Academic year: 2022/2023



Approval date: Issue:

Credit hours: 3

Course Syllabus

Bachelor

Course information

Course#	Course title		Prere	equisite		
0240101	General biology 1					
Course type			Class time	e	Room #	
□ University Requirement		🛛 Faculty Re	equirement			
🗆 Major Requirement		□ Elective		Mon, Wed: 9:45	- 11:15	61303
Compulsory						

Instructor Information

Name	Office No.	Phone No.	Office Hours	E-mail
Ayat Al-Azab	1018	2475	Sun, Tue: ٩:٤٥ – ١١:٠٠ Mon, Wed: ١٤:٠٠ –13:00	aalazab@philadelphia.edu.jo

Course Delivery Method

Course Delivery Method			
☑ Physical □ Online □ Blended			
Learning Model			
Precentage	Synchronous	Asynchronous	Physical

Course Description

This course involves studying the major concepts of biology including the structural and functional basics of the cell, metabolism and conversion of energy in respiration, cell division and DNA replication, transcription and translation, the basic principles of biotechnology in terms of hybridization, PCR and sequencing. In addition, the course covers the nervous, circularity and immunity systems in human.

Course Learning Outcomes

Number	Outcomes	Corresponding Program outcomes
	Knowledge	
K001	2.1 Distinguish between element and compound	Kp1
K002	3.2 Explain the major properties of water	Kp1
K003	4.2.1 Identify the major chemical functional groups	Kp1
K004	4.2.2 Understand the properties of each chemical group	Kp1
K005	5.1.1 List the different biological macromolecules found in	Kp1
	living organisms	
K006	5.1.2 Identify the monomers of each macromolecules	Kp1
K007	5.2.1Explain the importance of carbohydrate	Kp1

K008	5.2.2 Identify the linkage formed between two sugar	Kp1
K000	5.2.3 Compare between different polysaccharides (starch	Kn1
K009	s.z.s Compare between unrerent porysacchandes (starch,	крі
K010	5.3 Identify the structure of lipids	Kn1
K010	5.4 11 ist the functions of proteins and example of each one	Kp1
K012	5.4.2 Explain the formation of peptide bond	Kp1
K012 K013	5.4.3 Identify the different structures of protein	Kp1
K013	5.5.1 Describe the structure of puckie acids	Kp1
K014 K015	5.5.7 Describe the structure of huciele actus	Kp1
K015 K016	6.2.1 Distinguish between prokervotic and eukervotic cells	Kp1
K010 K017	6.2.2 Explain why there are both upper and lower limits to	крі
K 017	cell size. Be able to calculate the surface area to volume	Kn1
	ratio of a cube	крі
K018	6.2.3 Explain the advantage of compartmentalization in	
KUIO	outervotic calls	Kp1
K019	6.2.4 Compare and contract plant and animal cells	Kn1
K020	6.3.1 Describe the structure and function of the nuclear	K p1
1020	envelope, including the role of the pore complex	Kp1
K021	6.3.2 Briefly explain how the nucleus controls protein	
1021	synthesis in the cytoplasm	Kp1
K022	6.3.3 Explain the role of the nucleolus in protein synthesis	Kn1
K022	6.3.4 Distinguish between free and bound ribosomes in	K p1
1025	terms of location and function	Kp1
K024	6.4.1 L ist the components of the endomembrane system	
11024	and describe the structure and function of each component	Kp1
K025	6.4.2 Describe the path that a protein destined for the	
11025	organelles of the endomembrane system, the plasma	Kp1
	membrane or the outside of cell would follow.	iip i
K026	6.4.3 Compare the structure and functions of smooth and	
	rough ER.	Kpl
K027	6.4.4 Explain the significance of the <i>cis</i> and <i>trans</i> sides of	¥7. 1
	the Golgi apparatus.	Kpl
K028	6.4.5 Describe three examples of intracellular digestion by	17 1
	lysosomes.	Kpl
K029	6.4.6 Name three different kinds of vacuoles, giving the	IZ 1
	function of each kind.	Kpl
K030	6.5.1 Briefly describe the energy conversions carried out	V n 1
	by mitochondria / chloroplasts.	крі
K031	6.5.2 Describe the structure and function of mitochondria	Kn1
	and chloroplasts.	ĸpi
K032	6.5.3 Describe the evidence that mitochondria (and	Kn1
	chloroplasts) evolved from a prokaryotic endosymbiont.	крі
K033	6.5.4 Explain the roles of peroxisomes in eukaryotic cells.	Kp1
K034	6.6.1 Compare the structure, monomers, and functions of	Kn1
	microtubules, microfilaments, and intermediate filaments.	крт
K035	6.7.1 Describe the basic structure of a plant cell wall.	
	Distinguish between the primary cell wall, middle lamella,	Kp1
	and secondary cell wall.	
K036	6.7.2 Describe the structure and roles of the extracellular	Kn1
	matrix in animal cells.	
K037	6.7.3 Explain how the extracellular matrix may act to	.
	integrate changes inside and outside the cell. Describe the	Kp1
	proteins and glycoproteins involved in this process.	
K038	6./.4 Name the intercellular junctions found in plant and	T7 4
	animal cells and described the differences between the	Kpl
	intercellular junctions in animal tissues.	

K039	7.1.1 Explain the meaning of the statement that phospholipids and most other membrane constitutes (e.g.,	Kp1
K040	 7.1.2 Describe the fluidity of the components of a cell membrane and explain how membrane fluidity is 	Kp1
K041	influenced by temperature and membrane composition. 7.1.3 Explain how cholesterol resists changes in membrane	
K042	fluidity as temperatures change.	Kpl
K042	membrane proteins.	Kp1
K043	7.1.5 List six major functions of membrane proteins.	Kp1
K044	7.1.6 Explain the role of membrane carbohydrates in cell- cell recognition.	Kp1
K045	7.2.1 Explain how hydrophobic molecules cross cell	Kp1
K046	7.2.2 Distinguish between channel proteins and carrier	Kp1
K047	7.2.3 Explain how aquaporins facilitate the passage of water through membranes	Kp1
K048	7.3.1 Define diffusion. Explain why diffusion is a passive and spontaneous process.	Kp1
K049	7.3.2 Explain why a concentration gradient of a substance across a membrane represents potential energy.	Kp1
K050	7.3.3 Distinguish between solutions that are hypertonic, hypotonic, and isotonic to cell contents.	Kp1
K051	7.3.4 Define osmosis and predict the direction of water movement based on differences in solute concentrations.	Kp1
K052	7.3.5 Describe how living cells with and without cell walls regulate water balance.	Kp1
K053	7.3.6 Explain how transport proteins facilitate diffusion.	Kp1
K054	7.3.7 Distinguish between osmosis, facilitated diffusion, and active transport.	Kp1
K055	7.4.1 Describe the two forces that combine to produce an electrochemical gradient.	Kp1
K056	7.4.2 Explain how electrogenic pump creates voltage	Kp1
K057	7 4 3 Describe the process of cotransport	Kn1
K058	7.5.1 Explain how large molecule are transported across a	iip i
11000	cell membrane.	Kp1
K059	7.5.2 Distinguish between exocytosis and receptor- mediated endocytosis.	Kp1
K060	8.1.1 Explain the role of catabolic and anabolic pathways in cellular metabolism.	Kp1
K061	8.1.2 Distinguish between Kinetic and potential energy.	Kp1
K062	8.2.1 Distinguish between exergonic and endergonic reactions in terms of available energy change.	Kp1
K063	8.3.1 List three main kinds of cellular work and provide examples of each. Explain in general terms how cells obtain the energy to do cellular work.	Kp1
K064	8.3.2 Describe the structure of ATP and identify the major class of macromolecules to which ATP belongs.	Kp1
K065	8.3.3 Explain how ATP performs cellular work.	Kp1
K066	8.4.1 Describe the function of enzymes in biological systems.	Kp1
K067	8.4.2 Explain how enzyme structure determines enzyme	Kp1
K068	8.4.3 Explain the induced-fit model of enzyme function.	Kp1

K069	8.4.4 Describe the mechanisms by which enzymes lower	Kp1
	activation energy.	<u>F</u> -
K070	8.4.5 Explain how substrate concentration affects the rate	Kn1
	of an enzyme-catalyzed reaction.	
K071	8.4.6 Explain how temperature, pH, cofactors, and enzyme	Kn1
	inhibitors can affect enzyme activity.	11p1
K072	8.4.7 Distinguish between a competitive and	Kn1
	noncompetitive inhibitors.	11p1
K073	8.5.1 Describe how allosteric regulators may inhibit or	Kn1
	stimulate the activity of an enzyme.	11p1
K074	8.5.2 Explain how the binding of oxygen to hemoglobin	Kn1
	illustrates cooperativity.	
K075	8.5.3 Explain how feedback inhibition prevents a cell from	Kn1
	wasting chemical resources.	•••p1
K076	8.5.4 Describe how localization of enzymes within a cell	Kn1
	may help order metabolism.	11p1
K077	9.1.1 Distinguish, in general terms, between fermentation	Kn1
	and aerobic respiration.	11p1
K078	9.1.2 Define oxidation and reduction. Be able to identify	
	oxidizing agents and reducing agents as well as identify	Kn1
	which molecules are being oxidized/reduced in an	T tp1
	equation.	
K079	9.1.3 Explain in general terms how redox reactions are	Kn1
	involved in energy exchanges.	
K080	9.1.4 Describe the structure and the role of NAD ⁺ in	Kp1
	cellular respiration.	F -
K081	9.1.5 Explain, in general terms, the role of the electron	Kp1
	transport in cellular respiration.	1
K082	9.2.1 Name the three stages of cellular respiration and state	
	the region of the eukaryotic cell where each stage occurs.	
	State the net amount of ATP generated by each stage and	Kp1
	the type of phosphorylation that occurs. Be able label or	*
	draw a representation of centular respiration in a eukaryotic	
V092	0.2.2 Describe how the earbon skeleton of alwayse shanges	
K005	3.2.2 Describe now the carbon skeleton of glucose changes	Kn1
	as it proceeds through grycorysis (comparing grucose and	крт
K084	0.3.1 What is the energy yield of the citric acid cycle?	Kn1
K084 K085	9.5.1 What is the energy yield of the clube active cycle:	крт
K 005	and oxidative phosphorylation	Kp1
K086	0.4.2 In general terms, explain how the every sonic "slide"	
KUUU	of electrons down the electron transport chain is coupled to	Kn1
	the endergonic production of ATP by chemiosmosis	K p1
K087	9.4.3 Explain where and how the electron transport chain	
1007	creates a proton gradient	Kp1
K088	9.5.1 Distinguish between fermentation and anaerobic	
Rooo	respiration	Kp1
K080	9.5.2 State the basic function of fermentation	Kn1
K000	9.5.2 Compare the fate of pyruvate in alcohol fermentation	1321
11070	and lactic acid fermentation	Kp1
K091	9.6.1 Describe how food molecules other than alucose can	
11071	be oxidized to make ATP.	Kp1
K092	9.6.2 Explain how we regulate catabolic nathways	Kn1
K093	10.1 Explain the process of photosynthesis	Kn1
K094	10.2 Understand the reactions associated with	Kn1
	photosynthesis	••r•
K095	10.3 Understand Calvin cycle	Kn1
		r -

K096	16.1.1 Explain why researchers originally thought protein was the genetic material.	Kp1
K097	16.1.2 Explain how the experiments performed by the	
	following scientists provided evidence that DNA is the	
	genetic material:	
	a. Frederick Grifffith	Kp1
	b. Oswald Avery, Maclyn McCarty, and Colin Macleod	H p1
	c. Alfred Hershev and Martha Chase	
	d. Erwin Chargaff	
K098	16.1.3 Explain how Watson and Crick deduced the	
	structure of DNA and describe the evidence thy used.	¥7. 1
	Explain the significance of the research of Rosalind	Kpl
	Franklin.	
K099	16.1.4 Describe the structure of DNA. Explain the base-	17 1
	paring rule and describe its significance.	Kpl
K100	16.2.1 Describe the semiconservative model of replication	
	and the significance of the experiments of Matthew	Kp1
	Meselson and Franklin Stahl.	1
K101	16.2.2 Describe the process of DNA replication, including	¥7 1
	the role of the origins of replication and replication forks.	Kpl
K102	16.2.3 Explain the role of DNA polymerases in replication.	Kp1
K103	16.2.4 Explain the energy source drives the polymerization	TZ 1
	of DNA.	Kpl
K104	16.2.5 Distinguish between the leading strand and the	17 1
	lagging strand.	Kpl
K105	16.2.6 Explain how the lagging strand is synthesized even	
	though DNA polymerase can add nucleotides only to the 3'	Kp1
	end. Describe the significance of Okazaki fragments.	•
K106	16.2.7 Explain the role of DNA ligase, primer, primase,	V. 1
	helicase, topoisomerase, and single strand binding proteins.	крі
K107	16.2.8 Define "antiparallel" and explain why continuous	Kn1
	synthesis of both DNA strands is not possible.	ĸpi
K108	16.2.9 Explain the roles of DNA polymerase, mismatch	
	repair enzymes, and nuclease in DNA proofreading and	Kp1
	repair.	
K109	16.2.10 Describe the structure and function of telomers.	Kp1
K110	16.2.11 Explain the possible significance of telomerase in	Kn1
	germ cells and cancerous cells.	крт
K111	16.3.1 Compare bacterial chromosome and eukaryotic	Kn1
	chromosome.	крт
K112	16.3.2 Describe how the packing of chromatin changes	Kn1
	during the course of the cell cycle.	крі
K113	16.3.3 Distinguish between heterochromatin and	Kn1
	euchromatin.	npi
K114	17.1.1 Explain the reasoning that led Archibald Garrod to	Kn1
	suggest that genes dictate phenotypes through enzymes.	iip i
K115	17.1.2 Describe Beadle and Tatum's experiments with	
	<i>Neurospora</i> and explain the contribution they made to our	Kp1
	understanding of how genes control metabolism.	
K116	17.1.3 Distinguish between the "one gene-one enzyme"	
	hypothesis and the "one gene-one polypeptide" hypothesis	Kpl
77115	and explain why the original hypothesis was changed.	T 7 1
<u>K117</u>	1/.1.4 Explain how RNA differs from DNA.	Kpl
K118	1/.1.5 Briefly explain how information flows from gene to	Kp1
****	protein. Is the central dogma ever violated?	1
K119	17.1.6 Distinguish between transcription and translation.	Kpl

K120	17.1.7 Compare where transcription and translation occur in bacteria and in eukarvotes.	Kp1
K121	17.1.8 Define "codon" and explain the relationship	
	between the linear sequence of codons on mRNA and the	Kp1
	linear sequence of amino acids in a polypeptide.	1
K122	17.1.9 Explain the early techniques used to identify what	
	amino acids are specified by the triplets UUU, AAA,	Kp1
	GGG, and CCC.	ľ
K123	17.1.10 Explain why polypeptides begin with methionine	Vn1
	when they are synthesized.	ĸpi
K124	17.1.11 Explain what is means to say that the genetic code	Kn1
	is redundant and unambiguous.	крт
K125	17.1.12 Explain the significance of the reading frame	Kn1
	during translation.	крт
K126	17.1.13 Explain the evolutionary significance of nearly	Kn1
	universal genetic code.	крі
K127	17.2.1 Explain how RNA polymerase recognizes where	
	transcription should begin. Describe the role of the	Kp1
	promoter, the terminator and the transcription unit.	
K128	17.2.2 Explain the general process of transcription,	/
	including the three major steps of initiation, elongation and	Kpl
W100	termination.	
K129	17.3.1 Explain how RNA is modified after transcription in	Kp1
K120	eukaryotic cells.	1
K130	17.3.2 Define and explain the role of ribozyme. What three	IZ 1
	properties allow some RNA molecules to function as	Kpl
V121	17.2.2 Describe the functional and evolutionary	
K 151	significance of introns	Kp1
K132	17.3.4 Explain why due to alternative RNA splicing the	
K 152	number of different protein products an organism can	Kn1
	produce is much greater than its number of genes	крі
K133	17.4.1 Describe the structure and function of tRNA	Kn1
K134	17.4.2 Explain the significance of wobble.	Kp1
K135	17.4.3 Explain how tRNA is joined to the appropriate	
	amino acid.	Kpl
K136	17.4.4 Describe the structure and functions of ribosomes.	Kp1
K137	17.4.5 Explain the statement, "A ribosome can be regarded	I I
	as one colossal ribozyme".	Kpl
K138	17.4.6 Describe the process of translation (including	
	initiation, elongation, and termination) and explain which	Kn1
	enzymes, protein factors, and energy sources are needed	Kpi
	for each stage.	
K139	17.4.7 Describe the significance of polyribosomes.	Kp1
K140	17.4.8 Explain what determines the primary structure of a	
	protein and describe how a polypeptide must be modified	Kp1
	before it becomes fully functional.	
K141	17.4.9 Describe what determines whether a ribosome will	T 7 1
	be free in the cytosol or attached to the rough endoplasmic	Kpl
V 140	reticulum.	
K142	1/.5.1 Define "point mutations". Distinguish between	V a 1
	overpair substitutions and base-pair insertions. Give an	крі
K1/2	17.5.2 Distinguish between a missense and a personse	
K143	mutation	Kp1
K144	17.5.3 Why is an insertion or deletion more likely to be	
12177	deleterious than a substitution?	Kp1

K145	17.5.4 Define the term "mutation". Give an example of a	Kn1
	physical and a chemical agent of mutation.	Kpi
K146	17.5.5 Briefly, compare gene expression in bacteria,	
	archaea, and eukarya. In general, is archaeal gene	Kp1
	expression more similar to bacterial or eukaryotic gene	
17.1.47	expression?	
K147	17.5.6 Describe the historical evolution of the concept of a	Kp1
K1/18	10.1.1 Describe the structure of virus	Kn1
K140	19.1.2 List the different components of virus	Kp1
K150	19.7.2 East the different components of virus	Kp1
K151	20.1.1 Describe nucleic acid hybridization and how it is	
	widely used in DNA technology	Kpl
K152	20.1.2 Describe that DNA sequencing can be carried out	
	using dideoxy sequencing method in automated	Kp1
	sequencing machines.	
K153	20.1.3 Explain the fast and inexpensive next-generation	
	(high-throughput) techniques for sequencing DAN are	Kp1
	based on sequencing by synthesis.	
K154	20.1.4 Introduce the third sequencing methods including	Kp1
V155	nanopore technology	X
K155	20.1.5 Describe the natural function of restriction enzymes	Kn1
	technology	Kpi
	20.1.6 Explain how the creation of sticky ends by	
K156	restriction enzymes is useful in producing a recombinant	Kp1
	DNA molecule.	
17167	20.1.7 Outline the procedures for cloning a eukaryotic gene	IZ 1
K157	in bacterial plasmid.	Kpl
	20.1.8 Explain the rational for including a gene antibiotics	
K158	resistance and a gene that codes for a hydrolytic enzyme in	Kp1
	the plasmids.	
K159	20.1.9 Define and distinguish between genomic libraries	Kp1
V160	using plasmids, phages, and cDNA.	
K 100	20.1.10 Describe the fole of an expression vector.	Kpi
K161	instead of bacteria as bost for cloning or expression	Kn1
K 101	enkarvotic genes	K p1
W1 60	20.1.12 Describe the structure and function of a veast	¥7 1
K162	artificial chromosome (YAC).	Kpl
V162	20.1.13 Describe two techniques to introduce recombinant	V n 1
K 105	DNA into eukaryotic cells.	Kp1
K164	20.1.14 Describe the polymerase chain reaction (PCR) and	Kn1
	explain the advantages and limitations of this procedure.	
	Skills	
	Competencies	
C01	6.1 Four a cell and become acquainted with its	Cp1
	7.1 Learn how callular membranes control the passage of	
C02	substances often with transport proteins	Cp1
	7.2. Understand how the plasma membrane and its proteins	
C03	enable cells to survive and function.	Cp1
~ ^ :	8.1 Understand how matter and energy flow during life's	~ .
C04	processes.	Cpl
C05	8.2 Understand how energy flow is regulated.	Cp1
COF	9.1 Consider how cells harvest the chemical energy stored	Cn1
00	in organic molecules.	Срі

C07	9.2 Know how energy is used it to generate ATP.	Cp1
C08	9.3 Know the pathways of respiration and fermentation.	Cp1
C10	 16.1 Discovering how biologists deduced that DNA is the genetic material and how Watson and Crick worked out its structure. 16.2 Learn how a molecule of DNA is copied and how cells repair their DNA. 16.3 Explore how a DNA is packed together with proteins in a chromosome. 	Cp1
C11	17.1 Describe the flow of information from gene to protein.	Cp1
C12	17.2 Explain how genetic mutations affect organisms through their proteins.	Cp1
C13	17.3 Understand the processes of gene expression.	Cp1
C14	20.1 Describe the main techniques for sequencing and manipulating DNA.	Cp1

Learning Resources

Course textbook	Reece, J. B., Urry, L. A., Cain, M. L., Wasserman, S. A., Minorsky, P. V.,
	& Jackson, R. B. (2014). <i>Campbell biology</i> (Vol. 9). Boston: Pearson.
Supporting References	
Supporting websites	
Teaching Environment	⊠Classroom □ laboratory □Learning platform □Other

Meetings and subjects timetable

Week	Торіс	Learning Methods*	Tasks	Learning Material Text book
1	Introduction The chemical context of life 2.1. Matter consists of chemical elements in pure form and in combinations called compounds.	Lectures		P. 28-43
2	Water and life 3.1. Polar covalent bonds in water molecules result in hydrogen bonding 3.2. Four emerging properties of water contribute to Earth's suitability for life.	Lectures		P. 44-55

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	Carbon and the molecular diversity of life		
	4.2. Carbon atoms can form diverse molecules by		
	bonding to four other atoms. Exploring some biologically		
	important chemical groups.		
	The structure and the function of large biological		
	molecules		
	5.1. Macromolecules are polymers built from monomers		
	5.2. Carbohydrates serves as fuel and building material		
3.4	5.3. Lipids are a diverse group of hydrophobic Molecules	Lectures	P. 66-91
- , -	5.4. Proteins include a diversity of structures, resulting in		
	a wide range of functions		
	5.5 Nucleic acids store transmit and help express		
	hereditary information		
	A tour of the cell		
	6.2 Fukarvotic cells have internal membranes		
	compartmentalize the functions		
	6.2 The sukernotic cell's genetic instructions are housed		
	6.5. The eukaryotic cell's genetic instructions are noused		
	in the nucleus and carried out by the ribosomes		
	6.4. The Endomembrane system regulates protein traffic	.	D 00 105
5,6	and performs metabolic functions.	Lectures	P. 93-125
	6.5. Mitochondria and chloroplasts change energy from		
	one form to another		
	6.6. The Cytoskeleton is a network of fibers that organizes		
	structures and activities in the cell (In Brief).		
	6.7. Extracellular components and connections between		
	cells help coordinate cellular activities.		
	Membrane structure and function		
	7.1. Cellular membranes are fluid mosaics of lipids and		
	proteins.		
7	7.2. Membrane structures results in selective		
	permeability		
	7.3. Passive transport is diffusion of a substance across a	Lectures	P. 126-142
	membrane with no energy investment		
	7.4. Active transport uses energy to move solutes against		
	their gradients		
	7.5 Bulk transport across the plasma membrane occurs		
	by exocytosis and endocytosis		
	An introduction to metabolism		
	8.1. An Organism's metabolism transforms matter and		
	energy subject to the laws of thermodynamics.		
	8.2 The Free energy change of a reaction tells us whether		
	or not the reaction occurs spontaneously. Free energy and		
	metabolism: Equilibrium and Metabolism		P. 143-163
8	8.3 ATP powers cellular work by coupling everyonic to	Lectures	
	ondergonic reactions		
	8.4. Enzymas speed up matchelic reactions by lowering		
	anargy harriora		
	energy damers.		
	8.5. Regulation of enzyme activity netps control		
0	Midtorm ovom		
<u> </u>	Collular regnization and formantation		
	0.1 Cotabolic pothyeye vield energy by ovidining		
	organia fuala		
	0.2 Chucolucio homeosto chomical anorre ha anidicia		
10	9.2. Glycolysis harvests chemical energy by oxidizing	Lectures	P. 164-186
gl 9. co m	giucose to pyruvate		
	9.5. After pyruvate is oxidized, the citric acid cycle		
	completes the energy-yielding oxidation of organic		
	molecules		

	 9.4. During oxidation phosphorylation, chemiosmosis couples electron transport to ATP synthesis 9.5. Fermentation and anaerobic respiration enable cells to produce ATP without the use of oxygen 9.6. Glycolysis and citric acid cycle connect to many other metabolic pathways 		
11	 Photosynthesis 10.1. Photosynthesis converts light energy to the chemical energy of food 10.2. The light reactions convert solar energy to the chemical energy of ATP and NADPH 10.3. The Calvin cycle uses the chemical energy of ATP and NADPH to reduce CO₂ to sugar. 	Lectures	P.187-211
13	The molecular basis of inheritance 16.1. DNA is the genetic material 16.2. Many proteins work together in DNA replication and repair 16.3 A chromosome consists of a DNA molecule packed together with proteins.	Lectures	P. 314-334
14	Gene expression: From gene to protein 17.1. Genes specify proteins via transcription and translation 17.2. Transcription is the DNA-directed synthesis of RNA: <i>a closer look</i> 17.3. Eukaryotic cells modify RNA after transcription 17.4. Translation is the RNA-directed synthesis of a polypeptide: a <i>closer look</i> 17.5. Mutations of one or few nucleotides can affect protein structure and function	Lectures	P. 335-363
15	Viruses 19.1. A virus consists of a nucleic acid surrounded by a protein coat 19.2. Viruses replicate only in host cells.		P. 398-414
	DNA tools and biotechnology 20.1. DNA sequencing and DNA cloning are valuable tools for the genetic engineering and biological inquiry		 P.415-423
16	Final exam		

* includes: Lecture, flipped Class, project- based learning, problem solving based learning, collaborative learning

Course Contributing to Learner Skill Development

Using Technology
Communication skills
Application of concepts learnt

Assessment Methods and Grade Distribution

Assessment Methods	Grade Weight	Assessment Time (Week No.)	Link to Course Outcomes
Mid Term Exam	% 30	9	K001-K076 C01-C05
Various Assessments *	% 30	2,4,6,7,8,11,13,14,15	K001-K164 C01-C14
Final Exam	% 40	16	K001-K164 C01-C14
Total	%100		

* includes: quiz, in class and out of class assignment, presentations, reports, videotaped assignment, group or individual projects.

Alignment of Course Outcomes with Learning and Assessment Methods

Number	Learning Outcomes	Learning Method*	Assessment Method**
	Knowledge		
K001-K164	All outcomes	Lectures	Quizzes and exams
Skills			
	Competencies		
C01-C14	All outcomes	Lectures	Quizzes and exams

* includes: Lecture, flipped Class, project- based learning, problem solving based learning, collaborative learning

** includes: quiz, in class and out of class assignment, presentations, reports, videotaped assignment, group or individual projects.

Course Polices

Policy	Policy Requirements	
Passing Grade	The minimum passing grade for the course is (50%) and the minimum final mark	
	recorded on transcript is (35%).	
	• Missing an exam without a valid excuse will result in a zero grade to be assigned to the exam or assessment.	
Missing Exams	• A Student who misses an exam or scheduled assessment, for a legitimate reason, must submit an official written excuse within a week from the exam or assessment due date.	
	• A student who has an excuse for missing a final exam should submit the excuse to the dean within three days of the missed exam date.	
Attendance	The student is not allowed to be absent more than (15%) of the total hours	
	prescribed for the course, which equates to six lectures days (M, W) and seven	
	lectures (S,T,R). If the student misses more than (15%) of the total hours	
	prescribed for the course without a satisfactory excuse accepted by the dean of the	
	faculty, s/he will be prohibited from taking the final exam and the grade in that	
	course is considered (zero), but if the absence is due to illness or a compulsive	

	excuse accepted by the dean of the college, then withdrawal grade will be recorded.		
Academic	Philadelphia University pays special attention to the issue of academic integrity,		
Honesty	and the penalties stipulated in the university's instructions are applied to those who		
	are proven to have committed an act that violates academic integrity, such as:		
	cheating, plagiarism (academic theft), collusion, and violating intellectual property		
	rights.		

Program Learning Outcomes to be assessed in this Course

Number	Learning Outcome	Course Title	Assessment Method	Target Performance level
1	Kpl	Environmental	Quizzes and	
1	Крт	biotechnology	exams	
2	Cn1	Environmental	Quizzes and	
2	Срг	biotechnology	exams	

Description of Program Learning Outcome Assessment Method

Number	Detailed Description of Assessment	
Kp1	Quizzes and exams	
Cp1	Quizzes and exams	

Assessment Rubric of the Program Learning Outcome