		1. Understand the Cost-Benefit	Cost-benefit	Interactive	Simple
		Analysis method.	analysis (CBA)	lectures and	quizzes
		2. Knowing how to calculate the		related	
		indirect cost of the disease and		articles	
		indirect benefit of the			
		intervention/program			
		3. Using Huma Capital Method			
		(HCM).			
		4. Using HCM to calculate Daily			
		wage rate and Missed days to			
4	4	find out the indirect benefit of			
		the intervention/ management.			
		5. Describe in detail the			
		Willingness-to-Pay Method			
		(WTP): Hypothetical Scenario			
		& Bidding Vehicles			
		6. Formats for presenting Cost-			
		Benefit Analysis (CBA) When we			
		should select			
		7. Cost-Benefit or Cost-			
		Effectiveness Analysis?			
		1. Use of decision analysis to	Critical assessment	Interactive	Simple
		design economic evaluations	of economic	lectures and	quizzes
		2. Decision Analysis Structure or	evaluation	related	'
5	4	tree		articles	
		Define Cost of illness	Drug-focused versus	Interactive	Simple
		2. Knowing how to calculate Cost	disease-focused	lectures and	quizzes
		of illness	framework for	related	1
		3. Understand the difference	conducting	articles	
		between healthcare costs and	Pharmacoeconomic	articles	
6	4	the cost of illness			
		the cost of filless	analyses.		





COLLEGE OF MED	DICAL						4 77.	
		the students should be at	le to:	Introduction	to	Interactive	Simple	
		1. define epidemiology,	describe	epidemiology.		lectures	quizzes	
		basic terminology and	concepts			and related		
7	4	of epidemiology.				articles		
		2. identify types of data	sources.					
		3. Identify basic methods	of data					
		collection and interpr	etation.					
		Cost-Effectiveness projec	t can be	Project present	tation.		Presentation	
		assigned to teach stud	dents how				skills	
8	2	to understand the terr	ninologies					
		used in	published					
		Pharmacoeconomic st	udies.					
11. C	ourse l	Evaluation						
	4	points for quizzes, 5 points	for assignn	nents, 20 points	for midte	erm exam and 70	o points for the	
		final exam						
12. L	earning	g and Teaching Resources						
Requi	ired tex	tbooks (curricular books,	Bootman	Bootman JL, Townsend RJ, McGhan WF, (Eds.), Principles				
if any	)		Pharmacoeconomics, 2 <sup>nd</sup> ed., Harvey Whitney Books Compa					
			Cincinnati, Oh, latest edition					
			Renée J.G. Arnold. Pharmacoeconomics From Theory to Practice.					
			Second Edition, 2021. CRC Press, Boca Raton, FL, USA					
Main	referer	nces (sources)						
D								
		led books and references		/alue in Health Journal /alue in Health   Journal   ScienceDirect.com by Elsevier				
(scien	ilitic joi	urnals, reports)					<u>sevier</u>	
			Value in Health Journal Regional Issues  https://www.valuehealthregionalissues.com/					
					_			
Electronic References, Websites			Value in F	'alue in Health Journal and Value in Health Journal Regional				

Issues





4	_	N I
1	COURSE	Name:

#### Applied Therapeutics II

2. Course Code:

#### 555 ACIAt2

3. Semester / Year:

Second semester/Fifth Year

4. Description Preparation Date:

9- 2025

5. Available Attendance Forms:

On campus

6. Number of Credit Hours (Total) / Number of Units (Total)

2 Hours theory (30) /2 Units

7. Course administrator's name (mention all, if more than one name)

Name: Prof. Dr. Hayder Al-Tukmagi

E-mail: Tukmagi@bcms.edu.iq

#### 8. Course Objectives

#### **Course Objectives**

- It provides students with basic knowledge about pathophysiology, symptoms and aims of treatment for some cancers, endocrine, gynecological, psychiatric, and neurological disorders
- It provides students with basic knowledge about medications use, dose considerations, side effects, treatment algorithms and evaluation of therapeutic outcomes for the disorders

#### 9. Teaching and Learning Strategies

## Strategy

Lectures

Seminars Simple

quizzes

Brainstorming questions

#### 10. Course Structure

Week	Hours	Required Learning	Unit or	Learning	Evaluation
		Outcomes	subject name	method	method
1	1	1. Explain the regulation and	Adrenal gland	Lectures and	Simple
		physiological roles of hormones	disorders	Discussions	quizzes
		produced by the adrenal glands.			
		2. Recognize the clinical presentation of			
		adrenal insufficiency.			
		3. Describe the pharmacologic			
		management of acute and chronic			
		adrenal insufficiency.			
		4. Recommend therapy monitoring			
		parameters for adrenal insufficiency.			
		5. Recognize the clinical presentation of			





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			Cushing syndrome and the			
			physiological consequences of cortisol			
			excess.			
		6.	Describe the pharmacologic and			
			nonpharmacologic management of			
			Cushing syndrome.			
		7.	Recommend strategies to prevent the			
			development of hypercortisolism and			
			hypercortisolism.			
		8.	Recommend therapy monitoring			
		0.	parameters for Cushing syndrome.			
2	2	1.		Thyroid gland	Lectures	Simple
	2	1.	hypothalamic—pituitary— thyroid axis	disorders	and	quizzes
			and interaction among these	uisoruers	Discussions	quizzes
			_		Discussions	
		7	components.			
		۷.	Discuss the relationship between			
			serum thyroid- stimulating hormone			
			(TSH) levels and primary thyroid			
			disease, and advantages for the use of			
			TSH levels over other tests such as			
			serum T4 (thyroxine) and T3			
			(triiodothyronine) levels.			
		3.	Identify typical signs and symptoms of			
			hypothyroidism and consequences of			
			suboptimal treatment.			
		4.	Describe clinical use of levothyroxine			
			(LT4) in the treatment of			
			hypothyroidism.			
		5.	Discuss issues regarding LT4 product			
			bioequivalence and reasons for			
			maintaining patients on the same			
			product.			
		6.	Describe the management of			
			hypothyroidism and hyperthyroidism			
			in special populations, including			
			pregnant women.			
		7.	Identify typical signs and symptoms of			
			hyperthyroidism and consequences of			
			inadequate treatment.			
		8.	Discuss the pharmacotherapy of			
			hyperthyroidism, including advantages			
			and disadvantages of antithyroid drugs			
						ı





		versus radioactive iodine, adverse			
		effects, and patient monitoring.			
3	1	1. Describe the pathophysiology,	Alzheimer	Lectures and	Simple
		including genetic and environmental	disease	Discussions	quizzes
		factors that may be associated with AD			
		2. Detail the clinical presentation of the			
		typical patient with AD.			
		3. Explain how nonpharmacologic therap	y		
		is combined with pharmacologic			
		therapy for patients with AD.			
		4. Recognize and recommend treatment			
		options for disease- specific symptoms			
		as well as behavioral/ noncognitive			
		symptoms associated with AD.			
		5. Educate patients and/or caregivers			
		about the expected outcomes for			
		patients with AD and provide contact			
		information for support/advocacy			
		agencies.			
4	2	1. Recognize signs and symptoms of	Schizophrenia	Lectures	Simple
		schizophrenia	nia	and	quizzes
		2. Explain potential pathophysiologic		Discussions	
		mechanisms that are thought to			
		underlie schizophrenia.			
		3. Identify treatment goals for a patient			
		with schizophrenia.			
		4. Recommend appropriate antipsychoti			
		medications based on patient-specific			
		data.			
		5. Compare side effect profiles of			
		individual antipsychotics.			
		6. Educate patients and families about			
		schizophrenia, treatments, and the			
		importance of adherence to			
		antipsychotic treatment.			
5	2	1. Explain the etiology and	Depressive	Lectures	Simple
		pathophysiology of major depressive	disorders	and	quizzes
		disorder (MDD).		Discussions	
		2. Identify the signs and symptoms of			
		MDD.			
		3. Outline the treatment goals for a			
		patient with MDD.			
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		4. Recommend pharmacotherapy given a			
		specific patient with MDD.			
		5. Develop a monitoring plan for a			
		specific patient with MDD that			
		includes the assessment of efficacy as			
		well as adverse effects.			
		6. Educate patients and caregivers on the			
		proper use of antidepressant therapy.			
6	1	1. Explain the pathophysiologic	Anxiety	Lectures	Simple
		mechanisms underlying anxiety		and	quizzes
		disorders.		Discussions	
		2. Recognize common presenting			
		symptoms of generalized anxiety			
		disorder (GAD)			
		3. List treatment goals for patients with			
		GAD.			
		4. Identify appropriate lifestyle			
		modifications and over-the-counter			
		medication use in these patients.			
		5. Design a patient-specific			
		pharmacotherapy treatment plan for			
		patients.			
		6. Develop a monitoring plan for patients			
		with anxiety disorders.			
7	1	1. Describe the pathophysiology and	Sleep	Lectures	Simple
		characteristic features of insomnia.	disorders	and	quizzes
		2. Recommend and optimize appropriate	(insomnia)	Discussions	
		sleep hygiene and nonpharmacologic			
		therapies for the management and			
		prevention of sleep disorders.			
		3. Recommend and optimize appropriate			
		pharmacotherapy for insomnia.			
		4. Describe the components of the patient			
		care process to implement and assess			
		safety and efficacy of			
		pharmacotherapy for insomnia.			





8	1	1. Discuss the physiology of the female	Contracepti	Lectures	Simple
		reproductive system.	on	and	quizzes
		2. Compare the efficacy of oral		Discussions	
		contraceptives with that of other			
		methods of contraception.			
		3. State the mechanism of action of			
		hormonal contraceptives.			
		4. Discuss adverse effects, risks, and			
		contraindications associated with the			
		use of contraceptives and			
		recommend strategies for minimizing			
		or eliminating such risks.			
		5. Describe advantages and			
		disadvantages of various			
		contraceptives, including oral and			
		nonoral formulations.			
		6. Cite important drug interactions			
		that may occur with oral			
		contraceptives.			
		7. Provide appropriate patient education			
		regarding the use of oral and barrier			
		methods of contraception.			
		8. Discuss how emergency contraception			
		may be employed to prevent			
		unintended pregnancy.			
9	2	1. Explain the physiological changes	Hormone	Lectures	Simple
		associated with menopause.	replacemen	and	quizzes
		2. Identify the signs and symptoms	t therapy	Discussions	
		associated with menopause.	n nost		
		3. Determine the desired therapeutic	n post-		
		outcomes for patients taking	menopausa I women		
		menopausal hormone replacement	i women		
		therapy (MHRT).			
		4. Explain how to evaluate a patient for			
		the appropriate use of MHRT.			
		5. Recommend appropriate			
		nonpharmacologic and			
		pharmacologic interventions for			
		menopausal symptoms.			
		6. Design a monitoring plan to assess			
		the safety and effectiveness of			
		pharmacotherapy			





10	1	1.	Describe the underlying etiology of	Menstruati	Lectures	Simple
			dysmenorrhea, amenorrhea, and	on related	and	quizzes
			anovulatory bleeding.	disorders	Discussions	
		2.	Explain the physiological changes			
			associated with dysmenorrhea,			
			amenorrhea, and anovulatory			
			bleeding.			
		3.	Identify the signs and symptoms			
			associated with dysmenorrhea,			
			amenorrhea, and anovulatory			
			bleeding.			
		4.	Determine the desired therapeutic			
			outcomes for patients with			
			dysmenorrhea, amenorrhea, and			
			anovulatory bleeding.			
		5.	Recommend appropriate			
			nonpharmacologic and pharmacologic			
			interventions for dysmenorrhea,			
			amenorrhea, and anovulatory			
			bleeding.			
		6.	Design a monitoring plan to assess the			
			safety and effectiveness of			
			pharmacotherapy.			
11	2	1.	Describe the pathophysiology of	Cancer	Lectures	Simple
			cancer.	chemother	and	quizzes
		2.	Define the tumor, nodes, metastases	apy and	Discussions	
			(TNM) system of cancer staging.	treatment		
		3.	Define prevention and treatment			
			strategies for cancer.			
		4.	Outline actions for all healthcare			
			professionals to prevent medication			
	_		errors with cancer treatments.			
12	2	1.	Explain the pathophysiology of certain	Leukemias	Lectures	Simple
			types of leukemia.		and	quizzes
		2.	1 8 7 1		Discussions	
			laboratory disorders associated with			
		2	leukemias.			
		3.	, , ,			
			that would determine the most			
			appropriate chemotherapeutic			
			regimens for patients having leukemia.			
			іеикеппа.			





		4.	Describe the available treatment			
			options of certain types of leukemias			
		5.	Recognize the treatment			
			complications associated with the			
			therapy of leukemias.			
13	1	1.	Explain the risk factors associated with	Breast cancer	Lectures	Simple
			developing breast cancer.		and	quizzes
		2.	Recognize signs and symptoms		Discussions	
			related to early and late stages of the			
			disease.			
		3.	Distinguish between good and poor			
			prognostic factors.			
		4.	Determine treatment goals for early-			
			stage, locally advanced, and			
			metastatic breast cancers.			
		5.	Explain the available treatment			
			options for breast cancer.			
		6.	Describe the relevance of hormone,			
			HER2, and PD-1 receptors.			
		7.	Discuss the benefits and risks			
			associated with various therapies.			
14	1	1.	Identify risk factors associated with	Prostate	Lectures	Simple
14	1	1.	Identify risk factors associated with prostate cancer development.	Prostate cancer	Lectures and	Simple quizzes
14	1	1.	,			,
14	1		prostate cancer development.		and	,
14	1		prostate cancer development.  Appraise the prognostic- and patient-		and	,
14	1		prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine		and	,
14	1	2.	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.		and	,
14	1	2.	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic		and	,
14	1	2.	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types		and	,
14	1	<ol> <li>3.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.		and	,
14	1	<ol> <li>3.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects		and	,
14	1	<ol> <li>3.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for		and	,
14	1	<ol> <li>3.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for patients receiving androgen		and	,
14	1	<ol> <li>3.</li> <li>4.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for patients receiving androgen deprivation therapy for prostate cancer based on patient- specific factors and the prescribed regimen.		and	,
14	1	<ol> <li>3.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for patients receiving androgen deprivation therapy for prostate cancer based on patient- specific factors and the prescribed regimen.  Recognize the common adverse		and	,
14	1	<ol> <li>3.</li> <li>4.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for patients receiving androgen deprivation therapy for prostate cancer based on patient- specific factors and the prescribed regimen.  Recognize the common adverse effects and formulate a monitoring		and	,
14	1	<ol> <li>3.</li> <li>4.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for patients receiving androgen deprivation therapy for prostate cancer based on patient- specific factors and the prescribed regimen.  Recognize the common adverse effects and formulate a monitoring plan for patients receiving treatment		and	,
14	1	<ol> <li>3.</li> <li>4.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient-specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for patients receiving androgen deprivation therapy for prostate cancer based on patient-specific factors and the prescribed regimen.  Recognize the common adverse effects and formulate a monitoring plan for patients receiving treatment for metastatic prostate cancer.		and	,
14	1	<ol> <li>3.</li> <li>4.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient- specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for patients receiving androgen deprivation therapy for prostate cancer based on patient- specific factors and the prescribed regimen.  Recognize the common adverse effects and formulate a monitoring plan for patients receiving treatment for metastatic prostate cancer.  Provide recommendations for bone		and	,
14	1	<ol> <li>3.</li> <li>4.</li> </ol>	prostate cancer development.  Appraise the prognostic- and patient-specific data needed to determine appropriate treatment options.  Evaluate pharmacotherapeutic treatment options for different types of prostate cancer.  Recognize common adverse effects and formulate a monitoring plan for patients receiving androgen deprivation therapy for prostate cancer based on patient-specific factors and the prescribed regimen.  Recognize the common adverse effects and formulate a monitoring plan for patients receiving treatment for metastatic prostate cancer.		and	,





15	1	1. Introduction about common and	Adverse	Lectures	Simple
		problematic adverse effects of	effects of	and	quizzes
		chemotherapy	chemother	Discussions	
		2. Recognizing the clinically significant	ару		
		adverse effects			
		3. Explaining the preventive measures of			
		certain adverse effects			
		4. Discussing the available therapeutic			
		options of some adverse effects.			
16	2	1. Explain the pathophysiologic	Bipolar	Lectures	Simple
		mechanisms underlying bipolar	disorders	and	quizzes
		disorder.		Discussions	
		2. Recognize the symptoms of a manic			
		episode in patients with bipolar			
		disorder.			
		3. Identify common psychiatric			
		comorbidities of bipolar disorder.			
		4. List the desired therapeutic outcomes			
		for patients with bipolar disorder.			
		5. Identify the optimal use of medications			
		as first-line therapy in bipolar disorder,			
		including appropriate dosing.			
		6. Recommend drug therapy for acute			
		treatment of mania and depressive			
		episodes.			
		7. Recommend baseline and routine			
		monitoring for assessment of adverse			
		effects of medications used in the			
		treatment of bipolar disorder.			
		8. Identify general treatment differences			
		for agents used to treat bipolar disorder			
		in the pediatric population			
17	1	1. Identify the risk factors for colorectal	Colorecta	Lectures	Simple
		cancer.	l cancer	and	quizzes
		2. Outline preventive and screening		Discussions	
		strategies for individuals at average			
		and high risk for colorectal cancer.			
		3. Recognize the signs and symptoms of			
		colorectal cancer.			
		4. Describe the treatment options for			
		colorectal cancer based on patient-			
		specific factors, such as stage of			





			disease, age of patient, genetic			
			mutations, and previous treatment			
			received.			
		5.	Outline the pharmacological			
			principles for agents used to treat			
			colorectal cancer.			
		6.	Develop a monitoring plan to assess			
			the efficacy and toxicity of agents used			
			in colorectal cancer.			
		7.	Educate patients about the adverse			
			effects of chemotherapy that require			
			specific patient counseling.			
18	2	1.	Explain the routes of transmission for	Human	Lectures	Simple
			human immunodeficiency virus (HIV)	immunode	and	quizzes
			and its natural disease progression.	ficiency	Discussions	
		2.	Identify typical and atypical signs and	virus		
			symptoms of acute and chronic HIV			
			infection.			
		3.	Identify the desired therapeutic			
			outcomes for patients living with HIV.			
		4.	Recommend appropriate first-line			
			pharmacotherapy interventions for			
			patients with HIV infection.			
		5.	Describe the components of a			
			monitoring plan to assess			
			effectiveness and adverse effects of			
			pharmacotherapy for HIV infection.			
		6.	Educate patients about the state of the			
			disease, appropriate lifestyle			
			modifications, and drug therapy			
			required for effective treatment.			
	2	1.	Discuss the underlying	Lymphom	Lectures	Simple
			pathophysiologic mechanisms of the	a and	and	quizzes
			lymphomas and how they	multiple	Discussions	
		rel	ate to presenting symptoms of the	myeloma		
			disease.			
		2.	Differentiate the pathologic findings			
			of Hodgkin lymphoma (HL), follicular			
			indolent non-Hodgkin lymphoma			
			(NHL), and diffuse aggressive NHL			
			and how this information yields a			
			specific diagnosis.			





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		3.	Describe the general staging criteria					
			for the lymphomas and how it relates					
			to prognosis; evaluate the role of the					
			prognostic systems such as the					
			International Prognostic Score for HL,					
			the Follicular Lymphoma International					
			Prognostic Index (IPI), and the IPI for					
			diffuse, aggressive NHL.					
		4.	Compare and contrast the treatment					
			algorithms for early and advanced					
			stage disease for HL.					
		5.	Assess the role of autologous					
			hematopoietic stem-cell					
			transplantation for relapsed					
			lymphomas.					
		6.	Delineate the clinical course of					
			follicular indolent and diffuse					
			aggressive NHL and the implications					
			for disease classification schemes and					
			treatment goals.					
		7.	Outline the general treatment					
			approach to follicular indolent and					
			diffuse aggressive NHL for localized					
			and advanced disease.					
		8.	Interpret the current role for monoclona					
			antibody therapy in NHL					
20	1	1.	Explain the pathophysiology of	Endometri	Lectures	Simple		
			Endometriosis.	osis	and	quizzes		
		2.	Explain the signs/symptoms of		Discussions			
			Endometriosis.					
		3.	Outline the general treatment					
			approach					
11.	Course E	valu	ation			,		
Mid	term exa	m 2	5 marks, Quizzes and attendance 5 marl	cs, Final exam 70 m	narks			
12.	Learning	gano	d Teaching Resources					
Requ	Required textbooks (curricular books, if a Pharmacotherapy Handbook							
1- ACCP Updates in Therapeutics								
			2- Appl	ed therapeutics				
Recommended books and Review articles								
refer	ences (s	cien	tific journals, reports)					
Elect	ronic Re	fere	nces, Websites Med	scape				
_								





#### 1. Course Name:

#### Therapeutic drug monitoring

2. Course Code:

#### 556 PhTdm

3. Semester / Year:

Second semester/Fifth Year

4. Description Preparation Date:

2-2025

5. Available Attendance Forms:

On campus

6. Number of Credit Hours (Total) / Number of Units (Total)

2Hours theory +2-hour practical (60) /3Units

7. Course administrator's name (mention all, if more than one name)

Theory:

Name: Assistant lecturer Hussein Ali

Email: husseinali@bcms.edu.iq

Practical:

Name: Assistant lecturer Rana Kadum

Email: ranakadum@bcms.edu.iq

#### 8. Course Objectives

## **Course Objectives**

At the end of this unit, the student should be able to: recognize characteristics of drugs that make them good candidates for TDM, describe appropriate indications for TDM, understand the factors that may affect the measured concentrations list, and discuss the importance of information needed when requesting drug concentration interpret measured drug concentrations adjust dose based on TDM....

## 9. Teaching and Learning Strategies

Strategy	Lecture
	s
	Seminar
	Simple
	quizzes
	Brainstorming questions

Week	Hour	Required Learning Outcomes	Unit or subject	Learning	Evaluation
	S		name	method	method
	4	1. Discuss the goal of therapeutic drug	Chapter one:	Lectures,	Simple
		monitoring.	(Clinical	Discussions	quizzes
1-2		2. Discuss the need for therapeutic	Pharmacokinet		
1-2		drugs.	ic and		
		3. Identify the four principle biological	Pharmacodyna		
		events associated with	mic Concepts)		





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		<ul> <li>pharmacokinetics.</li> <li>4. Identify route(s) that drugs can be eliminated.</li> <li>5. Define the following: <ul> <li>a. TDM</li> <li>b. linear and nonlinear pharmacokinetics</li> <li>c. Pharmacokinetics parameters</li> <li>d. Half-life</li> <li>e. volume of distribution</li> <li>f. clearance</li> </ul> </li> </ul>			
3-4	4	<ol> <li>Discuss the applied equations that are used to measure the drug concentration</li> <li>Discuss the applied equations that are used to measure the individualized pharmacokinetic parameters</li> <li>Discuss the equations that are used to measure the dose and loading dose</li> </ol>	Chapter two: Clinical Pharmacokinet ic Equations and Calculations	Lectures, Discussions	Simple quizzes
	4	1.Discuss the effect of kidney, liver disease, and heart disease on the drug's pharmacokinetics.      2.Discuss the effect of obesity on the pharmacokinetics of the drug	Chapter Three: Drug dosing in special population	Lectures, Discussions	Simple quizzes
5-6	4	<ol> <li>Identify why we need to monitor drug concentration for aminoglycoside</li> <li>Determine the applied pharmacokinetics methods and equations to calculate the initial dose</li> <li>Determine the applied pharmacokinetics methods and equations to calculate the individualized dose</li> </ol>	Chapter four: Aminoglycoside	Lectures, Discussions	Simple quizzes
7-8	4	Identify why we need to monitor drug concentration for vancomycin     Determine the applied pharmacokinetics methods and equations to calculate the initial dose     Determine the applied pharmacokinetics methods and	Chapter five: vancomy cin	Lectures, Discussions	Simple quizzes





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		equations to calculate individualized dose	ıne			
		Identify why we need to monitor	or drug	Chapter	Lectures,	Simple
		concentration for digoxin	or drug	six:	Discussions	•
		2. Determine the applied		Digoxin	Discussions	quizzes
		pharmacokinetics methods and		Digoxiii		
9-10	3	equations to calculate the initial	dose			
5 .0	Ü	3. Determine the applied	2000			
		pharmacokinetics methods and				
		equations to calculate the				
		individualized dose				
		1. Identify why we need to monito	or drug	Chapter	Lectures,	Simple
		concentration for phenytoin		seven:	Discussions	quizzes
			pplied	phenytoin		,
		pharmacokinetics methods	and			
11-12	3	equations to calculate the initial	dose			
		3. Determine the a	pplied			
		pharmacokinetics methods	and			
		equations to calculate	the			
		individualized dose				
		1. Identify why we need to monito	or drug	Chapter eight:		
		concentration for valproic acid		valproic acid		
		2. Determine the a	applied			
		pharmacokinetics methods	and			
13-14	3	equations to calculate the initial	dose			
			pplied			
		pharmacokinetics methods	and			
		equations to calculate	the			
		individualized dose				
11. C	ourse E	valuation				
		exam + 20 Laboratory + 60 Final exa	m			
12. Learning and Teaching Resources						
		tbooks (curricular books, if any)		Applied Clinical Ph		, ,
		ces (sources)		Applied Clinical Pharmacokinetics by Larry		
	nmend			Applied therapeuti	CS	
		rnals, reports)				
Electr	onic Re	ferences, Websites		Review articles		





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1. Co	ourse Name:							
Advar	iced Pharmaceutic	al Analysis						
2. Co	ourse Code							
557 Cl	557 ChPApa							
3. Se	3. Semester / Year:							
Secon	Second Semester / Fifth Stage							
4. De	4. Description Preparation Date:							
9-202	5							
5. Av	ailable Attendance	Forms						
On car	npus							
6. Ni	umber of Credit Hou	ırs (Total) / Number of Units (Total)						
3 hour	s theory +2 hours p	ractical (75)/ 4 units						
7. Co	ourse administrator's	s name (if more than one name)						
Theo								
	e: Lecturer Dr. Imad							
	: <u>imad.muneeb@bc</u>	<u>ms.edu.iq</u>						
Pract								
	e: Assistant lecturer							
	: <u>hasan.fadhil@bcm</u>	is.edu.iq						
<b>6.</b> CC	ourse Objectives							
	<ul> <li>Learn about th</li> </ul>	e principles of spectrum						
	• Identify the dif	<ul> <li>Identify the different spectrum types (UV-FIS), IR, NMR, and Mass.</li> </ul>						
	Uses of the spectrum in identifying organic compounds and medicines.							
	Enable the student to gain the necessary skills and knowledge to use different							
Course	spectrometers in order to benefit from them after graduation in drug control							
Objectives	laboratories and pharmaceutical factories.							
	Gain the skills and knowledge necessary to operate advanced devices and							
	interpret the results to prepare qualified pharmacists capable of conducting							
	scientific research.							
Training in practical methods for different spectrometry								
9. Te	aching and Learning	g Strategies						
		1- Theoretical lectures covering all aspects of each method						
		2- Conducting reports and research on the applications of the						
		mentioned methods on chemical compounds and						
C:	<b>.</b>	pharmaceutical preparations						
Stra	ıegy	3- Display applied videos to help understand the material and						
		gain skill						

4- Use of methodological and supporting books

5- Holding scientific sessions in the form of discussions or





seminars

	10. Course Structure							
Week	Hours	Required Learning Outcomes	Unit or subject name	Learning method	Evaluation method			
1	3+2	UV / visible spectroscopy	Introduction & demonstration to visible spectrophotometry	Lectures Labs &	Paper exams			
2	3+2	UV / visible spectroscopy	Determination of KMnO4/Beers law	Lectures Labs &	Paper exams			
3	3+2	UV / visible spectroscopy	Unknown of KMnO4 + Quiz	Lectures Labs &	Paper exams			
4	3+2	Mass Spectrometry	Colorimetric assay of tetracycline using FeCl3	Lectures Labs &	Paper exams			
5	3+2	Mass Spectrometry	Unknown of tetracycline using FeCl3method + Quiz	Lectures Labs &	Paper exams			
6	3+2	Infrared Spectrometry	Determination of tetracycline in acidic medium	Lectures Labs &	Paper exams			
7	3+2	Infrared Spectrometry	Determination of tetracycline in basic medium	Lectures Labs &	Paper exams			
8			Midterm Exam					
9	3+2	Infrared Spectrometry	Colorimetric assay of streptomycin by maltol method	Lectures Labs &	Paper exams			
10	3+2	Infrared Spectrometry	Colorimetric assay of streptomycin by maltol method	Lectures Labs &	Paper exams			
11	3+2	Proton NMR Spectrometry	Unknown of streptomycin by maltol method + Quiz	Lectures Labs &	Paper exams			
12	3+2	Proton NMR Spectrometry	IR chart tutorial	Lectures Labs &	Paper exams			
13	3+2	C13 NMR	IR chart tutorial	Lectures	Paper exams			





		Spectrometry		Labs &	
14	3+2	C13 NMR Spectrometry	IR chart tutorial	Lectures Labs &	Paper exams
15			Review		

## 11. Course Evaluation

Mid-term written exams 17%

Daily preparation and daily and oral exams 3%

Practical side 20 marks

Final Exam 60 Marks arks

12. Learning and Teaching Resources	
Required textbooks (curricular books, if any)	Spectrometric Identification of Organic
	Compounds by Silverstein, Bassler and Morrill.
	Applications of absorption spectroscopy of
	organic compounds by Dyer JR.
Main references (sources)	like mention above
Recommended books and references (scientific	Organic Chemistry by McMurry; 5thed;
journals, reports)	Thomason learning CA, USA 2000
Web sites	Google scholar, ResearchGate





#### 1. Course Name:

## **Clinical Laboratory Training**

2. Course Code:

#### 558 ACICI

3. Semester / Year:

2<sup>nd</sup> semester/5<sup>th</sup> year

4. Description Preparation Date:

9-2025

#### 5. Available Attendance Forms:

Sheets signed by students (training in hospital)

## 6. Number of Credit Hours (Total) / Number of Units (Total)

4 hours Practical/ 2 unites

#### 7. Course administrator's name

Name: Kawther Faris

Email: <a href="mailto:kawther.kf@gmail.com">kawther.kf@gmail.com</a>
Name: Lecturer Dr. Nawfel Ayad
Email: <a href="mailto:nawfel.ayad@bcms.edu.iq">nawfel.ayad@bcms.edu.iq</a>
Name: Assistant lecturer Ibraheem Kais

Email: ibraheem.kais0@bcms.edu.iq

#### 8. Course Objectives

Course Objectives

Learning students about various Tests applied in hospital labs (Biochemical, Hematological, Bacteriological, Virological, General urine exam, general stool exam). Show them the normal values of each studied parameter and teach them how to explain abnormalities in association with clinical symptoms and diseases.

#### 9. Teaching and Learning Strategies

Strategy Explain work principles+ Applying the lab examinations + making weekly reports
+ written and practical quiz+ Visiting specific laboratories in general hospitals to
take a look on status situation in lab work field.

Week	Hours	Required Learning Outcomes	Unit or subject name	Learning method	Evaluation method
1	4	Importance of	Diagnostic test basis,	Practical	Exam
		laboratory tests, how	collecting and transporting		&
		to make sampling	specimen.		report
2	4	Glucose situation in	Biochemical test: fasting	practical	Exam
		the body	blood glucose, Post		&
			prandial glucose, oral		report
			glucose tolerance test.		





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3	4	Kidney function	Blood urea, blood	practical	Exam
			creatinine, creatinine		&
			clearance, uric acid.		report
4	4	Blood lipids situation	Cholesterol, lipoprotein,	practical	Exam &
			triglycerides.		report
5	4	Liver function	Blood proteins, bilirubin.	practical	Exam &
					report
6	4	Testing blood minerals	Calcium, inorganic	practical	Exam &
			phosphate, serum chloride.		report
7	4	Analysis of protein	Alkaline phosphatase, acid	practical	Exam &
		metabolism	phosphatase, alanine		report
			aminotransferase, aspartate		
			amino transferase, lactate		
			dehydrogenase, creatinine		
			phosphokinase.		
8	4	Virology test	Serological tests: VDRL,	practical	Exam &
			ASO-titer, hepatitis test.		report
9	4	Serological tests for	C-reactive protein test,	practical	Exam &
		infections	Rheumatic factor test, Ros		report
			Bengal test, typhoid fever		
			test (Widal test), Pregnancy		
			test, TORCH test.		
10	4	Urinalysis	General urine exam, urine	practical	Exam &
			specimen collection.		report
11	4	Stool analysis	General stool exam, stool	practical	Exam &
			specimen collection		report
12	4	Blood analysis	Hematological tests:	practical	Exam &
			RBC count, Hb, PCV, RBC		report
			indices, WBC count, Platelet		
			count.		
13	4	Blood analysis	Blood typing, COMB test,	Practical	Exam &
			Bleeding time, ESR.		report
14	4	Bacteriological and	Microbiological tests:	practical	Exam &
		sensitivity test	culture and sensitivity test,		report
			staining methods, enriched		
			media.		
			VITEK II system		
15	4	Applications of	Identifying the most	practical	Exam &
		Clinical Microbiology	prevalent lab techniques		report
			that can be used in		
			diagnostic microbiology		





	and correlate that with the	
	most clinical prevalent	
	infections	

#### 11. Course Evaluation

- 40 Marks Quest Practical: (10% Class activity and reports + 5% Oral exam. + 15% practical exam %10+written exam)
- 60 Marks final exam

100 Marks total

12. Learning and Teaching Resources

Required textbooks (curricular books, if any)

Oxford handbook of Clinical and Laboratory investigation. By: Drew Provan, 4<sup>th</sup> ed. 2018

Main references (sources)

Manual for laboratory training adopted by the department

Recommended books and references (scientific journals, reports...)

Laboratory tests in general practice. K reports 59 C. By: Gillet Pierr, et al 2007

Web sites

https://labtestsonline.org.uk





#### 1. Course Name:

#### **Drug Delivery Systems Design**

2. Course Code:

#### 559ACIDds

3. Semester / Year:

Second semester/Fifth Year

4. Description Preparation Date:

9-2025

5. Available Attendance Forms:

Students' signature on attendance sheet

6. Number of Credit Hours (Total) / Number of Units (Total)

2 hours Theoretical /2 units

7. Course administrator's name

Name: Lecturer Dr. Mohammed Jassim

Email: mohammed-jassim-neamah@bcms.edu.iq

#### 8. Course Objectives

#### **Course Objectives**

- 1. By the end of this course, students will be able to:
- 2. Explain the process of new drug development, including drug discovery, preclinical evaluation, early formulation studies, and regulatory submission of INDs, NDAs, and ICH guidelines.
- 3. Explore the principles and applications of pharmaceutical nanotechnology, including the design and use of liposomes, dendrimers, micelles, solid nanoparticles, and lipid-based delivery systems in modern therapeutics.
- 4. Describe the anatomical and physiological considerations for non-oral routes of drug delivery, including nasal, ocular, transdermal, and pulmonary routes.
- 5. Analyze formulation challenges and strategies for improving drug solubility, permeability, bioavailability, and patient adherence across various advanced delivery systems.
- 6. Evaluate the design, function, and clinical considerations of innovative delivery platforms such as patches, inhalers, eye drops, and nanoparticle-based systems.

#### 9. Teaching and Learning Strategies

Strategy	Lecturing
	Homework
	Quiz

Week	hours	Required Learning Outcomes	Unit or subject name	Learning method	Evaluation method
1-3		Introduction, Drug discovery and	New Drug	Presentation	Discussion
	C	drug design, Biologic	Development and	of lecture	Paper-
	6	characterization, Early	Approval Process	Interactive	based
		formulation studies, The	Chapter 2	discussions	exams





		investigational new drug	(Pharmaceutical	
		application, The new drug	Dosage Forms and	
		application, supplemental,	Drug Delivery Systems	
		abbreviated, and other	by Howard A. Ansel;	
		applications, international	11th edition, 2017)	
		conference on Harmonization of		
		technical Requirements for		
		registration of pharmaceuticals		
		for Human use.		
		Introduction, Applications of	Pharmaceutical	
		pharmaceutical nanotechnology,	Nanotechnology and	
		Polymer-drug conjugates,	Nanomedicines	
		Dendrimers, Micelle systems,	Chapter 46 (Aulton's	
4-7	7	Solid nanoparticles, Liposomes,	Pharmaceutics; The	
		bilayer vesicles and lipid	Design and	
		nanoparticles, Microcapsules	Manufacture of	
		and microspheres, Ongoing	Medicines; 6th	
		developments.	edition, 2022)	
		Introduction, Anatomy and	Nasal Drug Delivery	
		physiology, Drug delivery, Nasal	Chapter 40 (Aulton's	
		delivery systems	Pharmaceutics; The	
8-9	4		Design and	
			Manufacture of	
			Medicines; 6th edition,	
			2022)	
		Introduction, Anatomy and		
		physiology of the eye, Some		
		common ocular conditions and		
		pharmacological interventions,		
		Topical ophthalmic preparations,	Ocular Drug Delivery	
		formulating ophthalmic		
		preparations, Topical, liquid	Chapter 41(Aulton's	
		ophthalmic preparations, Barriers	Pharmaceutics; The	
10-11	4	to topical ocular drug absorption,	Design and	
		Increasing drug solubility and	Manufacture of	
		absorption in topical ophthalmic	Medicines; 6th edition,	
		preparations, Sterility of	2022)	
		ophthalmic preparations, Ocular		
		drug pharmacokinetics. Targeting		
		the posterior segment of the eye.		
		Problems with traditional and		
		new ocular drug delivery		





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		systems, Patient adherence and			
		instillation of eye drops.			
		Introduction, Factors affecting			
İ		percutaneous absorption,			
		Percutaneous absorption	Transdermals and		
		enhancers, Percutaneous	Transdermal Drug		
İ		absorption models, Design	Delivery Systems		
		features of transdermal drug			
42.42	4	delivery systems, Advantages and	Chapter 11		
12-13	4	disadvantages of TDDSs,	(Pharmaceutical		
		Examples of transdermal drug	Dosage Forms and		
		delivery systems, General clinical	Drug Delivery Systems		
		considerations in the use of	by Howard A. Ansel;		
		transdermal and TDDSs, Patches	11th edition, 2017)		
		(not systems), Tapes, Examples of			
		transdermal preparations.			
		Inhaled drug delivery,	Pulmonary Drug		
		Formulating and delivering	Delivery		
		therapeutic inhalation aerosols,	Chapter 39 (Aulton's		
14-15	4	Methods of aerosol size analysis.	Pharmaceutics; The		
	4		Design and		
			Manufacture of		
			Medicines; 6th edition,		
			2022)		
11. Cou	11. Course Evaluation				

## 11. Course Evaluation

- 30 Marks Theoretical assessment.
   (paper-based mid-term exam + quiz + attendance + seminar)
- 70 Marks paper-based theoretical final exam

Total 100 Marks

# 12. Learning and Teaching Resources Required textbooks (curricular books, if any) Aulton's Pharmaceutics; The Design and Manufacture of Medicines; 6th edition, 2022. 2) Pharmaceutical Dosage forms and Drug Delivery Systems by Howard A. Ansel; 11th edition, 2017. Main references (sources) Recommended books and references (scientific journals, reports...) Electronic References, Websites





#### 1. Course Name:

Pharmaceutical Biotechnology (Theoretical)

2. Course Code:

560 Phpb

3. Semester / Year:

2<sup>nd</sup> Semester/5<sup>th</sup> year

4. Description Preparation Date:

9-2025

5. Available Attendance Forms:

Students' signature on attendance sheet

6. Number of Credit Hours (Total) / Number of Units (Total)

1 hour Theoretical /1 units

7. Course administrator's name

Theory

Name: Assistant lecturer Rana Kadum

Email: ranakadum@bcms.edu.iq

8. Course Objectives

**Course Objectives** 

Identify the most common therapeutic peptides and proteins derived from biotechnological sources Knowing structure details, formulation requirements, and pharmacist role.

#### 9. Teaching and Learning Strategies

Strategy	Lecturing
	Homework
	Quiz

Week	Hours	Required Learning Outcomes	Unit or subject name	Learning method	Evaluation method
1	1	Importance and Definition of Biotechnology History of Biotechnology derived product	Biotechnology - introduction	Theoretical lectures.	Paper-based exams
2	1	Recombinant DNA biotechnology.	Formulation biotechnology prod (biopharmaceutical consideration)	Theoretical lectures.	Paper-based exams
3	1	Sterilization (chemical + physical Methods). Chemotherapy.	Microbial consideration- sterile pyrogen v decontamination	Theoretical lectures.	Paper-based exams





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		Types and specifications of	Excipients of parenteral	Theoretical	Paper-based
		excipients used in	products –soluble	lectures.	exams
4	1	biotechnological formulation	enhancer anti-adsorption		
			age buffer component		
			preservatives – osmotic		
			agents.		
		Types and specifications of	Excipients of parenteral	Theoretical	Paper-based
		excipients used in	products soluble	lectures.	exams
5	1	biotechnological formulation	enhancer anti-adsorption		
			age buffer component		
			preservatives – osmotic		
			agents.		
		Formulation requirements	Route of administration	Theoretical	Paper-based
6	1	according to route of	Parenteral route Oral route.	lectures.	exams
		administration			
7		<u> </u>	Mid-term exam		
		Formulation requirements	Route of administration	Theoretical	Paper-based
8	1	according to route of	Parenteral route Oral route	lectures	exams
		administration			
		Formulation requirements	Route of administration	Theoretical	Paper-based
9	1	according to route of	Parenteral route Oral route	lectures.	exams
	'	administration			
		Formulation requirements	Route of administration	Theoretical	Paper-based
10	1	according to route of	Alternative routes (nasal-	lectures.	exams
		administration	pulmonary rectal-buccal		
			transdermal	<b>T</b> 1 · ·	
		Formulation requirements	Route of administration	Theoretical	Paper-based
11	1	according to route of	Alternative routes (nasal-	lectures.	exams
		administration	pulmonary rectal-buccal		
		ADME 6 21 1	transdermal	TI . '	D 1 1
		ADME of peptides and	Pharmacokinetic of	Theoretical	Paper-based
12	1	proteins Assessments and	peptides and protein	lectures.	exams
12	1	relationships to	(Elimination of proteins		
		pharmacodynamics action	(proteolysis excretion- metabolism		
		ADME of popular and	Pharmacokinetic of	Theoretical	Danes based
		ADME of peptides and		lectures.	Paper-based
13	1	proteins Assessments and relationships to	peptides and protein (Elimination of proteins	iectures.	exams
13	'	·	(proteolysis excretion-		
		pharmacodynamics action	metabolism		
			metabonsiii		



Electronic References, Websites



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		ADME of peptides and	Pharmacokinetic of	Theoretical	Paper-based	
		proteins Assessments and	peptides and protein	lectures.	exams	
14	1	relationships to	(Elimination of proteins			
		Pharmacodynamics action	(proteolysis excretion-			
		,	metabolism			
15		Quest	ion and answers (Corse review)			
11. C	ourse Eva	luation				
	•	30 MarksTheoretical assess	ments; (paper-based mid-term e	xam)		
	•	70 Markspaper-based theor	etical final exam			
		100 Marks total				
12. Le	arning an	d Teaching Resources				
Requi	red textbo	ooks	1. pharmaceutical biotech	nology Cromm	elin, Robert D.	
			Syinder			
Main references (sources)			1. pharmaceutical biotechnology Crommelin, Robert D.			
			Syinder			
Recor	nmended	books and references				
(scien	tific journ	als, reports)				