# The Hashemite University







Deanship of Academic Development and International Outreach

عمادة التطوير الأكاديمي والتواصل الدولي

Syllabus: Hematopoietic System (181501207)
Second Year-Second Semester-2022/2023

COURSE INFORMATION			
Course Name: Hematopoietic System		Course Code: 181501207	
Semester:	Second (Spring) 2022/2023	Section: Preclinical Modules	
Department:	Pharmacology and Public Health	Core Curriculum: MD program	
Faculty:	Medicine		
Day(s) and Time	(s): 8:00AM - 3:00PM Sunday-Thursday	Credit Hours: 4	
(Teaching Period: <b>25.2.</b> 20 <b>24</b> – 13.3.20 <b>24</b> )		Prerequisites: NA	
Classroom:			
Theoretical lectures: Ibn Sina Complex 301, 302			
Practical sessions: labs of anatomy, physiology, microbiology			
and pathology, Ik	on Sina Complex		

#### **COURSE DESCRIPTION**

The Hematopoietic Module is an intensive, multidisciplinary, integrated 4 credit-hour course designed to provide medical students with essential basic science and clinical framework for topics related to normal function and abnormalities of the blood and lymphatic systems. The course encompasses lecture-based and laboratory sessions in the anatomy, physiology, pathology, pharmacology, biochemistry, and community medicine of the blood. Students are expected to be familiar with main disorders that affect the blood and lymphatic systems and their corresponding therapy approaches.

#### **DELIVERY METHODS**

The course will be delivered through a combination of active, in-class and online, learning strategies. These will include:

- PowerPoint lectures and active classroom-based discussion
- Live online-delivered lectures
- Relevant papers and reading materials
- E-learning resources: e-reading assignments, virtual meetings, and practice quizzes through Microsoft Teams.

Course Coordinator			
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	Office hours: Sunday: 10:00AM to 12:00AM and Wednesday:		
	10:00AM to 12:00PM		

## **REFERENCES AND LEARNING RESOURCES**

## \* Anatomy:

- Grey's Anatomy for Students by Richard Drake, 4<sup>th</sup> edition.
- Principles of Human Anatomy by Gerard J Tortora and Mark Nilsen, 14<sup>th</sup> edition.
- Clinical Anatomy for Medical Students. By R.S. Snell,  $\mathbf{5}^{\text{th}}$  edition.
- Before We Are Born, by K.L. Moore and T.V.N. Persaud, 10<sup>th</sup> edition.
- \* Physiology:
- Textbook of Medical Physiology by Gyton and Hall, latest edition.
- \* Biochemistry:
- Harper's Biochemistry by Robert K. Murray and Co., latest edition.
- \* Pharmacology:
- Lippincott's Illustrated Reviews: Pharmacology, 7th edition.
- \* Pathology:
- Basic Pathology by Kumar, Abbas and Aster, 10<sup>th</sup> edition.
- \* Microbiology:
- Medical microbiology. An introduction to infectious diseases. By Sheries. Latest edition.

# TOPICS DETAILS/ STUDENT LEARNING OUTCOMES MATRIX \*

Course Objectives				
				Method
A-Biomedical:  1. Describe the main components of blood,	(SUBJECTS& NUMBER OF LECTURES/	SUBJECT	Intended Learning Outcomes	-Online quizzes. -Exams
histology of bone marrow, lymph nodes and structure of spleen and their functions.  2. Understand the physiology of blood coagulation pathways and their contribution to thrombosis and bleeding disorders.  3. Identify abnormalities of red blood cells.  4. Understand the pathophysiology of anemia and its treatment.	*Topic 1: Structure and Function of Blood (11 lectures):  Anatomy: 2 Physiology: 6 Biochemistry: 3	Anatomy:  1. Anatomy and histology of blood (1)  2. Anatomy and histology of blood (2)	-Describe the composition of blood -Discuss the shape, size, color, structure, composition, number, and lifespan of erythrocytesUnderstand the structural-functional adaptation of erythrocytesDescribe the shape, size, structure, and differential count of granular and agranular leukocytesDescribe the histological features of neutrophils, eosinophils and basophilsDescribe the types of lymphocytes and their different basic functionsIdentify the light and electron microscopic features of monocytesDescribe the count, size, shape and lifespan of plateletsDescribe the light and electron microscopic structure of plateletsDescribe the histology of the bone marrow and haemopoiesis	
5. Familiarize with the malignant and non-malignant disorders of white blood cells.  6. Identify different types of leukemia and lymphoma, their pathological classification, clinical presentation and treatment.  7. Identify the microbial agents and diagnostic tests related to blood.  8. Identify the public health issues associated with anemia and malaria.		Physiology:  1. Blood Functions, Functions of plasma & RBCs characteristics.  2. RBCs functions and regulation of RBCs production  3. Blood Groups  4. Hemostasis and Blood Coagulation  5. Prevention of Blood Clotting and Lysis of Blood Clots 6. WBCs: Characterist ics and functions	-Understand the functions of bloodDescribe the constituents of plasma and how they are attributed to the general function of the plasmaState that RBCs are non-nucleated biconcave elastic discs, their number in peripheral bloodIdentify the sites of formation; discuss the normal percentage of reticulocytes of the whole circulating red blood cells and explain the causes of reticulocytosisHow RBCs are regulated, the effect of hypoxiaDescribe the role of iron, vitamin B12, and folic acid and describe the effect of their deficiencyUnderstand the basis of human blood typing into different blood groupsDescribe the ABO and Rh systems of blood groupingApply the knowledge given in the blood grouping system in blood transfusionDescribe the importance of crossmatching testsIdentify the complications of incompatible blood transfusion.	

8-Correlate the basic		-Describe the major types and causes of	
biomedical knowledge		anemia and polycythemia.	
to the clinical skills		-Define hemostasis and describe the three	
to the dimedians		steps involved in hemostasis.	
B-Critical thinking		-Understand the structure, function and	
skills:		life span of platelets.	
		-Understand the interaction of platelets,	
1-Observe, identify and		blood vessels and plasma coagulation	
predict health		factors in hemostasis.	
problems based on		-identify which coagulation factors are dependent on vitamin K and how vitamin	
previous experience		K modifies these coagulation factors.	
and make decisions		-Describe the fibrinolytic system and	
based on evidence		understand its role during hemostasis.	
rather than opinion		-Describe the mechanism of	
Taute and Spinion		anticoagulants and correlate to the	
2- Draw conclusions		coagulation pathways and components.	
about the collected		-Identify the pathophysiologic mechanisms	
data (inference).		of disease states caused by disturbed	
		hemostasis.	
3- Maintain good		-Discuss how to recognize different WBCs	
communication habits,		types and describe their site of production,	
such as active listening		life span and function.	
and respect.		-Differentiate between marginating and circulating pools of WBCs.	
·		-Understand the principle behind the total,	
4-Improve problem-		relative and absolute WBCs count.	
solving skills.		-Describe the properties of phagocytic	
		WBCs and physiological leukocytosis.	
5-Demonstrate		-Describe the tissue macrophages and the	
knowledge of		reticulo-endothelial system.	
resources and tools			
	Biochemistry:	-Understand and illustrate the steps of the	
lifelong learning 1	1. Erythrocyte	Embden-Meyerhof pathway, and how it	
	Metabolism (1)	helps regulate the reduction of	
2	2. Erythrocyte	methemoglobin back to hemoglobin. -Explain how the Embden-Meyerhof	
	Metabolism (2)	pathway relates to 2,3-DPG production.	
3	<ol><li>Hemoglobin and</li></ol>	-Understand and describe the steps of the	
	Hemoglobinopathi	hexose monophosphate shunt.	
	es	-Describe the HMP shunt function and	
		explain on a biochemical basis how this	
		shunt helps to protect red cells from	
		oxidative stress.	
		-Explain how G6PD deficiency causes	
		favism.	
		-List the normal hemoglobins found in fetal	
		and adult bloodDescribe the genetics of sickle cell anemia	
		and the precipitating factors by which	
		hemoglobin S causes sickling.	
		-Identify the different chromosomes	
		responsible for alpha-globin and beta-	
		globin synthesis.	
		-Describe the basic genetic differences	
		between alpha-thalassemia and beta-	
		thalassemia.	
		-Describe the genetic and hematologic	
		differences between alpha-thalassemia	

		trait, hemoglobin H disease, and hydrops fetalis.  -Describe the genetic and hematologic differences between beta—thalassemia minor and beta— thalassemia major.  -Identify normal and abnormal hemoglobins by electrophoresis.	
*Topic 2: Red Blood Cells Diseases and Treatment (lectures: 17)  Biochemistry: 1 Microbiology: 5 Community Medicine: 2 Pathology: 4 Pharmacology: 5	Microbiology 1. Epstein-Barr Virus (EBV) and parvovirus B19 2. Plasmodium and babesiosis 3. Trypanosomiasi s, visceral leishmaniasis and filariasis 4. Salmonella typhi, enteric fever and brucellosis 5. Yersinia pestis, and plague; Q- Fever and other rickettsia	Describe the reactions and rate-limiting steps implicated in heme synthesis  Describe the clinical consequences of the congenital deficiency of the enzymes involved in heme synthesis.  List and explain the intrinsic and extrinsic causes of hemolytic anemias.  List laboratory investigations that are used in the diagnosis of hemolytic anemias.  Describe the virology, epidemiology, pathogenesis, clinical presentation, and management of Epstein-Barr virus  Describe the virology, epidemiology, pathogenesis, clinical presentation, and management of parvovirus B19.  Describe the morphology, life cycle, epidemiology, pathogenesis, immunity, clinical presentations, diagnosis, management, and prevention of malaria  Describe the general characteristics, epidemiology, pathogenesis, clinical presentation, and management of eishmania.  Describe the general characteristics, epidemiology, pathogenesis, clinical presentation, and management of filaria.  Describe the general characteristics, epidemiology, pathogenesis, clinical presentation, and management of silaria.  Describe the general characteristics, epidemiology, pathogenesis, clinical presentation, and management of trypanosoma.  Describe the general characteristics, epidemiology, pathogenesis, clinical presentation, laboratory diagnosis and management of salmonella  Describe the general characteristics, epidemiology, pathology and virulence, clinical presentation, laboratory diagnosis and management of brucella.  Describe the general characteristics, epidemiology, pathogenesis, clinical presentation, laboratory diagnosis and treatment of brucella.  Describe the general characteristics, epidemiology, pathogenesis, clinical presentation, and management of Y. pestis.	

Community  Medicine  1. Blood-borne Infections (1)	Describe the general characteristics, epidemiology, classification, pathogenesis, clinical presentation and management of Rickettsia.  -Understand the definition of blood-born infections (BBI)Discuss the epidemiological aspects, mode of transmission, and prevention of the most	
2. Blood-borne Infections (2)	common BBIs including hepatitis B, hepatitis C, and HIV, AIDS.  Describe the risk of occupational exposure to blood.  Discuss the postexposure management of BBI.  -Understand the epidemiological importance of malaria -Identify the types of malaria species.  Discuss the life cycle, transmission, presentation, diagnosis, and complications of malaria.  -Familiarize with the epidemiology of malaria	
Pathology	Familiarize with the epidemiology of malaria in Jordan.	
1,2,3. Anemia (microcystic, normocytic, and macrocytic anemias)  4. Bleeding and Coagulation disorders	<ul> <li>- Understand the definition of anemia.</li> <li>- Discuss the classification of anemia according to the underlying mechanism or morphology (macrocytic, microcytic, and normocytic)</li> <li>- Understand the clinical presentation and approach for anemias</li> <li>- Start the discussion with microcytic anemias "iron def. anemia, AOCD and thalassemia)</li> <li>- Then, macrocytic anemia, "megaloblastic anemia."</li> <li>- Discuss thalassemia and understand the major types, the underlying genetics and the different presentations.</li> <li>- Discuss the pathophysiology of sickle cell disease and the precipitating factors of sickling.</li> <li>- Understand the consequences of sickling, including a plastic crisis.</li> <li>- Discuss the mode of inheritance and pathogenesis of G6PD</li> <li>Understand paroxysmal nocturnal hemoglobinuria.</li> <li>- Discuss immune and non-immune</li> </ul>	
	mediated hemolytic anemia -Understand the types of immune- mediated hemolytic anemia, including A- Warm antibody type and B-Cold antibody type	

	-Identify the causes of abnormal bleeding, including vascular disorders, thrombocytopenia, platelet function defects, and defective coagulation -Discuss the underlying etiologies of thrombocytopenia and describe both acute and chronic idiopathic thrombocytopenia -Describe the pathology of hemophilia A and B and von Willebrand disease -Discuss thrombotic thrombocytopenia purpura and microangiopathic hemolytic anemia -Define disseminated intravascular coagulopathy and it's clinical implications
Pharmacology 1. Drugs for the treatment of anemia (1) 2. Antiplatelets, anticoagulants and thrombolytics (1) 3. Antiplatelets, anticoagulants and thrombolytics (2) 4. Antiplatelets, anticoagulants and thrombolytics (3) 5. Chemotherapy for Malaria	-List the different approaches utilized for the treatment of anemia based on its classificationDescribe the main characteristics of iron preparations, their therapeutic indications, pharmacokinetics, and major adverse effectsDescribe the mechanism of action of folic acid and vitamin B12, their therapeutic indications and major adverse effectsUnderstand the role of erythropoietin in the treatment of anemia, therapeutic guidelines, and major adverse effectsList pharmacological therapy utilized for the treatment of neutropeniaDescribe the role of hydroxyurea in the treatment of sickle cell anemia, its mechanism of action and overall contribution to disease outcomeUnderstand the roles of the endothelium, platelets, and coagulation pathway in the development of arterial and venous thrombosisDelineate the pharmacological targets of the platelet plug formation process, including platelet activation, adhesion, and aggregationUnderstand the mechanisms of action, therapeutic uses, therapeutic guidelines, and major adverse effects of platelet aggregation inhibitorsList the hematological and non-hematological uses of aspirin -Understand the mechanisms of action, therapeutic uses, therapeutic guidelines, and major adverse effects of parenteral anticoagulantsUnderstand the mechanisms of action, therapeutic uses, therapeutic guidelines, and major adverse effects of oral anticoagulants (direct and indirect agents)Compare between heparins and warfarin in terms of mechanism of action, route of administration, onset, duration of action, drug interactions, teratogenic effects, and antidote.

		-Describe mechanisms of action, therapeutic uses, therapeutic guidelines, and major adverse effects of thrombolyticsUnderstand the different pharmacological approaches to treat bleeding -Understand the mechanism of action, pharmacokinetics, adverse effects, and clinical uses of antimalarial drugs -Describe the currently implemented guidelines for the treatment of complicated and uncomplicated malaria -Describe the drug regimens utilized for the prophylaxis against malaria	
*Topic 3: White Blood Cells Diseases and Treatment (Lectures: 9)  Anatomy: 2 Pathology: 5 Pharmacology: 2	Anatomy  1. Anatomy and histology of the lymphatic system (1)  2. Anatomy and histology of the lymphatic system (2)	-Understand the major components and function of the lymphatic system -Describe the origin and composition of lymph -Describe the structure of lymphatic vessels, trunks, and ducts -Explain the anatomy of the thoracic duct and right lymphatic ducts -Describe the anatomy and histology of the thymus -Describe the structure, histology, and function of lymph nodes -Identify the anatomy, histology, and function of the spleen and tonsils	
	Pathology  1. Neoplastic proliferation of WBCs, Acute Leukemia (ALL+AML)  2. Myeloproliferative Neoplasms (MPN) and Myelodysplastic Syndromes (MDS)  3. Lymphoid neoplasms, non-Hodgkin lymphomas  4. NHL, Multiple myeloma, and related plasma cell disorders  5. Lymphoid neoplasms, Hodgkin lymphoma	- Define leukopenia and discuss the pathogenesis, clinical features, and morphology of neutropenia/agranulocytosis - Define leukocytosis and discuss the causes based on the specific type of white cells affected Discuss in detail infectious mononucleosis, including definition, cause, pathogenesis, clinical features, morphology, and diagnosis Discuss the classification of hematologic malignancies Discuss the definition, epidemiology, pathogenesis, clinical features, morphology, and prognosis of precursor B & T cell neoplasms (ALL) Be familiar with the definition, incidence, pathogenesis, clinical features, and morphologic and immunophenotypic features of acute myeloid leukemia Discuss the WHO and FAB classifications of AML Talk about the clinical course and prognostic factors of AML Summarize the major differences between AML and ALL - Define myeloproliferative neoplasms and discuss their general features.	
		Discuss the pathogenesis, clinical features, differential diagnoses, laboratory findings, morphology, and clinical course of chronic myeloid leukemia (CML).	

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		- Discuss the genetic background, clinical	
		features, laboratory findings, morphology, and	
		clinical course of polycythemia vera and	
		essential thrombocythemia.	
		- Briefly discuss primary myelofibrosis.	
		- Define myelodysplastic syndrome (MDS).	
		- Discuss the pathogenesis, clinical features,	
		WHO classification, and morphology of MDS	
		- Discuss the normal lymph node morphology	
		- Discuss acute non-specific lymphadenitis	
		- Discuss chronic non-specific lymphadenitis by	
		concentrating on the different morphologic	
		patterns that occur.	
		- Define lymphoid neoplasms and mention the	
		WHO classification.	
		- Be familiar with the definition, clinical,	
		morphologic, and immunophenotypic	
		features of low and intermediate grades	
		peripheral B cell lymphomas, including	
		CLL/SLL, follicular lymphoma, MALT lymphoma,	
		and Mantle cell lymphoma.	
		- Discuss diffuse large B cell lymphoma and be	
		familiar with its clinical features, morphology,	
		subtypes, and prognosis	
		Describe the clinical features, types,	
		pathogenesis, morphologic features, and	
		prognosis of Burkitt lymphoma.	
		- Talk briefly about Hairy cell leukemia and T-cell	
		lymphoma/leukemia	
		Define plasma cell neoplasms	
		Describe the spectrum of plasma cell dyscrasias	
		Discuss multiple myeloma, including the	
		· · · · · ·	
		pathogenesis, clinical features, diagnosis, and morphologic and immunophenotypic features.	
		- Briefly talk about Monoclonal Gammopathy of	
		Undetermined Significance (MGUS).	
		- Discuss the morphologic and clinical features of	
		lymphoplasmacytic lymphoma	
		-Discuss the general characteristics,	
		classification, types, clinical features,	
		morphologic features, immunophenotype, and	
		prognosis of Hodgkin lymphoma.	
		- Describe in detail the morphologic and clinical	
		features of classical and non-classical Hodgkin	
		lymphoma.	
		- Be familiar with the staging system of Hodgkin	
-	Dhawaa aa la su	and non-Hodgkin lymphoma.	
	Pharmacology	-Understand the main steps involved the	
	1. Immunosuppre	activation of T lymphocytes	
	ssants	-Identify targetable molecular processes	
	2. Selected	involved in immunosuppression	
	chemotherapy	-Understand the role of	
	for the	immunosuppressants in organ	
	treatment of	transplantation and the treatment of	
	leukemia and	autoimmune diseases	
		-List the major pharmacotherapies utilized	
	lymphoma	for the induction of immunosuppression	

Γ		-List the major pharmacotherapies utilized	
		for the maintenance of	
		immunosuppression	
		-Understand the main differences between	
		monoclonal and polyclonal antibodies	
		-Understand the different phases of the	
		treatment of leukemia	
		-Understand the mechanisms of action,	
		therapeutic uses, therapeutic guidelines,	
		and major adverse effects of cytotoxic	
		chemotherapy utilized for the treatment of	
		hematological malignancies.	
		-Understand the mechanisms of action,	
		therapeutic uses, therapeutic guidelines,	
		and major adverse effects of targeted	
		therapies utilized for the treatment of	
		hematological malignancies.	
		-List the most frequently used	
		chemotherapy regimens for the treatment	
		of hematological malignancies.	
Practica	- '	-Identify the morphological characteristics	
Sessions	<u> </u>	of neutrophils, eosinophils, basophils,	
	Histology of	lymphocytes and monocytes under light	
	The	and electron microscopeIdentify the histological features of	
	Hematopoietic	-	
	System	platelets under light and electron microscope.	
		-Study the microscopic structure of bone	
		marrow.	
		-Identify the of different stages of	
		hemopoiesis under light microscope	
		-Identify the histological features of the	
		thymus	
		-Identify the histological characteristics of	
		lymph nodes	
		-Identify the gross anatomy of the spleen	
		and palatine tonsils.	
	Physiology Labs:	-Define Erythrocytes sedimentation Rate	
	1. RBCs: ESR,	(ESR), Demonstrate its measurement using	
	PCV; Blood	Westergren tube and identify the clinical	
	Grouping; and	significance	
	Blood indices	-Demonstrate the PCV test using the	
	2. Tests of	microhematocrite.	
	Hemostasis:	-Ask students to find their own blood group	
	Clotting,	and the percentage of each blood group of	
		the students attending the practical	
	Bleeding Time,	session.	
	capillary	-Understand the blood indices of the CBC	
	fragility and	test.	
	interpretation	-Use the capillary method to determine the	
	of lab results.	clotting time.	
		-Determine the bleeding time by using the	
		filter paperApply Hess test to assess the capillary	
		fragility	
		-Learn how to interpret laboratory tests on	
		coagulation profile.	

Pathology: 1. Blood Culture  Pathology: 1. Pathology of non-Hodgkin Lymphoma 2. Neoplastic diseases of WBCs	-Define bacteremia and determine its causes, types, and clinical course.  -Understand the indications, causes of contamination, approach to venipuncture, volume, number, and timing of blood culture.  -Explain the steps implicated in the laboratory processing and interpretation of blood culture  -Familiarize with case studies of bacteremia.  -Briefly describe the architecture of normal lymph node.  -Describe the gross and morphologic features of non-Hodgkin lymphoma types, the immunohistochemistry, and the flowcytometry.  - Briefly discuss the gross and microscopic features of the normal lymph node.
WBCS	features of the normal lymph node.  - Describe the morphologic features seen in acute leukemia and different chronic myeloid neoplasms.  - Be familiar with the morphologic features seen in the different types of non-Hodgkin and Hodgkin lymphomas.  - Describe the gross and microscopic features seen in plasma cell neoplasmS

## **ACADEMIC SUPPORT**

It is The Hashemite University policy to provide educational opportunities that ensure fair, appropriate, and reasonable accommodation to students who have disabilities that may affect their ability to participate in course activities or meet course requirements. Students with disabilities are encouraged to contact their Instructor to ensure that their individual needs are met. The University through its Special Need section will exert all efforts to accommodate for individual needs.

**Special Needs Section: Student Services and Care Unit** 

Tel: 053903333 ext. 4132 / 4583 / 5023 Location: Deanship of Students Affairs

Email: stydent@hu.edu.jo

## **COURSE REGULATIONS**

#### **Participation**

Class participation and attendance are important elements of every student's learning experience at The Hashemite University, and the student is expected to attend all classes. A student should not miss more than 15% of the classes during a semester. Those exceeding this limit of 15% will receive a failing grade regardless of their performance. It is a student's responsibility to monitor the frequency of their own absences. Attendance record begins on

the first day of class irrespective of the period allotted to drop/add and late registration. It is a student's responsibility to sign-in; failure to do so will result in a non-attendance being recorded.

In exceptional cases, the student, with the Instructor's prior permission, could be exempted from attending a class provided that the number of such occasions does not exceed the limit allowed by the University. The Instructor will determine the acceptability of an absence for being absent. A student who misses more than 25% of classes and has a valid excuse for being absent will be allowed to withdraw from the course.

## Plagiarism

Plagiarism is considered a serious academic offense and can result in your work losing marks or being failed. HU expects its students to adopt and abide by the highest standards of conduct in their interaction with their professors, peers, and the wider University community. As such, a student is expected not to engage in behaviors that compromise his/her own integrity as well as that of the Hashemite University.

Plagiarism includes the following examples, and it applies to all student assignments or submitted work:

- Use of the work, ideas, images or words of someone else without his/her permission or reference to them.
- Use of someone else's wording, name, phrase, sentence, paragraph or essay without using quotation marks.
- Misrepresentation of the sources that were used.

The Instructor has the right to fail the coursework or deduct marks where plagiarism is detected.

#### Late or Missed exams:

In all cases of assessment, students who fail to attend an exam, on the scheduled date without prior permission, and/or are unable to provide an accepted medical note, will automatically receive a failure grade for this part of the assessment.

## **Student Complaints Policy**

Students at The Hashemite University have the right to pursue complaints related to faculty, staff, and other students. The nature of the complaints may be either academic or non-academic. For more information about the policy and processes related to this policy, you may refer to the students' handbook.

## **COURSE ASSESSMENT**

#### Course Calendar and Assessment

Students will be graded through the following means of assessment, and their final grade will be calculated from the forms of assessment as listed below, with their grade weighting considered.

Assessment	Grade	Material	Date
Exam 1	40%	TBD	TBD
Exam 2	20%	Practical Labs	TBD
Final Exam	40%	Inclusive	TBD

## **Description of Exams**

Test questions will predominately come from the material presented in the lectures. The exam will consist of multiple-choice questions for the regular exams and short essay questions for makeup exams (for students with accepted excuses, only documented absences will be considered as per HU guidelines).

Grades are not negotiable and are awarded to the MD program according to the following criteria\*:

Letter Grade	Description	<b>Grade Points</b>
A+	Excellent	4.00
A		3.75
A-		3.50
B+	Very Good	3.25
В		3.00
B-		2.75
C+	Good	2.50
С		2.25
C-		2.00
D+	Pass	1.75
D	Pass	1.50
F	Fail	0.00
1	Incomplete	-

WEEKLY LECTURE SCHEDULE AND CONTENT DISTRIBUTION

<sup>\*</sup>Provided separately