Federal State Budgetary Educational Institution of Higher Education "Privolzhsky Research Medical University" Ministry of Health of the Russian Federation

APPROVED

Vice-Rector for E.S. Bogomolova 31 August 2021

WORKING PROGRAM

Name of the academic discipline: PHARMACEUTICAL CHEMISTRY

Specialty: 33.05.01 Pharmacy

Qualification: Pharmacist

Department: Pharmaceutical Chemistry and Pharmacognosy

Mode of study: full-time

Labor intensity of the academic discipline: 684 academic hours

Nizhny Novgorod 2021 The working program has been developed in accordance with the Federal State Educational Standard for the specialty 33.05.01 PHARMACY, approved by order of the Ministry of Science and Higher Education of the Russian Federation on March 27, 2018 N 219.

Developers of the working program:

O.A.Vorobeva, Associate Professor of the Department, PhD; D.S. Malygina, Associate Professor of the Department, Ph.D.

The program was reviewed and approved at the department meeting (protocol No. 1 of 08/29/2021)

Head of the Department of Pharmaceutical Chemistry and pharmacognosy, Ph.D. / O.V. Zhukova /

29 August 2021

AGREED Deputy Head of EMA ph.d. of biology

Alt Lovtsova L.V.

(signature)

29 August 2021

1. The purpose and objectives of mastering the academic discipline pharmaceutical chemistry.

The purpose of mastering the discipline: *participation in forming the relevant competencies UC-1,2, GPC-1,3,6, PC-4,7.*

Tasks of the discipline:

As a result of completing the discipline, the student should

Know:

- general methods for assessing the quality of medicinal products, the possibility of using each method depending on the method of obtaining medicinal products, the raw materials, the structure of medicinal substances, and the physical and chemical processes that may occur during storage and circulation of medicinal products;

- factors affecting the quality of medicines at all stages of circulation; determination of the main factors depending on the properties of medicinal substances (redox, ability to hydrolysis, polymerization); the possibility of preventing the influence of external factors on the good quality of medicines;

- chemical methods underlying the qualitative analysis of medicines; the main structural fragments of medicinal substances, according to which the identification of inorganic and organic medicinal substances is carried out; general and specific reactions to individual cations, anions and functional groups;

- chemical methods underlying the quantitative analysis of drugs; equations of chemical reactions taking place during acid-base, redox, precipitation, complexometric titration;

- the principles underlying the physicochemical methods of drug analysis;

- equipment and reagents for chemical analysis of medicines; requirements for reagents for testing for purity, identity and quantitation; equipment and reagents for physical and chemical analysis of medicinal substances; a schematic diagram of a refractometer, photocolorimeter, spectrophotometer, gas-liquid chromatography, high-performance liquid chromatography;

- the structure of regulatory documents regulating the quality of medicines; features of the structure of the pharmacopoeial article and the pharmacopoeial article of the enterprise;

- features of the analysis of individual dosage forms; concepts of disintegration, dissolution, strength; features of the analysis of soft dosage forms;

- physical and chemical constants of medicinal substances; methods for determining the melting point, rotation angle, specific absorption rate, boiling point;

- concept of validation; validation characteristics of qualitative and quantitative analysis methods;

general patterns of pharmacokinetics and pharmacodynamics of drugs; types of drug interactions and types of drug incompatibility;

- belonging of drugs to pharmacological groups, pharmacodynamics and pharmacokinetics of drugs, the most important toxic side effects, main indications and contraindications for use;

- normative documentation regulating the production and quality of medicines in pharmacies and pharmaceutical enterprises; basic requirements for dosage forms and indicators of their quality;

- nomenclature of industrial preparations;

- nomenclature of modern excipients and their properties, purpose.

Be able to:

- identify, prevent (if possible) pharmaceutical incompatibility;

- plan the analysis of medicines in accordance with their form according to regulatory documents and evaluate their quality according to the results obtained;

- prepare reagents, reference, titrated and test solutions, control them;

- to carry out the identification of medicinal substances by reactions to their structural fragments;

- determine the general indicators of the quality of medicinal substances: solubility, melting point, density, acidity and alkalinity, transparency, color, ash, weight loss upon drying;

- interpret the results of UV and IR spectrometry to confirm the identity of medicinal substances;

- use various types of chromatography in the analysis of medicinal substances and interpret its results;

- to establish the quantitative content of medicinal substances in the substance and dosage forms by titrimetric methods;

- to establish the quantitative content of medicinal substances in the substance and dosage forms by physical and chemical methods;

- carry out tests for the purity of medicinal substances and establish limits for the content of impurities by chemical and physico-chemical methods;

- perform analysis and quality control of pharmacy-made medicines in accordance with applicable requirements.

Possess:

- skills in interpreting the results of drug analysis to assess their quality; standard operating procedures for determining the order and execution of documents for the declaration of conformity of the finished product with the requirements of regulatory documents;

- skills in the use of chemical, biological, instrumental methods of analysis for the identification and determination of toxic, narcotic substances and their metabolites;

- methods of carrying out intra-pharmacy quality control of medicines;

- normative, reference and scientific literature for solving professional problems.

2. Position of the academic discipline in the structure of the General Educational Program of Higher Education (GEP HE) of the organization.

The discipline Pharmaceutical Chemistry refers to the core part (or *the part formed by the participants of educational relations*) of Block 1 of GEP HE (Academic discipline index).

The discipline is taught in 5, 6, 7, 8, 9 semesters/

2.2. The following knowledge, skills and abilities formed by previous academic disciplines are required for mastering the discipline: mathematics, physics, general and inorganic chemistry, physical and colloid chemistry, analytical chemistry, organic chemistry, biological chemistry.

2.3. Mastering the discipline is required for forming the following knowledge, skills and abilities for subsequent academic disciplines: *pharmacognosy*, *pharmaceutical* technology, toxicological chemistry.

3. Deliverables of mastering the academic discipline and metrics of competence acquisition

Mastering the discipline aims at acquiring the following universal (UC) or/and general professional (GPC) or/and professional (PC) competencies

		The content of the	Code and name of		of mastering the students show	the discipline,
№	Competen ce code	competence (or its	the competence			
		part)	acquisition metric	know	be able to	possess
1.	UC-1.	Able to realize critical analysis of prob- lem situations based on a systematic approach, develop strategy actions	UC-1.1. Analyzes the problem situa- tion as a system identifying its components and connections be- tween them UC-1.2. Identifies gaps in the information needed to solve a problem situation, and designs processes for their elimination UC-1.3. Critically assesses reliability of information sources, works with conflicting information from different sources UC-1.4. Develops and meaningfully argues the strategy of solving the problem situations based on the system and interdisciplinary approaches UC-1.5. Uses logi- cal and methodo-	 methodology of abstract thinking for systematization of processes and construction of cause-and-effect relationships; modern theoretical and experimental methods for the implementation of own and borrowed results of scientific research into practice. 	 abstract, analyze and synthesize the information received; highlight and to systematize the essential properties and connections of objects, to identify the main patterns of the objects under study; search, select and analyze information obtained from various sources in order to make the best decision at the modern scientific level, in accordance with professional tasks and the requirements of legal documents. 	 methods of self- control, abstract and analytical thinking; skills in analyzing methodological problems that arise in solving research and practical problems, including those in interdisciplinary areas; skills of presenting an independent point of view

			logical tools for critical evaluation of modern concepts of philosophical and social nature in its subject areas			
2.	UC-2.	Able to manage the pro- ject at all stages of its life cycle	UC-2.1. Formulates a project task on the basis of the set problems and a method of its solutions through the implementation of the project management UC-2.5. Moni- tors the progress of the project, corrects devia- tions, makes ad- ditional changes to the project implementation plan, clarifies zones of respon- sibilities of pro- ject participants	principles for developing a project implementation plan in the field of professional activity at all stages of its life cycle	develop a project implementation plan in the field of professional activity at all stages of its life cycle, providing for problem situations and risks	methods of planning and executing projects under conditions of uncertainty, managing the project (supporting the implementation of the project)
3.	GPC-1.	Able to use basic biological, physical- chemical, chemical, mathematical methods for the development, research and examination of medicines, the manufacture of medicinal products	GPC-1.1. Applies basic biological methods of analysis for the development, research and examination of pharmaceuticals and medicinal plant raw materials GPC-1.2. Applies basic physical- chemical analysis methods for the development, re- search and exam- ination of medici- nal products and medicinal plant raw materials GPC-1.3. Applies the basic methods of physical-	 •organization of a system of state control over the production and manufacture of drugs; • the main regulatory documents, production and manufacture, quality control, storage and use of medicines (domestic and international standards (GMP, GLP, GCP, GPP), pharmacopoeias, orders of the Ministry of Health of the Russian Federation, guidelines and instructions approved by the Ministry of Health of the Russian Federation, guidelines and instructions approved by the Ministry of Health of the Russian Federation) for examination using chemical, biological, physicochemical 	• apply chemical, biological, physico- chemical and other methods of analysis during the examination of medicines.	 ensuring the process of quality control of medicines with equipment and consumables; basic chemical, biological, physico-chemical and other methods of analysis during the examination of medicines.

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		chemical analysis in the manufacture of medicinal products GPC-1.4. Applies mathematical	and other methods; • pharmacopoeial methods of analysis used in the analysis of		
		methods and performs mathematical	medicinal products using chemical, biological, physicochemical		
		processing of data obtained during the development of medicines, as well	and other methods.		
		as research and examination of medicines and			
		medicinal plant raw materials			
GPC-3.	Able to carry out professional activities taking into account specific economic, environmental, social factors within the framework of the system of regulations of the medicine circulation sphere	GPC-3.1. Complies with norms and rules established by the authorized state authorities when solving the tasks of professional activity in the field of medicine circulation GPC-3.3. Performs labor actions taking into account their impact on the environment, preventing the occurrence of environmental hazards	 laws and legislative acts of the Russian Federation, normative and methodological materials of the Ministry of Health of Russia, regulating the procedure for conducting examinations provided for in the state registration of medicines; general principles of development, testing and registration of medicines; the basic principles, strategies, methods and procedures for quality control of medicines in the conditions of pharmaceutical organizations used in the course of examinations provided for in the state registration of medicines, in accordance with the requirements of the current regulatory and legislative 	 put into practice the basic principles of the system of quality control and safety of medicines in the conditions of pharmaceutical organizations; to organize and carry out the procedure for quality control of medicines at the level of their production, transportation and storage using methods of pharmacopoeial analysis. 	 skills in organizing and conducting quality control of medicines at the level of their production, transportation and storage; the main methods of pharmaceutical analysis provided for in the state registration of medicines; skills in carrying out preventive measures to ensure the quality of medicines at the level of their production, transportation and storage.
GPC-6.	Able to understand	GPC-6.2. Performs	framework. modern means of computing	use modern computer	methods of practical use
	the principles of modern information	an effective search for information	technology	technology and basic office applications	modern computers to search information
	technologies and	necessary to solve		And	processing and

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	use them to solve the tasks of professional activity	the tasks of professional activity using legal reference systems and professional pharmaceutical databases GPC-6.3. Uses specialized software for mathematical processing of observational and experimental data in solving problems of professional activity		graphic packages; evaluate way of implementing information systems and devices for solving task	fundamentals numerical methods for solving applied tasks
PC-4.	Able to participate in monitoring the quality, effectiveness and safety of medicines and medicinal plant raw materials	PC-4.1.Conductspharmaceuticalanalysisofpharmaceuticalsubstances,excipientsandmedicinesformedicaluseoffactoryproductioninaccordancewithqualitystandardsPC-4.2.Performsintra-pharmacyqualitycontrolqualitycontrolofmedicinesformedicinesformedicinesformedicinesformedicinesformedicinesformedicinesformedicinesformedicinalpharmacognosticanalysisofmedicinalpreparationsPC-4.4.Informsinaccordancewiththenon-compliancecomplianceofthemedicalusewiththeestablishedrequirementsoraboutthenon-complianceofthemedicinalproductformedicalusewiththenon-complianceofthenon-complianceofthenon-complianceofthenon-complianceofthenon-complianceofthenon-	 laws and legislative acts of the Russian Federation, regulatory and methodological materials of the Ministry of Health of Russia, regulating the procedure for quality control of medicines in the conditions of pharmaceutical organizations; methods of analysis used in the quality control of drugs in the conditions of pharmaceutical organizations; monitor drug quality assurance systems; the process of providing equipment and consumables for quality control in the conditions of 	 apply chemical, physico- chemical methods of intra-pharmacy quality of drugs in the conditions of pharmaceutical organizations; draw up documentation of the established form for the control of manufactured medicinal products in the conditions of pharmaceutical organizations; monitor drug quality assurance systems; provide the process of quality control in pharmaceutical organizations with equipment and consumables. 	• basic chemical and physico-chemical methods of intra- pharmacy quality control of drugs in the conditions of pharmaceutical organizations; • registration of documentation of the established sample for the control of manufactured drugs in the conditions of pharmaceutical organizations.

		data on the effec- tiveness and safety of the medicinal product with the data on the medici- nal product con- tained in the in- structions for its use			
PC-7.	Able to carry out operations related to the technological process in the production of medicines and their control	PC-7.5. Monitors the compliance of the raw materials and <u>excipients</u> used with the requirements of regulatory documentation	requirements of regulatory documentation for the raw materials and auxiliary materials used	carry out pharmacopoeial analysis of raw materials and auxiliary materials used	methods of quality control of raw materials and auxiliary materials used

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p /	Compete nce code	Section name of the discipline	The content of the section in teaching units
no.	nce code	of the discipline	
			Pharmaceutical chemistry as a science. The object of
			pharmaceutical chemistry. Methodology of
			pharmaceutical chemistry. The value of
			pharmaceutical chemistry in the preparation of a
			pharmacist. Tasks of pharmaceutical chemistry and
			ways to solve them together with chemical,
			biomedical and other disciplines. The place of
			pharmaceutical chemistry in the complex of
		C-1,2 Fundamentals of	pharmaceutical sciences.
			A brief historical outline of the development of
			pharmaceutical chemistry as a branch of pharmacy.
1.	GPC-	Pharmaceutical	Sources and methods for obtaining medicines:
	1,3,6	Analysis	isolation from natural raw materials; reproduction of
	PC-4,7	<i>j</i> -	physiologically active natural substances; synthesis
			based on metabolites and antimetabolites;
			biosynthesis; use of genetic engineering; fine organic
			synthesis. Computer modeling and prediction of the
			biological activity of new compounds.
			State principles and regulations governing the quality
			of medicines. Regulatory documentation and
			standardization of medicines. State Pharmacopoeia
			(SP), General Pharmacopoeia Articles (GPM),
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			Pharmacopoeia Articles (FS), Pharmacopoeia Article

of the Enterprise (FSP). Legislative nature of
pharmacopoeial articles. General characteristics of
ND (requirements, norms and methods of control).
The role of ND in improving the quality of medicines.
International and regional collections of unified
requirements and test methods for medicines,
European Pharmacopoeia, WHO International
Pharmacopoeia and other regional and national
pharmacopoeias.
Ensuring the quality of medicines. Organization of
quality control of medicines. GMP rules. Quality
control of medicines in production (industrial
enterprises and pharmacies). Quality control of
medicines during storage. The study of the expiration
dates of medicines. pharmacopoeial analysis.
Sampling procedure. Criteria for pharmacopoeial
analysis (selectivity, sensitivity, accuracy, analysis
time, amount of substance).
Subjective and objective criteria used to determine
the authenticity of a medicinal product. OFS "General
reactions to authenticity".
Chemical methods of authentication. Reactions to
cations, anions, functional groups and their use for the
qualitative analysis of drugs.
Establishing the authenticity of medicines by physical
constants (melting point, solidification point, boiling
point). Determination of solubility, degree of
whiteness, density and viscosity of drugs.
Establishing the authenticity of medicines using
instrumental methods (polarimetry, UV and IR
spectroscopy, GLC and HPLC, atomic absorption
spectroscopy, mass spectroscopy).
Test methods for purity. Possible reasons for the
appearance of impurities, their nature and character.
Unification and standardization of tests. Techniques
for determining the content of impurities based on the
degree of sensitivity of chemical reactions (reference
and non-reference methods).
Methods for quantitative and semi-quantitative
assessment of the content of impurities. Development
of requirements for drug purity testing. Quantitative
determination of impurities (chemical, physical,
physico-chemical methods).
Methods for quantitative analysis of drugs.
Prerequisites for choosing a method that allows to

			assess the content of the drug by functional groups that characterize its properties. Features of the
			quantitative analysis of pharmaceutical substances
			and drugs. Validation of analytical methods.
			Weight analysis (gravimetry).
			Method of acid-base titration in aqueous and non-
			aqueous media, complexometry, argentometry,
			bromatometry, iodometry, nitritometry.
			Determination of nitrogen in organic compounds.
			Optical methods: UV and IR spectrophotometry,
			NMR spectroscopy, photometry in the visible region
			of the spectrum, refractometry, polarimetry. Methods
			based on the emission of radiation: flame photometry,
			fluorimetry.
			Chromatographic methods: TLC, gas-liquid
			chromatography (GLC) and high performance liquid
			chromatography (HPLC), electrophoresis.
			Modern trends in the development of pharmaceutical
			analysis.
			Classification of medicinal products of inorganic
			compounds. Comparative assessment of quality
			requirements.
			Medicinal products of elements of group VII of the
			periodic system of elements. Iodine. Potassium and
			sodium chlorides, bromides, iodides. sodium fluoride.
			Hydrochloric acid.
			Medicinal products of elements VI, V and IV of
			groups of the periodic system of elements. Oxygen.
			Purified water, water for injections. Hydrogen
			peroxide solution, hydroperite (urea peroxide).
	UC-1,2	. .	Sodium thiosulfate, sodium metabisulphite. Sodium
2.	GPC-	Inorganic	bicarbonate, lithium carbonate, talc.
	1,3,6	medicines	Medicines of elements of groups II and III of the
	PC-4,7		periodic table of elements. Barium sulfate for
			fluoroscopy. Calcium chloride, calcium sulfate.
			Magnesium oxide, magnesium sulfate. Aluminum
			hydroxide, aluminum phosphate. Boric acid, sodium
			tetraborate.
			Medicines of bismuth, silver, copper, zinc. Bismuth
			nitrate basic. Zinc oxide, zinc sulfate. Silver nitrate,
			collargol (colloidal silver), protargol (silver
			proteinate). copper sulfate.
			Iron(II) compounds. Iron(II) sulfate. Complex
			compounds of iron (III) and platinum (IV). Maltofer,
			cisplatin.
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			radiopharmaceuticals. Prerequisites for the use of
			radioactive substances for diagnostic and therapeutic
			purposes. Features of standardization of
			radiopharmaceuticals. Sodium o-iodine hippurate.
			Halogen derivatives of hydrocarbons. Chloroethyl,
			halothane (halothane).
			Alcohols, aldehydes and ethers.
			Ethyl alcohol, glycerol (glycerin), polyethylene
			glycol, nitroglycerin, diethyl ether (medical ether and
			ether for anesthesia), formaldehyde solution.
			Carbohydrates (mono- and polysaccharides). Glucose,
			sucrose, lactose, glucosamine, chondroitin sulfate,
			starch, hydroxyethyl starch, hyaluronic acid.
			Derivatives of carbohydrates as excipients.
			Methylcellulose, carboxymethylcellulose.
			Carboxylic acids and their derivatives. Sodium
	UC-1,2	IC-1.2 Medicinal	acetate, calcium lactate, calcium gluconate, sodium
	GPC-	products of	citrate, sodium valproate, meldonium (mildronate),
3.	1,3,6	aliphatic and alicyclic structure.	sorbic acid.
	PC-4,7		Derivatives of uronic acids. Alginic acid.
	rC-4,/		-
			Lactones of unsaturated polyhydroxycarboxylic acids. Ascorbic acid.
			Amino acids and their derivatives. Glutamic acid,
			aminocaproic acid, gamma-aminobutyric acid
			(aminalon), methionine, cysteine, acetylcysteine,
			aspartame.
			Derivatives of polyaminopolycarboxylic acids.
			Tetacin-calcium (calcium sodium edetate).
			Piracetam, phenotropil as analogues of gamma-
			aminobutyric acid lactam.
			Proline derivatives: captopril, enalapril, lisinopril.
			Monocyclic terpenes: menthol, validol, terpinhydrate.
			Bicyclic terpenes: camphor, sulphocamphoric acid
			and its novocaine salt (sulfocamphocaine).
			Diterpenes: retinols and their derivatives (group A
			vitamins) as medicinal and prophylactic agents.
	UC-1,2		Statins. Lovastatin, simvastatin.
4.	GPC-	Terpenes and	Derivatives of cyclopentanperhydrophenanthrene.
4.	1,3,6	steroids.	
	PC-4,7		Cyclohexanolethylenehydrindane compounds.
			Calciferols (group D vitamins) as sterol conversion
			products. The mechanism of formation of vitamins
			ergocalciferol (D2) and cholecalciferol (D3).
			Cardenolides (cardiac glycosides). Structure and
			classification. Standardization. Biological and

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			physico-chemical methods for quantitative assessment of the activity of cardiac glycosides. Stability. Digitalis glycosides: digitoxin, digoxin.
			A number of strophanthidine: strophanthin K,
			preparations of lily of the valley.
			Corticosteroids. Mineral corticosteroids:
			Desoxycortone acetate (deoxycorticosterone acetate).
			Glucocorticosteroids: ortisone acetate, prednisolone,
			hydrocortisone acetate, dexamethasone, fluocinolone acetonide (sinaflan).
			Gestagens and their synthetic analogues.
			Progesterone, norethisterone, medroxyprogesterone acetate.
			Androgens. Testosterone propionate,
			methyltestosterone. Anabolic steroids: methandienone
			(methandrostenolone), methandriol
			(methylandrostenediol), nandrolone phenylpropionate
			(phenobolin), nandrolone decanoate (retabolil),
			Antiandrogens: Cyproterone acetate (Androcur).
			Estrogens. Estrone and estradiol as medicinal
			substances.
			Prerequisites for obtaining derivatives: ethinylestradiol, estradiol esters.
			Antiestrogens: tamoxifen, anastrozole (arimidex).
			Non-steroidal estrogen analogues: hexestrol
			(sinestrol), diethylstilbestrol.
			Phenols, quinones and their derivatives.
			Medicines of the phenol group: phenol, thymol,
			resorcinol, etamsylate, guaifenesin.
			Derivatives of naphthoquinones (group K vitamins):
			sodium menadione bisulfite (Vikasol). Aminophenol derivatives.
			Derivatives of n-aminophenol: paracetamol.
	UC-1,2		Derivatives of m-aminophenol: neostigmine methyl
5.	GPC-	Aromatic drugs	sulfate (prozerin).
	1,3,6		Aromatic acids and their derivatives. Benzoic acid,
	PC-4,7		sodium benzoate. Salicylic acid, sodium salicylate.
			Derivatives of p-hydroxybenzoic acid. Ethyl
			parahydroxybenzoate.
			Esters of salicylic acid. Acetylsalicylic acid.
			Derivatives of phenylpropionic acid. ibuprofen,
			ketoprofen.
			Derivatives of phenylacetic acid. Diclofenac sodium.

	Derivatives of butyrophenone. Haloperidol.
	aromatic amino acids.
	Derivatives of p-aminobenzoic acid: benzocaine
	(anesthesin), procaine hydrochloride (novocaine
	hydrochloride), tetracaine hydrochloride (dicaine).
	Diethylaminoacetanilides: trimecaine hydrochloride,
	lidocaine hydrochloride.
	Local anesthetics similar in structure: bupivacaine,
	articaine hydrochloride (ultracaine).
	Derivatives of p-aminobenzoic acid amide:
	procainamide hydrochloride (novocainamide),
	metoclopramide hydrochloride.
	Derivatives of p-aminosalicylic acid: sodium p-
	aminosalicylate.
	Derivatives of m-aminobenzoic acid: amidotrizoic
	acid and its sodium and N-methylglucamine salts
	(Triombrast for injections).
	Arylalkylamines and their derivatives. Biochemical
	prerequisites for obtaining medicinal substances in
	the series of phenylalkylamines. Ephedrine
	hydrochloride. Dopamine (dopamine). Epinephrine
	(adrenaline) and norepinephrine (norepinephrine),
	their salts. Isoprenaline hydrochloride, fenoterol,
	salbutamol, verapamil.
	Derivatives of hydroxyphenylaliphatic amino acids:
	levodopa, methyldopa.
	Derivatives of substituted aryloxypropanolamines (β -
	blockers): propranolol hydrochloride (anaprilin),
	atenolol, timolol, bisoprolol, fluoxetine.
	Aminodibromophenylalkylamines: bromhexine
	hydrochloride, ambroxol hydrochloride.
	Iodized derivatives of aromatic amino acids.
	Liothyronine (triiodothyronine), levothyroxine
	(thyroxine). Complex drug - thyroidin.
	Amides of benzenesulfonic acid. Sulfanilamide
	(streptocide).
	Sulfonamides substituted at the amide group
	(aliphatic and heterocyclic series): sodium
	sulfacetamide, co-trimoxazole, sulfadimethoxine,
	sulfalene.
	Sulfonamides substituted at the amide and aromatic
	amino groups. Phthalylsulfathiazole (phthalazol),
	salazopyridazine.
	Benzenesulfonic acid amide derivatives: furosemide,
	hydrochlorothiazide (dichlorothiazide, hypothiazide),

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			bumetanide.
			Derivatives of benzenesulfochloramide: chloramine
			B, halazon (pantocid).
			Derivatives of amides of sulfonic acids (substituted
			sulfonylurea) as antidiabetic agents. Carbutamide
			(Bucarban), glipizide (Minidiab), glibenclamide,
			gliclazide (Predian), gliquidone (Glurenorm).
			Non-aromatic antidiabetic drugs - biguanides:
			Metformin.
			Classification by action, chemical classification.
			quality requirements. Activity unit. Biological,
			chemical and physico-chemical methods of quality
			assessment. Standard samples of antibiotics.
			Beta lactamides.
			Penicillins. General characteristics and structure.
			Relationship between structure and biological action.
			Penicillins of natural origin: benzylpenicillin and
			preparations based on it, phenoxymethylpenicillin.
			Targeted semi-synthesis based on 6-aminopenicillanic
			acid (6-APA).
			Semi-synthetic penicillins: oxacillin sodium salt,
			ampicillin, carbenicillin disodium salt, amoxicillin.
			Beta-lactamase inhibitors: sulbactam, clavulanic acid.
			Combined preparations of penicillins: amoxiclav.
			Cephalosporins. Methods for obtaining
	UC-1,2		cephalosporins based on 7-aminocephalosporanic
6.	GPC-	Antibiotics	acid.
0.	1,3,6	1 millionoticis	I generation cephalosporins: cephalexin, cefazolin.
	PC-4,7		II generation cephalosporins: cefaclor, cefuroxime.
			III generation cephalosporins: ceftizoxime,
			cefotaxime.
			6th generation cephalosporins: cefmetazole,
			cefoxitim.
			Aminoglycoside antibiotics: streptomycin sulfate,
			kanamycin sulfate, gentamicin sulfate, amikacin.
			Derivatives of tetrahydropyrrole. Lincomycins:
			lincomycin hydrochloride, clindamycin.
			Macrolides and azalides: erythromycin, azithromycin.
			Tetracyclines. Tetracycline hydrochloride,
			oxytetracycline hydrochloride.
			Semi-synthetic analogues: doxycycline, metacycline.
			Aromatic nitro derivatives: chloramphenicol
			(levomycetin) - an aromatic antibiotic and its esters
			(stearate and succinate). Nimesulide.
7.	UC-1,2	Medicinal	Derivatives of 5-nitrofuran. Nitrofural, furagin,
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GPC-	products of	nifuratel, nifuroxazide (enterofuril).
1,3,6	heterocyclic	Furan derivatives. Amiodarone, griseofulvin.
PC-4	-	Benzopyran derivatives.
	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Chromane compounds as medicinal and prophylactic
		agents (group E vitamins - tocopherols). Tocopherol
		acetate.
		Benzo-gamma-pyrone derivatives: Cromoglycic acid
		(sodium cromoglycate).
		Phenylchromane compounds are flavonoids (group P
		vitamins). Rutoside (rutin), quercetin,
		dihydroquercetin, diosmin.
		Derivatives of pyrrole (vitamins of group B12).
		Cyanocobalamin, hydroxocobalamin, cobamamide.
		Derivatives of pyrrolizidine. Platifillina hydrotartrate,
		povidone (polyvinylpyrrolidone).
		Pyrazole derivatives. Phenazone (antipyrine),
		metamizole sodium (analgin), phenylbutazone
		(butadione), propyphenazone.
		indole derivatives. Reserpine, indomethacin, arbidol,
		vinpocetine.
		Ergoline derivatives (ergot alkaloids and their
		derivatives): nicergoline, ergometrine, ergotamine,
		methylergometrine, bromocriptine.
		imidazole derivatives. Pilocarpine hydrochloride,
		bendazole hydrochloride (dibazole), clonidine
		hydrochloride (clopheline), metronidazole,
		naphazoline nitrate (naphthyzine), clotrimazole,
		omeprazole and its S-isomer - esomeprazole
		(nexium), afobazole, domperidone (motilium),
		xylometazoline (galazolin).
		Histamine dihydrochloride.
		Antihistamines: diphenhydramine hydrochloride
		(diphenhydramine), chloropyramine, ranitidine,
		famotidine.
		1,2,4-triazole derivatives: fluconazole (Diflucan).
		Piperidine derivatives: trihexyphenidyl hydrochloride
		(cyclodol), ketotifen, loratadine, loperamide
		hydrochloride.
		Derivatives of dihydropyridine: nifedipine,
		amlopidine, nicardipine.
		Derivatives of pyridine-3-carboxylic acid: nicotinic
		acid, nicotinamide, nikethamide (nicotinic acid
		diethylamide), sodium salt of N-nicotinoyl-gamma-
		aminobutyric acid (picamilon), betahistine.
		Pyridine-4-carboxylic acid derivatives: isoniazid,

ftivazid, prothionamide, ethionamide.
Pyridinemethanol derivatives. Pyridoxine
hydrochloride (B6 vitamins), pyridoxal phosphate,
ethylmethylhydroxypyridine (emoxipin).
Tropane derivatives.
Tropane alkaloids and their synthetic analogues.
Atropine sulfate, scopolamine hydrochloride,
homatropine hydrobromide, tropacin, etc.
Derivatives of quinoline and isoquinoline.
Derivatives of 4-substituted quinoline. Quinine,
quinidine and their salts. Chloroquine phosphate
(Chingamine), Hydroxychloroquine sulfate
(Plaquenil).
Derivatives of 8-hydroxyquinoline: nitroxoline (5-
NOC), chlorquinaldol.
Fluoroquinolones: lomefloxacin, ofloxacin,
ciprofloxacin.
Benzylisoquinoline derivatives. Papaverine
hydrochloride and its synthetic analogue - drotaverine
hydrochloride.
Derivatives of phenanthrenisoquinoline. Morphine,
codeine and their salts.
Derivatives of morphine. Apomorphine
hydrochloride, ethylmorphine hydrochloride, glaucine
hydrochloride.
Synthetic analogues of morphine. Trimeperidine
hydrochloride (promedol), tramadol hydrochloride,
fentanyl.
Piperazine derivatives - cinnarizine.
Pyrimidine derivatives.
Derivatives of pyrimidine-2,4,6-trione (barbituric and
thiobarbituric acids). Phenobarbital, thiopental
sodium, benzonal (benzobarbital), hexobarbital
sodium (hexenal).
Derivatives of pyrimidine-2,4-dione. Methyluracil,
fluorouracil. Nucleosides: tegafur (ftorafur),
zidovudine (azidothymidine), stavudine.
Pyrimidine-4,6-dione derivatives: primidone
(hexamidine). Hydantoin derivatives. Phenytoin (difenin).
Purine derivatives.
The value of antimetabolites in the development of
-
new drugs. Xanthine derivatives: caffeine, theophylline,
theobromine, caffeine-sodium benzoate,

	1	r	
			aminophylline (eufillin), diprophylline, xanthinol
			nicotinate, pentoxifylline.
			Guanine derivatives. Acyclovir (Zovirax),
			Ganciclovir (Cymeven).
			Other purine derivatives: inosine (riboxin),
			allopurinol, mercaptopurine, azathioprine.
			Derivatives of pteridine and isoalloxazine.
			A group of folic acid derivatives. Folic acid and its
			analogues. Methotrexate.
			Derivatives of isoalloxazine (vitamin B2). Riboflavin,
			riboflavin mononucleotide.
			Derivatives of phenothiazine. Alkylamino
			derivatives: chlorpromazine hydrochloride
			(chlorpromazine), levomepromazine, trifluoperazine
			dihydrochloride (triftazine), fluphenazine decanoate,
			etc.
			Acyl derivatives: ethacizine, moracizine
			hydrochloride (ethmozine).
			Derivatives of benzodiazepines. Chlordiazepoxide,
			diazepam (sibazon), medazepam, nitrazepam,
			phenazepam, alprazolam, etc.
			Dibenzodiazepine derivatives: clozapine (azaleptin).
			Derivatives of 1,2-benzothiazine: piroxicam.
			Derivatives of 10,11-dihydrodibenzocycloheptene:
			amitriptyline.
			Derivatives of 1,5-benzothiazepine: diltiazem.
			Derivatives of iminostilbene: carbamazepine.
			Pyrimidinothiazole derivatives. Vitamins of group
			B1. Thiamine chloride and bromide, phosphothiamin,
			cocarboxylase, benfotiamine.
			Basics of metrology. Basic concepts. Metrological
			characteristics of the analysis results.
			Statistical processing of analysis results in accordance
			with the requirements of the Global Fund.
		Metrological	Types of analysis error. Errors in the analysis of
	UC-1,2	foundations of	physicochemical and chemical methods. Methods for
	GPC-	pharmaceutical	identifying systematic and random errors.
8.	1,3,6	analysis.	Validation evaluation of analysis methods. Validation
	PC-4,7	Validation	characteristics of the main types of methods.
	1,+, /	evaluation of	Establishment of the specificity of methods of
		analysis methods	qualitative and quantitative analysis, determination of
			foreign impurities. Linearity. Precision. Accuracy and
			correctness of analysis methods. Limit of detection
9.	UC-1,2	Standardization	and quantification. robustness. Legislation of the Russian Federation regulating the
9.	00-1,2	Stalidardization	Legislation of the Russian Federation regulating the

GPC-	and quality control	circulation of medicines.
1,3,6	of medicines.	
		State regulation of drug quality control
PC-4,7	Declaring the	The main directions of the modern concept of quality
	quality of	assurance of medicines. Rules for preclinical studies
	medicines	of the safety and efficacy of future drugs (GLP rules).
		Good Clinical Practice (GCP Practice).
		Organization of quality control of medicines. GMP
		rules. Quality control of medicines in production
		(industrial enterprises and pharmacies).
		Standardization of medicines as an organizational and
		technical basis for product quality management.
		Quality standards for medicines: OFS, FS, FSP, ND,
		orders of the Ministry of Health of the Russian
		Federation.
		Declaration of quality of medicines. Declaration
		procedure. The main stages of declaring the quality of
		medicines.
		Organization of quality control in the production of
		medicines at industrial enterprises and pharmacies.
		Quality control of medicines during storage. The
		study of the expiration dates of medicines.
		Methodological approach to the choice of methods
		for the analysis of industrial and pharmaceutical
		drugs.
		urugo.

5. Volume of the academic discipline and types of academic work

Type of educational work	Labor	intensity	Labor intensity (AH) in semesters		sters		
	volume vo						
	in credit	academic	5	6	7	8	9
	units (CU)	hours (AH)					
classroom work, including	10.9	390	92	84	84	64	66
Lectures (L)	2.6	92	22	14	20	18	18
Practicals (P)	8.3	298	70	70	64	46	48
Student's individual work (SIW)	7.1	258	52	60	60	44	42
Mid-term assessment							
exam	1	36					36
TOTAL LABOR INTENSITY	19	684	144	144	144	108	144

6. Content of the academic discipline

6.1. Sections of the discipline and types of academic work

ſ	p / no.	semester	Name	of	the	Types of academic work* (in AH)					
		number	section	of	the	L	Р			SIW	Total
			academic	,							

		the quality of medicines TOTAL	92		298			258	648
9	9	quality control of medicines. Declaring							
		Standardization and	14	-	24	-	-	18	56
8	9	pharmaceutical analysis. Validation evaluation of analysis methods							
		Metrological foundations of	4	-	24	-	-	18	46
7	7.8	Medicinal products of heterocyclic structure.	29	-	74	-	-	66	169
6	7	Antibiotics	9	-	28	-	-	30	67
5	6	Aromatic drugs	8	-	31	-	-	42	81
4	6	Terpenes and steroids.	6	-	11	-	-	18	35
3	5	aliphatic and alicyclic structure.							
		Medicinal products of	10	-	32	-	-	30	72
2	5	Inorganic medicines	10	-	30	-	-	18	58
1	5	FundamentalsofPharmaceuticalAnalysis	2	-	44	-	-	18	64
		discipline						10	

* - L – lectures; LP – laboratory practicum; P – practicals; S – seminars; SIW – student's individual work.

6.2. Thematic schedule of educational work types:

6.2.1 Thematic schedule of lectures

No.	Name of lecture topics			Volume in	AH	
p / p		5	6	7	8	9
1.	Pharmaceutical chemistry as a science.	2				
2.	Medicinal products of elements VI, V and IV of groups of the periodic system of elements.Oxygen. Purified water, water for injections. Hydrogen peroxide solution, hydroperite (urea peroxide). Sodium thiosulfate, sodium metabisulphite. Sodium bicarbonate, lithium carbonate, talc.	2				
3.	Medicinal products of elements of group VII of the periodic system of elements. Iodine. Potassium and sodium chlorides, bromides, iodides. sodium fluoride. Hydrochloric acid.	2				
4.	<i>radiopharmaceuticals</i> .Prerequisites for the use of radioactive substances for diagnostic and therapeutic purposes. Features of standardization of radiopharmaceuticals. Sodium o-iodine hippurate.	2				
5.	Medicines of elements of groups II and III of the periodic table of elements.Barium sulfate for fluoroscopy. Calcium chloride, calcium sulfate. Magnesium oxide, magnesium sulfate. Aluminum hydroxide, aluminum phosphate.	2				

	Boric acid, sodium tetraborate.				
E					
6.	Medicines of bismuth, silver, copper,				
	zinc.Bismuth nitrate basic. Zinc oxide, zinc				
	sulfate. Silver nitrate, collargol (colloidal				
	silver), protargol (silver proteinate). copper	2			
	sulfate.				
	<i>Iron(II) compounds</i> .Iron(II) sulfate. Complex				
	compounds of iron (III) and platinum (IV).				
	Maltofer, cisplatin.				
7.	Organic medicines. Classification,				
	nomenclature. Sources and methods of				
	obtaining. Analysis methods.	1			
	Halogen derivatives of				
	hydrocarbons. Chloroethyl, halothane				
	(halothane).			ļ	
8.	Alcohols, aldehydes and esters. Ethyl				
	alcohol, glycerol (glycerin), polyethylene	-			
	glycol, nitroglycerin, diethyl ether (medical	2			
	ether and ether for anesthesia), formaldehyde				
	solution.			 ļ	
9.	Carboxylic acids and their				
	derivatives. Sodium acetate, calcium lactate,	2			
	calcium gluconate, sodium citrate, sodium				
	valproate, meldonium (mildronate), sorbic acid.			ļ	
10.	Derivatives of uronic acids. Alginic acid.	_			
	Lactones of unsaturated	2			
	polyhydroxycarboxylic acids. Ascorbic acid.			 	
11.	Amino acids and their derivatives. Glutamic				
	acid, aminocaproic acid, gamma-aminobutyric				
	acid (aminalon), methionine, cysteine,				
	acetylcysteine, aspartame.				
	Derivatives of polyaminopolycarboxylic				
	acids. Tetacin-calcium (calcium sodium	3			
	edetate).				
	Piracetam, phenotropil as analogues of gamma-				
	aminobutyric acid lactam.				
	Proline derivatives: captopril, enalapril,				
	lisinopril.				
12.	Monocyclic terpenes: menthol, validol,		2		
	terpinhydrate.				
	Bicyclic terpenes: camphor, sulphocamphoric				
	acid and its novocaine salt (sulfocamphocaine).				
	Diterpenes: retinols and their derivatives				
	(group A vitamins) as medicinal and				
	prophylactic agents.				
	Statins.Lovastatin, simvastatin.				
13.	Derivatives of		1		
	cyclopentanperhydrophenanthrene.				
	Cyclohexanolethylenehydrindane				
	compounds.Calciferols (group D vitamins) as				
	sterol conversion products. The mechanism of				
	formation of vitamins ergocalciferol (D2) and				

	cholecalciferol (D3).			
14.	Corticosteroids. Mineral	1		
14.	<i>corticosteroids</i> : Desoxycortone acetate	1		
	(deoxycorticosterone acetate).			
	<i>Glucocorticosteroids</i> :cortisone acetate,			
	prednisolone, hydrocortisone acetate,			
	dexamethasone, fluocinolone acetonide			
	(sinaflan).			
15.	Gestagens and their synthetic analogues.	1		
13.	Progesterone, norethisterone,	1		
	medroxyprogesterone acetate.			
	Androgens. Testosterone propionate,			
	methyltestosterone.			
	<i>Anabolic steroid</i> :Methandienone			
	(Methandrostenolone), Methandriol			
	(Methylandrostenediol), Nandrolone			
	Phenylpropionate (Phenobolin), Nandrolone			
	Decanoate (Retabolyl),			
	Antiandrogens:cyproterone acetate			
	(Androcur).			
16.	<i>Estrogens</i> .Estrone and estradiol as medicinal	1		
10.	substances.	1		
	Prerequisites for obtaining derivatives:			
	ethinylestradiol, estradiol esters.			
	Antiestrogens:tamoxifen, anastrozole			
	(arimidex).			
	Non-steroidal estrogen analogues: hexestrol			
	(sinestrol), diethylstilbestrol.			
17.	aromatic compounds. General information	1		
17.	about the dependence of the chemical structure	1		
	and biological action in a number of aromatic			
	compounds. Sources and methods of obtaining.			
	General and private methods of analysis.			
	Phenols, quinones and their derivatives.			
	Medicines of the phenol group: phenol,			
	thymol, resorcinol, etamsylate, guaifenesin.			
	Derivatives of naphthoquinones (vitamins of			
	group K): menadione sodium bisulfite			
	(Vikasol)			
	Aminophenol derivatives.			
	Derivatives of n-aminophenol: paracetamol.			
	<i>m-aminophenol derivatives</i> : neostigmine			
	methyl sulfate (prozerin).			
18.	Aromatic acids and their derivatives. Benzoic	1		
	acid, sodium benzoate. Salicylic acid, sodium			
	salicylate.			
	Derivatives of p-hydroxybenzoic acid. Ethyl			
	parahydroxybenzoate.			
	<i>Esters of salicylic acid</i> . Acetylsalicylic acid.			
	Phenylpropionic acid derivatives. ibuprofen,			
	ketoprofen.			
	Derivatives of phenylacetic acid.Diclofenac			

	sodium.	[[
10	Derivatives of butyrophenone. Haloperidol.				
19.	aromatic amino acids.	2			
	Derivatives of p-aminobenzoic acid: benzocaine				
	(anesthesin), procaine hydrochloride (novocaine				
	hydrochloride), tetracaine hydrochloride				
	(dicaine). Diethylaminoacetanilides: trimecaine				
	hydrochloride, lidocaine hydrochloride.				
	Structurally related local				
	anesthetics: bupivacaine, articaine				
	hydrochloride (ultracaine).				
20.	Derivatives of p-aminobenzoic acid	1			
	amide:procainamide hydrochloride				
	(novocainamide), metoclopramide				
	hydrochloride.				
	Derivatives of p-aminosalicylic acid:sodium				
	p-aminosalicylate.				
	Derivatives of m-aminobenzoic				
	acid: amidotrizoic acid and its sodium and N-				
	methylglucamine salts (Triombrast for				
	injection).				
21.	Arylalkylamines and their derivatives.	1			
	Biochemical prerequisites for obtaining				
	medicinal substances in the series of				
	phenylalkylamines. Ephedrine hydrochloride.				
	Dopamine (dopamine). Epinephrine				
	(adrenaline) and norepinephrine				
	(norepinephrine), their salts. Isoprenaline				
	hydrochloride, fenoterol, salbutamol,				
	verapamil.				
22.	Derivatives of hydroxyphenyl-aliphatic	2			
	amino acids: levodopa, methyldopa.				
	Derivatives of substituted aryloxy-				
	propanolamines $(\beta$ -blockers):propranolol				
	hydrochloride (anaprilin), atenolol, timolol,				
	bisoprolol, fluoxetine.				
	Aromatic nitro derivatives: chloramphenicol				
	(levomycetin) is an aromatic antibiotic and its				
	esters (stearate and succinate). Nimesulide.				
	Aminodibromophenylalkylamines:				
	bromhexine hydrochloride, ambroxol				
	hydrochloride.				
	Iodized derivatives of aromatic amino acids.				
	Liothyronine (triiodothyronine), levothyroxine				
	(thyroxine). Complex drug - thyroidin.				
23.	Antibiotics. Classification by action, chemical		2		
	classification. quality requirements. Activity unit.				
	Biological, chemical and physico-chemical				
	methods of quality assessment. Standard samples				
	of antibiotics.				
24.	Beta lactamides.		2		
	<i>Penicillins.</i> General characteristics and		_		
<u>I</u>	- Stronthis, Sonoral Characteristics and			1	

	structure. Relationship between structure and				
	biological action.				
	Penicillins of natural origin: benzylpenicillin				
	and drugs based on it, phenoxymethylpenicillin.				
	Targeted semi-synthesis based on 6-				
	aminopenicillanic acid (6-APA).				
	Semi-synthetic penicillins: oxacillin sodium				
	salt, ampicillin, carbenicillin disodium salt,				
	amoxicillin.				
	Beta-lactamase inhibitors: sulbactam,				
	clavulanic acid.				
	Combined preparations of				
	penicillins:amoxiclav.				
25.	Cephalosporins. Methods for obtaining		2		
	cephalosporins based on 7-aminocephalosporanic				
	acid.				
	1st generation cephalosporins:cephalexin,				
	cefazolin.				
	<i>II generation cephalosporins</i> :cefaclor,				
	cefuroxime.				
	Third generation cephalosporins:ceftizoxime,				
	cefotaxime.				
	6th generation cephalosporins:cefmetazole,				
	cefoxitim.				
26.	Aminoglycoside antibiotics: streptomycin		2		
	sulfate, kanamycin sulfate, gentamicin sulfate,				
	amikacin.				
	<i>Derivatives of tetrahydropyrrole</i> . Lincomycins:				
27	lincomycin hydrochloride, clindamycin.		1		
27.	<i>macrolides and azalides</i> : erythromycin,		1		
29	azithromycin.		1		
28.	Heterocyclic compounds of natural and		1		
	synthetic origin. Study of natural biologically				
	active compounds of heterocyclic structure as				
	one of the ways to create new medicinal substances. Classification of heterocyclic				
	compounds. Application of general physical				
	and chemical patterns in the formation of				
	requirements for the quality of medicinal				
	substances and the choice of analysis methods.				
	oxygen-containing heterocycles.				
	Derivatives of 5-nitrofuran.Nitrofural,				
	furagin, nifuratel, nifuroxazide (enterofuril).				
	<i>Furan derivatives</i> . Amiodarone, griseofulvin.				
29.	Benzopyran derivatives.		1		
	<i>Chromine compounds</i> as medicinal and				
	prophylactic agents (group E vitamins -				
	tocopherols). Tocopherol acetate.				
	Benzo-gamma-pyrone derivatives:				
	Cromoglycic acid (sodium cromoglycate)				
30.	Phenylchromane compounds- flavonoids		1		
	(vitamins of the P group). Rutoside (rutin),				
-		•		•	•

	quercetin, dihydroquercetin, diosmin.			
31.	nitrogen-containing heterocycles.	3		
	<i>Pyrrole derivatives</i> (vitamins B12).			
	Cyanocobalamin, hydroxocobalamin,			
	cobamamide.			
	<i>Derivatives of pyrrolizidine</i> .Platifillina			
	hydrotartrate, povidone (polyvinylpyrrolidone).			
	<i>Pyrazole derivatives.</i> Phenazone (antipyrine),			
	metamizole sodium (analgin), phenylbutazone			
	(butadione), propyphenazone.			
32.	<i>indole derivatives.</i> Reserpine, indomethacin,	2		
52.	arbidol, vinpocetine.			
	<i>Ergoline derivatives</i> (ergot alkaloids and their			
	derivatives): nicergoline, ergometrine,			
	ergotamine, methylergometrine, bromocriptine.			
33.	<i>imidazole derivatives</i> . Pilocarpine	3		
55.	hydrochloride, bendazole hydrochloride	5		
	(dibazole), clonidine hydrochloride (clophelin),			
	metronidazole, naphazoline nitrate			
	(naphthyzine), clotrimazole, omeprazole and its			
	S-isomer - esomeprazole (nexium),			
	domperidone (motilium), xylometazoline			
	(galazolin), afobazole.			
	Histamine dihydrochloride.			
	Antihistamines: diphenhydramine			
	hydrochloride (diphenhydramine),			
	chloropyramine, ranitidine, famotidine.			
34.	<i>Piperidine derivatives</i> : trihexyphenidyl		1	
51.	hydrochloride (cyclodol), ketotifen, loratadine,		-	
	loperamide hydrochloride.			
	Derivatives of dihydropyridine: nifedipine,			
	amlopidine, nicardipine.			
35.	Derivatives of pyridine-3-carboxylic		1	
55.	<i>acid</i> :nicotinic acid, nicotinamide, nikethamide			
	(nicotinic acid diethylamide), N-nicotinoyl-			
	gamma-aminobutyric acid sodium salt			
	(picamilon), betahistine.			
36.	Derivatives of pyridine-4-carboxylic		1	
	<i>acid</i> :isoniazid, ftivazid, protionamide,			
	ethionamide.			
	Pyridinemethanol derivatives. Pyridoxine			
	hydrochloride (B6 vitamins), pyridoxal			
	phosphate, ethylmethylhydroxypyridine			
	(emoxipin).			
37.	Tropane derivatives.		1	
	Tropane alkaloids and their synthetic			
	analogues. Atropine sulfate, scopolamine			
	hydrochloride, homatropine hydrobromide,			
	tropacin, etc.			
38.	Derivatives of quinoline and isoquinoline.		1	
	Derivatives of 4-substituted quinoline.			
	Quinine, quinidine and their salts. Chloroquine			

		<u>г</u>	1	
	phosphate (Chingamine), Hydroxychloroquine			
	sulfate (Plaquenil).			
	8-hydroxyquinoline derivatives: nitroxoline			
	(5-NOC), chlorquinaldol.			
39.	Fluoroquinolones: lomefloxacin, ofloxacin,		1	
	ciprofloxacin.			
	Benzylisoquinoline derivatives. Papaverine			
	hydrochloride and its synthetic analogue -			
	drotaverine hydrochloride			
40.	Phenantrenisoquinoline derivatives and their		2	
	synthetic analogues.			
	Derivatives of			
	phenanthrenisoquinoline.Morphine, codeine			
	and their salts.			
	<i>Morphine derivatives.</i> Apomorphine			
	hydrochloride, ethylmorphine hydrochloride,			
	glaucine hydrochloride.			
	Synthetic analogues of			
	<i>morphine</i> .Trimeperidine hydrochloride			
	(promedol), tramadol hydrochloride, fentanyl.			
41.	Piperazine derivatives- cinnarizine.		1	
	Pyrimidine derivatives.			
	Derivatives of pyrimidine-2,4,6-trione			
	(barbituric and thiobarbituric			
	acids). Phenobarbital, thiopental sodium, benzonal			
	(benzobarbital), hexobarbital sodium (hexenal).			
42.	<i>Pyrimidine-2,4-dione derivatives.</i>		1	
	Methyluracil, fluorouracil. Nucleosides: tegafur			
	(ftorafur), zidovudine (azidothymidine),			
	stavudine.			
	<i>Pyrimidine-4,6-dione derivatives</i> : primidone			
	(hexamidine).			
	Hydantoin derivatives. Phenytoin (difenin).			
43.	<i>Pyrimidinothiazole derivatives.</i> Vitamins of		1	
15.	group B1. Thiamine chloride and bromide,			
	phosphothiamin, cocarboxylase, benfotiamine.			
44.	Purine derivatives.		1	
++.	The value of antimetabolites in the development			
	_			
	of new drugs.			
	<i>xanthine derivatives</i> : caffeine, theophylline,			
	theobromine, sodium caffeine benzoate,			
	aminophylline (euphylline), diprophylline,			
L	xanthinol nicotinate, pentoxifylline.			
45.	Guanine derivatives. Acyclovir (Zovirax),		2	
	Ganciclovir (Cymeven).			
	Other Purine Derivatives: inosine (riboxin),			
L	allopurinol, mercaptopurine, azathioprine.			
46.	Derivatives of pteridine and isoalloxazine.		1	
	A group of folic acid derivatives. Folic acid and			
	its analogues. Methotrexate.			
	Isoalloxazine derivatives(vitamin B2).			
	Riboflavin, riboflavin mononucleotide.			
			· ·	

		1				
47.	Derivatives of phenothiazine. Alkylamino				1	
	derivatives: chlorpromazine hydrochloride					
	(chlorpromazine), levomepromazine,					
	trifluoperazine dihydrochloride (triftazine),					
	fluphenazine decanoate, etc.					
	Acyl derivatives: ethacizine, moracizine					
	hydrochloride (ethmozine).					
48.	benzodiazepine derivatives. Chlordiazepoxide,				1	
	diazepam (sibazon), medazepam, oxazepam,					
	nitrazepam, phenazepam, alprazolam, etc.					
	Dibenzodiazepine derivatives: clozapine					
	(azaleptin).					
49.	<i>1,2-benzothiazine derivatives</i> : piroxicam.				1	
	Derivatives of 10,11-					
	dihydrodibenzocycloheptene: amitriptyline.					
	Derivatives of 1,5-benzothiazepine: diltiazem.					
	Derivatives of iminostilbene: carbamazepine.					
50.	Types of analysis error. Errors in the analysis					4
	of physicochemical and chemical methods.					
	Methods for identifying systematic and random					
	errors.					
51.	Validation evaluation of analysis methods.					2
	Validation characteristics of the main types of					
	methods. Establishment of the specificity of					
	methods of qualitative and quantitative analysis,					
	determination of foreign impurities. Linearity.					
	Precision. Accuracy and correctness of analysis					
	methods. Limit of detection and quantification.					
	robustness.					
52.	Organization of quality control of medicines.					6
	GMP rules. Quality control of medicines in					
	production (industrial enterprises and					
	pharmacies). Declaration of quality of					
	medicines.					
53.	Quality control of medicines during storage.					6
	The study of the expiration dates of medicines.					
	TOTAL (total 129 Ah)	22	14	20	18	18
I						

6.2.2. Thematic plan of practicals

No.	Name of topics of practical classes	Volume in AH				
p / p		5	6	7	8	9
1.	Introductory lesson. Goals and objectives of the					
	laboratory workshop. Safety precautions in the	2				
	chemical laboratory.					
2.	RD for medicines. General methods of pharmacopoeial analysis. Work with the methodical manual.	8				
3.	Determination of the quality of medicinal substances by appearance, color, transparency and degree of turbidity, solubility.	8				

				[
	Determination of weight loss on drying. Tablet				
	disintegration test.				
4.	Determination of the authenticity of inorganic	4			
	medicinal substances.	•			
5.	Determination of impurities of inorganic ions in				
	medicinal substances. Standard and non-	4			
	standard methods for the determination of	4			
	impurities. Solution of situational problems.				
6.	Preparation of reagents, indicators, buffer				
	solutions.Solution of situational problems.	4			
7.	Preparation of titrated solutions. Solution of				
	situational problems.	4			
8.	Analysis of purified water, water for injection,				
0.	water for injection in ampoules.	4			
9.	· · ·				
9.	Colloquium on the topic "General and private	4			
	methods for determining the quality of	4			
10	medicines"				
10.	Application of argentometry in the				
	pharmaceutical analysis of halogen-containing	4			
	medicinal substances of inorganic nature.				
11.	The use of permanganatometry in the				
	pharmaceutical analysis of drugs of compounds	4			
	of elements of groups VI and V of the periodic	-			
	system D.I. Mendeleev.				
12.	Acid-base titration method in pharmaceutical				
	analysis. Pharmacopoeial analysis of LP	4			
	compounds of elements of group III of the	4			
	periodic system D.I. Mendeleev.				
13.	Application of complexometry in				
	pharmaceutical analysis. Pharmacopoeial				
	analysis of drugs of compounds of elements of	4			
	groups V and II of the periodic system D.I.				
	Mendeleev.				
14.	Colloquium on the topic "Application of				
1	argentometry, permanganatometry,				
	complexometry, acid-base titration methods in	4			
	pharmaceutical analysis. Compounds I-III and	4			
15.	V-VII of PS groups.				
13.	Control work on practical skills. Solution of	4			
16	situational problems. Final testing.	A			
16.	Final coursework.	4	4		
17.	Qualitative analysis of organic medicinal		4		
	substances by functional groups.				
	Pharmacopoeial analysis of medicinal				
	substances of alcohols and their derivatives".				
18.	Analysis of medicinal substances, derivatives		4		
	of aldehydes.				
19.	Pharmacopoeial analysis of ether preparations.		4		
20.	Analysis of lactones of unsaturated		4		
	polyhydroxycarboxylic acids: Ascorbic acid.				
21.	Pharmacopoeial analysis of salts of carboxylic		4		
1			1		

	acids. Tablet analysis. GF requirements for the					
	quality of tablets.					
22.	Pharmacopoeial analysis of aliphatic amino		4			
	acid preparations.					
23.	Pharmacopoeial analysis of ester preparations.		4			
24.	Practical control work on the analysis of		4			
	organic drugs of the aliphatic series. Test					
	control by section.					
25.	Analysis of medicinal products of terpene		4			
	derivatives. Solution of situational problems.					
26.	Control work "Analysis of drugs of a steroid		4			
	structure." Solution of situational problems.		•			
	Test control by section.					
27.	Pharmacopoeial analysis of drugs from the		4			
27.			4			
	group of phenols.Solution of situational					
20	problems.		4			
28.	Pharmacopoeial analysis of drugs derived from		4			
	aromatic acids.Solution of situational problems.					
29.	Pharmacopoeial analysis of drugs derived from		4			
	aromatic amino acids.					
30.	Control work "Analysis of drugs of phenols,		4			
	aromatic acids and aromatic amino acids."					
	Solution of situational problems. Test control					
	by section.					
31.	Analysis of organic medicinal substances by		4			
	functional groups.					
32.	Pharmacopoeial analysis of organic medicinal		4			
	substances: benzenesulfonamides and their					
	derivatives.					
33.	Solution of situational problems on		2			
55.	pharmacopoeial methods of analysis.		-			
34.	Final control work "Medicines of aliphatic and		4			
54.	alicyclic structure" and "Medicines of aromatic		-			
	•					
35.	structure". Control work on practical skills.			9		
55.	Analysis of dosage forms of industrial and			9		
	pharmaceutical production. The use of					
	chemical and physico-chemical methods for the					
	analysis of dosage forms.			-		
36.	Educational and research work of students			5		
	(UIRS). Theoretical substantiation of methods					
	of analysis and experimental work on the					
	analysis of dosage forms of industrial and					
	pharmaceutical production					
37.	Pharmacopoeial analysis of antibiotics of the β -			5		
	lactamide group. Solution of situational					
	problems.					
38.	Pharmacopoeial analysis of antibiotics -			5		
	aminoglycosides and antibiotics of the					
	tetracycline group. Solution of situational					
	problems.					
39.	Pharmacopoeial analysis of			5		
		I		-	1	

	nitrophenylalkylamine derivatives			
	(levomycetin). Solution of situational problems.			
40.	Educational and research work of students	5		
	(UIRS). Theoretical substantiation of methods			
	of analysis and experimental work on the			
	analysis of dosage forms containing antibiotics.			
41.	Test work and seminar on the pharmacopoeial	5		
	analysis of antibiotics. Test control by section.			
42.	Analysis of drugs derived from furan. Solution	5		
	of situational problems.			
43.	Analysis of drugs derived from benzopyran and	5		
	pyrrole. Solution of situational problems.			
44.	Analysis of drugs of pyrazole derivatives.	5		
	Solution of situational problems.	-		
45.	Analysis of medicinal products of imidazole	5		
101	and benzimidazole derivatives. Solution of	J		
	situational problems.			
46.	Test work on the topic: Medicines derivatives	5		
	of five-membered heterocycles. Test control.	-		
47.	Analysis of medicinal products of pyridine-3-		3	
	carboxylic acid derivatives. Solution of			
	situational problems.			
48.	Analysis of drugs of pyridine-4-carboxylic acid		3	
	derivatives. Solution of situational problems.		C	
49.	Test work and seminar "Alkaloids, tropane		2	
.,.	derivatives, and their synthetic analogues." Test		-	
	control by section.			
50.	Analysis of medicinal products of quinoline		4	
	derivatives.			
51.	Analysis of drugs derived from isoquinoline		4	
	and benzylisoquinoline			
52.	Control work and seminar "Analysis of		2	
	medicines of quinoline and isoquinoline".			
	Solution of situational problems. Test control			
	by section.			
53.	Pharmacopoeial analysis of medicinal		4	
	substances of pyrimidine derivatives			
	(barbiturates, uracil derivatives).			
54.	Pharmacopoeial analysis of pyrimidinothiazole		4	
	derivatives.			
55.	Control work and seminar "Analysis of drugs		2	
	of pyrimidine and pyrimidinothiazole			
	derivatives". Solution of situational problems.			
	Test control by sections.			
56.	Pharmacopoeial analysis of drugs of purine		4	
	derivatives.			
57.	Pharmacopoeial analysis of drugs of pteridine		4	
	derivatives			
58.	Pharmacopoeial analysis of drugs derived from		4	
	isoalloxazine.			
59.	Pharmacopoeial analysis of drugs of		4	

	benzodiazepine derivatives.					
60.	Test work and seminar "Analysis of medicinal products of phenothiazine derivatives".				2	
	Solution of situational problems. Test control by section.					
61.	Types of analysis error. Errors in the analysis of physicochemical and chemical methods. Methods for identifying systematic and random errors.					10
62.	Statistical processing of pharmaceutical analysis methods.					10
63.	Quality control of medicines during storage. The study of the expiration dates of medicines.					10
64.	Incompatibility of medicinal substances and methods for its elimination.					10
65.	Control work and a seminar on the section.					8
	TOTAL (total 304 AH)	70	70	64	46	48

6.2.3. Types and topics of student's individual work (SIW)

No.	Types and topics of SIW		V	olume in	AH	
p / p		5	6	7	8	9
1.	Working with literary and other sources of	10	12	12	12	5
	information on the section under study					5
2.	Doing homework provided by the discipline	10	10	10	6	6
	program					6
3.	Writing an abstract (essay, report, scientific	-	-	-	-	20
	article) on a given problem					20
4.	Preparing for a business game	2	4	4	4	-
5.	Working with electronic educational resources	4	6	6	6	5
6.	The study of material submitted for independent	8	10	10	6	
	work					-
7.	Preparation for practical work	8	10	8	6	3
8.	Preparation for examinations and tests	10	8	10	4	3
	TOTAL (total 216 AH)	52	60	60	44	42

7. Types of assessment formats for ongoing monitoring and mid-term assessment

					Assessment formats			
No . p / p	semest er numbe r	Types of control	Name of section of academic discipline	Competen ce codes	types	number of test questio ns	numbe r of test task options	
1	2	3	4		5	6	7	
1.	5	Control of student's independe	Fundamentals of pharmaceutical analysis.	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	3	6	

		nt work					
2.	5	Control of student's independe nt work	Fundamentals of pharmaceutical analysis. Preparation of titrated solutions	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	4	13
3.	5	Control of student's independe nt work	Fundamentalsofpharmaceuticalanalysis.Determinationofimpurity contentand weight lossondrying	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	2	12
4.	5	Control of student's independe nt work.	inorganic drugs. Medicines of groups I and II of the Periodic system	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	2	6
5.	5	Control of student's independe nt work.	inorganic drugs. Medicines II and III groups of the Periodic system	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	3	6
6.	5	current control	inorganic drugs.	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	5	12
7.	5	Final control	inorganic drugs.	UC-1,2 GPC-1,3,6 PC-4,7	Test	240	Comput er testing (option is formed by random samplin g)
8.	5	Control of student's independe nt work.	Medicinal products of aliphatic and alicyclic structure. "Alcohols"	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	3	12
9.	5	current control	Medicinal products of aliphatic and alicyclic structure. "Alcohols and Esters"	UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing (option is formed by random samplin g)
10.	5	Control of student's independe nt work.	Medicinal products of aliphatic and alicyclic structure. "Carbohydrates and	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	3	12

			lactones"				
11.	5	current control	Medicinal products of aliphatic and alicyclic structure. "Aldehydes, carboxylic acids, lactones"	UC-1,2 GPC-1,3,6 PC-4,7	Test	40	Comput er testing (option is formed by random samplin g)
12.		Control of student's independe nt work.	Medicinal products of aliphatic and alicyclic structure.	UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing (option is formed by random samplin g)
13.	6	Final control	Terpenes and retinols. Steroid drugs.	UC-1,2 GPC-1,3,6 PC-4,7	Test	thirty	Comput er testing (option is formed by random samplin g)
14.	6	Final control	Terpenesandretinols.Steroiddrugs.	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	3	13
15.	6	Control of student's independe nt work.	Aromatic medicines. "Phenols, quinones and their derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing (option is formed by random samplin g)
16.	6	Control of student's independe nt work.	Aromatic medicines. Aromatic amino acids	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	3	5
17.	6	current control	Aromatic medicines. Aromatic amino acids	UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing

18.	6	Control of student's independe nt work.	Aromatic medicines. "Aromatic amines". Aromatic medicines.	UC-1,2 GPC-1,3,6 PC-4,7	Test	thirty	(option is formed by random samplin g) Comput er testing (option is formed by random samplin g)
19.	0	student's independe nt work.	"Arylalkylamines, nitrophenylalkylamin es".	UC-1,2 GPC-1,3,6 PC-4,7		20	Comput er testing (option is formed by random samplin g)
20.	6	Control of student's independe nt work.	Aromatic medicines. "sulfonamides".	UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing (option is formed by random samplin g)
21.	6	Final control	Aromatic medicines.	UC-1,2 GPC-1,3,6 PC-4,7	Test	50	Comput er testing (option is formed by random samplin g)
22.	7	Control of student's independe nt work.	Antibiotics. "β- lactam antibiotics"	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	3	12
23.	7	current	Antibiotics.	UC-1,2	Test	8	Comput

24.	7	control	"Analysis of derivatives of β- lactamides and aminoglycosides"	PC-4,7	Test	20	er testing (option is formed by random samplin g) Comput
24.	,	current control	Intrapharmacy control.	UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing (option is formed by random samplin g)
25.	7	Final control	Antibiotics.	UC-1,2 GPC-1,3,6 PC-4,7	Test	40	Comput er testing (option is formed by random samplin g)
26.	7	Control of student's independe nt work.	Medicinal products of heterocyclic structure. "Furan derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Test	10	Comput er testing (option is formed by random samplin g)
27.	7	Control of student's independe nt work.	Medicinal products of heterocyclic structure. "Pyrazole and imidazole derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Test	25	Comput er testing (option is formed by random samplin g)
28.	7	current control	Medicinal products of heterocyclic structure. "Furan	UC-1,2 GPC-1,3,6 PC-4,7	Test	4	Comput er testing

29.	8	current control	derivatives" Medicinal products of heterocyclic structure. "Derivatives of furan, benzopyran, pyrazole and imidazole"	UC-1,2 GPC-1,3,6 PC-4,7	Test	40	(option is formed by random samplin g) Comput er testing (option is formed by random samplin g)
30.	8	Control of student's independe nt work.	Heterocyclic drugs "Benzylisoquinoline and phenanthrenisoquinol ine derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Test	25	Comput er testing (option is formed by random samplin g)
31.	8	current control	Heterocyclic drugs "Benzylisoquinoline and phenanthrenisoquinol ine derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	4	6
32.	8	Control of student's independe nt work.		UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing (option is formed by random samplin g)
33.		Control of student's independe nt work.	Medicinal products of heterocyclic structure "Quinoline derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Test	thirty	Comput er testing (option is formed by random samplin g)

34.	8	current	Medicinal products	UC 12	Relate	8	6
		control	of heterocyclic structure "Quinoline and tropane derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	d Poll	-	
35.	8	current control	Medicinal products of heterocyclic structure "Pyridine, piperazine derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Test	40	Comput er testing (option is formed by random samplin g)
36.	8	Control of student's independe nt work.	Medicinal products of heterocyclic structure "Pyrimidine derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing (option is formed by random samplin g)
37.	9	Control of student's independe nt work.	Medicinal products of heterocyclic structure "Purine derivatives"	UC-1,2 GPC-1,3,6 PC-4,7	Test	20	Comput er testing (option is formed by random samplin g)
38.	9	current control	Medicinal products of heterocyclic structure "Derivatives of purine and pyrimidine	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	6	6
39.	9	Control of student's independe nt work.	Metrological foundations of pharmaceutical analysis. Validation evaluation of analysis methods	UC-1,2 GPC-1,3,6 PC-4,7	busine ss game	1	50
40.	9	current control	Incompatibility Issues	UC-1,2 GPC-1,3,6 PC-4,7	Relate d Poll	5	18

8. 8. Educational, methodological and informational support for mastering the academic discipline (printed, electronic publications, the Internet and other network resources)

No.	Name according to bibliographic requirements	Number	of copies
		At the department	In the library
1.	Huynh-Ba K. Handbook of Stability Testing in Pharmaceutical Development (Regulations, Methodologies, and Best Practices) Electronic resource Springer, 2009 390 p.	Electrical version	-
2.	Jouyban A. Handbook Of Solubility Data For Pharmaceuticals Electronic resource CRC Press, 2010 554 p.	Electrical version	-
3.	Putz M. V. (Ed.) Quantum Frontiers of Atoms and Molecules[Electronic resource] Nova Science Publishers, 2011 673 p.	Electrical version	-
4.	The British Pharmacopoeia 2012. – London: The Stationery Office on Behalf of the Medicines and Healthcare Products Regulatory Agency (MHRA) [Electronic resource].	Electrical version	-
5.	TheInternationalPharmacopoeia.4thEdition[Electronic resource]WHO PharmacopoeiaLibrary.2011.	Electrical version	-
6.	The United States Pharmacopeia (USP 32) and the 27th edition of the National Formulary (NF 27) [Electronic resource]. – Washington, DC: The United States Pharmacopeial Convention. 2009 815 p.	Electrical version	-
7.	TheJapanesePharmacopoeiaSixteenthEdition[Electronicresource].–Tokyo,TheCommittee on JapanesePharmacopoeia, 2011.2326 p.	Electrical version	-
8.	Pyatigorskaya N.V. and others. Rules for the organization of production and quality control of medicinal products from plant materials: textbook Electronic resource St. Petersburg: SpecLit, 2013 367 p.	Electrical version	-
9.	Pleteneva T.V. and others. Quality control of medicines: textbook Electronic resource M. : GEOTAR-Media, 2015 560 p Access mode: EBSStudent Advisor	Electrical version	-
10.	Order of the Ministry of Health of Russia dated October 26, 2015 N 751n "On approval of the rules for the manufacture and dispensing of drugs for medical use by pharmacy organizations, individual entrepreneurs licensed for pharmaceutical activities" Electronic resource	Electrical version	-

8.1. Key literature references

		Number	of copies
р / no.	no.		at the departme nt
1.	Belikov VG Synthetic and natural medicines : a brief guide / VG Belikov M.: Higher School, 1993 720 p.	2	-
2.	Laboratory work in pharmaceutical chemistry: Textbook / V. G. Belikov, I. Ya. Kul, G. I. Lukyanchikova, A. S. Saushkina and S. G. Tiraspolskaya; ed. E. N. Vergeichik and E. V. Kompantseva; Ed. organization Pyatigorsk State Pharmaceutical Academy 2nd ed., revised. and additional Pyatigorsk: B.I., 2003. (2003) - 342 p.	203	-
3.	Guide to laboratory studies in pharmaceutical chemistry : textbook / EN Aksenova and OP Andrianova ; ed. A. P. Arzamastsev 3rd ed., revised. and additional M .: Medicine, 2004. (2004) - 384 p.	20	-
4.	State Pharmacopoeia of the USSR: Issue 1: General methods of analysis 11th ed M.: Medicine, 1987 336 p.	32	-
5.	State Pharmacopoeia of the USSR: Issue. 2: General methods of analysis. Medicinal plant material 11th ed M.: Medicine, 2000 400 p.	5	-
6.	State Pharmacopoeia of the USSR: Issue. 2. General methods of analysis. Medicinal plant materials / redol. M. D. Mashkovsky, E. A. Babayan, A. N. Oboymakova, V. M. Bulaev, and V. A. Severtsev; Ed. organization Ministry of Health of the USSR 11th ed M.: Medicine, 1990 400 p.	25	-
7.	State Pharmacopoeia of the USSR 10th ed M.: Medicine, 1968 1080 p.	1	-
8.	Glushchenko N. N. Pharmaceutical chemistry: textbook / N. N. Glushchenko, T. V. Pleteneva and V. A. Popkov M. : Academy, 2004. (2004) - 384 s	118	-
9.	Soldatenkov A. T. Fundamentals of organic chemistry of medicinal substances / A. T. Soldatenkov, N. M. Kolyadina and I. V. Shendrik 3rd ed M.: Mir; M. : BINOM. Knowledge Laboratory, 2007. (2007) - 191 p.	1	-
10.	Granik V.G. Fundamentals of Medical Chemistry: Textbook / V.G. Granik M. : Vuzovskaya kniga, 2001. (2001) - 384 p.	1	-
11.	Slesarev V.I. Chemistry. Fundamentals of Living Chemistry: A Textbook for High Schools / V.I. Slesarev St. Petersburg. : Himizdat, 2000 768 p.	1	-

8.3. Electronic educational resources for teaching academic subjects8.3.1. Internal Electronic Library System of the University (IELSU)

Name of the	Brief description (content)	Access conditions	Number of users
electronic resource			
Internal	Proceedings of the teaching	J 1	Not limited
Electronic Library	staff of the Academy:	on the Internet,	

System of the University (IELSU)textbooks and text monographs, col scientific papers, articles, dissertat abstracts of dissertat patents.	login and password entific
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8.4.2. Electronic educational	resources acquired b	by the University

No. p / p	Name of the electronic	Brief description (content)	Access conditions	Number of users
1.	resource DB "Medicine. Healthcare (HPE)" (EBS "Student Consultant")	Educational literature + additional materials (audio, video, interactive materials, test tasks) for higher medical and pharmaceutical education	from any computer on the Internet, using an individual login and password	Not limited
2.	Electronic library system «BookUp»	Educational and scientific medical literature of Russian publishing houses, incl. translations of foreign publications	from university computers; from any computer on the Internet using an individual login and password Subscribed editions are available for reading.	Not limited
3.	Electronic Medical Library "Doctor's Consultant"	National guidelines in all areas of medicine, clinical guidelines, textbooks, monographs, atlases, pharmaceutical reference books, audio and video materials, ICD-10 and ATC, recent publications in foreign journals with brief annotations in Russian	from any computer on the Internet, using an individual login and password	Not limited
4.	Domestic electronic periodicals	Medical periodicals	from the university computers on the platform of the SCIENTIFIC electronic library eLIBRARY.RU Subscribed editions are available for reading.	Not limited
5.	DB Medline	Foreign full-text	from university	Not limited

			2	
	Complete	database of articles	computers; from any	
		from scientific	computer on the	
		periodicals and	Internet, using an	
		collections of medical	individual login and	
		and natural science	password	
		topics	-	
6.	Springer	Full-text scientific	from university	Not limited
	Electronic	publications (journals,	computers	
	Collection	books, articles,	-	
		scientific protocols,		
		conference materials,		
		etc.) in the natural		
		sciences, medical		
		sciences and the		
		humanities		
7.	Electronic	Books and periodicals	from university	Not limited
	collection	of the publishing house	computers	
	"Freedom" on the	"Elsevier" in the natural	1	
	Science Direct	sciences, medicine and		
	platform	humanities		
8.	DB Scopus	International Science	from university	Not limited
	-	Citation Abstract	computers	
		Database	1	
9.	DB Web of	International Science	from university	Not limited
	Science Core	Citation Abstract	computers; from any	
	Collection	Database	computer on the	
			Internet, using an	
			individual login and	
			password	
10.	DB Questel	Questel Patent	from university	Not limited
	Orbit	Database	computers	
			1	

8.4.3 Open access resources

No . p / p	Name of the electronic resource	Brief description (content)	Access conditions
1	Federal Electronic Medical Library (FEMB)	Includes electronic analogues of printed publications and original electronic publications that have no analogues recorded on other media (dissertations, abstracts, books, magazines, etc.).	from any computer on the Internet
2.	Scientific electronic library eLIBRARY.RU	The largest Russian information portal in the field of science, technology, medicine and education, containing abstracts and full texts of scientific articles and	from any computer on the Internet.

		publications.	
3.	Scientific electronic library of open access CyberLeninka	Full texts of scientific articles with annotations published in scientific journals in Russia and neighboring countries.	from any computer on the Internet
4.	Russian State Library (RSL)	Abstracts for which there are copyright agreements with permission for their open publication	from any computer on the Internet
5.	Reference and legal system "Consultant Plus"	Federal and regional legislation, judicial practice, financial advice, legislative comments, etc.	from any computer on the Internet

9. Material and technical support for mastering an academic discipline

- 9.1. List of premises for classroom activities for the discipline
- 1. Lecture room classroom
- 2. Classrooms for practical classes, seminars, intermediate certification in the same place.
- 3. Scientific laboratories for conducting practical exercises and laboratory workshops
- 9.2. List of equipment for classroom activities for the discipline
- 1. Multimedia complex (laptop, projector, screen)

overhead multimedia projector Vega Focus 400 GLS (101042910) - 1 pc.,

multimedia projector BenQ NB 6110 (101042596) - 1 pc.,

Notebook HP Pavilion Notebook 15-ab234ur (101341033) - 1 pc.;

Laptop ASUS Z99H (101041277) – 1 pc.;

screen - 2 pcs.

- 2. A set of electronic presentations on lecture topics.
- 3. Instruments and equipment:

spectrophotometer UNICO 1200 (101043138) – 1 pc.,

RN-meter millivoltmeter RN-150M (101043000) - 1 pc.,

Liquid chromatograph LC-10AVP (101043413) – 1 pc.,

spectrophotometer Specord S100 Bio (101043137) - 1 pc.

Fourier IR spectrophotometer IRAffinity-1S (101241054) – 1 pc.,

Rotary evaporator LEKI RE 52AA (101041294) - 1 pc.,

Scales EK-400N (101041435) - 1 pc.,

Spectrophotometer UV-1800 scanning 2-beam (101240610) - 1 pc.,

Analytical balance ATX-224 (101240947) - 1 pc.,

Infrared Fourier spectrophotometer (101040380) - 1 pc.,

Water purification system MILLIPORE Elix-3 (101041324) – 1 pc., AA-7000F atomic

absorption spectrophotometer (101340100) - 1 pc., Liquid chromatograph LC-20AD

Prominence (101240611) – 1 pc., Chromatographic column C 18 (101040683)) - 1 PC.

Computer Pentium 4 (101041937) - 1 pc.,

Printer HP LJ 1010 laser (101042738) - 1 pc.,

Refrigerator 2-chamber Atlant XM-4012-000 (101065445) - 1 pc.

Exhaust cabinet 1460*700*2100 (101260844) - 1 pc.,

Exhaust cabinet 1800*700*2100 (101260842, 101260843) - 2 pcs.,

fume hood 1460*700*2100 (101261000) – 1 pc.,

information stand (101261001 and 101260845) - 2 pcs.

information stand (101261002, 101261003) - 2 pcs.

9.3. A set of licensed and freely distributed software, including domestic production

Ite m no.	Software	number of licenses	Type of software	Manufactur er	Number in the unified register of Russian software	Contract No. and date
1	Wtware	100	Thin Client Operating System	Kovalev Andrey Alexandrovi ch	1960	2471/05-18 from 28.05.2018
2	MyOffice is Standard. A corporate user license for educational organizations, with no expiration date, with the right to receive updates for 1 year.	220	Office Application	LLC "NEW CLOUD TECHNOL OGIES"	283	without limitation, with the right to receive updates for 1 year.
3	LibreOffice		Office Application	The Document Foundation	Freely distributed software	
4	Windows 10 Education	700	Operating systems	Microsoft	Azure Dev Tools for Teaching Subscripti on	
5	Yandex. Browser		Browser	«Yandex»	3722	
6	Subscription to MS Office Pro for 170 PCs for FGBOU VO "PIMU" of the Ministry of Health of Russia	170	Office Applica- tion	Microsoft		23618/HN10 030 LLC "Softline Trade" from 04.12.2020

10. List of changes to the working program (to be filled out by the template)

Federal State Budgetary Educational Institution of Higher Education "Privolzhsky Research Medical University" Ministry of Health of the Russian Federation (FSBEI HE "PRMU" of the Ministry of Health of Russia)

Department of *Name of the department*

CHANGE REGISTRATION SHEET

working program for the academic discipline *NAME OF THE ACADEMIC DISCIPLINE*

Field of study / specialty / scientific specialty:

Training profile: _____

(name) - for master's degree programs

Mode of study: _____

full-time/mixed attendance mode/extramural

Position	Number and name of the program section	Contents of the changes made	Effective date of the changes	Contributor's signature
1				

Approved at the department meeting	
Protocol Noof	20

Head	of the	De	partment
Head	of the	De	partment

department name, academic title

signature

print name

(code, name)