

# SYLLABUS OF THE DISCIPLINE "BIOLOGICAL CHEMISTRY"

1. General information				
Name of the faculty	Department of the Foreign Students (Dentistry Faculty)			
Educational program (branch,	22 Public Health,			
specialty, level of higher education,	221 Dentistry, second (master's) level of the higher			
form of training)	education, daily form			
Academic year	2022-2023			
Name of the discipline, code	Biological chemistry			
(electronic address is on the web-	Code OK 12			
site of the Danylo Halytsky Lviv	Kaf_biochemistry@meduniv.lviv.ua			
National Medical University)				
Department (name, address, tel, e-	Department of Biological Chemistry			
mail)	79010, Lviv, 54, Pekarska Street			
	tel. +38 (032) 275 76 02			
	Kaf_biochemistry@meduniv.lviv.ua			
Head of the Department ( <i>e-mail</i> )	Lesya Kobylinska, PhD, DrSci, professor			
	kobylinska_lesya@meduniv.lviv.ua			
Year of training (when the discipline	Second year of training			
is taught)				
Semester (when the discipline is	III-IV			
taught)				
Туре	Mandatory			
Staff (names, scholar degrees, e-	Iryna Fomenko, PhD, DrSci, professor			
mail)	irynafomenkolviv@gmail.com			
	Lesya Kobylinska, PhD, DrSci, associate professor			
	kobylinska_lesya@meduniv.lviv.ua			
	Natalya Denysenko, PhD, assistant professor			
	denysenko.natalka@gmail.com			
Erasmus yes/no (availability of the	No			
discipline for students in the framework of the <i>Erasmus</i> +)				
,	Lagua Kabulingka DhD DrSai professor			
A person, responsible for the syllabus (receiving comments	Lesya Kobylinska, PhD, DrSci, professor kobylinska_lesya@meduniv.lviv.ua			
regarding syllabus, e-mail)	KODYIIIISKa_lesya@meduniv.iviv.ua			
Number of ECTS credits	5			
	Total 150 h			
Number of hours ( <i>lectures/practical classes/students independent work</i> )	(16 lectures / 59 practical classes / 75 students individual			
classes/students independent work)	work)			
Language of training	English			
Information on consultations	Consultations are carried out according to the approved plan			
information on consultations	once per week during the academical year.			
	Consultations before exam are carried out according to the			
	approved plan by lectors.			
Address, telephone and working	-			
schedule of the Department				
	ort annotation to the course			
General characteristics, short course description, peculiarities, advantages				

Teaching of *Biological chemistry* at the Department of the Foreign Students (Dentistry Faculty) in Danylo Halytsky Lviv National Medical University is provided during the second year of studying.

The subject of study in the educational discipline "Biological Chemistry" is the chemical composition of living organisms (the human body) and the biochemical transformations to which the molecules that makeup them are subject. Biological chemistry is based on students' study of medical biology, biophysics, medical chemistry (bioorganic, bioinorganic, physical and colloidal chemistry), and morphological disciplines and are integrated with these disciplines; lays the foundations for students to study molecular biology, genetics, physiology, pathology, general and molecular pharmacology, toxicology and propaedeutics of clinical disciplines, which involves the integration of teaching with these disciplines and the formation of skills to apply knowledge of biological and bioorganic chemistry, primarily biochemical processes that take place in the body of a healthy and sick person, in the process of further education and professional activity; lays the disease, control over the effectiveness of the use of medicines and measures aimed at preventing the occurrence and development of pathological processes.

# 3. Aim and scope of the course

**1. The aim of the discipline** is to study biomolecules and molecular organization of cell structures, general patterns of enzymatic catalysis and biochemical dynamics of transformation of major classes of biomolecules (amino acids, carbohydrates, lipids, nucleotides, porphyrins, etc.), molecular biology and genetics of informational macromolecules. , ie molecular mechanisms of heredity and realization of genetic information, hormonal regulation of metabolism and biological functions of cells, biochemistry of special physiological functions.

**2. Learning objectives:** To determine the structure of bioorganic compounds and the functions they perform in the human body; the reactivity of the main classes of biomolecules, which provides their functional properties and metabolic transformations in the body; biochemical mechanisms of pathological processes in the human body; features of diagnostics of a physiological condition of an organism and development of pathological processes on the basis of biochemical researches; connection of features of structure and transformations in an organism of bioorganic compounds as bases of their pharmacological action as medicines; basic mechanisms of biochemical action and principles of directed application of different classes of pharmacological agents; biochemical and molecular bases of physiological functions of cells, organs and systems of the human body; functioning of enzymatic processes occurring in membranes and organelles to integrate metabolism in individual cells; norms and changes in biochemical parameters used to diagnose the most common human diseases; on the beginning of biochemical processes of metabolism and its regulation in ensuring the functioning of organs, systems and the whole human body.

Analyze the compliance of the structure of bioorganic compounds with the physiological functions they perform in the human body. Interpret the features of the physiological state of the organism and the development of pathological processes on the basis of laboratory tests. Analyze the reactivity of carbohydrates, lipids, amino acids, which provides their functional properties and metabolic transformations in the body. Interpret the features of the structure and transformations in the body of bioorganic compounds as the basis of their pharmacological action as drugs. Interpret the biochemical mechanisms of pathological processes in the human body and the principles of their correction. Explain the main mechanisms of biochemical action and the principles of targeted use of different classes of pharmacological agents. Explain the biochemical and molecular basis of physiological functions of cells, organs and systems of the human body. Analyze the results of biochemical studies and changes in biochemical and enzymatic parameters used to diagnose the most common human diseases. Classify the results of biochemical studies and changes in biochemical processes of metabolism and its regulation in ensuring the functioning of organs, systems and the whole human body.

**3.** Competences and learning outcomes, the formation of which provides the study of the discipline (*general competence -GC*):

- Ability to abstract thinking, analysis and synthesis.
- Knowledge and understanding of the subject area and understanding of professional activity.
- Ability to apply knowledge in practical activities.
- Ability to communicate in the national language both orally and in writing.
- Ability to communicate in English.
- Skills in using information and communication technologies.
- Ability to search, process and analyze information from various sources.
- Ability to identify, pose and solve problems.
- Ability to be critical and self-critical.
- Ability to work in a team.
- The ability to act socially responsibly and consciously.
- special (professional, subject):
- Ability to collect medical information about the patient and analyze clinical data.
- Ability to interpret the results of laboratory and instrumental research.
- Ability to diagnose: determine preliminary, clinical, final, accompanying diagnosis, emergency conditions.

### 4. Course prerequisites

Biological chemistry as a discipline:

- 1. based on the knowledge of students obtained on the basis of the State Standard of Education
- in such disciplines as medical biology, inorganic and organic chemistry, human anatomy, histology;provides a high level of general medical training;
- 3. lays the foundation for students to further master their knowledge of specialized theoretical and clinical professional and practical dental disciplines

and chinical professional and practical dental disciplines					
5. Program results of learning					
Learning outcome code	List of learning results     The content of the learning outcome	Reference to the competency matrix code			
The code is created when filling the syllabus (category: Kn – knowledge, general competencies, PL- professional competencies)	Learning outcomes determine that the student must know, understand and be able to perform, after completing the discipline. Learning outcomes follow from the set learning goals. To enroll in the discipline, it is necessary to confirm the achievement of each learning outcome.	Symbol of the Program Learning Outcome Code in the Higher Education Standard			
Kn-1	Know the structure of bioorganic compounds and the functions they perform in the human body.	PL-2			
Kn-2	Know the reactivity of the main classes of biomolecules, which provides their functional properties and metabolic transformations in the body.	PL-3			
Kn-3	To know the biochemical mechanisms of pathological processes in the human body.	PL-4			
Kn-4	To know about the peculiarities of the diagnosis of the physiological state of the organism and the development of pathological processes on the basis of biochemical studies.	PL-6			
Kn-5	To know about the peculiarities of the structure and transformations in the body of bioorganic compounds as the basis of their pharmacological action as drugs.	PL-9			

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	Know the basic mechanisms of	PL-10
Kn-6	biochemical action and the principles of	
<i>Kn-0</i>	targeted use of different classes of	
	pharmacological agents.	
	Know the biochemical and molecular	PL-14
Kn-7	basis of physiological functions of cells,	
	organs and systems of the human body.	
	Know the functioning of enzymatic	PL-15
	processes occurring in membranes and	
Kn-8	organelles to integrate metabolism in	
	individual cells.	
	Know the norms and changes in	PL-17
Kn-9	biochemical parameters used to diagnose	1 L-1/
Kh-3	the most common human diseases.	
	To know the beginnings of biochemical	
<i>V</i> <sub></sub> 10	processes of metabolism and its regulation	
Kn-10	in ensuring the functioning of organs,	
	systems and the whole human body.	
	Skills	[
	Analyze the compliance of the structure of	
Sk-1	bioorganic compounds with the	
57-1	physiological functions they perform in	
	the human body.	
	Interpret the features of the physiological	
GL 0	state of the organism and the development	
Sk-2	of pathological processes on the basis of	
	laboratory tests.	
	Analyze the reactivity of carbohydrates,	
~ ~ ~	lipids, amino acids, which provides their	
<i>Sk-3</i>	functional properties and metabolic	
	transformations in the body.	
	Interpret the features of the structure and	
	transformations in the body of bioorganic	
Sk-4	· ·	
	1	
	pharmacological action as drugs.	
GL 5	Interpret the biochemical mechanisms of	
Sk-5	pathological processes in the human body	
	and the principles of their correction.	
	Explain the main mechanisms of	
Sk-6	biochemical action and the principles of	
58.0	targeted use of different classes of	
	pharmacological agents.	
	Explain the biochemical and molecular	
Sk-7	basis of physiological functions of cells,	
	organs and systems of the human body.	
	Analyze the results of biochemical studies	
	and changes in biochemical and enzymatic	
Sk-8	parameters used to diagnose the most	
	common human diseases	
SI- 0	Classify the results of biochemical studies	
Sk-9	and changes in biochemical and enzymatic	
	parameters used to diagnose the most	

	common human diseases.	
	Interpret the importance of biochemical	
Sk-10	processes of metabolism and its regulation	
	in ensuring the functioning of organs,	
	systems and the whole human body.	
	Autonomy and responsibility	
AR-1	Be responsible for the timely acquisition	
	of modern knowledge.	
	Be responsible for the timely acquisition	
AR-2	of basic general and professional	
	knowledge.	
AR-3	Be responsible for the timeliness of	
	decisions in these situations.	
AR-4	Responsible for the quality of the tasks	
AR-5	Be responsible for the timely acquisition	
AR-5	of knowledge and handling of information	
AR-6	Be responsible for the quality of work.	
AR-7	Be responsible for your civic position and	
AR-7	activities	
	Be responsible for literacy in professional	
AR-8	communication.	
Сотр	etencies and profeccional competencies	
	Ability to abstract thinking, analysis and	
GC-1	synthesis.	
	Knowledge and understanding of the	
GC -2	subject area and understanding of	
	professional activity.	
<i>GC-3</i>	Ability to apply knowledge in practice.	
	Ability to communicate in the national	
<i>GC-5</i>	language both orally and in writing.	
GC-5	Ability to communicate in English.	
	Skills in the use of information and	
GC-6	communication technologies.	
	Ability to search, process and analyze	
<i>GC</i> -7	information from various sources.	
	Ability to identify, pose and solve	
<i>GC-9</i>	problems.	
GC-10	*	
	The ability to be critical and self-critical.	
GC-11	Ability to work in a team.	
GC-13	The ability to act socially responsibly and	
	consciously.	
<i>PC-1</i>	Ability to collect medical information	
	about the patient and analyze clinical data.	
<i>PC-2</i>	The ability to interpret the results of	
	laboratory and instrumental research.	
	Ability to diagnose: determine	
<i>PC-3</i>	preliminary, clinical, final, accompanying	
	diagnosis, emergency conditions.	
	Format and contents of the course	
Course format	eye	
(specify full-time or part-time)		
Kind of occupations	Number of hours	Number of groups
lectures	16	10

practical			59			10
seminars						
individu	al stident work		75			10
		7	Toping and content of the course			
Code	Topic	/.	Topics and content of the course Learning content	Code	of the	Lecturer
of classes type	Topic		Learning content	results	of the	Lecturer
L-1	Biochemistry as a science. Enzymes. Regulation of enzymatic processes. Enzymology.	and phy enz acti stuc	acquaint students with the subject tasks of biochemistry. Describe the rsicochemical properties of ymes, the mechanism of their on and regulation. To acquaint lents with enzyme diagnostics, yme therapy and enzymopathy	Kn-1 Kn-3 Kn-4 Kn-5 Kn-6 Kn-8 AR-1 AR-2 GC-1 GC-6		Prof. Kobylinska L.I.
L-2	Metabolism of carbohydrates, its regulation and changes in pathology	of c cart path glya glua of c the dev	acquaint students with the process carbohydrate digestion, the use of pohydrates in various metabolic nways, anaerobic and aerobic colysis, glycogen metabolism , coneogenesis, hormonal regulation carbohydrate metabolism. Describe causes, mechanism of elopment, diagnostic criteria for petes	Kn-1 Kn-2 Kn-3 Kn-4 Kn-7 Kn-8 Kn-9 Kn-10 Sk-3 Sk-5 AR-1 AR-2 GC-1 GC-6		Prof. Fomenko I.S.
L-3	Metabolism of lipids, its regulation and changes in pathology	of li the lipid syn con oxid path	acquaint students with the process ipid digestion in the digestive tract, functions of simple and complex ds, metabolic transformations: thesis-decomposition of simple and nplex lipids, the processes of dation and synthesis of fatty acids, nological processes - obesity, ttosis, atherosclerosis	Kn-1 Kn-2 Kn-3 Kn-4 Kn-7 Kn-8 Kn-9 Kn-10 Sk-3 Sk-5 AR-1 AR-2 GC-1 GC-6		Prof. Kobylinska L.I.
L-4	Metabolism of amino acids. General pathways of amino acid turnover. Metabolism of ammonia: urea	spec met dest proc	acquaint students with general and cific ways of amino acid cabolism; by the formation and truction of ammonia, pathological cesses of amino acid metabolism the ornithine cycle of urea	Kn-1 Kn-2 Kn-3 Kn-4 Kn-7 Kn-8 Kn-9		Prof. Fomenko I.S.

	synthesis and its disorders. Hereditary enzymopathias of distinct amino acids.		Kn-10 Sk-3 Sk-5 AR-1 AR-2 GC-1 GC-6	
L-5	Basic molecular biology and genetics	To acquaint students with the synthesis and breakdown of purine and pyrimidine nucleotides, regulation of these processes, and pathological changes that may occur when these exchanges are disturbed. To acquaint students with the processes of replication, transcription and broadcasting.	Kn-1 Kn-2 Kn-3 Kn-4 Kn-7 Kn-7 Kn-7 Kn-7 Kn-7 Kn-9 Kn-10 Sk-3 Sk-5 AR-1 AR-2 GC-1 GC-6	Prof. Fomenko I.S.
L-6	Biochemistry of hormones: molecular mechanisms of hormone action; pathology of endocrine action.	Introduce students to the general characteristics of hormones, types of receptors, the mechanism of action of hormones of protein-peptide nature and steroid hormones, to characterize the features of each hormone and pathological processes that occur in the absence / excess of a hormone	Kn-1 Kn-2 Kn-3 Kn-4 Kn-7 Kn-10 Sk-2 Sk-10 GC-1 GC-6	Prof. Kobylinska L.I.
L-7	Biochemistry of blood. Coagulation and fibrinolytic systems. Pathobiochemistr y of blood.	To acquaint students with the process of vascular-platelet and coagulation hemostasis, the mechanism of fibrinolysis, to characterize the anticoagulant system, features and structure of the immune system. Explain the mechanism of development of blood coagulation disorders and immunodeficiency states	Kn-1 Kn-2 Kn-3 Kn-4 Kn-7 Kn-8 Kn-9 Kn-10 Sk-3 Sk-5 AR-1 AR-2 AR-5 GC-1 GC-6	Prof. Fomenko I.S.
L-8	Biochemical functions of liver. Biochemistry of jaundices; biotransformation of foreign substances in	To acquaint students with the classification of xenobiotics and features of their metabolism in the human body. Describe the stages of disposal of foreign substances. Biochemical reactions conjugation oftoxic substances	Kn-1 Kn-4 Kn-6 Kn-7 Kn-8 Kn-9 Kn-10 Sk-4	Prof. Kobylinska L.I.

	liver.		Sk-6	
			AR-1	
			AR-2	
			AR-5	
			GC-1	
			GC-6	
P-1	Objectives and	Piological chemistry as a science. The	Kn-1	According
P-1	Objectives and	Biological chemistry as a science. The		According
	assignments of	place of biochemistry among other	Kn-2	to the time-
	biochemistry.	medical and biological disciplines.	Kn-7	table
	Aims and	Objects of study and tasks of	AR-1	
	methods of	biochemistry. The leading role of	AR-2	
	biochemical	biochemistry in establishing the	AR-4	
	investigations.	molecular mechanisms of pathogenesis	AR-5	
		of human diseases. Connection of	AR-6	
		biochemistry with other biomedical	GC-1	
		sciences. Medical biochemistry.	GC-2	
		Clinical Biochemia. Biochemical	GC-3	
		laboratory diagnostics.	GC-11	
		Structural and functional components		
		of cells, their biochemical functions.		
		Classes of biomolecules. Their		
		hierarchy and origin. Basic and		
DO		methods of biochemical research.	77 1	A 1'
P-2	Physico-chemical	Physicochemical properties of	Kn-1	According
	properties and	enzymes: surface charge of the	Kn-8	to the
	structure of	molecule, solubility, thermodynamic	Sk-1	time-table
	enzyme proteins.	stability of protein-enzyme molecules,	AR-1	
	Mechanism of	precipitation, denaturation, interaction	AR-2	
	action and	with ligands and its functional	AR-4	
	kinetics of	significance.	AR-5	
	enzymatic	Simple and complex proteins-enzymes,	AR-6	
	catalysis.	prosthetic groups of complex proteins-	GC-1	
		enzymes (cofactors, coenzymes).	<i>GC-2</i>	
		Structure of enzymes: active,	GC-3	
		regulatory (allosteric) centres.	GC-6	
		Levels of structural organization of	GC-7	
		enzymes. Multienzyme complexes,	GC-11	
		enzymetic ensembles, multifunctional	PC-2	
			10-2	
		enzymes, their advantages. Nomenclature and classification of		
		enzymes. Types of reactions catalyzed		
		by individual classes of enzymes.		
		Mechanism of action and kinetics of		
		enzymatic reactions: dependence of the		
		reaction rate on temperature, pH of the		
		medium, substrate concentration.		
		Specificity of enzymes.		
		Intracellular localization of enzymes,		
		tissue (organ) specificity of enzymes.		
		Saliva enzymes.		
		Isoenzymes, multiple molecular forms		
		of enzymes. Principles and methods of		
		detecting enzymes in biological		
		objects. Units of activity and amount		

D 2	Study of the	of enzymes.	V. 1	A appendin -
P-3	Study of the	Activators and inhibitors of enzymes:	Kn-1	According
	regulation of	examples and mechanisms of their	Kn-8	to the time-
enzymatic processes. Medical	enzymatic	action.	Sk-1	table
	processes.	Types of enzyme inhibition: reversible	AR-1	
	Medical	(competitive, non-competitive) and	AR-2	
	enzymology.	irreversible.	AR-4	
		Regulation of enzymatic processes.	AR-5	
		Ways and mechanisms of regulation:	AR-6	
		allosteric enzymes, covalent	GC-1	
		modification of enzymes, proteolytic	GC-2	
		activation of enzymes (limited	GC-2 GC-3	
		5 、	GC-6	
		proteolysis).		
		Cyclic nucleotides (cAMP, cGMP) as	<i>GC-7</i>	
		regulators of enzymatic reactions and	GC-11	
		biological functions of the cell.	<i>PC-2</i>	
		Enzymopathies are congenital		
		(hereditary) defects in the metabolism		
		of carbohydrates, amino acids,		
		porphyrins, and purines.		
		Enzyme diagnosis of pathological		
		processes and diseases.		
		Enzymotherapy is the use of enzymes,		
		their activators and inhibitors in		
		medicine.		
P-4	Study of the role	Classification of coenzymes according	Kn-1	According
-4	of cofactors and	to their chemical nature and the type of	Kn-1 Kn-8	to the
		• •		
	coenzyme	reaction they catalyze.	<i>Sk-1</i>	time-table
	vitamins in the	Vitamins as irreplaceable biologically	AR-1	
	manifestation of	active components necessary for the	AR-2	
	catalytic activity	human body. The history of the	AR-4	
	of enzymes. The	discovery of vitamins. Development of	AR-5	
	role of water- and	vitaminology in Ukraine.	AR-6	
	fat-soluble	Causes of exo- and endogenous hypo-	GC-1	
	vitamins in the	and vitamin deficiency.	GC-2	
	metabolism of	Vitamins B1 and B2, their structure,	GC-3	
	living organisms.	biological role, sources for humans,	GC-6	
	n ving organisms.	daily requirement. Signs of	GC-7	
		hypovitaminosis.	GC-11	
		• 1	PC-2	
		Structure, properties of vitamin H and	rt-2	
		pantothenic acid. Their participation in		
		metabolism, main sources, daily need.		
		The role of CoA in exchange		
		processes.		
		Antianemic vitamins (B12, folic acid),		
		their structure, participation in		
		metabolism, human sources, daily		
		need, signs of hypovitaminosis.		
		Vitamins B6 and PP, their structure,		
		biological role, human sources, daily		
		requirement, signs of hypovitaminosis.		
		Vitamins C and P, their structure,		
		biological role, sources for humans,	1	1
		daily requirement. Functional		

P-5	Metabolic pathways and bioenergetics. Tricarboxylic acid cycle and its regulation and energetic balance	relationship between vitamin P and vitamin C. Manifestations of deficiency in the human body. Use of water-soluble vitamins in dental practice. Vitamins of group D, structure, biological role, daily need, sources for humans, signs of hypo- and hypervitaminosis, vitamin deficiency. Vitamin A, structure, biological role, daily need, sources for humans, signs of hypo-hypervitaminosis. Vitamins E, F, structure, biological role, sources for humans, mechanism of action, daily requirement, signs of deficiency, application in medicine. Antihemorrhagic vitamins (K2, K3) and their water-soluble forms, structure, biological role, sources for humans, mechanism of action, daily requirement, signs of deficiency, application in medicine. Use of fat-soluble vitamins in dental practice. Provitamins, antivitamins. Mechanism of action and application in practical medicine. Vitamin-like substances, their structure and role. The concept of metabolism and energy. Characteristics catabolic, anabolic and amfibolic metabolic pathways and their significance. Exergonic and endergonic biochemical reactions; the role of ATP and other macroergic phosphates in their conjugation. Intracellular location metabolic pathways in the cell. The methods of studying metabolism. Catabolic metabolic pathways of biomolecules: proteins, carbohydrates, lipids, their characteristics.	Kn-2 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3 GC-3 GC-6	According to the time-table
		localization of enzymes TCA cycle, TCA cycle reactions; characterization of enzymes and nucleotides TCA cycle, substrate phosphorylation reaction of TCA cycle, the impact of allosteric modulators on the regulation of TCA cycle, energy balance Citric acid cycle). Anaplerotic and amphibolic reactions of TCA cycle.		
P-6	Molecular basis of bioenergetics.	Biological oxidation reactions; types of reactions (dehydrogenase, oxidase, oxygenase) and their biological	Kn-1 Kn-2 Kn-6	According to the

·			r	,
		significance. Tissue respiration.	<i>Kn-7</i>	time-table
		Pyridine-dependent dehydrogenases.	Kn-8	
		The structure of NAD+ and NADPH+.	Kn-10	
		Their importance in oxidation and	Sk-1	
		reduction reactions.	Sk-10	
		Flavin-dependent dehydrogenases.	GC-1	
		Building of FAD and FMN. Their role	GC-2	
		in oxidation and reduction reactions.	GC-3	
		Cytochromes and their role in tissue	GC-6	
		respiration. The structure of their		
		prosthetic group.		
		The sequence of the components of the		
		mitochondrial respiratory chain.		
		Molecular complexes of the inner		
		membranes of mitochondria.		
		Oxidative phosphorylation: points of		
		conjugation of electron transport and		
		phosphorylation, coefficient of		
		oxidative phosphorylation.		
		Chemiosmotic theory of oxidative		
		phosphorylation, mitochondrial ATP		
		synthetase.		
		Inhibitors of electron transport in the		
		respiratory chain of mitochondria.		
		1 1		
		Uncouplers of electron transport and oxidative phosphorylation in the		
		1 1 2		
D 7	Ctor Jon of	mitochondrial respiratory chain.	V 1	A
P-7	Study of	Digestion of carbohydrates in the	Kn-1 Km 2	According
P-7	anaerobic	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of	Kn-2	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their	Kn-2 Kn-6	-
P-7	anaerobic	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of	Kn-2 Kn-6 Kn-7	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the	Kn-2 Kn-6 Kn-7 Kn-8	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine.	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose).	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body.	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose.	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolysis.	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolysis. Glycolytic oxido-reduction: substrates	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolysis. Glycolytic oxido-reduction: substrates of phosphorylation, and shuttle	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolysis. Glycolytic oxido-reduction: substrates	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolysis. Glycolytic oxido-reduction: substrates of phosphorylation, and shuttle mechanisms glycolytic oxidation of NADH. Alcohol fermentation,	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolysis. Glycolytic oxido-reduction: substrates of phosphorylation, and shuttle mechanisms glycolytic oxidation of	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolysis. Glycolytic oxido-reduction: substrates of phosphorylation, and shuttle mechanisms glycolytic oxidation of NADH. Alcohol fermentation,	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolytic oxido-reduction: substrates of phosphorylation, and shuttle mechanisms glycolytic oxidation of NADH. Alcohol fermentation, enzymatic reactions. Reactions are	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
P-7	anaerobic oxidation of	Digestion of carbohydrates in the digestive tract: glycolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of carbohydrate hydrolysis products in the small intestine. Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane transport of hexoses, absorption of glucose and galactose). Glucose as an important metabolite of carbohydrate metabolism: a general scheme of sources and ways of conversion of glucose in the body. Anaerobic oxidation of glucose. Reaction sequence and enzymes of glycolysis. Glycolytic oxido-reduction: substrates of phosphorylation, and shuttle mechanisms glycolytic oxidation of NADH. Alcohol fermentation, enzymatic reactions. Reactions are common and different for glycolysis	Kn-2 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-10 GC-1 GC-2 GC-3	to the
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	alternative pathways of monosaccharide metabolism.	Enzymes, coenzymes and the sequence of reactions in the multienzyme pyruvate dehydrogenase complex. Comparative characteristics of bioenergetics of aerobic and anaerobic oxidation of glucose. The Pasteur effect. Pentose phosphate pathway of glucose oxidation: process scheme and biological significance. Metabolic ways of conversion of fructose and galactose; hereditary enzymopathies of their metabolism.	Kn-7 Kn-8 Kn-9 Kn-10 Sk-1 Sk-3 Sk-7 Sk-10 AR-1 AR-2 AR-4 AR-2 AR-4 AR-5 AR-6 GC-1 GC-2 GC-6 GC-7 GC-11	time-table
P-9	Study of glycogen catabolism and biosynthesis. Biosynthesis of glucose - gluconeogenesis.	Glycogen biosynthesis: enzymatic reactions, physiological significance. Regulation of glycogen synthetase activity. The phosphorolytic pathway of glycogen breakdown in the liver and muscles. Regulation of glycogen phosphorylase activity. The mechanism of reciprocal regulation of glycogenolysis and glycogenesis due to cascade cAMP-dependent phosphorylation of enzyme proteins. The role of epinephrine, glucagon, and insulin in the hormonal regulation of glycogen metabolism in muscle and liver. Genetic disorders of glycogen metabolism (glycogenosis and aglycogenosis). Gluconeogenesis: substrates, enzymes, reactions and physiological significance of the process. Relationship between glycolysis and gluconeogenesis (Cori cycle). Glucose-lactate, glucose- alanine cycles.	PC-2 Kn-1 Kn-2 Kn-7 Kn-8 Kn-9 Kn-10 Sk-1 Sk-3 Sk-7 Sk-7 Sk-10 AR-1 AR-2	According to the time-table
P-10	Studies on mechanisms of metabolic and humoral regulation of carbohydrate metabolism. Diabetes mellitus.	Biochemical processes that ensure a stable level of glucose in the blood. The role of different pathways of carbohydrate metabolism in the regulation of blood glucose levels. Hormonal regulation of carbohydrate metabolism (insulin - structure, mechanism of action, role in carbohydrate metabolism; adrenaline and glucagon - mechanisms of their regulatory action on carbohydrate metabolism).	Kn-1 Kn-2 Kn-7 Kn-8 Kn-9 Kn-10 Sk-1 Sk-2 Sk-3 Sk-5 Sk-7 Sk-8	According to the time-table

		Characteristics of normo-, hyper-,	Sk-9	
		hypoglycemia and glucosuria.	Sk-10	
		Insulin-dependent and non-insulin-	GC-1	
		dependent forms of diabetes.	GC-2	
		Characteristics of biochemical	GC-6	
		disorders in diabetes.	<i>GC-7</i>	
		Biochemical tests for the assessment of	GC-11	
		diabetes mellitus.	<i>PC-1</i>	
		Violation of carbohydrate metabolism	<i>PC-2</i>	
		during fasting.		
P-11	Catabolism and	Digestion of lipids in the digestive	Kn-1	According
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	Intracellular	Disorders of lipid digestion in the	Kn-10	
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	0	thermoregulation, biosynthetic).	AR-1	
		Participation of lipids in the	AR-2	
		construction and operation of cell	AR-4	
		membranes. Liquid-mosaic model of	AR-5	
		biomembranes. Liposomes, their use in	AR-6	
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		Circulatory transport and deposition of	GC-2	
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		Lipoproteinlipase of endothelial tissue.	GC-7	
		Catabolism of triacylglycerols in	GC-11	
		adypocytes, the sequence of reactions,	PC-2	
		the mechanisms regulating the activity	102	
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		triacylglycerols. Neurohumoral		
		regulation of lipolysis: role of		
		epinephrine, norepinephrine,		
		glucagone, insulin.		
		Metabolism of sphingolipids. Genetic		
		anomalies of sphingolipid metabolism		
		- sphingolipidoses. Lysosomal		
		diseases.		
P-12	Beta-oxidation	Reactions of beta-oxidation of fatty	Kn-1	According
	and biosynthesis	acids: localization of the process;	Kn-2	to the
	of fatty acids.	activation of fatty acids; the role of	Kn-7	time-table
	Metabolism of	carnitine in the transport of fatty acids	Kn-8	time-table
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		enzymatic reactions and energy cost of	Kn-10	
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		Glycerol oxidation: enzymatic	Sk-2	
		reactions, bioenergetics.	Sk-2 Sk-3	
		Biosynthesis of higher fatty acids:	Sk-5	
		localization of the process; metabolic	Sk-7	
		sources of fatty acid synthesis; stages	<i>Sk-8</i>	
		of synthesis of saturated fatty acids;	Sk-9	
		or synthesis of subtracted fully delus,	510 2	1

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		the value of acyl-transporting protein,	GC-1	
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		bodies: enzymatic reactions of ketone		
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		value; metabolism of ketone bodies in		
		conditions of pathology; mechanisms		
		of excessive increase in the content of		
		ketone bodies in diabetes and		
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P-13	Biosynthesis and	Cholesterol biosynthesis in the human	Kn-1	According
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		development, the role of genetic	<i>Sk-8</i>	
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P-14	General ways of	Digestion of proteins in the digestive	Kn-1	According
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		proteolytic enzymes of the pancreas and small intestine, their mechanism of action. Absorption of protein hydrolysis products in the small intestine. Decay of proteins in the large intestine. Ways of formation and maintenance of the pool of free amino acids in the human body. General ways of conversion of free amino acids. Types of amino acid deamination reactions and their final products. Mechanism of oxidative deamination of amino acids. L- and D-amino acid oxidases. Their enzymatic activity, specificity of action. Glutamate dehydrogenase: structure of the enzyme, mechanism of the glutamate dehydrogenase reaction, biological significance. Transamination of amino acids, substrates for transamination reactions. The mechanism of the transamination reaction. Transaminases. Localization of transaminases in organs and tissues. The clinical and diagnostic value of determining the activity of transaminases. Decarboxylation of amino acids. Decarboxylases. Formation of biogenic amines (□- aminobutyric acid, histamine, serotonin, dopamine). Decarboxylation of amino acids in the process of protein	AR-5 AR-6 GC-1 GC-2 GC-6 GC-7 GC-11 PC-2	
P-15	Detoxification of ammonia and urea biosynthesis. Biosynthesis of glutathione and creatine.	decay in the intestine. Oxidation of biogenic amines. Ways of ammonia formation. Ammonia toxicity and mechanisms of its neutralization. Circulatory transport of ammonia (glutamine, alanine). Urea biosynthesis: localization of the ornithine cycle; enzymatic reactions; sources of ammonia; energy supply. Genetic defects of enzymes (enzymopathy) of urea synthesis. Glutathione: structure, biosynthesis, biological functions, role in the exchange of organic peroxides. Biosynthesis and biological role of creatine and creatine phosphate, formation of creatinine, clinical and biochemical significance of disorders of their metabolism.	Kn-1 Kn-2 Kn-7 Kn-8 Kn-9 Kn-10 Sk-1 Sk-2 Sk-3 Sk-5 Sk-7 Sk-8 Sk-9 Sk-10 GC-1 GC-2 GC-6 GC-7 GC-11	According to the time-table

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P-16	Specialized	General pathways of metabolism of	Kn-1	According
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		Exchange of sulfur-containing amino	GC-1	
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		Arginine exchange; biological role of	GC-6	
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		Tryptophan metabolism: kynurenine	GC-11	
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		tyrosine, sequence of enzymatic		
		reactions. Hereditary enzymopathies.		
P-17	Biosynthesis and	Purine nucleotide biosynthesis: scheme	Kn-1	According
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	determination of	(retroinhibition).	Kn-10	
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	metabolism.	dTMP biosynthesis as antitumor	<i>Sk-7</i> <i>Sk-8</i>	
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		Catabolism of purine nucleotides.	Sk-10	
		Hereditary disorders of uric acid	GC-1	
		metabolism. Clinical and biochemical	GC-2	
		characteristics of hyperuricemia, gout,	GC-6	
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		Scheme catabolism of pyrimidine	GC-11	
		nucleotides.	<i>PC-1</i>	
			<i>PC-2</i>	
P-18	Replication of	DNA replication: biological	Kn-1	According
	DNA and	significance, semi-conservative	Kn-2	to the
	transcription of	mechanism of replication.	Kn-7	time-table
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	the mechanisms of mutations,	DNA replication in prokaryotes and eukaryotes. RNA transcription: RNA	Kn-9 Kn-10	

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P-19	Investigation of molecular and cellular mechanisms of action of protein and peptide hormones on target cells. Mechanism of hormonal action of amino acid derivatives and biogenic amines.	Hormones: general characteristics; the role of hormones and other bioregulators in the system of intercellular integration of human body functions. Classification of hormones and biomodulator; compliance with the structure and mechanisms of action of hormones. The reaction of target cells to the action of hormones. Membrane (ionotropic, metabotropic) and cytosolic receptors. Biochemical systems of intracellular transmission of hormonal signals: G- proteins, secondary mediators (cAMP, Ca <sup>2+</sup> / calmodulin, IF <sub>3</sub> , DAG, protein kinase C, A), their role. Hypothalamic hormones - liberins and statins. Functional connection between the hypothalamus and the pituitary gland. Anterior pituitary hormones: somatotropin (STG), prolactin. Pathological processes associated with dysfunction of these hormones. Hormones of the posterior pituitary gland. Vasopressin and oxytocin: structure, biological functions. Pancreatic hormones. Insulin - structure, biosynthesis and secretion; effects on the metabolism of	Kn-1 Kn-2 Kn-3 Kn-4 Kn-5 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-2 Sk-3 Sk-4 Sk-5 Sk-7 Sk-8 Sk-7 Sk-8 Sk-10 AR-1 AR-2 AR-4 AR-2 AR-4 AR-5 GC-1 GC-2 GC-3 GC-6 GC-11 PC-1 PC-2 PC-2 PC-3	According to the time-table

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		carbohydrates, lipids, amino acids and proteins. Growth-stimulating effects of		
		insulin. Glucagon. Chemical nature		
		and biological action of the hormone.		
		Catecholamines (epinephrine,		
		norepinephrine, dopamine): structure,		
		biosynthesis, biological effects,		
		biochemical mechanisms of action.		
P-20	Molecular and	Steroid hormones nomenclature	Kn-1	According
	cellular	classification. Mechanism of action of	Kn-2	to the
	mechanisms of	steroid hormones.	Kn-3	time-table
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	hormones upon	Biochemical effects of corticosteroids.	Kn-6	
	target cells.	Glucocorticoids; the role of cortisol in	Kn-7	
	Eicosanoids.	the regulation of gluconeogenesis; anti-	Kn-8	
		inflammatory properties of	Kn-10	
		glucocorticoids. Itsenko-Cushing's	Sk-1	
		disease.	Sk-2	
		Mineralocorticoids; the role of aldosterone in the regulation of water-	Sk-3 Sk-4	
		salt metabolism; aldosteronism. Steroid	Sk-4 Sk-5	
		hormones of the gonads. Female sex	Sk-7	
		hormones: estrogens - estradiol,	Sk-7 Sk-8	
		estrone (C $_{18}$ -steroids), progesterone	Sk-10	
		$(C_{21}$ -steroids); biochemical	AR-1	
		effects; connection with the phases of	AR-2	
		the menstrual cycle; regulation of	AR-4	
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		hormones (androgens) - testosterone,	GC-1	
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		Thyroid hormones. Structure and	C-11	
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		Biological effects of $T_4$ and $T_3$ .	<i>PC-2</i>	
		Pathology of the thyroid gland;	<i>PC-3</i>	
		features of metabolic disorders in conditions of hyper- and		
		conditions of hyper- and hypothyroidism. Mechanisms of		
		endemic goiter and its prevention.		
		Eicosanoids: structure, biological and		
		pharmacological properties. Aspirin		
		and other nonsteroidal anti-		
		inflammatory drugs as inhibitors of		
		prostaglandin synthesis.		
P-21	Intermediate	Porphyrins: nomenclature; structure of	Kn-1	
	products of heme	hemoglobin and myoglobin. Reactions	Kn-2	
	biosynthesis and	of heme biosynthesis. Regulation of	Kn-3	
	their	the process.	Kn-4	
	accumulation in	Hereditary disorders of porphyrin	Kn-7	
	porphyrias. The	metabolism (porphyria). Hemoglobin:	Kn-8	
	structure and	properties, types of hemoglobin.	Kn-9	
	properties of	Hemoglobin derivatives. Mechanisms	Sk-1	

	hemoglobin and its derivatives. Pathological forms of hemoglobin - hemoglobinopath y, thalassemia.	of hemoglobin participation in the transport of oxygen and carbon dioxide. The participation of hemoglobin in the regulation of the acid-base state of the blood. Pathological forms of human hemoglobins. Hemoglobinoses: hemoglobinopathy and thalassemia.	Sk-2 Sk-5 Sk-6 Sk-7 Sk-9 Sk-10 AR-1 AR-2 AR-4 AR-5 GC-1 GC-2 GC-3 GC-3 GC-6 C-11 PC-1 PC-1 PC-2 PC-3	
P-22	Proteins and non- protein nitrogen- containing blood components. Blood buffer systems.	General characteristics of the protein composition of blood. Factors influencing the content of proteins in blood plasma: hyper-, hypo- and dysproteinemia, paraproteinemia. Albumins and globulins. Electrophoresis of blood plasma proteins. Proteins of the acute phase of inflammation and their clinical and biochemical characteristics. Enzymes of blood plasma: value in enzymodiagnosis of diseases of organs and tissues. Kallikrein-kinin system of blood and tissues. Medicines are antagonists of kinin formation. Non-protein organic compounds of blood plasma. Inorganic components of blood plasma. Blood buffer systems. Violation of the acid-base balance in the body (metabolic and respiratory acidosis, alkalosis).	PC-3 Kn-1 Kn-2 Kn-3 Kn-4 Kn-5 Kn-6 Kn-7 Kn-8 Kn-10 Sk-1 Sk-2 Sk-3 Sk-4 Sk-2 Sk-3 Sk-4 Sk-5 Sk-7 Sk-8 Sk-7 Sk-8 Sk-10 AR-1 AR-2 AR-4 AR-2 AR-4 AR-5 GC-1 GC-2 GC-3 GC-6 GC-11 PC-1 PC-2 PC-3 PC-13	According to the time-table
P-23	Blood coagulation, anticoagulant and fibrinolytic systems of blood. Biochemistry of	Functionalandbiochemicalcharacteristicsofthehemostasissystem in the human body;vascular-plateletandcoagulationhemostasis.Bloodcoagulationsystem;characteristicsofcoagulationfactors.	Kn-2 Kn-3 Kn-4 Kn-7 Kn-7 Kn-8	According to the time-table

	immune	Cascade mechanism of activation and	Kn-9	
	reactions.	function of blood coagulation; intrinsic	Sk-1	
	Immunodeficienc	and extrinsic blood coagulation	Sk-2	
	y.	pathways. Role of vitamin K in	Sk-5	
	<i>J</i> .	reactions of hemocoagulation	Sk-6	
		(carboxylation of glutamic acid	Sk-7	
		residues, its role in Ca binding).	Sk-9	
		Medical preparations as vitamin K	Sk-10	
		agonists and antagonists. Hereditary	AR-1	
		and acquired disorders of vascular-	AR-1 AR-2	
		platelet and coagulation hemostasis.	AR-2 AR-4	
		Anticoagulation system of blood,	AR-4 AR-5	
		functional characteristics of its	GC-1	
		components. Fibrinolytic system of		
		blood: stages and factors of fibrinolysis. Medicinal influencing		
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		fibrinolytic process. Activators and	GC-11 PC-1	
		inhibitors of plasmin. Drugs that affect		
		the processes of fibrinolysis.	<i>PC-2</i>	
		Plasminogen activators and plasmin	<i>PC-3</i>	
		inhibitors.		
		Immunoglobulins: structure, biological		
		functions, mechanisms of regulation of		
		immunoglobulin synthesis.		
		Biochemical characteristics of distinct		
		immunoglobulin classes of human		
		blood. Mediators and hormones of		
		immune system: interleukins,		
		interferons, protein and peptide factors		
		of cell growth and proliferation.		
		Factors of complement system.		
		Classical and alternative pathways of		
		complement activation Biochemical		
		mechanisms of immunodeficiencies:		
		primary (hereditary) and secondary		
		immunodeficiencies.		
P-24	Investigation of	The role of the liver in the metabolism	Kn-1	According
	end products of	of bile pigments. Hemoglobin	Kn-2	to the
	heme catabolism.	catabolism, bilirubin conversion.	Kn-3	time-table
	Pathobiochemistr	Pathobiochemistry of jaundices;	Kn-4	
	y of jaundices.	hemolytic (prehepatic),	Kn-7	
	-	parenchimatous (hepatic), occlusive	Kn-7	
		(posthepatic). Enzymatic congenital	Kn-8	
		jaundices: • Crigler-Najjar syndrome	Kn-9	
		as a consequence of insufficient	Sk-1	
		activity of UDP-glucuronyl-	<i>Sk-2</i>	
		transferase. • Gilbert disease –	Sk-5	
		pathology caused by combined	Sk-6	
		disorder of synthesis of bilirubin	Sk-7	
		diglucuronide and absorption of	Sk-9	
		bilirubin from blood by liver cells	Sk-10	
		("absorption jaundice"). • Dubin-	AR-1	
		Johnson syndrome – jaundice caused	AR-2	
			AR-4	
	l	by disorder of transport of bilirubin		

			1	
		diglucuronide from liver cells to bile	AR-5	
		("excretory jaundice"). Enzymatic	GC-1	
		jaundices of neonates and methods of	<i>GC-2</i>	
		their prevention.	GC-3	
		Porphyrins: nomenclature, reactions of	GC-6	
		biosynthesis of prothoporphyrin IX;	GC-11	
		heme production. Regulations of	<i>PC-1</i>	
		porphyrins synthesis. Hereditary	<i>PC-2</i>	
		disorders of porphyrin metabolism	<i>PC-3</i>	
		(porphyria).		
P-25	Biotransformation	Homeostatic role of the liver in the	Kn-1	According
1 20	of xenobiotics	metabolism of the whole organism.	Kn-2	to the
	and endogenous	Biochemical functions of hepatocytes.	Kn-3	
	toxins.	Detoxification function of liver;	Kn-3 Kn-4	time-table
	toxills.	biotransformation of xenobiotics and	Kn-4 Kn-7	
		endogenous toxins. Reactions of	Kn-7	
		microsomal oxidation; inducers and	Kn-8	
		inhibitors of microsomal	Kn-9	
		monoxygenases. Biological role of	<i>Sk-1</i>	
		cytochrome P-450. Electron transport	<i>Sk-2</i>	
		chains in the membranes of the	Sk-4	
		endoplasmic reticulum of hepatocytes.	Sk-5	
		Conjugation reactions in hepatocytes:	Sk-6	
		biochemical mechanisms, functional	Sk-7	
		significance.	Sk-9	
			Sk-10	
			AR-1	
			AR-2	
			AR-4	
			AR-5	
			GC-1	
			GC-2	
			GC-3	
			GC-6	
			GC-0 GC-11	
D 26	Watan and	Water calt match aligns in the		Assauling
P-26	Water and	Water-salt metabolism in the	Kn-1 Km-2	According
	mineral	body.Intra-cellular and extracellular	Kn-2	to the
	metabolism.	water. Metabolism of water, sodium,	Kn-3	time-table
	Normal and	potassium. The role of the kidneys in	Kn-4	
	pathological	the regulation of volume, electrolyte	Kn-7	
	constituents of	composition and pH of body fluids.	<i>Kn-7</i>	
	urine.	Biochemical mechanisms of urinary	Kn-8	
		renal function (filtration, reabsorption,	Kn-9	
		secretion and excretion). Biochemical	Sk-1	
		characteristics of renal clearance and	<i>Sk-2</i>	
		renal threshold, their diagnostic	Sk-5	
		value. Renin-angiotensin system of the	Sk-6	
		kidneys. Biochemical mechanisms of	Sk-7	
		renal hypertension. Antihypertensive	Sk-9	
		drugs are angiotensin-converting	Sk-10	
		enzyme inhibitors. Physico-chemical	AR-1	
		properties of urine: quantity, color,	AR-2	
		odor, transparency, reaction (pH), its	AR-2 AR-4	
			AR-4 AR-5	
		dependence on the composition of	ал-э	

food. The role of the kidneys and lungs $GC-1$ in maintaining the acid-base state of $GC-2$ the body. Ammonium genesis. $GC-3$ The biochemical composition of $GC-6$ human urine is normal and under $GC-11$ conditionsofpathological $PC-1$ processes. Clinicalanddiagnostic $PC-2$	
the body. Ammonium genesis.GC-3The biochemical composition of human urine is normal and under conditions of processes. Clinical and diagnosticGC-11PC-2	
The biochemical composition of human urine is normal and under conditions of processes. Clinical and diagnosticGC-6 PC-1 PC-2	
human urine is normal and under <i>GC-11</i> conditions of pathological <i>PC-1</i> processes. Clinical and diagnostic <i>PC-2</i>	
conditionsofpathologicalPC-1processes. ClinicalanddiagnosticPC-2	
processes. Clinical and diagnostic PC-2	
value of urine composition analysis. <i>PC-3</i>	
P-27 Investigation of General characteristics of morphology <i>Kn-2</i> Acco	rding
biochemical and biochemical composition of <i>Kn-3</i> to the	;
components of connective tissue. Biochemical features Kn-4 time-	table
connective tissue. of intercellular substance of connective Kn-7	uore
tissue. Proteins of connective tissue Kn-7	
fibers - collagen. Proteins of Kn-8	
connective tissue fibers – collagens. Kn-9	
Biosynthesis of collagen and formation Sk-1	
of fibrilar structures. Breakdown of Sk-2	
collagen. Structure and properties of Sk-5	
non-collagen proteins (elastin, large Sk-6	
and small proteoglycans). Noncollagen Sk-7	
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(fibronectin, integrins, laminin, Sk-10	
vitronektyn, tenastsyn, <i>AR-1</i>	
thrombospondin). Complex AR-2	
carbohydrates of the main amorphous AR-4	
matrix of connective tissue - $AR-5$	
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of participation of glycosaminoglycan <i>GC-3</i>	
molecules (hyaluronic acid, GC-6	
chondroitin-, dermatan-, keratan <i>GC-11</i>	
sulfates) in the construction of the <i>PC-1</i>	
main substance of connective tissue. PC-2	
Distribution of various PC-3	
glycosaminoglycans in human organs	
and tissues. Biochemical mechanisms	
1 5	
collagenosis, their clinical and biochemical diagnosis	
biochemical diagnosis.	
Organization and chemical structure of hone tissue Riochemical mechanisms	
bone tissue. Biochemical mechanisms	
of bone formation and physiological	
regeneration. Regulation of	
metabolism in bone tissue: systemic	
and local factors, markers of bone	
metabolism. Bone tissue response to	
dental implants.	1.
P-28 The biochemical Regulation of salivation. The Kn-1 Acco	-
composition and mechanism of saliva formation. <i>Kn-2</i> to the	;
functions of Functions of human saliva (digesting, <i>Kn-3</i> time-	table
saliva. protective, anticariesQuantity indixes Kn-4	
of saliva secretion in health and Kn-5	
pathology. Density and viscosity of the <i>Kn-6</i>	
saliva, pH in health and pathology. Kn-7	

r				
		Organic substances of saliva - proteins and enzymes, their role in ensuring the functions of saliva. Changes in the pathology of the oral cavity and the body as a wholeNoprotein nitrogenous components of saliva, carbohydrates and lipids. Hormones of the saliva, their role in regulation of metabolic processes in oral cavity and human organism in generalInorganic constituents of the saliva (trace and macroelements), changes in disorders of the oral cavity. Protective mechanisms of saliva in smoking.	Kn-8 Kn-10 Sk-1 Sk-2 Sk-3 Sk-4 Sk-5 Sk-7 Sk-8 Sk-7 Sk-8 Sk-7 Sk-8 Sk-10 AR-1 AR-2 AR-4 AR-2 AR-4 AR-5 GC-1 GC-2 GC-3 GC-3 GC-6 GC-11 PC-1 PC-2 PC-3	
P-29	Biochemical composition of tooth tissues: organic and mineral components. Amelogenesis.	General characteristics of the chemical composition of tooth tissues (enamel, dentin, cement, pulp). Inorganic compounds of enamel; apatites, their properties and biological role. Organic substances of enamel (specific proteins, peptides, carbohydrates, lipids). Special features of dentine composition, its structure and functional organization. Cementum. Pulp - features of biochemical composition and metabolism. Amelogenesis. The processes of mineralization - demineralization - the basis of mineral metabolism of tooth tissues. Enamel permeability. The role of vitamins A, D, E, K, C in the regulation of mineralization of tooth tissues. Hormonal regulation of tooth tissue mineralization processes. Superficial formations on the teeth under normal conditions (cuticle, pellicle) and pathology (plaque and tartar). Biochemical changes in tooth tissues during caries. The value of fluoride for caries and fluorosis. The composition of gingival fluid and its changes in periodontal pathology.	Kn-2   Kn-3   Kn-4   Kn-7   Kn-7   Kn-7   Sk-7   Sk-6   Sk-7   Sk-9   Sk-10   AR-1   AR-2   AR-4   AR-5   GC-1   GC-2   GC-3   GC-6   GC-11   PC-1   PC-2   PC-3   PC-7   PC-13	According to the time-table
SIW-1	History of biochemistry; development of biochemical	Make a periodic table of stages of development of biochemistry in Ukraine	AR-1 AR-2 AR-4 AR-5	According to the time- table

	research in		GC-6	
	Ukraine.		GC-7	
SIW-2	Connection of	Give examples of the relation between	AR-1	According
	biochemistry with	biochemistry and other biomedical	AR-2	to the time-
	other biomedical	sciences.	AR-4	table
	sciences. Medical	List the tasks of medical and clinical	AR-5	lucit
	biochemistry.	biochemistry, as well as laboratory	GC-6	
	Clinical	diagnostics	GC-7	
	biochemistry.		007	
	Biochemical			
	laboratory			
	diagnostics.			
SIW-3	The contribution	To compile a periodic table of	AR-1	According
51 W-5	of scientists of the	development of the Department of	AR-1 AR-2	to the time-
		1 1	AR-2 AR-4	to the time-
	Department of	Biochemistry and describe the activities of the heads of the	AR-4 AR-5	table
	Biochemistry, Lviv National			
		department in these periods	GC-6	
	Medical		<i>GC-7</i>	
	University in the			
	development of			
	biological			
CINI 4	chemistry.			A 1'
SIW-4	Principles of	Describe the principles of collecting	AR-1	According
	collection and	material for laboratory tests,	AR-2	to the time-
	storage of	describe their use for diagnostic	AR-4	table
	material for	purposes. List and systematize	AR-5	
	laboratory	mistakes that are allowed for research	GC-6	
	research. Errors in		GC-7	
SIW-5	research.	Malas a dable of an arrive of the dama in	<i>V</i> 7	A
51W-5	Salivary enzymes:	Make a table of enzymes that are in	Kn-7	According
	their specificity	saliva, indicating their functions and	Kn-8	to the time-
	and role.	type of specificity	AR-1	table
			AR-2	
			AR-4	
			AR-5	
			GC-6	
0.000.0			GC-7	
SIW-6	Levels of	Describe the primary, secondary,	<i>Kn-7</i>	According
	structural	tertiary and quaternary structure of	Kn-8	to the time-
	organization of	5 / 51	AR-1	table
	enzymes.	of bonds. Define the concepts of	AR-2	
	Multienzyme	multienzyme complexes, enzymatic	AR-4	
	complexes,	ensembles, multifunctional enzymes,	AR-5	
	enzymatic	give examples	GC-6	
	ensembles,		GC-7	
	multifunctional			
	enzymes, their			
<u>anı: -</u>	advantages.			
SIW-7	Principles and	Describe the main methods of enzyme	Kn-2	According
	methods of	detection. Name the main principles of	Kn-8	to the time-
	detecting	determining the activity of enzymes.	<i>Sk-1</i>	table
	enzymes in	Define the concepts of calor,	AR-1	
	biological objects.	international unit, specific activity,	AR-2	
	Units of activity	molar activity	AR-4	

	and amount of enzymes.		AR-5 GC-6	
CIW 0		Provide a chronological table of the	<i>GC-7</i>	Asserting
SIW-8	Vitamins as irreplaceable biologically active components necessary for the human body. The history of the discovery of vitamins. Development of vitaminology in Ukraine.	Provide a chronological table of the development of vitaminology in the world and in Ukraine	AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW-9	Causes of exo- and endogenous hypo- and vitamin deficiency.	List the factors that lead to the occurrence of hypovitaminosis, hypervitaminosis, avitaminosis	Kn-3 Kn-4 Kn-9 Sk-2 Sk-4 Sk-5 AR-2 AR-5 GC-1 GC-2 GC-3 PC-1	According to the time- table
SIW- 10	Use of water- and fat-soluble vitamins in dental practice.	Fill in the vitamin application table, indicating the name of the vitamin and the pathochemical process in the oral cavity that it affects	Kn-3 Kn-4 Kn-5 Kn-6 Sk-4 Sk-6 AR-1 AR-2 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 11	Provitamins, antivitamins. Mechanism of action and application in practical medicine.	Define the key terms: provitamins and antivitamins. Give examples and describe the mechanism of action.	Kn-1 Kn-5 Kn-8 Kn-9 Sk-1 Sk-4 AR-1 AR-2 AR-5 GC-3 GC-7 PC-1	According to the time- table
SIW- 12	Vitamin-like substances, their	Define the term "vitamin-like substance". Write the structure of	Kn-1 Kn-5	According to the time-

	structure and role.	vitamin-like substances, indicate the mechanism of action and the biological role of each	Kn-8 Kn-9 Sk-1 Sk-4 AR-1 AR-2 AR-5 GC-3 GC-7	table
SIW- 13	Methods of studying metabolism.	Present a table of subcellular structures isolated under the conditions of fractionation of tissue homogenates by	PC-1 Kn-7 Kn-8 Kn-9	According to the time- table
		the method of differential centrifugation	Kn-10 Sk-1 Sk-7 Sk-10 AR-1 AR-2 AR-4 AR-5 AR-6 GC-1	
SIW- 14	Intracellular localization of metabolic pathways, compartmentaliza tion of metabolic processes in the cell.	Make a table of the location of certain metabolic pathways in cell organelles, indicating the enzymes that provide these pathways.	Kn-2 Kn-7 Kn-8 Kn-10 Sk-1 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 15	The structure of NAD+ and NADPH+. Their importance in oxidation and reduction reactions. Building of FAD and FMN. Their role in oxidation and reduction reactions.	Write the structural formulas of NAD+ and NADPH+, FAD, FMN. Specify the oxidized and reduced forms of active structures and explain the mechanism of transfer of reducing equivalents.	Kn-1 Kn-2 Kn-5 Kn-8 Kn-10 Sk-4 Sk-10 AR-1 AR-5 GC-7	According to the time- table
SIW- 16	Hereditary enzymopathies of digestive processes (insufficiency of disaccharidases, violation of membrane	Indicate the relationship between the protein (apoenzyme) and the non- protein part in the composition of dehydrogenases.	Kn-3 Kn-4 Sk-3 Sk-5 Sk-9 AR-1 AR-5 GC-6	According to the time- table

	4			
	transport of		<i>PC-1</i>	
	hexoses,		<i>PC-2</i>	
	absorption of		<i>PC-3</i>	
	glucose and			
CINI	galactose).			
SIW-	The Pasteur	Give examples of metabolic processes	Kn-2	According
17	effect.	in which they participate.	<i>Kn-8</i>	to the time-
			Sk-1	table
			AR-1	
			AR-2	
			AR-4	
			AR-5	
			GC-6	
			<i>GC-7</i>	
SIW-	Glucose-alanine	Introduce a diagram of the glucose-	Kn-1	According
18	cycle.	alanine cycle with a description of the	Kn-2	to the time-
		principle of its operation	Kn-8	table
			Kn-10	
			AR-1	
			AR-2	
			AR-4	
			AR-5	
			GC-6	
			<i>GC-7</i>	
SIW-	Pentose	Present the scheme of the pentose	Kn-1	According
19	phosphate	phosphate pathway of glucose use in	Kn-2	to the time-
	pathway of	the human body (describe the	Kn-3	table
	glucose oxidation:	oxidation stage and the stage of	Kn-8	
	process reaction	isomeric transformations with the	Kn-10	
	scheme	indication of metabolites and	<i>Sk-3</i>	
		enzymes). Specify the biological	Sk-5	
		significance of the pentose phosphate	Sk-10	
		pathway of glucose use in the human	AR-1	
		body. Hereditary deficiency of	AR-2	
		glucose-6-phosphate dehydrogenase.	AR-4	
		Describe the clinical manifestations of	AR-5	
		enzymopathy and describe the	GC-1	
		biochemical causes of their occurrence.	GC-2	
			<i>GC-4</i>	
			<i>GC-5</i>	
			GC-6	
			<i>GC-7</i>	
<u>ann</u>			<i>PC-3</i>	
SIW-	The role of	Make a table of the mechanisms of	Kn-1	According
20	adrenaline,	action of these hormones on specific	Kn-2	to the time-
	glucagon and	enzymes of glycogen metabolism	<i>Kn-8</i>	table
	insulin in the		Kn-10	
	hormonal		AR-1	
	regulation of		AR-2	
	glycogen		AR-4	
	metabolism in		AR-5	
	muscles		GC-6	
			<i>GC-7</i>	
SIW-	Characteristics of	Define concepts, describe the reasons	Kn-2	According

21	normo hunor	for their occurrence	Kn-3	to the time-
21	normo-, hyper-,	for their occurrence		
	hypoglycemia and		Kn-4	table
	glucosuria, causes		<i>Sk-1</i>	
	of their		<i>Sk-2</i>	
	occurrence.		AR-1	
			AR-2	
			AR-4	
			AR-5	
			GC-6	
			GC-7	
SIW-	Biological	Describe the functions of lipids,	Kn-1	According
22	functions of	indicate which compounds are	Kn-2	to the time-
	simple and	provided.	Kn-8	table
	complex lipids in	provided.	Kn-10	tuble
	the human body		AR-10	
	_		AR-1 AR-2	
	(spare, energy,		AR-2 AR-4	
	participation in			
	thermoregulation,		AR-5	
	biosynthetic).		GC-6	
			<i>GC-7</i>	
SIW-	Participation of	Recreate the classification of lipids.	Kn-7	According
23	lipids in the		Kn-8	to the time-
	construction and		<i>Sk-3</i>	table
	functioning of		Sk-7	
	biological		AR-1	
	membranes of		AR-2	
	cells. Liquid-		AR-4	
	mosaic model of		AR-5	
	biomembranes.		AR-6	
	bioinemoranes.		GC-1	
			<i>GC-4</i>	
			GC-5	
			GC-6	
			<i>GC-7</i>	
SIW-	Disorders of lipid	Schematically characterize the	Kn-3	According
24	digestion in the	biological functions of simple and	Kn-4	to the time-
	digestive tract	complex lipids in the human body. To	<i>Sk-2</i>	table
	(steatorrhea, its	characterize the structural organization	<i>Sk-3</i>	
	types).	of biomembranes, to schematically	Sk-5	
	51 /	depict the liquid-mosaic structure of	Sk-9	
		the membrane. Describe the main	AR-1	
		functions of membranes.	AR-5	
			GC-1	
			GC-1 GC-6	
			PC-1	
			<i>PC-2</i>	
			<i>PC-3</i>	
SIW-	Liposomes, their	Describe the structure of liposomes,	Kn-1	According
25	structure and	explain their vector action in the	Kn-2	to the time-
	vectors of use in	process of use	Kn-5	table
	medicine.		Kn-6	
			AR-1	
			AR-2	
			AR-4	
	<u> </u>		AN-4	

			105	
			AR-5	
			<i>GC-6</i>	
~~~~			<i>GC-7</i>	
SIW-	Lysosomal	Make a table of lysosomal diseases,	Kn-3	According
26	diseases: causes,	indicating the name, enzyme and	Kn-4	to the time-
	clinical and	clinical manifestations.	AR-1	table
	biochemical		AR-2	
	characteristics.		AR-4	
			AR-5	
			GC-6	
			<i>GC-7</i>	
SIW-	Pathological	Describe the causes, clinical and	Kn-3	According
27	processes of lipid	biochemical characteristics of obesity.	Kn-4	to the time-
	metabolism that		AR-1	table
	lead to the		AR-2	
	development of		AR-4	
	obesity.		AR-5	
	, i i i i i i i i i i i i i i i i i i i		GC-6	
			GC-7	
SIW-	Disorders of lipid	To define the term "diabetes mellitus",	Kn-3	According
28	metabolism in	to name the causes of lipid metabolism	Kn-4	to the time-
20	diabetes.	disorders in diabetes mellitus and their	AR-2	table
	diabetes.	manifestations.	AR-4	tuoie
		mannestations.	AR-5	
			GC-6	
			GC-0 GC-7	
			PC-1	
			PC-2	
			PC-2 PC-3	
SIW-	Formation	Show the chamistry of the recetions of	К <i>n</i> -6	Assarding
		Show the chemistry of the reactions of		According
29	mechanism and	the formation of hydrochloric acid,	Kn-8	to the time-
	role of	5	Sk-4	table
	hydrochloric acid.	types of acidity, its quantitative	Sk-6	
	Acidity of gastric	indicators. Name the drugs that are	Sk-9	
	juice and forms of	used to stimulate the release of	AR-1	
	its expression.	hydrochloric acid	AR-2	
	Quantitative		AR-4	
	indicators in		AR-5	
	normality and		<i>GC-6</i>	
	pathology by the		<i>GC-7</i>	
	pH-metry		<i>PC-1</i>	
	method.		<i>PC-2</i>	
	Mechanisms of		<i>PC-3</i>	
	stimulation of			
	release of			
	hydrochloric acid.			
SIW-	Decay of proteins	Define the concept of "decay of	Kn-1	According
30	in the large	proteins in the large intestine". Write	Kn-2	to the time-
	intestine.	reactions for the formation of indole,	Kn-3	table
		skatole, cresol, putrescine, cadaverine.	Sk-3	
			AR-1	
			AR-2	
			AR-5	
			GC-1	
			501	1

			GC-2 GC-4 GC-5 GC-6 GC-7 PC-2	
SIW- 31	Transaminases. Localization of transaminases in organs and tissues. Clinical and diagnostic value of determining the activity of transaminases.	Describe transaminases, indicating the name, function, localization in the organs. Describe the clinical and diagnostic value based on the determination of the de Ritis coefficient	Kn-2 Kn-3 Kn-4 Kn-8 Kn-9 Sk-8 AR-1 AR-2 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 32	Genetic defects of enzymes (enzymopathy) of urea synthesis.	In the form of a table to describe the genetic defects of the ornithine cycle, indicating the name of the pathology, enzyme, clinical and biochemical characteristics	Kn-3 Kn-4 Kn-8 Sk-8 AR-1 AR-2 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 33	The role of tetrahydrofolate (H4-folate) in the transfer of single- carbon fragments, dihydrofolate reductase inhibitors as antitumor agents.	Write the chemistry of reactions, describe the role of tetrahydrofolate, explain the mechanism of action of dihydrofolate reductase and the effect of inhibitors on it. Name them.	Kn-8 Kn-9 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 34	Participation of coenzyme forms of vitamin B <sub>12</sub> in amino acid metabolism.	Give the chemistry of reactions, indicate the role of vitamin B 12	Kn-8 Kn-9 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 35	Clinical and biochemical significance of creatine and creatine phosphate metabolism disorders.	Give the chemistry of the formation reactions, indicate the pathochemical changes in violation of the formation of creatine and creatine phosphate	Kn-3 Kn-4 Kn-8 Sk-8 AR-1 AR-2 AR-4 AR-5	According to the time- table

			GC-6 GC-7	
SIW- 36	Orotaciduria: causes, types, clinical and biochemical characteristics.	Define the concept, indicate the causes of orotaciduria type 1 and 2, clinical manifestations and pathochemical changes	Kn-3 Kn-4 Kn-8 Sk-8 AR-1 AR-2 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 37	Inhibitors of transcription and translation in prokaryotes and eukaryotes: antibiotics and interferons - their use in medicine; diphtheria toxin.	In the form of a table to give a description of antibiotics, indicating the name and the specific process that is inhibited. Describe interferons and diphtheria toxin, indicating the origin and mechanism of action	Kn-5 Kn-6 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 38	Gene (point) mutations: role in the occurrence of enzymopathies and hereditary human diseases. Biochemical mechanisms of action of chemical mutagens.	Describe the role of point mutations in the occurrence of enzymopathies. Give examples of such pathologies. Give examples of chemical mutagens	Kn-3 Kn-4 Sk-10 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 39	The reaction of target cells to the action of hormones. Membrane (ionotropic, metabotropic) and cytosolic receptors.	Define the concept of hormone and target cell Describe the features of the structure and localization of membrane and cytosolic receptors	Kn-2 Kn-10 Sk-1 AR-1 AR-2 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 40	Aspirin and other nonsteroidal anti- inflammatory drugs as inhibitors of prostaglandin synthesis.	To give the mechanism of influence of aspirin on formation of proinflammatory postaglandins. Give examples of other NSAIDs	Kn-3 Kn-5 Kn-6 Sk-6 AR-1 AR-2 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table

SIW- 41	Blood plasma enzymes: importance in enzymatic diagnosis of diseases of organs and tissues.	Fill in the table, indicating the pathological processes and enzymes that can be used to diagnose	Kn-9 Kn-10 Sk-2 Sk-8 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 42	Immunoglobulins : structure, biological functions, mechanisms of regulation of immunoglobulin synthesis. Biochemical characteristics of certain classes of human immunoglobulins.	Fill a table indicating the structure of , biological functions, mechanisms of regulation of the synthesis of antibodies , their localization.	Kn-2 Sk-1 Sk-7 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 43	Biochemical mechanisms of urinary renal function (filtration, reabsorption, secretion and excretion).	Describe in detail the stages of urine formation, indicating the location of each	Kn-1 Kn-2 Kn-4 Kn-7 Kn-10 Sk-1 Sk-7 Sk-10 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table
SIW- 44	Physico-chemical properties of urine: quantity, color, odor, transparency, reaction (pH), its dependence on the composition of food.	Describe the quantity, color, odor, clarity, reaction (pH) of urine, its dependence on the food	Kn-4 Sk-2 Sk-10 AR-1 AR-2 AR-4 AR-5 GC-6 GC-7	According to the time- table

SIW-	Bone tissue	Describe the stages of bone changes in	Kn-4	According
45	response to dental	response to dental implants	Sk-1	to the time-
	implants.		Sk-5	table
			Sk-7	
			<i>Sk-8</i>	
			AR-1	
			AR-2	
			AR-4	
			AR-5	
			GC-6	
			<i>GC-7</i>	
SIW-	Protective	Describe the protective mechanisms	Kn-2	According
46	mechanisms of	that occur in the oral cavity during	Kn-3	to the time-
	saliva when	smoking and in chronic smokers	Kn-10	table
	smoking.		AR-1	
			AR-2	
			AR-4	
			AR-5	
			GC-6	
			GC-7	
SIW-	The value of	Describe the mechanism of caries	Kn-2	According
47	fluoride for caries	under conditions of insufficient	Kn-3	to the time-
	and fluorosis.	fluoride intake and the mechanism of	Kn-10	table
		fluorosis under conditions of excess	Sk-5	
		fluoride. Submit the chemistry of the	<i>Sk-6</i>	
		reactions	AR-1	
			AR-2	
			AR-4	
			AR-5	
			<i>C-6</i>	
			<i>C</i> -7	
SIW-	Myofibril	Fill in the table. Giving a comparative	Kn-1	According
48	proteins: myosin,	description of thick and thin filaments	<i>Sk-1</i>	to the time-
	actin,	of muscle tissue.	<i>Sk-7</i>	table
	tropomyosin,		AR-1	
	troponin.		AR-2	
	Molecular		AR-4	
	organization of		AR-5	
	thick and thin		GC-6	
	filaments of		<i>GC-7</i>	
CINY	muscle tissue.		12 1	A 1'
SIW-	Muscle	To characterize the extractive	Kn-1	According
49	extractives,	substances of muscles, nitrogenous and	Kn-2	to the time-
	nitrogenous and	non-nitrogenous, their chemical nature	<i>Sk-1</i>	table
	non-nitrogenous,	and significance. Write the structure of	Sk-10	
	their chemical	anserine and carnosine, indicate their	AR-1	
	nature and role.	role.	AR-2	
			AR-4	
			AR-5	
			Ar-6	
			GC-1	
			GC-2	
			GC-4	
			GC-5	

			GC-6 GC-7	
SIW- 50	Molecular mechanisms of muscle contraction: modern ideas about the interaction of muscle filaments. The role of Ca2+ ions in the regulation of contraction and relaxation of skeletal and smooth muscles.	Schematically depict and describe the molecular mechanisms of skeletal and smooth muscle fiber contraction. Explain the role of Ca2+ ions in the regulation of the contraction-relaxation process.	Kn-1 Kn-7 Kn-8 Kn-10 Sk-1 Sk-7 Sk-7 Sk-10 AR-1 AR-2 AR-4 AR-5 GC-1 GC-2 GC-4 GC-5 GC-6 GC-7	According to the time- table
SIW- 51	Bioenergetics of muscle tissue. Macroergic compounds of muscles. Structure, formation and role of ATP, creatine phosphate, creatine phosphokinase, sources of ATP in muscles; the role of creatine phosphate in providing energy for muscle contraction.	Draw the structural formula of ATP, describe the mechanisms of ATP formation in muscles and its role in the bioenergetics of muscle tissue. Draw the structural formula of creatine phosphate, write the creatine kinase reaction and explain their role in the bioenergetics of muscle tissue. Explain the peculiarities of bioenergetic processes in the myocardium.	GC-7   Kn-1   Kn-7   Kn-8   Kn-9   Kn-10   Sk-1   Sk-7   Sk-8   Sk-10   AR-1   AR-2   AR-4   AR-5   GC-1   GC-2   GC-4   GC-5   GC-6   GC-7	According to the time- table
SIW- 52	Biochemical changes in muscles during pathology.	Explain the biochemical changes in the heart muscle during myocardial infarction. Describe biochemical changes in muscles in myopathies, muscular dystrophies and metabolic disorders in skeletal muscles during aging.	Kn-3 Kn-4 Kn-8 Kn-9 Sk-2 Sk-5 Sk-8 Sk-9 AR-1 AR-2 AR-4 AR-5 GC-1 GC-2 GC-4 GC-5	According to the time- table

			GC-6 GC-7 PC-1 PC-2 PC-3	
SIW- 53	Features of the biochemical composition and metabolism of the brain: chemical composition of the brain, neurospecific proteins and lipids (gangliosides, cerebrosides, cholesterol), features of the amino acid composition of the brain, the role of the glutamic acid system.	Describe the chemical composition of the brain. Describe neurospecific proteins (neuroalbumins, neuroglobulins, neuroscleroproteins, etc.) and lipids (gangliosides, cerebrosides, cholesterol). To describe the amino acid composition of the brain; the role of the glutamic acid system; GABA - shunt.	Kn-1   Kn-7   Kn-8   Kn-9   Kn-10   Sk-1   Sk-7   Sk-8   Sk-10   AR-1   AR-2   AR-4   AR-5   GC-1   GC-2   GC-4   GC-5   GC-6   GC-7	According to the time- table
SIW- 54	Energy exchange in the human brain.	Explain the meaning of aerobic oxidation of glucose; to describe the changes in energy metabolism in the conditions of physiological sleep and anesthesia.	Kn-1 Kn-7 Kn-8 Kn-10 Sk-1 Sk-7 Sk-10 AR-1 AR-2 AR-4 AR-5 GC-1 GC-2 GC-4 GC-5 GC-6 GC-7	According to the time- table
SIW- 55	Biochemistry of neurotransmitters (acetylcholine, norepinephrine, dopamine, serotonin, excitatory and inhibitory amino acids), their role in the transmission of nerve impulses and memory	Give the characteristics and biological role of each neurotransmitter. Name the excitatory and inhibitory amino acids and explain their role in the transmission of nerve impulses. Describe receptors for neurotransmitters and physiologically active compounds. 8.1. Describe the molecular organization and functioning of receptors for the neurotransmitters acetylcholine, serotonin, dopamine, norepinephrine, and amino acids	Kn-1 Kn-7 Kn-8 Kn-10 Sk-1 Sk-7 Sk-10 AR-1 AR-2 AR-4 AR-5 GC-1 GC-2	According to the time- table

	regulation.	(excitatory and inhibitory).	GC-4	
		(	GC-5	
			GC-6	
			<i>GC-7</i>	
SIW-	Receptors for	Describe the molecular organization	Kn-1	According
56	neurotransmitters	and functioning of receptors for the	Kn-7	to the time-
	and	neurotransmitters acetylcholine,	Kn-8	table
	physiologically	serotonin, dopamine, norepinephrine,	Kn-10	
	active	and amino acids (excitatory and	Sk-1	
	compounds.	inhibitory).	Sk-7	
			Sk-10	
			AR-1	
			AR-2	
			AR-4	
			AR-5	
			GC-1	
			<i>GC-2</i>	
			<i>GC-4</i>	
			GC-5	
			GC-6	
			<i>GC-7</i>	
SIW-	Peptidergic	Name the representatives of opoid	Kn-1	According
57	system of the	peptides, indicate the immediate	Kn-7	to the time-
	brain.	predecessors, give the functional	Kn-8	table
		characteristics of enkephalins,	Kn-10	
		endorphins, dynorphins.	Sk-1	
			<i>Sk-7</i>	
			Sk-10	
			AR-1	
			AR-2	
			AR-4	
			AR-5	
			GC-1	
			GC-2	
			<i>GC-4</i>	
			GC-5 GC-6	
			GC-0 GC-7	
SIW-	Disruption of the	List the receptors of opoid peptides and	Kn-2	According
58	exchange of brain	indicate the biochemical mechanisms	Kn-2 Kn-3	to the time-
50	mediators and	of their functioning.	Kn-3 Kn-4	to the time-
	modulators in	or alon runeuolillig.	Sk-2	labic
	mental disorders.		Sk-2 Sk-5	
	montal abordors.		AR-1	
			AR-1 AR-2	
			AR-2 AR-4	
			AR-4 AR-5	
			GC-6	
			GC-7	
			PC-2	
SIW-	Biochemical	Describe the clinical and biochemical	Kn-2	According
58	mechanisms	characteristics of alcoholism, drug	Kn-3	to the time-
	underlying human	addicts, Alzheimer's disease,	Kn-4	table
	neuropsychiatric	multiplesclerosis, Parkinson's disease,	<i>Sk-2</i>	

diseases	epilepsy. Submit in the form of a table,	Sk-5	
(alcoholism, drug	indicating the name of the pathology,	AR-1	
addiction,	pathochemical causes, manifestations	AR-2	
Alzheimer's		AR-4	
disease, multiple		AR-5	
sclerosis,		GC-6	
Parkinson's		GC-7	
disease, epilepsy).		<i>PC-2</i>	

It is necessary to provide the system of the classes organization, the use of interactive methods, educational technologies that are used for the transfer of knowledge and skills.

#### 8. Verification of learning results

### **Current control**

is carried out during training sessions and aims to check the assimilation of students of educational material (it is necessary to describe the forms of current control during training sessions). Forms of assessment of current educational activities should be standardized and include control of theoretical and practical training. The final grade for the current educational activity is set on a 4-point (national) scale.

Code of the learning result	Code of	Code of the learning result	Code of the learning
C	the	C	result
	learning		
	result		
		Types of	A grade of
		educational activities	"excellent" is given
		of students are:	to a student who took
		a) lectures	an active part in
		b) practical classes	discussing the most
		c) individual work of	difficult questions on
		students (SIW)	the topic of the lesson,
		Thematic plans of lectures,	gave at least 19-
		practical classes, SIW ensure	20 correct answers to
		the implementation in the	standardized test tasks,
		educational process of all	answered written tasks
		topics included in the content	without errors,
72 1	7 1	of the program.	completed practical
Kn-1	L-1,		work and drew up a
Kn-2 Kn-3	L-2, L-3,	The lecture course consists of 8 lectures. The topics of	protocol.
Kn-3 Kn-4	L-3, L-4,	the lecture course reveal the	A grade of <b>"good"</b> is given to a student who
Kn-4 Kn-5	L-4, L-5,	problematic issues of the	took part in the
Kn-5 Kn-6	L-J, L-6,	relevant sections of medical	discussion of the most
Kn-7	L-0, L-7,	biology and parasitology.	difficult questions on
Kn-8	L-8,	During the lectures, the	the topic, gave at least
Kn-9	L-9,	students formed the	17-18 correct answers
Kn-10	L-10	theoretical basic knowledge	to standardized test
<i>Sk-1</i>		will ensure there is a	tasks, made some
<i>Sk-2</i>		motivational component of	minor mistakes in
<i>Sk-3</i>		general and tentative stage	answering written
<i>Sk-4</i>		mastering scientific	tasks, did practical
<i>Sk-5</i>		knowledge during	work and drew up a
<i>Sk-6</i>		independent work. In the	protocol.
<i>Sk-7</i>		lecture course maximum	A student who did not
Sk-10		used various teaching tools -	participate in the
A B -1		multimedia presentations,	discussion of the most

		• 1 1 / 1 / 1	1:00:1				
AR-2		video lectures, educational	difficult questions on				
GC-1		films, slides.	the topic, gave at least				
<i>GC-2</i>		Practical classes are aimed	14-16 correct answers				
GC-3		at controlling the	to standardized test				
		assimilation of theoretical	tasks, made significant				
Kn-1		material, the formation of	mistakes in answering				
Kn-2		practical skills and abilities,	written tasks,				
Kn-3		as well as the ability to	performed practical				
Kn-4		analyze and apply the	work and drew up a				
Kn-5	<i>P-1, P-2,</i>	acquired knowledge to solve	protocol received a				
Kn-6	<i>P-3, P-4,</i>	practical problems.	grade of				
Kn-7	<i>P-5, P-6,</i>	Each session begins with a	"satisfactory".				
Kn-8	<i>P-7, P-8,</i>	test control (20 tests) to	A grade of				
Kn-9	<i>P-9</i> ,	assess baseline knowledge	"unsatisfactory" is				
Kn-10	<i>P-10</i> ,	and determination with	given to a student who				
Sk-1	<i>P-11</i> ,	tupenya readiness of students	did not participate in				
Sk-1 Sk-2	<i>P-12</i> ,	to classes. The teacher	the discussion of the				
<i>Sk-2</i> <i>Sk-3</i>	P-13,	determines the purpose of the	most difficult				
<i>Sk-3</i> <i>Sk-4</i>	P-14,	lesson and creates a positive	questions on the topic,				
<i>Sk-4</i> <i>Sk-5</i>	P-15, P-16,	cognitive motivation;	gave less than 14				
Sk-5 Sk-6	P-10, P-17,	answers questions from	0				
	<i>P-17</i> , <i>P-18</i> ,	1					
Sk-7	<i>P-19</i> ,	students that arose during the	standardized test tasks,				
Sk-8	<i>P-20</i> ,	VTS on the topic of the	made gross mistakes				
<i>Sk-9</i>	<i>P-21</i> ,	lesson.	in answering written				
<i>Sk-10</i>	<i>P-22</i> ,	The main stage of the lesson	tasks or did not answer				
AR-1	<i>P-23</i> ,	is to perform practical work.	them at all, did not do				
AR-2	<i>P-24</i> ,	At the final stage of	practical work and				
AR-3	<i>P-25</i> ,	the lesson in order to assess	without drawing up a				
AR-4	<i>P-26</i> ,	the student's mastery of the	protocol.				
AR-5	<i>P-27</i> ,	topic he					
AR-6	<i>P-28</i> ,	is asked to answer three					
GC-1	<i>P-29</i> ,	theoretical questions. The					
GC-2	<i>P-30</i> ,	teacher sums up the lesson					
GC-3	<i>P-31</i> ,	gives students tasks for					
GC-6	<i>P-32</i> ,	independent work, indicating					
<i>GC-7</i>	<i>P-33,</i> <i>P-34,</i>	the main issues the next topic					
GC-9	P-34, P-35	and offers a list of					
GC-10	1-33	recommended literature.					
GC-11		The duration of the practical					
GC-13		lesson is 2 academic hours.					
<i>PC-1</i>							
<i>PC-2</i>							
<i>PC-3</i>							
	F	Final control					
General evaluation system		ion in the work during the seme	ster / exam - 60% / 40%				
,	-	point scale					
Rating scales		l 4-point scale, multi-point (200	-point) scale. ECTS				
	rating scale						
Conditions of admission to the	All types of work provided by the curriculum must						
final control	be completed and all topics submitted for current control must						
	be included . The student has received at least 72 point s on						
	current progress						
Type of final control		Exam	Enrollment criteria				
Examination							
	The						

	standardized and includes control	of points is 80.
	of theoretical and practical training.	The minimum number
	Jan	of points is 50
	Exam evaluation criteria	
Exam	1. written answers to 40 standard test	Criteria for
	tasks, each of which has one correct	evaluating test tasks:
	answer out of five proposed (format	Less than 25 MCQs -
	A). 40 minutes are allocated for writing	"unsatisfactory";
	the test control (at the rate of 1 test for	25 - 30 MCQs -
	1 min);	"satisfactory";
	2. written standardized answers to 5	31 - 36 MCQs -
	problems in the form of chains of	"good";
	transformations of bioorganic	37 - 40 MCQs -
	compounds, the list of certain	"excellent".
	biochemical indicators, the filled	The correct answer to
	tables, drawing up of schemes, writing	1 test is 1 point.
	of equations of chemical reactions (1 -	The minimum number
	4 questions); description of the	of points for 40 tests is
	principles of methods and clinical and	25 points.
	diagnostic value of determining	The maximum number
	biochemical parameters (5	of points for 40 MCQs
	questions). The duration of the exam	is 40 points.
	is 95 minutes.	Criteria for
		evaluating
		theoretical tasks:
		Each of the five
		theoretical tasks is
		evaluated from 6 to 8
		points:
		Less than 5 points -
		"unsatisfactory" - the
		student made gross
		mistakes in answering written tasks or did not
		give answers to them
		at all;
		5 points -
		"satisfactory" - the
		student made
		significant mistakes in
		answering written
		theoretical tasks
		(including practical
		skills);
		7 points - "good" -
		the student made some
		minor mistakes in
		answering written
		theoretical tasks
		(including practical
		skills), or did not fully
		substantiate them;
		8 points - "excellent"
		- the student has

comprehensively and
deeply mastered the
curriculum; has full
theoretical knowledge
and practical skills.
The minimum number
of points for 5
1
theoretical questions
is 25 points.
The maximum number
of points for 5
theoretical questions is
_
40 points.
The maximum
number of
points that a student
can score when
taking the exam is
80.
The minimum
number of points in
the exam - not less
than 50.

*The maximum number of points* that a student can score for the current academic activity for admission to the exam is 120 points.

*The minimum number of points* that a student must score for the current academic activity for admission to the exam is 72 points.

*The calculation of the number of points* is based on the grades obtained by the student on a 4-point (national) scale during the study of the discipline, by calculating the arithmetic mean (CA), rounded to two decimal places. The resulting value is converted into points on a multi-point scale as follows:

disciplines culminating in the exam									
4-point	200		4.95	119		4.79	115	4.62	111
scale	point		4.91	118		4.75	114	4.58	110
	scale		4.87	117		4.7	113	4.54	109
5	120		4.83	116		4.66	112	4.5	108
4.45	107		3.95	95		3.58	86	3.2	77
4.29	103		3.91	94		3.41	82	3.04	73
4.12	99		3.74	90		3.37	81	3.0	72

Recalculation of the average grade for current activity in a multi-point scale for

*Students' individual work* is assessed during the current control of the topic in the relevant lesson. Assimilation of topics that are submitted only for independent work is controlled during the final control.

The grade for the discipline that ends with the exam is defined as the sum of the points

for current educational activity (not less than 72) and points for the exam (not less than 50).

Points from the discipline are independently converted into both the ECTS scale and the 4-point (national) scale. ECTS scale scores are not converted to a 4-point scale and vice versa.

The scores of students studying in one specialty, taking into account the number of points earned in the discipline are ranked on the ECTS scale as follows:

ECTS assessment	Statistical indicator
А	The best 10% of students
В	The next 25% of students
С	The next 30% of students
D	The next 25% of students
E	The last 10% of students

Ranking with assignments of grades "A", "B", "C", "D", "E" is carried out for students of this course who study in one specialty and have successfully completed the study of the discipline.

Discipline scores for students who have successfully completed the program are converted into a traditional 4-point scale according to the absolute criteria, which are given in the table below:

Points in the discipline	Score on a 4-point scale
From 170 to 200 points	5
From 140 to 169 points	4
From 139 points to the minimum number of points that a student must score	3
Below the minimum number of points that a student must score	2

The ECTS score is not converted to the traditional scale, as the ECTS scale and the fourpoint scale are independent.

The objectivity of the assessment of students' learning activities is checked by statistical methods (correlation coefficient between ECTS assessment and assessment on a national scale).

# Criteria for assessing the objective structured practical (clinical) exam / Complex of practice-oriented exam Master's thesis

# 9. Course policy

The policy of the course is determined by the system of requirements for the student in the study of the discipline "Biological Chemistry" and is based on the principles of academic integrity. Students are explained the value of acquiring new knowledge, the need for independent performance of all types of work, tasks provided by the work program of this discipline. Lack of references to used sources, fabrication of sources, writing off, interference in the work of other students are examples of possible academic dishonesty. Detection of signs of academic dishonesty in the student's work is the basis for its non-enrollment by the teacher, regardless of the extent of plagiarism or deception. Literary sources may be provided by the teacher exclusively for educational purposes without the right to transfer to third parties. Students are encouraged to use other literature sources not provided by the recommended list.

10. Literature

Mandatory

Main:

1. Gubsky Yu. Bioorganic and biological chemistry. Book 2. Biological chemistry. Second edition.

Medicine 2021. 500 p.

- 2. Harper's Illustrated Biochemistry 30<sup>th</sup> edition / V. W. Rodwell et al.; NY: McGraw-Hill Education, 2015. 817 p.
- 3. Satyanarayana U., Chakrapani U. Biochemistry. Fifth edition, N.Delhy: Elsevier, copublished with Book and Allied, 2017. 788 p.
- 4. Gubsky Yu. Biological Chemistry. Nova Knyha, Vinnytsia, 2017. 487 p.
- 5. Lippincott Illustrated Reviews: Biochemistry. Denise R. Ferrier. Seventh edition. Wolters Kluwer, 2017. 2224 p.
- 6. MCQs in biochemistry 2<sup>nd</sup> edition / A. Ya. Sklyarov et al.: Lviv: Danylo Halytsky Lviv National Medical University Press, 2020. 319 p.
- 7. MCQs in biochemistry / A. Ya. Sklyarov et al.: Lviv: Danylo Halytsky Lviv National Medical University Press, 2012. 308 p.

# **Optional:**

- 1. Textbook of Biochemistry for Medical Students by Vasudevan D.M., Sreekumari S., Kannan Vaidyanathan. Seventh edition. Jaypee Brothers Medical Pub, 2013. 791 p.
- 2. Chatterjea M.N., Rana Shinde. Textbook of Medical Biochemistry. Eighth edition. Jaypee Brothers Medical Pub, 2012. 894 p.
- 3. Nelson D.L., Cox M.M. Lehninger Principles of Biochemistry. Seventh edition. W.H. Freeman and Company, New EPYOrk, 2017. 1328 p.
- 4. Trudy McKee, James R. McKee. Biochemistry. The molecular basis of life. Sixth edition. Oxford University Press, 2015. 928 p.
- 5. Peter Ronner. Netter's Essential Biochemistry. Elsevier, 2018. 482 p.

# Information resources:

1. Centre of testing - base of licenced test tasks Krock-1 http://testcentr.org.ua/

# 1. Equipment, material and technical supply

# Methodical supply:

- 1. Working educational program of the discipline;
- 2. Multimedia lectures supply;
- **3.** Lectures theses;
- 4. Methodical recommendations for the lecturers;
- **5.** Educational platform Misa;
- **6.** Study guides;
- 7. Methodical guides to the practical classes for students;
- 8. Methodical instruction for the students independent work;
- 9. Test and control tasks to practical classes;
- **10.** Questions and tasks for the summary control (exam); Laboratory equipment (PEC, SP, centrifuges, laboratory utensils, biochemical analyzers)

Information resource - http://misa.meduniv.lviv.ua/

Testing center - database of licensed test tasks Step-1 http://testcentr.org.ua

# 12. Additional information

Responsible for the educational work with foreign students – Prof. Iryna Fomenko Practical classes and lectures are delivered in the Department classrooms at the address: Lviv, 54, Pekarska Street, Chemical building, ground floor.

Web-site of the Department - *e-mail:* <u>Kaf\_biochemistry@meduniv.lviv.ua</u>

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