Head (deputy head)					
Federal Accreditation Service					
signature	initials, last name				
Annex to the accredita	tion certificate				
№ <u>RA.RU.21ФВ02</u>	2				
from « »	20				
on <u>243</u> pages, page <u>1</u>					

SCOPE ACCREDITATION OF TESTING CENTRE

FEDERAL STATE BUDGETARY INSTITUTION «THE RUSSIAN STATE CENTER FOR ANIMAL FEED AND DRUG STANDARDIZATION AND QUALITY» Address: Zvenigorodskoye shosse 5, 123022, Moscow, Russia

143511, Russia, Moscow region, Istra district, Manikhino laboratory facility 1, laboratory facility 2, laboratory facility 4, office building 2

N p/p	Documents setting rules and methods for research (tests), measurements	Name of object	OKPD 2 CODE	EAEU CN of FEA	Defined characteristic (indicator)	Range of measurement
1	2	3	4	5	6	7
		Zvenigorodskoye sl	nosse 5, 123022, N	Moscow		
1	GOST 34136	Food raw materials and food	10.11	0201-0205	spiramycin	(2 - 320) mcg/kg
		products: meat and meat products	10.12		erythromycin	(10 - 320) mcg/kg
			10.13		tilmicosin	(1 - 160) mcg/kg
					tylosin	(1 - 160) mcg/kg
					tylvalosin	(5 - 160) mcg/kg
				_	valnemulin	(1 - 160) mcg/kg
				_	tiamulin	(1 - 160) mcg/kg
				_	tulathromycin	(1 - 160) mcg/kg
					clarithromycin	(1 - 160) mcg/kg
					lincomycin	(1 - 160) mcg/kg
					clindamycin	(1 - 160) mcg/kg
					pirlimycin	(1 - 160) mcg/kg
		Food raw materials and food	10.11	0206-0208	spiramycin	(20 - 3200) mcg/kg
		products: offal	10.12		erythromycin	(10 - 320) mcg/kg
			10.13		tilmicosin	(10 - 1600) mcg/kg
				_	tylosin	(1 - 160) mcg/kg
				_	tylvalosin	(5 - 160) mcg/kg
					valnemulin	(5 - 800) mcg/kg
					tiamulin	(10 - 1600) mcg/kg
				_	tulathromycin	(20 - 3200) mcg/kg
				_	clarithromycin	(1-160) mcg/kg
					lincomycin	(15 - 2400) mcg/kg
					clindamycin	(15 - 2400) mcg/kg
					pirlimycin	(10 - 1600) mcg/kg

1	2	3	4	5	6	7 7
		Food raw materials and food	10.51	0401-0406	spiramycin	(2 - 320) mcg/kg
		products: milk and dairy products	10.52		erythromycin	(10 - 320) mcg/kg
					tilmicosin	(1 - 160) mcg/kg
					tylosin	(5 - 160) mcg/kg
					tylvalosin	(1 - 160) mcg/kg
					valnemulin	(20 - 160) mcg/kg
					tiamulin	(1 - 160) mcg/kg
					tulathromycin	(1 - 160) mcg/kg
					clarithromycin	(1 - 160) mcg/kg
					lincomycin	(1,5 - 240) mcg/kg
					clindamycin	(1 - 160) mcg/kg
					pirlimycin	(1 - 160) mcg/kg
2	GOST 34140	Food products, food raw materials	01.11	1001-1008,	mycophenolic acid	(20 - 2000) mcg/kg
		for cereals, fodder, food raw	01.12 10.91	1101-1108, 1201-1207	15-acetyl deoxynivalenol	(100 - 2000) mcg/kg
		materials for cereals and oilseeds,	10.91	1201-1207	3-acetyl deoxynivalenol	(100 - 2000) mcg/kg
		compound feed	10.52		agroclavine	(10 - 1000) mcg/kg
					alternariol	(10 - 2000) mcg/kg
					alternariolamethyl ether	(20 - 2000) mcg/kg
					aflatoxin g1	(1 - 200) mcg/kg
					aflatoxin g2	(1 - 200) mcg/kg
					aflatoxin B1	(1 - 200) mcg/kg
					aflatoxin B2	(1 - 200) mcg/kg
					beauvericin	(50 - 10000) mcg/kg
					wortmannin	(20 - 2000) mcg/kg
					gliotoxin	(100 - 2000) mcg/kg
					griseofulvin	(20 - 2000) mcg/kg
					deoxynivalenol	(100 - 10000) mcg/kg
					deoxynivalenol-3-glucoside	(100 - 2000) mcg/kg
					deepoxy-deoxynivalenol	(200 - 2000) mcg/kg
					diacetoxyscirpenol	(10 - 2000) mcg/kg
					zearalenone	(20 - 4000) mcg/kg
					kojic acid	(10000 - 20000) mcg/kg
1					meleagrine	(20 - 2000) mcg/kg
					moniliformin	(20 - 2000) mcg/kg
					neosolaniol	(10 - 2000) mcg/kg
					nivalenol	(100 - 10000) mcg/kg
					NT-2 toxin	(10 - 2000) mcg/kg
					ochratoxin b	(1 - 200) mcg/kg
					ochratoxin a	(1 - 200) mcg/kg
					paxillin	(20 - 200) mcg/kg

1	2	3	4	5	6	243 page
1	<u> </u>	3	4	5		/
					patulin	(1000 - 2000) mcg/kg
					penicillin acid	(20 - 2000) mcg/kg
					roquefortine c	(10 - 2000) mcg/kg
					roridin a	(100 - 2000) mcg/kg
					stachybotrys lactam	(10 - 2000) mcg/kg
					sterigmatocystin	(10 - 2000) mcg/kg
					т-2 tetraol	(100 - 2000) mcg/kg
					т-2 toxin	(10 - 2000) mcg/ml
					т-2 triol	(20 - 2000) mcg/kg
					tentoxin	(20 - 2000) mcg/kg
					tenuazonic acid	(20 - 2000) mcg/kg
					fusarenon x	(500 - 10000) mcg/kg
					fusarium acid	(100 - 20000) mcg/kg
					fumagillin	(100 - 2000) mcg/kg
					Fumonisin b3	(100 - 10000) mcg/kg
					Fumonisin в1	(100 - 20000) mcg/kg
					Fumonisin в2	(100 - 20000) mcg/kg
					cyclopiazonic acid	(20 - 2000) mcg/kg
					citreoviridin	(100 - 2000) mcg/kg
					citrinin	(50 - 10000) mcg/kg
					ergocornin	(20 - 2000) mcg/kg
3	GOST 33978	Unprocessed food products: meat	10.11	0201-0208,	6-propyl-2-thiouracil	(2 - 30) mcg/kg
		(including poultry), offal (liver);	10.12	1001-1008,	6- methyl -2-thiouracil	(2 - 30) mcg/kg
		compound feed; animal urine	10.13	1101-1108	2-thiouracil	(2 - 30) mcg/kg
			01.11		6-phenyl -2-thiouracil	(2 - 30) mcg/kg
			01.12 10.91			
			10.91		2-mercaptobenzimidazole	(0,4 - 30) mcg/kg
4	GOST 33934	Meat, including poultry, offal, meat	10.11	0201-0208		
	33731	and meat products	10.12			
		and meat products	10.13		zinc bacitracin	(0,02 - 100) mg/kg
5	METHODOLOGICAL GUIDELINES	Fish	10.20	0301-0305	netobimin	(5 - 1000) mcg/kg
	A-1/044				albendazole	(1 - 1000) mcg/kg
	Methodical guidelines for the arbitration				albendazole-2-	
	determination of anthelminthics in fish				aminosulfone	(1 - 1000) mcg/kg
	by a method of high-performance liquid				albendazole	(1 - 1000) mcg/kg
	chromatography with a mass				albendazole	(1 - 1000) mcg/kg
	spectrometer detector				aminomebendazole	(1 - 1000) mcg/kg
					aminooxybendazole	(1 - 1000) mcg/kg
					aminotriclabendazole	(1 - 1000) mcg/kg
					anniourciauciluazuie	(1 - 1000) IIICg/kg

1	2	3	4	5	6	7 243 page
					aminoflubendazole	(1 - 1000) mcg/kg
					hydroxymebendazole	(1 - 1000) mcg/kg
					hydroxytyabendazole	(1 - 1000) mcg/kg
					cambendazole	(1 - 1000) mcg/kg
					ketotriclabendazole	(1 - 1000) mcg/kg
					closantel	(1 - 1000) mcg/kg
					clorsulon	(1 - 1000) mcg/kg
					levamisole	(1 - 1000) mcg/kg
					mebendazole	(1 - 1000) mcg/kg
					morantel	(1 - 1000) mcg/kg
					netobimin	(1 - 1000) mcg/kg
					niclosamide	(1 - 1000) mcg/kg
					nitroxynil	(1 - 1000) mcg/kg
					oxibendazole	(1 - 1000) mcg/kg
					oxyclozanide	(1 - 1000) mcg/kg
					oxfendazole	(1 - 1000) mcg/kg
					oxfendazole	(1 - 1000) mcg/kg
					parbendazole	(1 - 1000) mcg/kg
					pyrantel	(1 - 1000) mcg/kg
					Praziquantel	(1 - 1000) mcg/kg
					rafoxanide	(1 - 1000) mcg/kg
					thiabendazole	(1 - 1000) mcg/kg
					triclabendazole	(1 - 1000) mcg/kg
					triclabendazole	(1 - 1000) mcg/kg
					triclabendazole	(1 - 1000) mcg/kg
					febantel	(1 - 1000) mcg/kg
					fenbendazole	(1 - 1000) mcg/kg
					flubendazole	(1 - 1000) mcg/kg
6	METHODOLOGICAL GUIDELINES	Food raw materials and food	10.11	0201-0208,	bacitracin a	(5-500) mcg/kg
	A-1/045	products: meat and meat products,	10.12 10.13	0401-0408	bacitracin в	(1-100) mcg/kg
	Methodical guidelines for the arbitrage	offal (liver, kidneys), eggs, milk,	01.47		colistin a	(5-500) mcg/kg
	determination of residual polypeptide	dairy products,	10.51		colistin B	(3,75-375) mcg/kg
	antibiotics in animal products by high-		10.52		polymyxin в1	(5-500) mcg/kg
	performance liquid chromatography with a mass spectrometer detector				polymyxin в2	(2,5-250) mcg/kg
	a mass spectrometer detector				virginiamycin s1	(5-500) mcg/kg
					virginiamycin m1	(5-500) mcg/kg
					actinomycin d	(5-500) mcg/kg
L			10.11	0001 000	novobiocin	(5-500) mcg/kg
7	GOST 33971	Food raw materials and food	10.11	0201-0208	quinoxaline-2-carboxylic	(0,5-8) mcg/kg
		products: meat, meat products,	10.12		acid	(0,0 0)

1	2	3	4	5	6	7 243 page
		offal (liver, kidneys)	10.13		3-methylquinoxaline-2- carboxylic acid	(0,5-8) mcg/kg
					1,4-bisdeoxycarbadox	(0,5-8) mcg/kg
8	GOST 34137	Food raw materials and food	10.11	0201-0208,	cefacetrile	(5-500) mcg/kg
		products: meat (all types of	10.12	0401-0408,	cephalexin	(5-500) mcg/kg
		animals), including poultry, offal,	10.13 01.47	1601-1602	cephalonium	(5-500) mcg/kg
		meat products, semi-finished	10.51		cefoperazone	(5-500) mcg/kg
		products, eggs and their products,	10.52		cefkinom	(5-500) mcg/kg
		milk, dairy products			cephapirin	(5-500) mcg/kg
					desacetyl cephapirin	(5-500) mcg/kg
					cefadroxil	(5-500) mcg/kg
					cefsulodin	(5-500) mcg/kg
					cefotaxime	(5-500) mcg/kg
					ceftibuten	(5-500) mcg/kg
					cefpodoxime	(5-500) mcg/kg
					cefpirome	(5-500) mcg/kg
					cefotiam	(5-500) mcg/kg
					cefaclor	(5-500) mcg/kg
					cefetamet	(5-500) mcg/kg
					cefepime	(5-500) mcg/kg
					ceftiofur and metabolites (desfuroil	
					ceftiofur, desfuroil ceftiofur	(30-3000) mcg/kg
					cysteine disulfide)	
9	GOST 34139	Food raw materials and food	10.11 10.12	0201-0205, 0401-0408	azaperol	(1-500) mcg/kg
		products: meat (all kind of animal),	10.12	0401-0408	azaperone	(1-500) mcg/kg
		including poultry meat, meat	10.13		propionyl-promazine	(10-500) mcg/kg
		products, milk, Dairy products	10.52		haloperidol	(1-500) mcg/kg
					fluphenazine	(1-500) mcg/kg
					carazolol	(1-500) mcg/kg
					acepromazine	(1-500) mcg/kg
					xylazine	(1-500) mcg/kg
					triflupromazine	(1-500) mcg/kg
					detomidine	(1-500) mcg/kg
					medetomidine	(1-500) mcg/kg
					chlorpromazine	(1-500) mcg/kg
					meperidine	(1-500) mcg/kg
					diazepam	(1-500) mcg/kg
					metoprolol	(1-500) mcg/kg
					romifidine	(1-500) mcg/kg

1	2	3	4	5	6	7
					promazine	(10-500) mcg/kg
		Food raw materials and food	10.11	0206-0208	azaperol	(1-500) mcg/kg
		products: offal	10.12		azaperone	(1-500) mcg/kg
			10.13		propionyl-promazine	(10-500) mcg/kg
					haloperidol	(1-500) mcg/kg
					fluphenazine	(1-500) mcg/kg
					carazolol	(1-500) mcg/kg
					acepromazine	(1-500) mcg/kg
					xylazine	(1-500) mcg/kg
					triflupromazine	(1-500) mcg/kg
					detomidine	(1-500) mcg/kg
					medetomidine	(1-500) mcg/kg
					chlorpromazine	(10-500) mcg/kg
					meperidine	(1-500) mcg/kg
					diazepam	(1-500) mcg/kg
					metoprolol	(1-500) mcg/kg
					romifidine	(10-500) mcg/kg
					promazine	(10-500) mcg/kg
10	METHODOLOGICAL GUIDELINES	Food products: meat (all kind of	10.11	0201-0208,	Mass fraction:	
	A-1/049	animal), including poultry meat,	10.12	0301-0305,	2-hydroxymethyl-1-methyl-	(1-1000) mcg/kg
	Methodical guidelines for the	offal, meat products, fish, fish	10.13 10.20	0401-0408, 0409000000,	5-nitroimidazole	
	comprehensive detection of broad	products, eggs and processed egg	10.20	1001-1008,	sulfachloropyridazine	(1-1000) mcg/kg
	spectrum xenobiotics in food and feed	product, honey, milk, Dairy	10.52	1101-1106,	Sulfaethoxypyridazine	(1-1000) mcg/kg
	based on ultra-high performance liquid	products, feed	01.11	1201-1207	cephapirin	(5-1000) mcg/kg
	chromatography with high resolution		01.12		cefotiam	(5-1000) mcg/kg
	mass spectrometric detection		10.91		ceftibuten	(5-1000) mcg/kg
			10.92		desfuroilceftiofur	(5-1000) mcg/kg
					2-aminoflubendazole	(1-1000) mcg/kg
					aminotriclabendazole	(1-1000) mcg/kg
					netobimin	(5-1000) mcg/kg
					sulfachloropyridazine	(1-1000) mcg/kg
					sulfadoxine	(1-1000) mcg/kg
					sulfasalazine	(1-1000) mcg/kg
					pefloxacin	(1-1000) mcg/kg
					dapsone	(1-1000) mcg/kg
					nafcillin	(1-1000) mcg/kg
					thiamphenicol	(1-1000) mcg/kg
					decoquinate	(1-1000) mcg/kg
					oxyphenbutazone	(1-1000) mcg/kg
					3-methylquinoxaline-2-	(1-1000) mcg/kg

1		2	A			243 page
	2	3	4	5	6	7
					carboxylic acid	
					quinoxaline-2-carboxylic	(1-1000) mcg/kg
					acid	
					tobramycin	(1-1000) mcg/kg
					clarithromycin	(1-1000) mcg/kg
					pirlimycin	(1-1000) mcg/kg
					tulathromycin	(1-1000) mcg/kg
					2-mercaptobenzimidazole	(1-1000) mcg/kg
					clenpenterol	(1-1000) mcg/kg
					tulobuterol	(1-1000) mcg/kg
					a-zeranol	(1-1000) mcg/kg
					17-a-trenbolone	(1-1000) mcg/kg
					β-nortestosterone	(1-1000) mcg/kg
					methylboldenone	(1-1000) mcg/kg
					methyltestosterone	(1-1000) mcg/kg
					ampicillin	(1-1000) mcg/kg
					metronidazole	(1-1000) mcg/kg
					h dimetridazole	(1-1000) mcg/kg
					penicillin G	(1-1000) mcg/kg
					penicillin V	(1-1000) mcg/kg
					sulfamethazine	(1-1000) mcg/kg
					sulfamethox-pyridoxine	(1-1000) mcg/kg
					sulfonamide	(1-1000) mcg/kg
					sulfaquinoxaline	(1-1000) mcg/kg
					tinidazole	(1-1000) mcg/kg
					florfenicol	(1-1000) mcg/kg
					chloramphenicol	(0,2-1000) mcg/kg
					leucocrystal violet	(1-1000) mcg/kg
					brilliant green	(1-1000) mcg/kg
					leukomalachite green	(1-1000) mcg/kg
1					cefadroxil	(5-1000) mcg/kg
1					cefaclor	(5-1000) mcg/kg
1					cefepime	(5-1000) mcg/kg
					cefetamet	(5-1000) mcg/kg
1					cefotaxime	(5-1000) mcg/kg
					cefpirome	(5-1000) mcg/kg
					cefpodoxime	(5-1000) mcg/kg
					cefsulodin	(5-1000) mcg/kg
					flumequine	(1-1000) mcg/kg
					albendazole	(1-1000) mcg/kg

1	2	3	4	5	6	243 page 7
					closantel	(1-1000) mcg/kg
					clorsulon	(1-1000) mcg/kg
					niclosamide	(1-1000) mcg/kg
					oxibendazole	(1-1000) mcg/kg
					oxfendazole	(1-1000) mcg/kg
					febantel	(1-1000) mcg/kg
					flubendazole	(1-1000) mcg/kg
					detomidine	(1-1000) mcg/kg
					carazolol	(1-1000) mcg/kg
					medetomidine	(1-1000) mcg/kg
					sotalol	(1-1000) mcg/kg
					oxytetracycline	(1-1000) mcg/kg
					chlortetracycline	(1-1000) mcg/kg
					dinitrocarbonylide	(1-1000) mcg/kg
					lasalocid a	(1-1000) mcg/kg
					tinidazole	(1-1000) mcg/kg
					nicarbazin	(1-1000) mcg/kg
					4-acetamidoantipirin	(1-1000) mcg/kg
					vedaprofen	(1-1000) mcg/kg
					deoxycarbadox	(1-1000) mcg/kg
					hygromycin b	(1-1000) mcg/kg
					dihydrostreptomycin	(1-1000) mcg/kg
					kanamycin a	(1-1000) mcg/kg
					clindamycin	(1-1000) mcg/kg
					spiramycin	(1-1000) mcg/kg
					tilmicosin	(1-1000) mcg/kg
					tylosin	(1-1000) mcg/kg
					erythromycin	(1-1000) mcg/kg
					brombuterol	(1-1000) mcg/kg
					hydroxy-methylclenbuterol	(1-1000) mcg/kg
					zilpaterol	(1-1000) mcg/kg
					isoxsuprine	(1-1000) mcg/kg
					clenbuterol	(1-1000) mcg/kg
					ritodrine	(1-1000) mcg/kg
					terbutaline	(1-1000) mcg/kg
					fenoterol	(1-1000) mcg/kg
					testosterone	(1-1000) mcg/kg
					hexestrol	(1-1000) mcg/kg
					6α-methylprednisolone	(1-1000) mcg/kg

1	2	3	4	5	6	243 page 7
					prednisolone	(1-1000) mcg/kg
					amikacin	(50-1000) mcg/kg
					paromomycinum	(50-1000) mcg/kg
					sulfathiazole	(1-1000) mcg/kg
					sulfamonomethoxine	(1-1000) mcg/kg
					cephaloridine	(5-1000) mcg/kg
					cephalothin	(5-1000) mcg/kg
					cefradine	(5-1000) mcg/kg
					cefazolin	(5-1000) mcg/kg
					cefamandole	(5-1000) mcg/kg
					cefoxitin	(5-1000) mcg/kg
					ceftriaxone	(5-1000) mcg/kg
					ceftazidime	(5-1000) mcg/kg
					cefixime	(5-1000) mcg/kg
					cefodizime	(5-1000) mcg/kg
					cefkinom	(5-1000) mcg/kg
					cefonicid	(5-1000) mcg/kg
					ceforanide	(5-1000) mcg/kg
					ceftizoxime	(5-1000) mcg/kg
					cefoselis	(5-1000) mcg/kg
					cefprozil	(5-1000) mcg/kg
					cefdinir	(5-1000) mcg/kg
					cephalonium	(5-1000) mcg/kg
					piperacillin	(5-1000) mcg/kg
					ceftizoxime	(5-1000) mcg/kg
					cephatrizine	(5-1000) mcg/kg
					cefazedone	(5-1000) mcg/kg
					ceftezole	(5-1000) mcg/kg
					cefotetan	(5-1000) mcg/kg
					cefbuperazone	(5-1000) mcg/kg
					cefminox	(5-1000) mcg/kg
					cefcapene	(5-1000) mcg/kg
					cefdaloxime	(5-1000) mcg/kg
					cefditoren	(5-1000) mcg/kg
					cefpimizole	(5-1000) mcg/kg
					cefteram	(5-1000) mcg/kg
					cefozopran	(5-1000) mcg/kg
					desacetylcephalothin	(5-1000) mcg/kg
					cypermethrin	(5-1000) mcg/kg

						243 pages
1	2	3	4	5	6	7
					bifenthrin	(5-1000) mcg/kg
					permethrin	(5-1000) mcg/kg
					deltamethrin	(5-1000) mcg/kg
					diazinon	(5-1000) mcg/kg
					haloperidol	(5-1000) mcg/kg
					dactinomycin	(5-1000) mcg/kg
					gramicidin a	(5-1000) mcg/kg
					gramicidin c	(5-1000) mcg/kg
					polymyxin B	(5-1000) mcg/kg
					colistin	(5-1000) mcg/kg
					vancomycin	(5-1000) mcg/kg
					bacitracin	(5-1000) mcg/kg
					3-amino-2-oxazolidinone	(1-1000) mcg/kg
					crystal violet	(1-1000) mcg/kg
					pefloxacin	(5-1000) mcg/kg
					sulfamethoxazole	(1-1000) mcg/kg
					sulfamethizole	(1-1000) mcg/kg
					florfenicol	(1-1000) mcg/kg
					valnemulin	(5-1000) mcg/kg
					tiamulin	(5-1000) mcg/kg
					narasin	(1-1000) mcg/kg
					oxacillin	(5-1000) mcg/kg
11	METHODOLOGICAL GUIDELINES A	Feed, feed raw materials, Food	01.11	0901-0909,	Mass fraction:	
	1/050	products of vegetable products	01.12	1001-1008,	15-acetyldeoxynivalenol	(100-2000) mcg/kg
	Methodical guidelines for the		01.19.1	1101-1108, 1201-1212	15-monoacetoxyscirpenol	(20-2000) mcg/kg
	multicomponent determination of			1201-1212	3-acetyldeoxynivalenol	(100-2000) mcg/kg
	mycotoxins in feed, feed raw material				3-nitropropionic acid	(1000-10000) mcg/kg
	and food products by high-performance				Fujimycin FK 506	(50-5000) mcg/kg
	liquid chromatography with mass				HC-toxin	(50-5000) mcg/kg
	spectrometric detection method				NG 012	(1000-10000) mcg/kg
					A 23187	(50-5000) mcg/kg
					Averantin	(1000-10000) mcg/kg
					Agistatin E	(1000-10000) mcg/kg
					Agroclavine	(10-1000) mcg/kg
					Alamethicin F50	(1000-10000) mcg/kg
					Altenuene	(50-5000) mcg/kg
					Altenusin	(50-5000) mcg/kg
					Alternatiol	(10-1000) mcg/kg
					Alternatiolmethylether	(20-2000) mcg/kg
					Altersolanol	(50-5000) mcg/kg

1	2	3	4	5	6	7 7
					Anisomycin	(50-5000) mcg/kg
					Anomalin A	(1000-10000) mcg/kg
					Apicidin	(50-5000) mcg/kg
					Ascomycin	(50-5000) mcg/kg
					Aspergillimide	(50-5000) mcg/kg
					Aspirochlorine	(50-5000) mcg/kg
					Aspartic acid	(1000-10000) mcg/kg
					Atpenin	(50-5000) mcg/kg
					Austdiol	(500-10000) mcg/kg
					Aflatoxin B1	(1-200) mcg/kg
					Aflatoxin B2	(1-200) mcg/kg
					Aflatoxin G1	(1-200) mcg/kg
					Aflatoxin G2	(1-200) mcg/kg
					Aflatoxin M1	(1-200) mcg/kg
					Aflatoxin M2	(1-200) mcg/kg
					Beauvericin	(50-10000) mcg/kg
					Butyrolactone II	(1000-10000) mcg/kg
					Verrukofortin	(1000-10000) mcg/kg
					Viomellein	(1000-10000) mcg/kg
					Viridicatin	(500-10000) mcg/kg
					Wortmannin	(20-2000) mcg/kg
					Geldanamycin	(500-10000) mcg/kg
					Helvolic acid	(1000-10000) mcg/kg
					Geodin	(500-10000) mcg/kg
					Gibberellic acid	(1000-10000) mcg/kg
					Hypothemycin	(1000-10000) mcg/kg
					Gliotoxin	(100-2000) mcg/kg
					Griseofulvin	(20-2000) mcg/kg
					Daunorubicin	(1000-10000) mcg/kg
					Deoxynivalenol	(100-10000) mcg/kg
					Deoxynivalenol-3-	
					glycoside	(100-2000) mcg/kg
					Dechlorogriseofulvin	(20-2000) mcg/kg
					Deepoxydeoxynivalenol	(20-2000) mcg/kg
					Diacetoxyscirpenol	(10-2000) mcg/kg
					Dynactin	(100-10000) mcg/kg
					Zearalenone	(20-4000) mcg/kg
					Izofusidienol A	(200-10000) mcg/kg
					Irgasan	(200-10000) mcg/kg
					Calphostin C	(200-10000) mcg/kg

1	2	3	4	5	6	243 pages, 7
					Kojic acid	(10000-20000) mcg/kg
					Cochliodinol	(50-5000) mcg/kg
					Xantomegnin	(200-10000) mcg/kg
					Curvularin	(200-10000) mcg/kg
					Macrosporin	(200-10000) mcg/kg
					Malformine C	(50-5000) mcg/kg
					Markfortin C	(50-5000) mcg/kg
					Meleagrine	(20-2000) mcg/kg
					Mevinolin	(50-5000) mcg/kg
					Mycophenolic acid	(20-2000) mcg/kg
					Myriocin	(200-10000) mcg/kg
					Monactin	(200-10000) mcg/kg
					Moniliformin	(20-2000) mcg/kg
					Neoxaline	(50-5000) mcg/kg
					Neosolaniol	(10-2000) mcg/kg
					Nivalenol	(100-10000) mcg/kg
					Nigericin	(200-10000) mcg/kg
					Nidulin	(200-10000) mcg/kg
					Nonactin	(200-10000) mcg/kg
					Nornidulin	(200-10000) mcg/kg
					NT-2 toxin	(10-2000) mcg/kg
					Oligomycin A	(200-10000) mcg/kg
					Oligomycin B	(200-10000) mcg/kg
					Ophiobolin A	(200-10000) mcg/kg
					Ochratoxin A	(1-200) mcg/kg
					Ochratoxin B	(1-200) mcg/kg
					Paxillin	(20-200) mcg/kg
					Paraherquamide A	(50-5000) mcg/kg
					Patulin	(1000-2000) mcg/kg
					Penitrem A	(50-5000) mcg/kg
					Penicillin acid	(20-2000) mcg/kg
					Piranonigrin A	(50-5000) mcg/kg
					Pyrenophorol	(100-10000) mcg/kg
					Pyripyropene A	(100-10000) mcg/kg
					Pseurotin A	(50-5000) mcg/kg
					Puromycin	(50-5000) mcg/kg
					Radicicol	(200-10000) mcg/kg
					Rapamycin	(200-10000) mcg/kg
					Roquefortine C	(10-2000) mcg/kg
					Roridin A	(20-2000) mcg/kg

1	2	3	4	5	6	243 pages,
					Rugulosin	(100-10000) mcg/kg
					Secalonic acid	(1000-10000) mcg/kg
					Semivioxanthin	(200-10000) mcg/kg
					Setosusin	(200-10000) mcg/kg
					Skyrin	(200-10000) mcg/kg
					Staurosporine	(200-10000) mcg/kg
					Stachybotrys lactam	(10-2000) mcg/kg
					Sterigmatocystin	(10-2000) mcg/kg
					Sulochrin	(200-10000) mcg/kg
					T-2 tetraol	(100-2000) mcg/kg
					T-2 toxin	(10-2000) mcg/kg
					T-2 triol	(20-2000) mcg/kg
					Taxol	(200-10000) mcg/kg
					Tentoxin	(20-2000) mcg/kg
					Tenuazonic acid	(20-2000) mcg/kg
					Terrein	(200-10000) mcg/kg
					Territrem B	(200-10000) mcg/kg
					Thiolutin	(200-10000) mcg/kg
					Tryptophol-OH	(200-10000) mcg/kg
					Trichostatin A	(200-10000) mcg/kg
					Fiscion	(200-10000) mcg/kg
					Famopsin A	(200-10000) mcg/kg
					Fusarenon X	(500-10000) mcg/kg
					Fusarium acid	(100-20000) mcg/kg
					Fumagillin	(100-2000) mcg/kg
					Fumitremorgin C	(50-5000) mcg/kg
					Fumonisin B1	(100-20000) mcg/kg
					Fumonisin B2	(100-20000) mcg/kg
					Fumonisin B3	(100-20000) mcg/kg
					Chaetoglobosin A	(200-10000) mcg/kg
					Hetocin	(200-10000) mcg/kg
					Chrysophanol	(200-10000) mcg/kg
					Cyclopeptide A	(200-10000) mcg/kg
					Cycloheximide	(50-5000) mcg/kg
					Cyclopenin	(50-5000) mcg/kg
					Cyclopenol	(50-5000) mcg/kg
					Cyclopiazonic acid	(20-2000) mcg/kg
					Cyclosporine D	(200-10000) mcg/kg
					Cyclosporine H	(200-10000) mcg/kg
					Cyclosporine A	(200-10000) mcg/kg

-1	2		T 4	-		243 pages,
1	2	3	4	5	6	
					Cyclosporine C	(200-10000) mcg/kg
					Cytochalasin A	(50-5000) mcg/kg
					Cytochalasin B	(50-5000) mcg/kg
					Cytochalasin D	(50-5000) mcg/kg
					Cytochalasin E	(50-5000) mcg/kg
					Cytochalasin H	(50-5000) mcg/kg
					Cytochalasin J	(50-5000) mcg/kg
					Citreoviridin	(100-2000) mcg/kg
					Citrinin	(50-10000) mcg/kg
					Equisetin	(50-5000) mcg/kg
					Emodin	(200-10000) mcg/kg
					Enniatin A	(50-5000) mcg/kg
					Enniatin A1	(50-5000) mcg/kg
					Enniatin B	(50-5000) mcg/kg
					Enniatin B1	(50-5000) mcg/kg
12	METHODOLOGICAL GUIDELINES	Fish, non-fish objects	10.20	0301-0308	Mass fraction:	(2000-40000) mcg/kg
	A-1/051				domoic acid	(2000-40000) meg/kg
	Methodical guidelines for the				Okadaic acid	(12,5-625) mcg/kg
	determination of phycotoxins in food				dinophysis toxin-1	(12,5-625) mcg/kg
	products by high-performance liquid				dinophysis toxin-2	(2,5-125) mcg/kg
	chromatography with mass spectrometric				pectenotoxin-2	(10-500) mcg/kg
	detection				brevetoxin	(10-500) mcg/kg
					Yessotoxin	(10-500) mcg/kg
					1a-homotoxin	(10-500) mcg/kg
					13-desmethyl spirolide C	(10-500) mcg/kg
					13,19-	(10-500) mcg/kg
					didesmethyl spirolide C	
					20-methylspirolide g	(10-500) mcg/kg
					azaspiracid-1	(1-50) mcg/kg
					azaspiracid-2	(1-50) mcg/kg
					azaspiracid-3	(1-50) mcg/kg
					azaspiracid-4	(1-50) mcg/kg
					azaspiracid-5	(1-50) mcg/kg
					saxitoxin	(40-1600) mcg/kg
					neosaxitoxin	(40-1600) mcg/kg
					decarbamoyl-saxitoxin	(40-1600) mcg/kg
					decarbamoyl-neosaxitoxin	(40-1600) mcg/kg
					gonyautoxin-2	(40-1600) mcg/kg
					gonyautoxin-3	(2,9-288) mcg/kg
					gonyautoxin-5	(40-1600) mcg/kg

1	2	3	4	5	6	243 pages
					gonyautoxin-6	(40-1600) mcg/kg
					decarbamoyl-	(40-1600) mcg/kg
					gonyautoxin-2	
					decarbamoyl-	(1.9.176)
					gonyautoxin-3	(1,8-176) mcg/kg
					N-sulfo-carbamoyl-	(40-1600) mcg/kg
					gonyautoxin-2	
					N-sulfo-carbamoyl-	(2,2-224) mcg/kg
					gonyautoxin-3	, , ,
	METHODOLOGICAL GUIDELINES	Honey	01.44.21	0409000000	nystatin	(5 - 500) mcg/kg
	A-1/052				clotrimazole	(0,1 - 10) mcg/kg
	Methodical guidelines for the				rifampicin	(1 -100) mcg/kg
13	determination of xenobiotics in honey by				fumagillin	(5 - 500) mcg/kg
	high performance liquid chromatography				colchicine	(1 -100) mcg/kg
	with mass spectrometric detection				dapsone	(1 -100) mcg/kg
					clothianidin	(1 -100) mcg/kg
					imidacloprid	(1 -100) mcg/kg
14	GOST 34141	Food products, feed, food raw	10.11.1-	0201-0210	mass fraction:	
		materials	10.11.3	0201-0210 0302-0308;	arsenic	(0.010 - 500) mg/kg
			10.11.5 10.12; 10.13	0401-0406	cadmium	(0.005 - 100) mg/kg
			10.2; 10.41.12	0409;1001;100 3;1005;1101;	Mercury	(0,002-20) mg/kg
			10.5	1102	lead	(0,010 – 500) mg/kg
			10.91.10.110	1501-1517 1604-1605;	icac	(0,010 300) mg/kg
			10.91.10.120 10.91.10.180	2304;2306;230		
				9		
15	GOST 34462	Food products, feed, food raw	10.2; 10.61.1;	0301-0308;		
		materials	10.20.22.120; 10.20.1;	1001-1008; 1101-1109;	mass fraction:	
			10.20.1;	2301-2309	inorganic arsenic	(0.03 - 10.0) mg/kg
			10.91.10.110	2301 2309	morganic arsenic	
			10.91.10.180			
16	METHODOLOGICAL	Muscle tissue of meat, including	10.11	0201-2010	mass fraction:	(10,0 - 100) mg/kg
	RECOMMENDATIONS 55-14 Methods	poultry	10.12		iron	
	of measurement of mass concentrations		10.13 10.20		cadmium	(0,05 - 0,5) mg/kg
	of chemical elements in muscle tissues		10.20		calcium	(100 - 1000) mg/kg
	(in meat) of animals and poultry by		10.42		cobalt	(0,01 - 0,1) mg/kg
	inductively coupled plasma mass		10.51		magnesium	(100 - 5000) mg/kg
	spectrometry method FR.1.31.2015.21645		10.52			(0,5 - 5,0) mg/kg
	TK.1.31.2013.21043		10.85.11 10.85.12		manganese	
			10.65.12		copper	(0,5 - 5,0) mg/kg

1	2	3	1	5	6	243 pages 7
1	2	3	4] 3	6	
			10.86 10.89		arsenic	(0,05 - 0,5) mg/kg
			10.89		nickel	(0,01 - 0,1) mg/kg
					lead	(0.05 - 0.5) mg/kg
					selenium	(0,5 - 5,0) mg/kg
					strontium	(0,1 - 1,0) mg/kg
					chrome	(0.05 - 0.5) mg/kg
					zinc	(5,0 - 100) mg/kg
17	METHODOLOGICAL GUIDELINES	Bio-substrates, drugs and	21.20.23.190	3003	mass fraction:	(0.001.20)
	4.1.1483-03	biologically active additives	21.20.23.193		aluminium	(0,001-20) mC/kg
	Methodical guidelines. Determination of		21.20.23.199 21.1		iron	(0,1-500) mC/kg
	chemical elements content in diagnosed		21.10		potassium	(1-5000) mC/kg
	biosubstrates, drugs and biologically		21.10		cadmium	(0,0001-0,5) mC/kg
	active additives by mass spectrometry				calcium	(2-2000) mC/kg
	with inductively coupled argon plasma				cobalt	(0,0001-0,5) mC/kg
	(approved by the Chief State Sanitary				magnesium	(0,001-500) mC/kg
	Doctor of the Russian Federation on				manganese	(0,0001-2) mC/kg
	29.06.2003)				arsenic	(0,0005-0,5) mC/kg
					Mercury	(0,0001-1) mC/kg
					copper	(0,0001-50) mC/kg
					natrium	(1-1000) mC/kg
					nickel	(0,0001-2) mC/kg
					lead	(0,0001-10) mC/kg
					phosphorus	(5-5000) mC/kg
					chrome	(0,001-10) mC/kg
					zinc	(0,001-500) mC/kg
					selenium	(0,0005-2) mC/kg
18	GOST P 56219	Drinking, natural and sewage water	10.86.10.300	-	Mass concentration	$(1 - 1000) \text{ mcg/dm}^3$
			10.86.10.310		silver	
			36.00.11. 36.00.11.000		aluminium	$(5 - 5000) \text{ mcg/dm}^3$
			36.00.11.000		arsenic	$(1 - 1000) \text{ mcg/dm}^3$
					boron	$(10 - 10000) \text{ mcg/dm}^3$
					barium	$(0.5 - 1000) \text{ mcg/dm}^3$
					beryllium	$(0.5 - 1000) \text{ mcg/dm}^3$
					calcium	$(10 - 10000) \text{ mcg/dm}^3$
					cadmium	$(0,1 - 500) \text{ mcg/dm}^3$
					cobalt	$(0,2 - 1000) \text{ mcg/dm}^3$
					chrome	$(1 - 1000) \text{ mcg/dm}^3$
					copper	$(1 - 1000) \text{ mcg/dm}^3$
					potassium	(50 - 10000) mcg/dm ³

1	2	3	4	5	6	7 243 pages,
					lithium	$(1 - 1000) \text{ mcg/dm}^3$
					magnesium	$(1 - 1000) \text{ mcg/dm}^3$
					manganese	$(3 - 5000) \text{ mcg/dm}^3$
					molybdenum	$(0.5 - 1000) \text{ mcg/dm}^3$
					natrium	$(10 - 10000) \text{ mcg/dm}^3$
					nickel	$(1 - 1000) \text{ mcg/dm}^3$
					phosphorus	(5 - 5000) mcg/dm ³
					lead	$(0.1 - 500) \text{ mcg/dm}^3$
					antimony	$(0,2 - 1000) \text{ mcg/dm}^3$
					selenium	$(10 - 10000) \text{ mcg/dm}^3$
					tin	$(1 - 1000) \text{ mcg/dm}^3$
					strontium	$(0,3 - 1000) \text{ mcg/dm}^3$
					tellurium	$(2 - 5000) \text{ mcg/dm}^3$
					thallium	$(0,1 - 500) \text{ mcg/dm}^3$
					vanadium	$(1 - 1000) \text{ mcg/dm}^3$
					Tungsten	$(0,3 - 1000) \text{ mcg/dm}^3$
					zinc	$(1 - 1000) \text{ mcg/dm}^3$
19	GOST P 58144	Distilled water	20.13.52.120	2853901000	substances reducing	compliant/non compliant
	p. 8.2;8.4-8.7-8.9-8.11.			2853001000	KMnO4	
	8.12.					
	8.14.				pН	un.pH (0-14)
	8.15.				specific electrical	(10 ⁻⁴ - 10) Cm/m
20	GOOD ALOGE	5.11	10.06.10.200		conductance	
20	GOST 31867 p. 4	Drinking water	10.86.10.300 10.86.10.310	-	ion sulfate	$(0.5 - 50) \text{ mg/dm}^3$
			36.00.11.		ion chloride	$(0.5 - 50) \text{ mg/dm}^3$
			36.00.11.000			
21	GOST 33045	Drinking, natural and sewage water	10.86.10.300		Ammonia and ammonium	
	p.5		10.86.10.310		ions (total)	$(0.1 - 3.00) \text{ mg/dm}^3$
	p, 6	7	36.00.11.		nitrite ions	$(0,003 - 0,3) \text{ mg/dm}^3$
22	GOST P 57162	Deigling costs	36.00.11.000 10.86.10.300	_		$(0.01 - 10) \text{ mg/dm}^3$
22	GOS1 P 3/102	Drinking water	10.86.10.310	_	aluminium iron	$(0.01 - 10) \text{ mg/dm}^3$ $(0.04 - 25) \text{ mg/dm}^3$
			36.00.11.			$(0.04 - 2.5) \text{ mg/dm}^3$
			36.00.11.000		copper lead	$(0,001 - 3) \text{ mg/dm}^3$ $(0,002 - 5) \text{ mg/dm}^3$
					zinc	$(0,002 - 5) \text{ mg/dm}^3$
23	GOST 24240	Feed, compound feed, animal	10.91.10.110	2309		(0,001 - 30) mg/am ²
23	GOST 34249	compound feed	10.91.10.110	2307	mass fraction:	(0,1 - 5,0) mg/kg
		•			chrome	
24	GOST 33411	Raw materials and food products	10.11; 10.12	0201-0208,	mass fraction:	(0.01, 50.0)
			10.13; 10.20 10.51; 10.52	0301-0305, 0401-0408,	arsenic	(0,01 - 50,0) mg/kg
			10.51, 10.52	0401-0408,		

1	2	3	4	5	6	243 pages, 7
			01.11; 01.12 10.91; 10.92	0409000000		
25	GOST 33412	Raw materials and food products	10.11; 10.12 10.13; 10.20 10.51; 10.52 01.11; 01.12 10.91; 10.92	0201-0208, 0301-0305, 0401-0408, 0409000000	mass fraction: Mercury	(0,002 - 5,000) mg/kg
26	GOST 33413	Raw materials and food products	10.11; 10.12 10.13; 10.20 10.51; 10.52 01.11; 01.12 10.91; 10.92	0201-0208, 0301-0305, 0401-0408, 0409000000	mass fraction: tin	(25,0 - 1000,0) mg/kg
27	GOST ISO 14377	Condensed milk canned	10.51.51.111	0401-0406	mass fraction: tin	(0,25 - 5,0) mg/kg
28	GOST ISO/TS 6733	Milk and Dairy products	10.51	0401-0406	mass fraction: lead	(0,001 - 200) mg/kg
29	GOST P 54639	Food and feed for animals	10.11; 10.12 10.13; 10.20 10.51; 10.52 01.11; 01.12 10.91; 10.92	0201-0208, 0301-0305, 0401-0408, 0409000000 2102 1101-1104	mass fraction: Mercury	(0,0025 - 5) mg/kg
30	GOST EN 15505	Food products	10.11; 10.12 10.13; 10.20 10.51; 10.52	0201-0208, 0301-0305, 0401-0408,	mass fraction: natrium	(1500 -15000) mg/kg
			01.11; 01.12 10.91; 10.92	0409000000	magnesium	(250 - 1000) mg/kg
31	GOST 9793	Meat and meat products	10.13.14.610 10.11	0201-0210	humidity	(1,0-85,0)%
32	GOST ISO 6092	Powdered milk	10.51.2	0401	titratable acidity (practical method)	pH 8,0 – 10,0
33	GOST ISO/TS 17837	Processed cheese products	10.51.40.217	0406	nitrogen content and calculation of total protein	(0,5-99,0)%
34	GOST P 57221p.7	Feed yeast			mass fraction of ash	(0,1-99,0)%
	p.8 p. 9		10.91.10.151	2102 1101-1104	mass fraction of raw protein Bernstein mass fraction of protein	(0,5-99,0)% (0,5-99,0)%
35	GOST 32052 p.8.7	Food additives. Lecithin E322		2923	mass fraction of substances, insoluble in toluene	(0, 1 - 0,30)%
	p.8.8		-		mass fraction of substances, insoluble in acetone	(60-95)%

			,	•		243 pages
1	2	3	4	5	6	7
	p.8.9				mass fraction of moisture and volatile	(0,01 – 95,00)%
36	GOST 2081 p. 7.4.2	Carbamide			mass fraction of nitrogen by distillation method	(45 – 47)%
	p.7.6		-	-	mass fraction of free ammonia	(0,01 - 0,04)%.
	p. 7.7				mass fraction of water	(0,05-0,5)%.
37	GOST P 55063, p 7.7	Cheese and melting cheeses	10.51.4 10.51.40.170	0406	mass fraction of moisture and dry matter	(3,0-70,0) %
38	GOST 25011, p.6	Meat andmeat products	10.13.14.610 10.11	0201-0210	Kjeldahl mass fraction of protein	(1,0-55,0) %
39	STB ISO 8968-1-2008	Whole and Skim Milk	10.51.56.420	0401	nitrogen by the Kjeldahl method	(0,5-15,0)%
40	GENERAL PHARMACOPOEIA ARTICLE 1.2.3.0011.15	Drugs		-	Kjeldahl nitrogen in organic compounds	(0,1-99,0)%
41	GOST P 56374	Feed, compound feed, feed raw materials, feed additives	10.91	2309	mass fraction of cations: ammonium potassium sodium magnesium calcium	(0,01 - 40,00)%
42	GOST P 56375	Feed, compound feed, feed raw materials, feed additives	10.91	2309	mass fraction: chloride ions sulfate-ions	(0,005 - 60,00)% 0,005 -70,00)%
					nitrate-ions	(0,002 -1,00)%
					phosphate-ions	(0,005 - 80,00)%
43	GOST P 57124	Feed, compound feed, animal compound feed, feed additives	10.91	2309	mass fraction choline chloride	(0,01 - 100)%.
44	GOST 33500	Dairy raw materials, drinking milk and cream	10.51	0201-0210	mass concentration of phosphate-ions	$(5-1500) \text{ mg/dm}^3$
45	GOST EN 12014-4	Meat products, vegetables	_{10.11}	0201-0210	mass fraction of nitrate	(50-2500) mg/kg
			01.13; 10.3	0701-0714	mass fraction of nitrite	(40 - 300) mg/kg
46	GOST P 55569 and other normative documents approved	Feed, compound feed, feed additives, animal compound feed	10.91 10.92	2309	mass fraction: alanine	(0,25-10) %
	in the established order that specify the	r			arginine	(0,5-10) %
	application of the research (testing) method, measurements that establish				asparagine acid and asparagine in total	(0,5-10) %
	requirements for feed and feed additives				valine	(0,5-10) %
	registered in the established order and				histidine	(0,5-10) %
	included in the State registries of feeds				glycine	(0,25-10) %
	and feed additives of the Eurasian Economic Union member states				glutamine acid and glutamine in total	(0,5-10) %
					leucine and isoleucine in total	(0,25-10) %

1	2	3	4	5	6	7 243 pages,
					methionine	(0,25-10) %
					proline	(0,25-10) %
					lysine	(0,25-20) %
					serine	(0,25-10) %
					tyrosine	(0,25-10) %
					threonine	(0,5-10) %
					phenynalanine	(0,25-10) %
					cystine	(0,1-10,0)%
47	GOST 34258	Drugs for veterinary use, feed	10.91	2309	mass concentration:	
		additives.			vitamin B ₁	(60 - 4800) mg/kg
					vitamin B2	(25 - 2000)mg/kg
					vitamin PP	(60 - 4800) mg/kg
					vitamin B6	(25 - 2000) mg/kg
					vitamin B5	(125 - 10000) mg/kg
					vitamin B9	(25 - 2000) mg/kg
					vitamin B12	(25 - 2000) mg/kg
					vitamin H	(25 - 2000) mg/kg
48	GOST 31483	Premixes	10.91	2309	mass fraction:	(0,1-5,0)g/kg
		Vitamin additives			B1(thiamine chloride)	
					B2(riboflavin)	(0,1-5,0)g/kg
					B3(pantothenicacid)	(1,0-25,0)g/kg
					B5 (nicotine acid)	(2,0–100,0)g/kg
					B5 (niacinamide)	(0,1-5,0)g/kg
					B6 (pyridoxine)	(0,2-10,0)g/kg
					B9 (folic acid)	(0,1-5,0)g/kg
					C (Ascorbic acid)	(2,0-50,0)g/kg
49	GOST 31643	Juice products	10.32	-	mass fraction:	
					ascorbic acid	(5-1000)mg/dm ³ , (mln ⁻¹)
50	GOST 33527	Dairy products for child nutrition	10.86	1901	weight fraction of mono	(0,5% -10,0) %.
				0401	and disaccharides	(2.0 05.0) 0/
51	GOST 34178,	Melted spreads and mixtures,	10.42	210690980	mass fraction milk fat in	(3,0 – 85,0) %
	Annex B	milk and dairy products	10.51.30	4	fat phase	
52	GOST 31754, p.7	Animal and vegetable oils and	10.41	1501-1518	mass fraction of	(1 - 50) %
	-	fats	10.42		isolated trans-isomers	
					of fatty acids	
53	GOST P 55483	Meat and meat products	10.11	0201-0210	mass fraction individual	(0,03 – 98) %-
			10.13.1	1501-1502	fatty acids	
54	GOST P 56373	Feed and feed additives	10.91	2309	mass fraction of organic	(0,03% - 10,00)%
					acids:	
					oxalic	

1	2	3	4	5	6	7 7
					formic	(0,15% - 80,00)%
					fumaric	(0,005% - 80,00)%
					succinic	(0,05% - 80,00)%
					malic	(0,05% - 80,00)%
					citric	(0,05% - 80,00)%
					acetic	(0,10% - 80,00)%
					propanoic	(0,10% - 80,00)%
					lactic	(0,12% - 80,00)%
					benzoic	(0,005% - 50,00)%
					sorbic	(0,025% - 50,00)%
					butyric	(0,05% - 50,00)%
55	GOST 34164	food products, food raw materials in	10.51.11		mass fraction of fuccilin	(0,5 - 62,5) mcg/kg
		terms of meat, poultry, eggs, egg	10.51.22		metabolite	, , ,
		powder, egg melange, milk, fish,	10.11.11		(semicarbazide)	
		honey	10.11.12			
			10.11.13	0201		
			10.11.31	0203		
			10.11.32	0207		
			10.11.33	0401		
			10.12.10	0407		
			10.12.20	040900000		
			01.47.21.000 10.89.12.110	0		
			03.11.20	0306		
			01.49.21.110			
			03.12.20			
			03.12.20			
			03.22.10			
56	GOST 34209	feed, compound feed, animal	10.91.10.180	1001	total pleuromutilin	(0-48)ng/kg
		compound feed	10.91.10.186	1003	content (tiamulin and	(= -=/ == / ==
		*	10.91.10.183	1005	valnemulin)	
			10.91.10.181	1102	,	
			10.91.10.184	1101		
			10.91.10.182	2304		
			10.91.10.185	2306		
			10.91.10.189			
			10.91.10.188			
			10.91.10			
			10.92.10			
	GOST 34284	meat (all kind of animal), feed, feed	10.91.10.180	2309	total β-agonists content	$(0.03-4.21) \text{ ng/cm}^3$

1	2	3	4	5	6	7 7
57		additives, as well as biological objects of animal origin	10.91.10.186 10.91.10.183	0201 0202	boldenone	(0,03-4,43) ng/cm ³
		objects of annhar origin	10.91.10.181	0202	total corticosteroid content	(0,04-4,61) ng/cm ³
			10.91.10.182		nandrolone	$(0.07-8.70) \text{ ng/cm}^3$
			10.91.10.185 10.91.10.189		ractopamine	(0,03-4,43) ng/cm ³
			10.91.10.188		stanozolol	(0,03-4,36)ng/cm ³
			10.91.10 10.92.10		total stilbene content	(0,04-5,32)ng/cm ³
			10.11.11		trenbolone	(0,03-3,69)ng/cm ³
			10.11.12 10.11.13		zeranol	(0,02-4,74)ng/cm ³
			10.11.31			
			10.11.32 10.11.33			
			10.12.10			
50	GOST 34285		10.12.20	0201		
58	GOS1 34283	meat (all kind of animal), including poultry meat, milk, honey	01.49.21 10.51.11	0201 0203	sulfadiazine	
		poundy mean, man, noney	10.51.21	0207	sulfadimetoxin	
			10.51.22	0401	sulfaquinoxaline	Positive\negative
			10.11.11	040900000	sulfamethazine	
			10.11.12	0	sulfamethoxazole	
			10.11.13 10.11.31		sulfathiazole	
			10.11.31		sulfisoxazole	
			10.11.33		sulfapyridine	
			10.12.10		sulfamerazine	
			10.12.20		sulfamonomethoxine	
					sulfamethox-pyridoxine	
					sulfachloropyridazine	
					dapsone	
					sulfadoxine	
					trimethoprim	
					quinolones ceftiofur	
					thiamphenicol	
					streptomycin	
					tylosin	

1	2	3	4	5	6	7 243 pages
					tetracyclines	
					aoz	
					amoz	
					agd	
					sem	
					spiramycin	
					apramycin	
					bacitracin	
					neomycin	
					tobramycin	
					tylosin в	
					spectinomycin	
					amikacin	
					lincosamides	
					erythromycin	
					streptomycin	
					virginiamycin	
					Nitroimides of ash	
					chloramphenicol	
					cephalexin	
					beta-lactam type	
					antibiotics	
					cefuroxime	
					Benzimide of ash	
					aminbenzimide of ash	
					levamisole	
					avermectins	
					thiabendazole	
					moxidectin	
50	METHODOLOGICAL CURRELINES		01.40.21	0201	triclabendazole	
59	METHODOLOGICAL GUIDELINES	food products, raw materials such as	01.49.21	0201	mass fraction of	
	A-1/042 Methods for massuring the mass fraction	meat, poultry and eggs, milk, honey	10.51.11 10.51.21	0203 0207	pleuromutilins	
	Methods for measuring the mass fraction of pleuromutilins in animal products by		10.51.21	0401		
	solid-phase competitive immunoenzyme		10.31.22	0401		
	analysis		10.11.11	040300000		(10-1200) mcg/kg
	mini y 515		10.11.31-			
			10.11.31			
			10.12.10			
			10.12.20			

1	2	3	4	5	6	7 243 pages.
60	METHODOLOGICAL GUIDELINES	Meat, milk, eggs, feed	01.49.21	0201	mass fraction of	meat (0,009-0,3) mg/kg
	4.1.3379-16		10.51.11	0203	bacitracin	milk(0,011-0,2) mg/kg
	Determination of residues of bacitracin		10.51.21	0207		eggs(0,011-0,3) mg/kg
	in animal products by enzyme		10.51.22	0401		feed(0,092-0,8) mg/kg
	immunoassay		10.11.11-	0407		
			10.11.13	2309		
			10.11.31-			
			10.11.33			
			10.12.10			
			10.12.20			
			10.91.10.180-			
			10.91.10.186			
			10.91.10.189			
			10.91.10.188			
			10.91.10			
			10.92.10			
61	METHODOLOGICAL GUIDELINES	Meat, milk and dairy products, eggs,	01.49.21	0201	total content of	(1-240) mcg/kg
	A-1/048	offal, honey	10.51.11	0203	lincosamides(lincomycin	
	Method of screening determination of		10.51.21	0207	and clindamycin)	
	the residual content of lycosamides in		10.51.22	0210		
	animal products by the method of solid-		10.11.11-	0401		
	phase competitive immunoenzyme		10.11.13	0407		
	analysis		10.11.31-	040900000		
			10.11.33	0		
			10.12.10			
			10.12.20			
			10.11.20			
62	METHODOLOGICAL GUIDELINES	Meat, milk, eggs	01.49.21	0201	mass fraction	
	4.1.1912-04		10.51.11	0203	chloramphenicol	$(0-40,5) \text{ ng/cm}^3$
	Determination of residual amounts of		10.51.21	0207	(Levomycetin)	
	Levomycetin (Chloramphenicol,		10.51.22	0401		
	Chlormecithin) in animal products by		10.11.11-	0407		
	high-performance liquid		10.11.13			
	chromeatography and immunoassay.		10.11.31-			
	p. 5		10.11.33			
			10.12.10			
			10.12.20			
63	GOST P 57025	Fish, crustacean, mollusca	10.20.11	0302	total content of	System testing 1 (0-0,2) ng/cm ³
			10.20.13	0303	triphenylmethane dyes	System testing 2 (0,0-3,2)
			10.20.14	0304		ng/cm ³
			10.20.15	0306		
			10.20.31	0307		

1	2	3	4	5	6	7 7
			10.20.32			
64	GOST EN 12856	Food products	10.11 – 10.89	2001- 2009 2201-2202	mass fraction of artificial sweetener: Acesulfame potassium aspartame saccharine	(10 - 3000) mg/kg or mg/dm ³
65	GOST 34138	Milk and Dairy products, meat and poultry products	10.11 – 10.13 10.41.1 10.41.6 10.42; 10.51	0201- 0210 0401-0406	mass concentration of macrocyclic lactones	(0,0005 – 0,25) mg/kg
66	GOST 31644	Fruits and vegetables juice products, Fruits and vegetables juice products for child nutrition	10.32 10.86.10.230 – 10.86.10.249	2009	mass concentration 5- hydroxymethyl-furfural	$(1,0-50,0) \text{ mg/dm}^3$
67	GOST 34228	Fruits and vegetables juice products	10.32	2009	mass concentration of preservatives: 4- hydroxybenzoic acid benzoic acid sorbic acid methyl-4- hydroxybenzoates ethyl-4- hydroxybenzoates n-propyl-4- hydroxybenzoates n-butyl-4- hydroxybenzoates	(10,0 – 320,0) mg/dm ³ (10,0 – 320,0) mg/dm ³
68	GOST ISO 14501	Milk and Powdered milk	10.51.11.110 – 10.51.11.119 10.51.21 10.51.22.110 – 10.51.22.122	0402	content determination of aflatoxin m1 milk Powdered milk	(0,008 - 0,100) mcg/l (0,08 - 0,10) mcg/kg
69	GOST ISO 9231	Milk and dairy products	10.51 10.52	0401-0406	mass fraction: benzoic acid sorbic acid	(5 – 2000) mg/kg (5 – 1000) mg/kg
70	ISO 18329	Milk and dairy products	10.51 10.52	0401-0406	mass fraction of furosine	(5 - 5000) mg/100 g
71	ISO 11868	Milk	10.51.11 10.51.21 10.51.22.110 – 10.51.22.122	0401-0406	mass concentration of lactulose	(200 – 1500) mg/l

1	2	3	4	5	6	7 243 pages,
72	METHODOLOGICAL GUIDELINES A 1/053 Methodical guidelines for determining polybrominated pollutants in animal products, feed and feed additives	Food raw materials Feed Feed additives	10.1 10.2 10.4 10.5 10.9	0401-0408 0201-2010 0301-0308 1001-1008 1102;1101 2304;2306 2309	mass concentration of polybrominated compounds	(0,004 - 8) mcg/kg
73	METHODOLOGICAL GUIDELINES A 1/054 Methodical guidelines for the determination of pesticides in honey by gas-liquid chromeatogrphy with mass spectrometric detection	Honey	10.8	0409	mass concentration of pesticides	(0,005 – 0,5) mg/kg
74	GOST 32193	Feed Compound feed	10.9 10.91.10.180	1001-1008 2304; 2306 2309	mass fraction organophosphorus of pesticides	(0,01 - 1,0) mcg/g
75	METHODOLOGICAL GUIDELINES A-1/043	Feed Compound feed Soybean Raw material of vegetable products	10.9 10.91.10.180 01.11	1001-1008 2304;2306 2309; 1201 0701-0714 1001-1008 1101-1109 1201-1213	mass fraction of glyphosate, gluphosinate and aminomethyl- phosphonic acid	(0,1 - 10) mg/kg
76	METHODOLOGICAL GUIDELINES A 1/055 Methodical guidelines for determining	Honey	01.49.21	0409	mass fraction of glyphosate of glufosinate	(0,05-2) mg/kg (0,5-20) mg/kg
	the content of glufosinate, glyphosate and its metabolite by high-performance liquid chromeatography with high- resolution time-of-flight mass spectrometer detector in honey				aminomethyl-phosphonic acid	(0,1-2) mg/kg
77	METHODOLOGICAL GUIDELINES A 1/056 Methodical guidelines for the determination of polyfluorinated contaminants using high performance liquid chromeatography with mass spectrometric detection	Fish and non-fish objects	10.20.1 10.20.3	0301-0308	mass fraction of organofluorine compounds	(0,5 - 100) mcg/kg
78	GOST 34449	Food products, food raw materials, feed, feed additives.	10.1 10.2 10.4 10.5 10.9	0401-0408 0201-2010 0301-0308 1001-1008 1102;1101 2304; 2306 2309	mass fraction of dioxin	(1.0 - 30.0) ng/kg
79	ISO 18363-1	Animal and vegetable fats and oils	10.4	1501- 1518	Mass fraction 3-MCPD and glycidol	(0.003 - 3) mg/kg

1	2	3	4	5	6	7 7
80	CT PK ISO 18363-1	Animal and vegetable fats and oils	10.4	1501- 1518	Mass fraction 3-MCPD and glycidol	(0.003 - 3) mg/kg
81	STATE PHARMACOPOEIA OF THE	Drugs.	21.1	3003 –	Appearance (description)	-
	REPUBLIC OF BELARUS section 1		21.10	3004	Color (description)	-
	«General information» and other normative documents approved in the		21.10.1 21.10.20.120	from 4201	Odor (description)	-
	established order, specifying the		21.10.20.120		Consistency (description)	-
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124		Solubility	Pass test/ fail test (if necessary, specify conditions)	
82	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.3.4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs.	21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10 21.20.10.158 21.20.10.159 21.20.21.130 21.20.21.130		Odor (description)	-
83	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions; Suspensions and emulsions: Drops; Tinctures and Extracts; Powders and granules; Pharmaceutical substances; Lyophilizate		21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130		Transparency and degree of turbidity of liquids (description)
84	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.2 and other normative documents		02.30.40.140 15.12.11		Degree of liquids coloration / color / colour / colour solution /	-

1	2	3	4	5	6	7 7
	approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				coloration / solution coloration (description)	
85	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				pH / activity (concentration) hydrogen ions / pH / pH solution	from 0 to 14
86	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Density/ Relative density/ density at 20 °C / density at 25 °C	700 – 1840 kg/m ³ 0,001 – 3,000 mg/cm ³ 0,0001 – 3,000 mg/cm ³
87	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.17 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Nominal Volume / recoverable volume / filling volume / filling volume of bottle / drug volume in bottle	0,1 – 1000 ml (cm3; l; dm3); 80 -150 % of nominal; Compliant/ Non compliant; Pass test/ Fail test

1	2	3	4	5	6	7 243 pages,
88	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.			Refractive index / Quantitative determination	1,3 – 1,7; 0,0001 – 500 g/ml; mg/ml; g/l; mg/l; g/cm ³ ; mg/cm ³
89	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.9 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions; Suspensions and emulsions: Drops (eye): Tinctures and Extracts; Pharmaceutical substances			Viscosity	0,0001–100000 mm2/c; Ps; cPs; PAHs; MPAHs; m2/c; St; cSt;
90	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.10 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					

	2	3	4	5	6	7
91	STATE PHARMACOPOEIA OF THE	Drugs:	21.1	3003-3004	Moisture content / Water	0,01 – 100%
	REPUBLIC OF BELARUS art. 2.2.13	Solutions;	21.10		content	
	and other normative documents	Powders and granules	21.10.32			
	approved in the established order,	(microgranules, pellets):	21.10.5			
	specifying the application of the research	- for preparation (solution for	21.10.51.120			
	(testing) method, measurements,	injection, for oral use, drops);	21.10.51.121			
	establishing requirements for drugs	- for oral use;	21.10.51.122			
	registered in the established order and	- for external use;	21.10.51.123			
	included in State registers of drug for	- for local application;	21.10.51.124			
	veterinary use of the Eurasian Economic	Lyophilizate;	21.10.51.125			
	Union member states	Capsules;	21.10.51.126			
		Ointments;	21.10.51.129			
0.2	CELEBRATE DIA DIA CODORIA OF EILE	Tablets and dragee;	21.10.52.110			
	STATE PHARMACOPOEIA OF THE	Pharmaceutical substances	21.10.53			
	REPUBLIC OF BELARUS art. 2.5.12		21.10.53.120			
	and other normative documents		21.10.54			
	approved in the established order,		21.10.54.110			
	specifying the application of the research		21.10.54.120			
	(testing) method, measurements,		21.10.54.130			
	establishing requirements for drugs		21.10.54.140			
	registered in the established order and		21.10.54.150			
	included in State registers of drug for		21.10.54.160			
	veterinary use of the Eurasian Economic		21.10.54.170			
	Union member states		21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.21.130			
			21.20.21.139			
93	STATE PHARMACOPOEIA OF THE	Pharmaceutical substances	21.1	3003	Melting temperature/	25 – 400 °C
	REPUBLIC OF BELARUS art. 2. 2.14		21.10		melting point	
	and other normative documents					
	approved in the established order,					
	specifying the application of the research					
	(testing) method, measurements,					
	establishing requirements for drugs					
	registered in the established order and					
	included in State registers of drug for					
	veterinary use of the Eurasian Economic					
	Union member states					
	STATE PHARMACOPOEIA OF THE	Pharmaceutical substances			Sulfated ash	0,001 - 10,000%
	REPUBLIC OF BELARUS art. 2.4.14	Medicinal plant raw material and				

1	2	3	4	5	6	7 7
	and other normative documents	collections.				
	approved in the established order,					
	specifying the application of the research					
	(testing) method, measurements,					
	establishing requirements for drugs					
	registered in the established order and					
	included in State registers of drug for					
	veterinary use of the Eurasian Economic					
	Union member states					
95	STATE PHARMACOPOEIA OF THE				Total ash	0,001 - 10,000%
	REPUBLIC OF BELARUS art. 2.4.16					
	and other normative documents					
	approved in the established order,					
	specifying the application of the research					
	(testing) method, measurements,					
	establishing requirements for drugs					
	registered in the established order and					
	included in State registers of drug for					
	veterinary use of the Eurasian Economic					
	Union member states					
96	STATE PHARMACOPOEIA OF THE	Drugs.	21.1	3003-3004	Quantitative	0,1-10000 mkg/kg (g; 100g; ml;
	REPUBLIC OF BELARUS art. 2.2.20	Medicinal plant raw material and	21.10		determination /	cm3; 1; dm3; 100ml; tablet.;
	and other normative documents	collections.	21.10.1		quantitative content /	capsule; pipette; syringe; bottle.;
	approved in the established order,	Pharmaceutical substances.	21.10.20.120		mass fraction / mass	plate; suppository; stick;
	specifying the application of the research		21.10.32		concentration	package, bag);
	(testing) method, measurements,		21.10.5			0,000001-10000 mg/kg (g; 100g;
	establishing requirements for drugs		21.10.51.120			ml; cm3; l; dm3; 100ml; tablet.;
	registered in the established order and		21.10.51.121			capsule; pipette; syringe; bottle.;
	included in State registers of drug for		21.10.51.122			plate; suppository; stick;
	veterinary use of the Eurasian Economic		21.10.51.123			package, bag);
	Union member states		21.10.51.124			0,00001-1000 g/kg (g; 100g; ml;
97	STATE PHARMACOPOEIA OF THE		21.10.51.125			cm3; 1; dm3; 100ml; tablet;
	REPUBLIC OF BELARUS art. 2.5.8		21.10.51.126			capsule; pipette; syringe; fl.
	and other normative documents		21.10.51.129			plate; suppository; stick;
	approved in the established order,		21.10.52.110			package, bag)
	specifying the application of the research		21.10.53			0,01 -10 mg KOH/g (cm3;g);
	(testing) method, measurements,		21.10.53.120			0,0001-150%; 0,0001-150% weight;
	establishing requirements for drugs		21.10.54			0,0001-150% weight; 0,0001-150% volume;
	registered in the established order and		21.10.54.110			not found
	included in State registers of drug for		21.10.54.120			
	veterinary use of the Eurasian Economic		21.10.54.130			
	Union member states		21.10.54.140			

1	2	3	4	5	6	7 7
98	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.23 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.130 21.20.21.139 02.30.40.140			0,01-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,000001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-1000 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag) not found
99	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.25 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Authenticity Quantitative determination / quantitative content / mass fraction / mass concentration	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository);

1	2	3	4	5	6	7
100	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.5.11 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination / quantitative content / mass fraction / mass concentration	0,00001-150%; 0,00001-150% weight; 0,00001-150% volume; 1,0 - 200,0 % of declared; not found 0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,000001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-10,0 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag) 0,0001-150%; 0,0001-150%; 0,0001-150% weight; 0,0001-150% volume; not found
101	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.58 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE				Quantitative determination (quantitative content; mass fraction; mass concentration) of active substance	(0,002 – 500) mg/kg, mg/dm ³ 0,1-10000 mkg/kg (g; 100g; ml;
102	REPUBLIC OF BELARUS art. 2.5.50 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements,				determination / quantitative content / mass fraction / mass concentration	cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,000001-10000 mg/kg (g; 100g;

1	2	3	4	5	6	243 pages
1		3	'	<u> </u>	U U	1 2 1 1 2 100 1 11
103	establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE	Drugs.			Authenticity	ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-1000 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag) 0,0001-150%; 0,0001-150% weight; 0,0001-150% volume; not found Compliant/ Non compliant; Pass
103	REPUBLIC OF BELARUS art. 2.2.28 and other normative documents approved in the established order, specifying the application of the research	Medicinal plant raw material and collections. Pharmaceutical substances.			·	test/ fail test (if necessary, specify conditions)
	specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g;
						100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%;

1	2	3	4	5	6	7 7
						0,00001-150% weight; 0,00001-150% volume; 1,0 - 200,0 % of declared; not found
					Foreign matter / related compounds	0,01 - 20% 0,01 - 20% of active substance (if necessary, specify conditions)
		Drugs: Solutions: - for injection;			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
		Suspensions and emulsions: - for injection; Drops (eye)			Quantitative determination of antimicrobial preservatives / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-10000 mg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-20%; 0,00001-20% weight; 0,00001-20% volume; 1,0 - 200,0 % of declared; not found
		Drugs: Solutions: - for oral use;			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
		- for injection; Suspensions and emulsions: - for oral use; - for injection; Powders and granules (microgranules, pellets): - for oral use. Tinctures and Extracts. Medicinal plant raw material and collections. Pharmaceutical substances			Quantitative determination of aromatic compounds / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-1000 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag);

1	2	3	4	5	6	7 7
104	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.29 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.			Authenticity Quantitative determination (quantitative content; mass fraction; mass concentration) of active substance	0,1 – 100000000 IU/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150%, 0,00001-150% weight,

1	2	3	4	5	6	7 7
					Foreign matter / related compounds	0,00001-150% volume, 1,0 – 200,0 % of declared; not found 0,01 - 20% 0,01 - 20% of active substance (if necessary, specify
		Drugs: Solutions: - for injection;			Authenticity	conditions) Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
		Suspensions and emulsions: - for injection; Drops (eye)			Quantitative determination of antimicrobial preservatives / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-10000 mg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-20%; 0,00001-20% weight; 0,00001-20% volume; 1,0 - 200,0 % of declared; not found
		Drugs: Solutions: - for oral use;			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
		- for injection; Suspensions and emulsions: - for oral use; - for injection; Powders and granules (microgranules, pellets): - for oral use. Tinctures and Extracts; Pharmaceutical substances. Medicinal plant raw material and collections.			Quantitative determination of aromatic compounds / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-1000 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag) 0,1 – 1000000000 IU/kg (g; 100g; ml;

1	2	3	4	5	6	7 7
		Drugs: Solutions: - for oral use; - for injection; Suspensions and emulsions: - for oral use; - for injection; Powders and granules (microgranules, pellets): - for oral use. Tinctures and Extracts; Syrups; Balsams; Pharmaceutical substances; Pharmaceutical substances of vegetable products. Medicinal plant raw material and collections.			Authenticity; Quantitative determination of antioxidants / quantitative content / mass fraction / mass concentration	cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions); 0,1-10000 mkg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-10000 mg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-20%; 0,00001-20% weight; 0,00001-20% volume; 1,0 – 200,0 % of declared; not found
		Drugs: Solutions: - for oral use; - for injection; Suspensions and emulsions: - for oral use; - for injection; Powders and granules (microgranules, pellets): - for oral use. Pharmaceutical substances			Authenticity Quantitative determination of organic acids / quantitative content / mass fraction / mass concentration	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-1000 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet;

1	2	3	4	5	6	7 7
						capsule; pipette; syringe; fl. plate; suppository; stick; package, bag) 0,1 – 100000000 IU/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
105	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 5.10 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.			Foreign matter / related compounds	0,01 - 20% 0,01 - 20% of active substance (if necessary, specify conditions)
106	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.27 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
107	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.3.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and					

1	2	3	4	5	6	7 7
	included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
108	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.24 and other normative documents	Drugs: Solutions; Drops;			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	approved in the established order, specifying the application of the research (testing) method, measurements,	Aerosols and sprays; Suspensions and emulsions; Powders and granules;			Quantitative determination / quantitative content /	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.;
	establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Suppositories (sticks) Tablets, dragee, briquettes, pastilles; Capsules; Ointments; Lyophilizate;			mass fraction / mass concentration	plate; suppository; stick; package; bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.;
		Pharmaceutical substances. Medicinal plant raw material and collections.				plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet;
						capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g;
						100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g;
						100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%,
						0,00001-150%weight, 0,00001-150% volume, 1,0 – 200,0 % of declared;
109	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.40 and other normative documents				Authenticity	not found Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)

1	2	3	4	5	6	7 7
	approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 1000000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 - 200,0 % of declared; not found
110	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.2.48 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Authenticity Quantitative determination / quantitative content / mass fraction / mass concentration	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag);

1	2	3	4	5	6	
1	<u> </u>	<u> </u>	4	<u> </u>	0	/
111	STATE PHARMACOPOEIA OF THE	Drugs.			Weight loss during	0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found 0,001 – 50,0 %
	REPUBLIC OF BELARUS art. 2.2.32 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Medicinal plant raw material and collections. Pharmaceutical substances.			drying / Drying method / mass fraction of humidity / humidity	
112	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.4.24 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.			Residual organic solvents	10 -5000ppm (mg/kg; mcg/g); 0,00001 – 10%

1	2	3	4	5	6	7 243 pages,
113	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 5.4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.4.27 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.4.8 (Method A, Method B) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Content of heavy metals (cadmium, lead, arsenic, Mercury)	(0,002 – 500) mg/kg, mg/dm ³
116	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.5.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and	Drugs: Solutions; Ointments; Drops (eye); Extracts	21.10.51.120 21.10.52.110 21.10.54 21.10.54.180 21.20.1 21.20.10 02.30.40.140	3004	Acid value	0,01 -10 mg KOH/ml (cm3;g)

1	2	3	4	5	6	7 7
	included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
117	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.5.5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Peroxide value	0,01-50 mmol O2/kg
118	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.5.33 (Method 1, Method 5, Method 7) and other normative documents approved in the established order, specifying the application of the research (testing)	Drugs: Solutions; Lyophilizate.	21.20 21.20.1	3004	Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination of protein / quantitative content / mass fraction / mass concentration	0,2 - 2,0 mg/ml
119	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.6.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for external use (when applied to wounds); - for intracisternal injection; - for infusion; Suspensions and emulsions: - for injection; - for intracisternal injection; Drops (eye); Ointments (Gels, creams, liniments, pastes); - for external use; - for intracisternal injection;	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110	3003-3004	Sterility	Sterile/non-sterile; pass test/fail test

1	2	3	4	5	6	7 7
		- eye;	21.10.53			
		Powders and granules	21.10.53.120			
		(microgranules, pellets):	21.10.54			
		- for preparation solution for	21.10.54.110			
		injection;	21.10.54.120			
		- for external use (when applied to	21.10.54.130			
		wounds)	21.10.54.140			
120	STATE PHARMACOPOEIA OF THE	Drugs:	21.10.54.150		Microbiological purity:	Pass test/fail test (if necessary,
	REPUBLIC OF BELARUS art. 2.6.12	Solutions:	21.10.54.160		Total number of aerobic	specify conditions)
	and other normative documents	- for oral use;	21.10.54.170		bacteria;	
	approved in the established order,	- for external use;	21.10.54.180		Total number of fungi;	
	specifying the application of the research	- for intrauterine administration;	21.10.54.190		Enterobacteria, etc.	
	(testing) method, measurements,	- for local application;	21.20.1		gram-negative bacteria;	
	establishing requirements for drugs	Suspensions and emulsions:	21.20.10		E. coli;	
	registered in the established order and	- for oral use;	21.20.10.158		Pseudomonas	
	included in State registers of drug for	- for external use;	21.20.10.159		aeruginosa;	
	veterinary use of the Eurasian Economic	- for intrauterine administration;	21.20.10.213		Staphylococcus	
	Union member states	- for local application;	21.20.21.130		aureus;	
121	STATE PHARMACOPOEIA OF THE	Drops:	21.20.21.139		Salmonella	
	REPUBLIC OF BELARUS art. 2.6.13	- Ear;	02.30.40.140			
	and other normative documents	- nasal;				
	approved in the established order,	- for local application;				
	specifying the application of the research	- for oral use;				
	(testing) method, measurements,	Ointments (Gels, creams, liniments,				
	establishing requirements for drugs	pastes);				
	registered in the established order and	- for external use;				
	included in State registers of drug for	- for local application;				
	veterinary use of the Eurasian Economic	Powders and granules				
	Union member states	(microgranules, pellets):				
122	STATE PHARMACOPOEIA OF THE	- for preparation (solution for oral				
	REPUBLIC OF BELARUS art. 2.6.31	use, drops);				
	and other normative documents	- for oral use;				
	approved in the established order,	- for external use;				
	specifying the application of the research	- for local application;				
	(testing) method, measurements,	Aerosols and sprays;				
	establishing requirements for drugs	Capsules;				
	registered in the established order and	Suppositories (sticks);				
	included in State registers of drug for	Tablets, dragee, briquettes, pastilles;				
	veterinary use of the Eurasian Economic	Tinctures and Extracts:				
	Union member states	- for oral use;				

1	2	3	4	5	6	7 243 pages,
123	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 5.1.4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 5.1.8 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	- for external use; - for local application; Syrups; Balsams; System: - Intravaginal administration Pharmaceutical substances				
125	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.7.2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for intracisternal injection; - for local application; Suspensions and emulsions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for intracisternal injection; - for local application; Drops; Ointments; Powders and granules (microgranules, pellets): - for preparation (solution for	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53 21.10.54 21.10.54.110 21.10.54.120	3003 – 3004	Determination of the antimicrobial activity of antibiotics by diffusion in agar/ Quantitative determination / quantitative content	0,001-1000000 mkg/kg (mg; g; 10 mg; 100g; ml; cm3; 10 ml; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle; plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (mg; g; 10 g; 100 g; ml; cm3; l; dm3; 10 ml; 100 ml; tabl; capsule; pipette; syringe; bottle; plate; suppository; stick; packing; bag); 0,00001-1000,0 g/kg (g; 10 g; 100 g; ml; 10 ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle; plate; suppository; stick; package, bag) 0,1 – 100000000 IU/kg (mg; g; 10 g; 100g; ml; 10 ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle, plate, suppository);

1	2	3	4	5	6	7
126	STATE PHARMACOPOEIA OF THE	injection, for oral use, drops); - for oral use; - for external use; - for local application; Lyophilizate; Tablets; Pharmaceutical substances. Drugs:	21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140		Bacterial endotoxins	0,1 – 100000000 Unit/kg (mg; g; 10 g; 100g; ml; 10 ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository) 0,00001-150%, 1,0 – 200,0 % of declared; not found Pass test/fail test; Compliant/ Non
	REPUBLIC OF BELARUS art. 2.6.14 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Solutions: - for injection; - for infusion; Suspensions and emulsions: - for injection; Powders and granules: - for preparation solution for injection. Pharmaceutical substances		21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140		
127	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.2 and other normative documents approved in the established order, specifying the application of the research	Drugs: Tablets, dragee, briquettes, pastilles; Capsules; Suppositories; Powders and granules (microgranules, pellets); System: - Intravaginal administration Drugs: Tablets, dragee, briquettes, pastilles; Capsules; Suppositories; Powders and granules.	21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120	3004	Disintegration	0,5 -120 minutes; Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)

1	2	3	4	5	6	7
129	(testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements,	Drugs: Tablets, dragee, briquettes, pastilles; Capsules; Suppositories; Powders and granules (microgranules, pellets);	4 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 02.30.40.140	5	Dissolution Average mass / mass uniformity	0,1 – 120% of declared; Compliant/ Non compliant; Pass test/ fail test 0,1 – 10 kg; 0,001 – 500 g; 1,0 – 5000 mg; 0,01 – 50 % of average mass; Compliant/ Non compliant; Pass test/ fail test
	establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	System: - Intravaginal administration Drugs: Tablets, dragee, briquettes, pastilles; Capsules; Suppositories; Powders and granules; Drops				
131	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic	Drugs: Tablets and dragee, briquettes, pastels; Capsules; Suppositories; Powders and granules (microgranules, pellets); Drops (eye); Ointments; Cornea (eye);	21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129		Dosage uniformity	Compliant/ Non compliant (if necessary, specify conditions); Pass test/ fail test (if necessary, specify conditions)

1	2	3	4	5	6	7
132	Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.40 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS monograph 0672 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic	Aerosols and sprays; System: - Intravaginal administration Drugs: Tablets and dragee, briquettes, pastels; Capsules; Suppositories; Powders and granules (microgranules, pellets); Drops (eye); Ointments; Cornea (eye);	4 21.10.52.110 21.10.53 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 02.30.40.140	5	6	7 7
	included in State registers of drug for	Ointments;			Sedimentary stability	0,5 – 120 minutes; Compliant/
		Suspensions; Drops (eye)			Resuspension ability	Non compliant; Pass test/ fail test 0,5 – 120 minutes; Compliant/ Non compliant; Pass test/ fail test

1	2	3	4	5	6	7 7
134	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.27 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Drops: - eye; - Ear; - nasal; - sublingual; - for local application; - for oral use; Aerosols and sprays; Ointments; Cornea (eye)			Dosage mass uniformity / mass uniformity	0,01 – 50 % of average mass; Compliant/ Non compliant; Pass test/ fail test
135	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.10 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for local application; Tinctures and Extracts: - for oral use; - for external use; - for local application; Syrups	21.10.51.120 21.10.52.110 21.10.54 21.10.54 21.20.1 21.20.10 02.30.40.140	3004	Quantitative determination of ethyl alcohol / quantitative content / mass fraction / mass concentration / Volume concentration	0,01-96 % (mass, volume); g/l (dm3, cm3, ml); mg/ml (cm3)
136	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.12 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.35 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and	Drugs: Powders and granules.	21.10.32 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1	3004	Determination of fractional composition / particle size distribution/ particle size	45 μm – 11,2 mm

1	2	3	4	5	6	7 7
	included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.20.10			
138	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.38 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					10.100001 (3
139	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.34 (Method 1) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bulk density	10-10000 kg/m ³ ; 0,01 – 10 g/ml
140	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.9.20 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Liquid and solid parenteral dosage forms, eye dosage forms	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120	3004	Mechanical inclusions / presence of mechanical inclusions	Absent/ Present; Pass test/ fail test (if necessary, specify conditions)

1	2	3	4	5	6	7 243 pages.
			21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.213 21.20.21.130 21.20.21.139			
141	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS monograph 0132 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Ointments	21.10.54 21.10.54.180 21.20.1 21.20.10	3004	Uniformity	Uniform/non uniform; Compliant/ Non compliant; Pass test/ fail test
142	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS monograph 1433 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products. Feed additives	21.1 21.10 21.10.53 21.10.53.120 21.20.1 21.20.10 02.30.40.140 10.91.10.230	3003 – 3004; 2309	Content of pharmacologically active substances or biological activity	10 ⁻⁸ – 10,0 %; 10 ⁻⁸ – 0,1 %/tab (bottle); 10 ⁻⁸ – 10,0 mg/g (mg; cm3; ml; dm3; l); 10 ⁻³ – 100 mcg/g (mg; cm3; ml; dm3; l)
	veterinary use of the Eurasian Economic Union member states; and other normative documents approved in the established order that specify the application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and included in the State registries	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products.			Impurity content	0,1-20%; Pass test/ fail test; Presence/absence (if necessary, specify conditions)

1	2	3	4	5	6	7 7
	of feeds and feed additives of the Eurasian Economic Union member states					
143	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS monograph 0765 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; and other normative documents approved in the established order that specify the application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and included in the State registries of feeds and feed additives of the Eurasian Economic Union member states	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products. Feed additives			Content of pharmacologically active substances or biological activity	10 ⁻⁸ – 10,0 %; 10 ⁻⁸ – 0,1 %/tab(bottle); 10 ⁻⁸ – 10,0 mg/g (mg; cm3; ml; dm3; l); 10 ⁻³ – 100 mcg/g (mg; cm3; ml; dm3; l)
144	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.8.16 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA OF THE	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products.			Mass fraction of dry residue / dry residue	0,001 – 10% 0,001 – 50%
	REPUBLIC OF BELARUS art. 2.8.17 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs				drying / humidity determination / mass fraction of humidity / humidity	5,551

1	2	3	4	5	6	7 7
	registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
146	STATE PHARMACOPOEIA OF THE REPUBLIC OF BELARUS art. 2.8.2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Impurity content	0,1-20%; Pass test/ fail test; Presence/absence (if necessary, specify conditions)
147	STATE PHARMACOPOEIA, XIII edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.1.0001.15 and other normative documents approved in the established order, specifying the	Drugs: Solutions; Drops; Aerosols and sprays; Suspensions and emulsions; Powders and granules;	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5	3003-3004	Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Suppositories (sticks) Tablets, dragee, briquettes, pastilles; Capsules; Ointments; Lyophilizate; Pharmaceutical substances. Medicinal plant raw material and collections.	21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53 21.10.54.120 21.10.54.130 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.170 21.10.54.170		Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g;

						243 pages									
1	2	3	4	5	6	7									
148	STATE PHARMACOPOEIA, XIII edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.1.0002.15 and other normative documents approved in the		21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139		Authenticity	100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 1000000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,0000; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 - 200,0 % of									

1	2	3	4	5	6	243 pages
1		3	+		0	declared; not found
149	STATE PHARMACOPOEIA, XIII edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.1.0009.15 and other normative documents approved in the established order, specifying the				Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found

1	2	3	4	5	6	7 7
150	STATE PHARMACOPOEIA, XIII edition, GENERAL PHARMACOPOEIA ARTICLE.1.1.0015.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Powders and granules.	21.1 21.10 21.10.32 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1	3003-3004	Determination of fractional composition / particle size distribution/ particle size	45 μm – 11,2mm
151	STATE PHARMACOPOEIA, XIII edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0012.15 (Method 1, Method 5, Method 7) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions; Lyophilizate.	21.20 21.20.1	3004	Quantitative determination of protein / quantitative content / mass fraction / mass concentration	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,2 - 2,0 mg/ml
152	STATE PHARMACOPOEIA, XIII edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0019.15 and other normative documents approved in the	Drugs: Tinctures; Extracts; Solutions	21.10.53 21.10.53.120 21.20.1 21.20.10 02.30.40.140	3003-3004	Mass fraction of dry residue / dry residue	0,001 – 10%

1	2	3	4	5	6	7 243 pages.
150	established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
153	STATE PHARMACOPOEIA, XIII edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0021.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
154	STATE PHARMACOPOEIA, XIII edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0006.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for intracisternal injection; - for local application; - for infusion; Suspensions and emulsions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for intracisternal injection; - for local application; Drops: - eye; - Ear; - nasal; - sublingual	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53 21.10.54 21.10.54.110 21.10.54.130	3003-3004	Color / colour / colour solution / coloration / solution coloration (description)	-

1	2	3	4	5	6	7 243 pages
		- for local application;	21.10.54.140			
		- for oral use.	21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.158			
			21.20.10.159			
			21.20.10.213			
			21.20.21.130			
			21.20.21.139			
			02.30.40.140			
155	USP, (201) and other normative	Drugs.	21.1	3003 –	Authenticity	Compliant/ Non compliant; Pass
	documents approved in the established	Medicinal plant raw material and	21.10	3004		test/fail test (if necessary,
	order, specifying the application of the	collections.	21.10.1			specify conditions)
	research (testing) method,	Pharmaceutical substances.	21.10.20.120			
	measurements, establishing requirements		21.10.32			
	for drugs registered in the established		21.10.5			
	order and included in State registers of		21.10.51.120			
	drug for veterinary use of the Eurasian		21.10.51.121			
	Economic Union member states		21.10.51.122			
			21.10.51.123			
156	USP, (203) and other normative		21.10.51.124			
	documents approved in the established		21.10.51.125			
	order, specifying the application of the		21.10.51.126			
	research (testing) method,		21.10.51.129			
	measurements, establishing requirements		21.10.52.110			
	for drugs registered in the established		21.10.53			
	order and included in State registers of		21.10.53.120			
	drug for veterinary use of the Eurasian		21.10.54			
	Economic Union member states		21.10.54.110			
157	USP, (1064) and other normative		21.10.54.120			
	documents approved in the established		21.10.54.130			
	order, specifying the application of the		21.10.54.140			
	research (testing) method,		21.10.54.150			
	measurements, establishing requirements		21.10.54.160			
	for drugs registered in the established		21.10.54.170			
	order and included in State registers of		21.10.54.180			
	drug for veterinary use of the Eurasian		21.10.54.190			

1	2	3	4	5	6	7 243 pages.
	Economic Union member states		21.20.1			
			21.20.10			
1.70			21.20.10.158			
158	USP, (621) and other normative		21.20.10.159			
	documents approved in the established		21.20.10.213			
	order, specifying the application of the		21.20.21.130			
	research (testing) method,		21.20.21.139			
	measurements, establishing requirements		02.30.40.140			
	for drugs registered in the established					
	order and included in State registers of					
	drug for veterinary use of the Eurasian					
150	Economic Union member states	D	21.10.5	2004	N. 1 . 1 . 1 /	A1
159	USP, (790) and other normative	Drugs:	21.10.5	3004	Mechanical inclusions /	Absent/ Present; Pass test/ fail
	documents approved in the established	Liquid and solid parenteral dosage	21.10.51.120		presence of mechanical inclusions	test (if necessary, specify
	order, specifying the application of the	forms, eye dosage forms	21.10.51.121		inclusions	conditions)
	research (testing) method,		21.10.51.122			
	measurements, establishing requirements		21.10.51.123			
	for drugs registered in the established		21.10.51.124 21.10.51.125			
	order and included in State registers of		21.10.51.125			
	drug for veterinary use of the Eurasian Economic Union member states		21.10.51.126			
	Economic Union member states		21.10.51.129			
			21.10.52.110			
			21.10.53			
			21.10.54			
			21.10.54.110			
			21.10.54.110			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.213			
			21.20.21.130			
			21.20.21.139			
160	USP, (1) and other normative documents	Drugs:	21.10.51.120	3004	Resuspension ability	0,5 – 120 minutes
	approved in the established order,	Suspensions;	21.10.52.110		1	,
	specifying the application of the research	Drops (eye)	21.10.54			

1	2	3	4	5	6	7 243 pages
	(testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.10.54.180 21.20.1 21.20.10			
161	USP, (1151) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
162	USP, (755) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Drops: - eye; - Ear; - nasal; - sublingual; - for local application; - for oral use; Suspensions and emulsions; Aerosols, sprays, foams; Tinctures and Extracts: - for oral use; - for external use; - for local application; Ointments; Syrups; Balsams; Powders and granules	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.125 21.10.51.126 21.10.51.129 21.10.51.129 21.10.53 21.10.53 21.10.53 21.10.54 21.10.54 21.10.54.110 21.10.54.130 21.10.54.150 21.10.54.150 21.10.54.170 21.10.54.170 21.10.54.170 21.10.54.180 21.10.54.190	3004	Weight (volume) of package contents	0,1 – 25000 ml (cm3; 1; dm3); 0,1 – 10 kg; 0,001 – 500 g; 1,0 – 5000 mg; Compliant/ Non compliant; Pass test/ fail test

1	2	3	4	5	6	7 7
			21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140			
163	USP, (603) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Aerosols, sprays, foams	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120 21.10.54 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	3004	Number of doses per package	0-1000; Compliant/non compliant; Pass test/ fail test

1	2	3	4	5	6	7 7
			02.30.40.140			
164	USP, (561) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and	21.1 21.10 21.10.53 21.10.53.120 21.20.1	3003 – 3004; 2308	21.10 3004; 21.10.53 2308 Prese 21.10.53.120 specifi	0,1-20%; Pass test/ fail test; Presence/absence (if necessary, specify conditions)
	for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	collections. Pharmaceutical substances of vegetable products.	21.20.10 02.30.40.140		Weight loss during drying / humidity determination / mass fraction of humidity	0,001 – 50%
165	USP, (565) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Mass fraction of dry residue / dry residue	0,001 – 10%
166	USP, (611) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for local application; Tinctures and Extracts: - for oral use; - for external use; - for local application; Syrups	21.10.51.120 21.10.52.110 21.10.54 21.10.54.180 21.20.1 21.20.10 02.30.40.140	3004	Quantitative determination of ethyl alcohol / quantitative content / mass fraction / mass concentration / Volume concentration	0,01-96 % (mass, volume); g/l (dm3, cm3, ml); mg/ml (cm3)

1	2	3	4	5	6	7 7
167	USP, (401) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions; Ointments; Drops (eye); Extracts	21.10.51.120 21.10.52.110 21.10.54 21.10.54.180 21.20.1 21.20.10 02.30.40.140	3004	Acid value	0,01 -10 mg KOH/ml (cm3;g)
168	USP, (541) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for local application; - for infusion; Drops: - eye; - Ear; - nasal; - for local application; - for oral use; Powders and granules (microgranules, pellets); Tablets; Ointments; Aerosols and sprays; Tinctures and Extracts; Syrups; Balsams; Drug checker; Cord. Pharmaceutical substances.	21.1 21.10 21.10.20.120 21.10.51.120 21.10.54 21.10.54.180 21.20.1 21.20.10 21.20.10.213 21.20.10.158 21.20.10.159 21.20.21.130 21.20.21.139 21.20.21.4 21.20.14.000 02.30.40.140	3004	Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,000001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-10,0 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag) 0,01 -10 mg KOH/g (cm3;g); 0,0001-150%; 0,0001-150%; weight; 0,0001-150% volume; not found
169	USP, (786) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements	Drugs: Powders and granules.	21.10.32 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123	3004	Determination of fractional composition / particle size distribution/ particle size	45 μm – 11,2mm

1	2	3	4	5	6	7 7
170	for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states USP, (811) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states USP, (616) Metod 1 and other normative documents approved in the established	3	4 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1	5	Bulk density	
172	order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states USP, (1103) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; Suspensions and emulsions: - for injection; Powders and granules (microgranules, pellets): - for preparation (solution for injection, drops); Lyophilizate	21.10.52.110 21.20.1 21.20.10 21.20.10.213	3004	Authenticity, Quantitative determination (quantitative content, mass fraction; mass concentration) of active substance	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; package; bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; package; bag); 0,1 – 1000000000 IU/kg (mg; g;
173	USP, (631) and other normative documents approved in the established	Drugs: Solutions:	21.1 21.10	3003-3004	Degree of liquids coloration / color / colour	100g; ml; cm3; l; dm3; 100 ml); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml); not found

1	2	3	4	5	6	7
	order, specifying the application of the	- for injection;	21.10.1		/ colour solution /	
	research (testing) method,	- for oral use;	21.10.20.120		coloration / solution	
	measurements, establishing requirements	- for external use;	21.10.32		coloration	
	for drugs registered in the established	- for intrauterine administration;	21.10.5		(description)	
	order and included in State registers of	- for intracisternal injection;	21.10.51.120			
	drug for veterinary use of the Eurasian	- for local application;	21.10.51.121			
	Economic Union member states	- for infusion;	21.10.51.122			
		Suspensions and emulsions:	21.10.51.123			
		- for injection;	21.10.51.124			
		- for oral use;	21.10.51.125			
		- for external use;	21.10.51.126			
		- for intrauterine administration;	21.10.51.129			
		- for intracisternal injection;	21.10.52.110			
		- for local application;	21.10.53			
		Drops:	21.10.53.120			
		- eye;	21.10.54			
		- Ear;	21.10.54.110			
		- nasal;	21.10.54.120			
		- sublingual	21.10.54.130			
		- for local application;	21.10.54.140			
		- for oral use.	21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.158			
			21.20.10.159			
			21.20.10.213			
			21.20.21.130			
			21.20.21.139			
			02.30.40.140			
174	USP, (197) and other normative	Drugs:	21.1	3003-3004	Authenticity	Compliant/ Non compliant; Pass
	documents approved in the established	Solutions;	21.10			test/ fail test (if necessary,
	order, specifying the application of the	Drops;	21.10.1			specify conditions)
	research (testing) method,	Aerosols and sprays;	21.10.20.120			
	measurements, establishing requirements	Suspensions and emulsions;	21.10.32			

1	2	3	4	5	6	7
175	for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states USP, (854) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Powders and granules; Suppositories (sticks) Tablets, dragee, briquettes, pastilles; Capsules; Ointments; Lyophilizate; Pharmaceutical substances. Medicinal plant raw material and collections.	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.140 21.10.54.150 21.10.54.140 21.10.54.150 21.10.54.190 21.20.10 21.20.10 21.20.10 21.20.10 21.20.10.158 21.20.21.130 21.20.21.139 02.30.40.140	5	Quantitative determination / quantitative content / mass fraction / mass concentration Authenticity Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% wolume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet;

1	2	2	1		6	243 pages
1	2	3	4	5	6	
176	USP, (1119) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Authenticity Quantitative determination / quantitative content / mass fraction / mass concentration	package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-10000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 10000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; lo0 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository);

1	2	3	4	5	6	7 7
177	USP, (1120) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		4	5	Authenticity Quantitative determination / quantitative content / mass fraction / mass concentration	7 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag);
						package, odg), 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found

1	2	3	4	5	6	7
178	EP, art. 2.2.27 and other normative	Drugs.	21.1	3003 -	Authenticity	Compliant/ Non compliant; Pass
	documents approved in the established	Medicinal plant raw material and	21.10	3004	,	test/ fail test (if necessary,
	order, specifying the application of the	collections.	21.10.1	from 4201		specify conditions)
	research (testing) method,	Pharmaceutical substances.	21.10.20.120			
	measurements, establishing requirements		21.10.32			
	for drugs registered in the established		21.10.5			
	order and included in State registers of		21.10.51.120			
	drug for veterinary use of the Eurasian		21.10.51.121			
	Economic Union member states		21.10.51.122			
179	EP, art. 2.2.6 and other normative	Drugs.	21.10.51.123		Refractive index /	1,3-1,7;
	documents approved in the established	Medicinal plant raw material and	21.10.51.124		Quantitative	0,0001 - 500 g/ml; mg/ml; g/l;
	order, specifying the application of the	collections.	21.10.51.125		determination	mg/l; g/cm ³ ; mg/cm ³
	research (testing) method,	Pharmaceutical substances.	21.10.51.126			
	measurements, establishing requirements		21.10.51.129			
	for drugs registered in the established		21.10.52.110			
	order and included in State registers of		21.10.53			
	drug for veterinary use of the Eurasian		21.10.53.120			
	Economic Union member states		21.10.54			
			21.10.54.110			
			21.10.54.120			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.158			
			21.20.10.159			
			21.20.10.213			
			21.20.21.130			
			21.20.21.139			
			02.30.40.140			
105		-	15.12.11	2001	3	
180	EP, art. 2.9.20 and other normative	Drugs:	21.10.5	3004	Mechanical inclusions /	Absent/ Present; Pass test/ fail
	documents approved in the established	Liquid and solid parenteral dosage	21.10.51.120		presence of mechanical	test (if necessary, specify
	order, specifying the application of the	forms, eye dosage forms	21.10.51.121		inclusions	conditions)
	research (testing) method,		21.10.51.122			
	measurements, establishing requirements		21.10.51.123			
	for drugs registered in the established		21.10.51.124			

1	2	3	4	5	6	243 pages
	order and included in State registers of		21.10.51.125			
	drug for veterinary use of the Eurasian		21.10.51.126			
	Economic Union member states		21.10.51.129			
			21.10.52.110			
			21.10.53			
			21.10.53.120			
			21.10.54			
			21.10.54.110			
			21.10.54.120			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.213			
			21.20.21.130			
			21.20.21.139			
181	EP, art. 2.9.12 and other normative	Drugs:	21.10.32	3004	Determination of	45 μm – 11,2 mm
101	documents approved in the established	Powders and granules.	21.10.51.120		fractional composition /	11,2
	order, specifying the application of the	1 0 N dots and grandles.	21.10.51.121		particle size distribution/	
	research (testing) method,		21.10.51.122		particle size	
	measurements, establishing requirements		21.10.51.123		Financia seri	
	for drugs registered in the established		21.10.51.124			
	order and included in State registers of		21.10.51.125			
	drug for veterinary use of the Eurasian		21.10.51.126			
	Economic Union member states		21.10.51.129			
182	EP, art. 2.9.35 and other normative		21.10.54.110			
102	documents approved in the established		21.10.54.120			
	order, specifying the application of the		21.10.54.130			
	research (testing) method,		21.10.54.140			
	measurements, establishing requirements		21.10.54.150			
	for drugs registered in the established		21.10.54.160			
	order and included in State registers of		21.10.54.170			
	drug for veterinary use of the Eurasian		21.10.54.180			
	Economic Union member states		21.10.54.190			
183	EP, art. 2.9.38 and other normative		21.20.1			
	documents approved in the established		21.20.10			
	order, specifying the application of the					

1	2	3	4	5	6	7 7 243 pages,
	research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
184	EP, art. 2.9.34 (Method 1) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bulk density	10-10000 kg/m ³ ; 0,01 – 10 g/ml
185	EP, art. 2.8.16 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products.	21.1 21.10 21.10.53 21.10.53.120 21.20.1 21.20.10 02.30.40.140	3003 – 3004	Mass fraction of dry residue / dry residue	0,001 – 10%
186	EP, art. 2.5.8 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53	3003-3004	Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml;

1	2	3	4	5	6	7 243 pages,
			21.10.54 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140			tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
187	EP, art. 2.2.2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for intracisternal injection; - for local application; - for infusion; Suspensions and emulsions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for intracisternal injection; - for local application; Drops: - eye; - Ear; - nasal; - sublingual - for local application; - for oral use.	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.125 21.10.51.125 21.10.51.126 21.10.51.129 21.10.53 21.10.53 21.10.53 21.10.54 21.10.54.110 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.150 21.10.54.160	3003-3004	Degree of liquids coloration / color / colour / colour solution / coloration / solution coloration (description)	-

documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states Suppositories (sticks) Tablets, dragee, briquettes, pastilles; Lyophilizate; Pharmaceutical substances. Medicinal plant raw material and collections. Medicinal plant raw material and collections. Solutions; 21.10.20.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.125 21.10.51.126 21.10.51.126 21.10.51.126 21.10.51.127 21.10.51.129 21.10.51.129 21.10.52.110 21.10.53.120 21.10.54.140 21.10.54.160 Solutions; 21.10 determination / quantitative content / mass fraction	1	2	3	4	5	6	7
21.10.54.170 21.10.54.180 21.10.54.190 tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%,	188	EP, art. 2.2.24 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian	Drugs: Solutions; Drops; Aerosols and sprays; Suspensions and emulsions; Powders and granules; Suppositories (sticks) Tablets, dragee, briquettes, pastilles; Capsules; Ointments; Lyophilizate; Pharmaceutical substances. Medicinal plant raw material and	21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140 21.1 21.10 21.10.2 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.125 21.10.51.126 21.10.51.129 21.10.53 21.10.53 21.10.53 21.10.54 21.10.54 21.10.54.140 21.10.54.150 21.10.54.150 21.10.54.170 21.10.54.170 21.10.54.170 21.10.54.170 21.10.54.170 21.10.54.170 21.10.54.170		Authenticity Quantitative determination / quantitative content / mass fraction / mass	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository);

-	2		4	_		243 pages,
1	2	3	4	5	6	7
			21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140			declared; not found
189	EP, art 2.2.40 and other normative documents approved in the established order, specifying the application of the				Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 - 200,0 % of declared; not found

1	2	3	4	5	6	7 243 pages,
190	EP, art. 2.2.48 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian				Authenticity Quantitative determination / quantitative content /	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl
	Economic Union member states				quantitative content / mass fraction / mass concentration	capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% wolume, 1,0 – 200,0 % of declared; not found
191	BP, Appendix III A and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)

1	2	3	4	5	6	7
	drug for veterinary use of the Eurasian Economic Union member states					
192	BP, Appendix V E and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53 21.10.53.120 21.10.54 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139	3003 - 3004	Refractive index / Quantitative determination	1,3 – 1,7; 0,0001 – 500 g/ml; mg/ml; g/l; mg/l; g/cm³; mg/cm³
193	BP, Appendix XIII B and other normative documents approved in the established order, specifying the application of the research (testing)	Drugs: Liquid and solid parenteral dosage forms, eye dosage forms	02.30.40.140 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122	3004	Mechanical inclusions / presence of mechanical inclusions	Absent/ Present;; Pass test/ fail test (if necessary, specify conditions)

1	2	3	4	5	6	7 243 pages
	method, measurements, establishing		21.10.51.123			
	requirements for drugs registered in the		21.10.51.124			
	established order and included in State		21.10.51.125			
	registers of drug for veterinary use of the		21.10.51.126			
	Eurasian Economic Union member		21.10.51.129			
	states		21.10.52.110			
			21.10.53			
			21.10.53.120			
			21.10.54			
			21.10.54.110			
			21.10.54.120			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.213			
			21.20.21.130			
			21.20.21.130			
			21.20.21.137			
194	BP, Appendix XVII A and other	Drugs:	21.10.32	3004	Determination of	45 μm – 11,2 mm
	normative documents approved in the	Powders and granules.	21.10.51.120		fractional composition /	•
	established order, specifying the		21.10.51.121		particle size distribution/	
	application of the research (testing)		21.10.51.122		particle size	
	method, measurements, establishing		21.10.51.123			
	requirements for drugs registered in the		21.10.51.124			
	established order and included in State		21.10.51.125			
	registers of drug for veterinary use of the		21.10.51.126			
	Eurasian Economic Union member		21.10.51.129			
	states		21.10.54.110			
195	BP, Appendix XVII B and other		21.10.54.120			
	normative documents approved in the		21.10.54.130			
	established order, specifying the		21.10.54.140			
	application of the research (testing)		21.10.54.150			
	method, measurements, establishing		21.10.54.160			
	requirements for drugs registered in the		21.10.54.170			
	established order and included in State		21.10.54.170			
	registers of drug for veterinary use of the		21.10.54.190			
	registers of drug for veterinary use of the		21.10.54.130			

1	2	3	4	5	6	7 243 pages
	Eurasian Economic Union member states		21.20.1 21.20.10			
196	BP, Appendix XVII S, Metod 1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bulk density	10-10000 kg/m³; 0,01 – 10 g/ml
197	BP, General notices, Part II and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Extracts; Powders and granules; Pharmaceutical substances.	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.51.129 21.10.52.110 21.10.53 21.10.53 21.10.54 21.10.54 21.10.54.120 21.10.54.130 21.10.54.150 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10	3003 – 3004	Solubility	Pass test/ fail test (if necessary, specify conditions)

1	2	3	4	5	6	7 243 pages
			21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140			
198	BP, Appendix XI P and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products.	21.1 21.10 21.10.53 21.10.53.120 21.20.1 21.20.10 02.30.40.140	3003 – 3004; 2308	Mass fraction of dry residue / dry residue	0,001 – 10%
199	BP, Appendix XI Q and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Weight loss during drying / humidity determination / mass fraction of humidity / humidity	0,001 - 50%
200	BP, Herbal Drug Extracts and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; and other normative documents approved in the established order that specify the application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and included in the State registries of feeds and feed additives of the	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products. Feed additives	21.1 21.10 21.10.53 21.10.53.120 21.20.1 21.20.10 02.30.40.140 10.91.10.230	3003 – 3004; 2308-2309	Content of pharmacologically active substances or biological activity	10 ⁻⁸ – 10,0 %; 10 ⁻⁸ – 0,1 %/tab(bottle); 10 ⁻⁸ – 10,0 mg/g (mg; cm3; ml; dm3; l); 10 ⁻³ – 100 mcg/g (mg; cm3; ml; dm3; l)

				T	Г	243 pages
1	2	3	4	5	6	7
	Eurasian Economic Union member					
	states					
201	BP, Herbal Drugs and other normative	Drugs:	-			
201	documents approved in the established	Tinctures and Extracts;				
	order, specifying the application of the	Syrups;				
	research (testing) method,	Balsams.				
	measurements, establishing requirements	Medicinal plant raw material and				
	for drugs registered in the established	collections.				
	order and included in State registers of	Pharmaceutical substances of				
	drug for veterinary use of the Eurasian	vegetable products.				
	Economic Union member states;	Feed additives				
	and other normative documents	Drugs:			Impurity content	0,1-20%;
	approved in the established order that	Tinctures and Extracts;				Pass test/ fail test;
	specify the application of the research	Syrups;				Presence/absence (if necessary,
	(testing) method, measurements that	Balsams.				specify conditions)
	establish requirements for feed and feed	Medicinal plant raw material and				
	additives registered in the established	collections.				
	order and included in the State registries	Pharmaceutical substances of				
	of feeds and feed additives of the	vegetable products.				
	Eurasian Economic Union member					
202	states	Discourse	_		Town of a sentent	0.1.2007
202	BP, Appendix XI D and other normative documents approved in the established	Drugs: Tinctures and Extracts;			Impurity content	0,1-20%; Pass test/ fail test;
	order, specifying the application of the	Syrups;				Presence/absence (if necessary,
	research (testing) method,	Balsams.				specify conditions)
	measurements, establishing requirements	Medicinal plant raw material and				specify conditions)
	for drugs registered in the established	collections.				
	order and included in State registers of	Pharmaceutical substances of				
	drug for veterinary use of the Eurasian	vegetable products.				
	Economic Union member states	5 r r				
203	BP, Appendix VII (Method A, Method	Drugs.	21.1	3003-3004	Content of heavy metals	(0.002 - 500) mg/kg, mg/dm ³
	B, Limit Test for Heavy Metals in	Medicinal plant raw material and	21.10		(cadmium, lead, arsenic,	
	Herbal Drugs and Herbal Drug	collections.	21.10.1		Mercury)	
	Preparations) and other normative	Pharmaceutical substances.	21.10.20.120			
	documents approved in the established		21.10.32			
	order, specifying the application of the		21.10.5			
	research (testing) method,		21.10.51.120			
	measurements, establishing requirements		21.10.51.121			
	for drugs registered in the established		21.10.51.122			
	order and included in State registers of		21.10.51.123			
	drug for veterinary use of the Eurasian		21.10.51.124			

1	2	3	4	5	6	7 243 pages
	Economic Union member states		21.10.51.125			
			21.10.51.126			
			21.10.51.129			
			21.10.52.110			
			21.10.53			
			21.10.53.120			
			21.10.54			
			21.10.54.110			
			21.10.54.120			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.158			
			21.20.10.159			
			21.20.10.213			
			21.20.21.130			
			21.20.21.130			
			02.30.40.140			
204	BP, Appendix X B and other normative	Drugs:	21.10.51.120	3004	Acid value	0,01 -10 mg KOH/ml (cm3;g)
204	documents approved in the established	Solutions;	21.10.51.120	3004	Acid value	
	order, specifying the application of the	Ointments;	21.10.54			
	research (testing) method,	Drops (eye);	21.10.54.180			
	measurements, establishing requirements	Extracts	21.20.1			
	for drugs registered in the established	Extracts	21.20.10			
	order and included in State registers of		02.30.40.140			
	drug for veterinary use of the Eurasian		02.30.40.140			
	Economic Union member states					
205	BP, Appendix X F and other normative				Peroxide value	0,01-50 mmol O2/kg
203	documents approved in the established				1 CIUNIUE VAIUE	0,01-30 Hillof O2/kg
	order, specifying the application of the					
	research (testing) method,					
	measurements, establishing requirements					
	for drugs registered in the established					
	order and included in State registers of					
	drug for veterinary use of the Eurasian					
	Economic Union member states					

1	2	3	4	5	6	7
207	BP, Appendix VI and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states BP, Appendix VIII B (Potentiometric Titration, Determination of Primary Aromatic Amino-nitrogen) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53 21.10.54 21.10.54 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.150 21.10.54.190 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213	5 3003-3004	Quantitative determination / quantitative content / mass fraction / mass concentration	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,000001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-10,0 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag) 0,01 -10 mg KOH/g (cm3;g); 0,0001-150%, 0,0001-150% weight, 0,0001-150% volume, not found
			21.20.21.130 21.20.21.139 02.30.40.140			
208	BP, Appendix XIV B and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State	Drugs: Solutions: - for injection; Suspensions and emulsions: - for injection; Powders and granules (microgranules, pellets):	21.10.52.110 21.20.1 21.20.10 21.20.10.213	3004	Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; package; bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; package; bag); 0,00001-1000 g/kg (g; 100 g; ml;

1	2	3	4	5	6	7 7
	registers of drug for veterinary use of the Eurasian Economic Union member states	- for preparation (solution for injection, drops); Lyophilizate				cm3; l; dm3; 100ml; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml); not found
209	BP, AppendixV III F and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for local application; Tinctures and Extracts: - for oral use; - for external use; - for local application; Syrups	21.10.51.120 21.10.52.110 21.10.54 21.10.54.180 21.20.1 21.20.10 02.30.40.140	3004	Quantitative determination of ethyl alcohol / quantitative content / mass fraction / mass concentration / Volume concentration	0,01-96 % (mass, volume); g/l (dm3, cm3, ml); mg/ml (cm3)
210	BP, Appendix II A and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions; Drops; Aerosols and sprays; Suspensions and emulsions; Powders and granules; Suppositories (sticks) Tablets, dragee, briquettes, pastilles; Capsules; Ointments; Lyophilizate; Pharmaceutical substances. Medicinal plant raw material and collections.	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53 21.10.54.120 21.10.54.130 21.10.54.130 21.10.54.140 21.10.54.150	3003-3004 2308	Authenticity Quantitative determination / quantitative content / mass fraction / mass concentration	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 00,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 1000000000 Unit/kg (mg; g; 0,1 - 1000000000 Unit/kg (mg; 0,1 - 10000000000 Unit/kg (mg; 0,1 - 1000000000 Unit/kg (mg; 0,1 - 100000

1	2	3	4	5	6	7 7
211	BP, Appendix II H and other normative		21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130		Authenticity	100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Pass
	documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.20.21.139 02.30.40.140		Quantitative determination / quantitative content / mass fraction / mass concentration	test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 100000000 IU/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; table; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; table; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 - 200,0 % of declared; not found

1	2	3	4	5	6	7
212	BP, Appendix IV B and other normative	Drugs:	21.1	3003-3004	Degree of liquids	-
	documents approved in the established	Solutions:	21.10		coloration / color / colour	
	order, specifying the application of the	- for injection;	21.10.1		/ colour solution /	
	research (testing) method,	- for oral use;	21.10.20.120		coloration / solution	
	measurements, establishing requirements	- for external use;	21.10.32		coloration	
	for drugs registered in the established	- for intrauterine administration;	21.10.5		(description)	
	order and included in State registers of	- for intracisternal injection;	21.10.51.120		_	
	drug for veterinary use of the Eurasian	- for local application;	21.10.51.121			
	Economic Union member states	- for infusion;	21.10.51.122			
		Suspensions and emulsions:	21.10.51.123			
		- for injection;	21.10.51.124			
		- for oral use;	21.10.51.125			
		- for external use;	21.10.51.126			
		- for intrauterine administration;	21.10.51.129			
		- for intracisternal injection;	21.10.52.110			
		- for local application;	21.10.53			
		Drops:	21.10.53.120			
		- eye;	21.10.54			
		- Ear;	21.10.54.110			
		- nasal;	21.10.54.120			
		- sublingual	21.10.54.130			
		- for local application;	21.10.54.140			
		- for oral use.	21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.158			
			21.20.10.159			
			21.20.10.213			
			21.20.21.130			
			21.20.21.139			
			02.30.40.140			
213	BP, Appendix VIII P (Method 1,	Drugs:	21.20	3004	Authenticity	Compliant/ Non compliant; Pass
	Method 5, Method 7) and other	Solutions;	21.20.1			test/ fail test (if necessary,
	normative documents approved in the	Lyophilizate.				specify conditions)
	established order, specifying the				Quantitative	0,2-2,0 mg/ml
	application of the research (testing)				determination of protein /	
	method, measurements, establishing				quantitative content /	
	requirements for drugs registered in the				-	

1	2	3	4	5	6	7 243 pages
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				mass fraction / mass concentration	
214	STATE PHARMACOPOEIA, XIV edition, GENERAL	Drugs. Medicinal plant raw material and	21.1 21.10	3003 – 3004	Appearance (description)	-
	PHARMACOPOEIA	collections.	21.10.1	from 3305	Color (description)	-
	ARTICLE.1.1.0001.18 and other	Pharmaceutical substances.	21.10.20.120		Odor (description)	-
	normative documents approved in the	Drugs.	21.10.32		Consistency (description)	-
	established order, specifying the	Medicinal plant raw material and	21.10.5			
	application of the research (testing)	collections.	21.10.51.120			
	method, measurements, establishing requirements for drugs registered in the	Pharmaceutical substances.	21.10.51.121 21.10.51.122			
	established order and included in State		21.10.51.122			
	registers of drug for veterinary use of the		21.10.51.124			
	Eurasian Economic Union member		21.10.51.125			
	states		21.10.51.126			
215	STATE PHARMACOPOEIA, XIV	Drugs:	21.10.51.129		Appearance (description)	-
	edition, GENERAL	Tablets, dragee, briquettes, pastilles;	21.10.52.110		Color (description)	-
	PHARMACOPOEIA	Capsules;	21.10.53		Determination of talc,	0-5 %
	ARTICLE.1.4.1.0015.15 and other	Suppositories;	21.10.53.120		aerosil, titanium dioxide	
	normative documents approved in the established order, specifying the	Powders and granules (microgranules, pellets)	21.10.54 21.10.54.110		and other auxiliary	
	application of the research (testing)	(inicrogranules, penets)	21.10.54.110		substances	
	method, measurements, establishing		21.10.54.130			
	requirements for drugs registered in the		21.10.54.140			
	established order and included in State		21.10.54.150			
	registers of drug for veterinary use of the		21.10.54.160			
	Eurasian Economic Union member		21.10.54.170			
	states		21.10.54.180			
216	STATE PHARMACOPOEIA, XIV		21.10.54.190 21.20.1		Appearance (description)	-
	edition, GENERAL PHARMACOPOEIA		21.20.1			
	ARTICLE.1.4.1.0013.15 and other		21.20.10			
	normative documents approved in		21.20.10.159			
	accordance with the established		21.20.10.213			
	procedure, specifying the application of		21.20.21.130			
	the Methodology of research (tests),		21.20.21.139			
	measurements, establishing requirements		02.30.40.140			
	for drugs registered in accordance with		15.12.11			
	the established procedure and included					
	in the State for veterinary use of the					

1	2	3	4	5	6	7 7
	Eurasian Economic Union member					
	states					
217	STATE PHARMACOPOEIA, XIV				Appearance (description)	-
	edition, GENERAL					
	PHARMACOPOEIA					
	ARTICLE.1.4.1.0010.15 and other					
	normative documents approved in the					
	established order, specifying the				Dissolution time	0.5 - 120 min;
	application of the research (testing)					Pass test/ fail test
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states					
218	STATE PHARMACOPOEIA, XIV				Appearance (description)	-
	edition, GENERAL					
	PHARMACOPOEIA					
	ARTICLE.1.4.1.0004.18 and other					
	normative documents approved in the				Size	45 μm – 11,2 mm
	established order, specifying the					7
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the				Disintegration	0.5 - 120 min;
	established order and included in State					Pass test/ fail test (if necessary,
	registers of drug for veterinary use of the					specify conditions)
	Eurasian Economic Union member					
	states					
219	STATE PHARMACOPOEIA, XIV				Appearance (description)	-
	edition, GENERAL					
	PHARMACOPOEIA					
	ARTICLE.1.4.1.0011.18and other					
	normative documents approved in the					
	established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states					
		1			1	

1	2	3	4	5	6	7 7
220	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0027.18and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Drops: - eye; - Ear; - nasal; - for local application; - for oral use;	21.1 21.10 21.10.20.120 21.10.51.120 21.10.54 21.10.54.180 21.20.1 21.20.10 21.20.10.213 21.20.10.158 21.20.10.159 21.20.21.130 21.20.21.130 21.20.21.139 21.20.14	3004	Appearance (description)	-
221	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0031.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Lyophilizate	21.1 21.10 21.20.21.130 21.20.21.139	3004	Appearance (description) Dissolution time	0,5 – 120 min; Pass test/ fail test (if necessary, specify conditions)
222	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0041.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions	21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.14 21.20.14.000	3004	Appearance (description) Foaming capacity	- 10 -700 mm

1	2	3	4	5	6	7 7
223	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0014.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member	Drugs: Suspensions Drops (eye) Solutions	21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.51.129	3004	Appearance (description) Sedimentary stability Resuspension ability Needle penetration/Injectable Solution Stratification	- 0,5 – 120 min; Pass test/ fail test (if necessary, specify conditions) 0,5 – 120 min; Pass test/ fail test (if necessary, specify conditions) from 0,2 seconds to 10 minutes; Pass test/ fail test 0,5 – 120 min;
224	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0017.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Emulsions	21.10.53 21.10.53.120 21.10.54.110 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140		(delamination) Appearance (description) Stratification (delamination)	Pass test/ fail test (if necessary, specify conditions) - 0,5 - 120 min; Pass test/ fail test (if necessary, specify conditions)
225	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0008.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member	Ointments (Gels, creams, liniments, pastes)	21.10.54 21.10.54.180 21.20.1 21.20.10	3004	Appearance (description) Color (description) Odor (description) Consistency (description) Dosage mass uniformity (mass uniformity) Dosage uniformity Uniformity	

1	2	3	4	5	6	7 7
	states					
226	STATE PHARMACOPOEIA, XIV	Drugs.	21.1	3003 –	pH/activity	from 0 to 14
	edition, GENERAL	Medicinal plant raw material and	21.10	3004	(concentration) hydrogen	
	PHARMACOPOEIA	collections.	21.10.1		ions/pH/concentration	
	ARTICLE.1.2.1.0004.15 and other	Pharmaceutical substances.	21.10.20.120		index of hydrogen ions	
	normative documents approved in the		21.10.32			
	established order, specifying the		21.10.5			
	application of the research (testing)		21.10.51.120			
	method, measurements, establishing		21.10.51.121			
	requirements for drugs registered in the		21.10.51.122			
	established order and included in State		21.10.51.123			
	registers of drug for veterinary use of the		21.10.51.124			
	Eurasian Economic Union member		21.10.51.125			
	states		21.10.51.126			
			21.10.51.129			
			21.10.52.110			
			21.10.53			
			21.10.53.120			
			21.10.54			
			21.10.54.110			
			21.10.54.120			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10.158			
			21.20.10.159			
			21.20.10.213			
			21.20.21.130			
			21.20.21.139			
			02.30.40.140			
			20.20.14			
			20.20.14.000			
227	STATE PHARMACOPOEIA, XIV	Drugs.	21.1	3003 -	Residual organic	10 -5000ppm (mg/kg; mcg/g);
	edition, GENERAL	Medicinal plant raw material and	21.10	3004	solvents	0,00001 – 10%
	PHARMACOPOEIA	collections.	21.10.1			

1	2	3	4	5	6	7 7
228	ARTICLE.1.1.0008.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0010.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Pharmaceutical substances.	21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.123 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.54 21.10.54 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10 21.20.10 21.20.10 21.20.10.158 21.20.10.159 21.20.21.130 21.20.21.130 21.20.21.130 21.20.21.130 21.20.21.139 02.30.40.140		Weight loss during drying/Drying method/mass fraction of humidity	0,001 -50%
229	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.2.0005.15 and other normative documents approved in the established order, specifying the	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances. Drugs. Medicinal plant raw material and	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5	3003 – 3004 from4201 from 3808	Foreign matter (related compounds) Authenticity	0,01 - 20% 0,01 - 20% of active substance (if necessary, specify conditions) Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)

method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states Pharmaceutical substances. 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.125 21.10.51.126 21.10.51.126 21.10.51.129 21.10.51.129 21.10.51.120	1	2	3	4	5	6	7
Drugs: Drugs: Solutions: - for injection; Drops (eye) Drops (eye) Drops (eye) Drugs: Drops (eye) Drops (eye) Drops (eye) Drugs: Drops (eye) Drugs: Drops (eye) Drugs: Drops (eye) Drugs: Drugs: Drops (eye) Drugs: Drops (eye) Drugs: Drops (eye) Drugs: Drops (eye) Drugs: Drugs: Drops (eye) Drugs: Drops (eye) Drugs: Drugs: Drops (eye) Drugs:	met requ esta regi Eur	plication of the research (testing) ethod, measurements, establishing quirements for drugs registered in the tablished order and included in State gisters of drug for veterinary use of the urasian Economic Union member	Collections. Pharmaceutical substances. Drugs: Solutions: - for injection; Suspensions and emulsions: - for injection;	21.10.51.120 21.10.51.121 21.10.51.123 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10 21.20.10.158 21.20.10.158 21.20.10.159 21.20.21.139 02.30.40.140 15.12.11 20.20.14	5	Quantitative determination / quantitative content / mass fraction / mass concentration Authenticity Quantitative determination of antimicrobial preservatives / quantitative content /	0,1-10000 mkg/kg (g; 100 g; ml cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle. plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; mcm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle. plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,0001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Passtest/ fail test (if necessary, specify conditions) 0,1-10000 μg/ml (cm3; l; dm3; 100 ml; pipette; syringe; bottle.); 0,00001-20%, 0,00001-20%, 0,00001-20% weight,

1	2	3	4	5	6	7 243 pages
		Drugs: Solutions: - for oral use;			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
		- for injection; Suspensions and emulsions: - for oral use; - for injection; Powders and granules (microgranules, pellets): - for oral use. Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances. Pharmaceutical substances of vegetable products.			Quantitative determination of antioxidants/ quantitative content / mass fraction / mass concentration	0,1-10000 mkg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-10000 mg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-20%, 0,00001-20% weight, 0,00001-20% volume, 1,0 – 200,0 % of declared; not found
		Drugs: Solutions: - for oral use;			Authenticity;	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
		- for injection; Suspensions and emulsions: - for oral use; - for injection; Powders and granules (microgranules, pellets): - for oral use. Pharmaceutical substances			Quantitative determination of organic acids/ quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository);
						0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml;

1		2	4	-	-	243 pages
1	2	3	4	5	6	/
230	STATE PHARMACOPOEIA, XIV	Drugs.			Authenticity	tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found Compliant/ Non compliant; Pass
	edition, GENERAL PHARMACOPOEIA	Medicinal plant raw material and collections.				test/ fail test (if necessary, specify conditions)
	ARTICLE.1.2.1.2.0004.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Pharmaceutical substances.			Quantitative determination / quantitative content / mass fraction / mass concentration	o0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150%, 0,00001-150% volume, 1,0 – 200,0 % of declared;
						not found

1	2	3	4	5	6	7
		Drugs: Solutions: - for injection; Suspensions and emulsions: - for injection; Drops (eye)			Authenticity Quantitative	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 µg/ml (cm3; 1; dm3; 100
		Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120	3003 – 3004	determination of antimicrobial preservatives / quantitative content / mass fraction / mass concentration	ml; pipette; syringe; bottle.); 0,00001-10000 mg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-20%, 0,00001-20% weight, 0,00001-20% volume, 1,0 – 200,0 % of declared; not found
231	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.2.0003.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120		Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
232	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.1.0003.15 and other		21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170		Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions
	normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139		Optical density Quantitative determination / quantitative content / mass fraction / mass concentration	0,0001 – 3,0 E.O.P. 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule;

1	2	3	4	5	6	7 7
			02.30.40.140			pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,0001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
233	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.1.0008.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination / quantitative content / mass fraction / mass concentration	(0,002 – 500) mg/kg, mg/dm ³
234	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0017.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances.	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.125 21.10.51.125 21.10.51.125	3003 – 3004 from 3808	Refractive index / Quantitative determination	1,3 – 1,7; 0,0001 – 500 g/ml; mg/ml; g/l; mg/l; g/cm ³ ; mg/cm ³

	T		 			243 pages
1	2	3	4	5	6	7
			21.10.52.110			
			21.10.53			
			21.10.53.120			
			21.10.54			
			21.10.54.110			
			21.10.54.120			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.10			
			21.20.10.159			
			21.20.10.133			
			21.20.10.213			
			21.20.21.130			
			02.30.40.140			
			20.20.14			
			20.20.14			
225	CTATE DILADMA CODOCIA VIV	Danaga	21.10.5	3004	Mechanical inclusions	Absent/ Present; Pass test/ fail
235	STATE PHARMACOPOEIA, XIV edition, GENERAL	Drugs:		3004	Mechanical inclusions	
	PHARMACOPOEIA	Liquid and solid parenteral dosage	21.10.51.120			test (if necessary, specify
		forms, eye dosage forms	21.10.51.121			conditions)
	ARTICLE.1.4.2.0005.18 and other		21.10.51.122			
	normative documents approved in the		21.10.51.123			
	established order, specifying the		21.10.51.124			
	application of the research (testing)		21.10.51.125			
	method, measurements, establishing		21.10.51.126			
	requirements for drugs registered in the		21.10.51.129			
	established order and included in State		21.10.52.110			
	registers of drug for veterinary use of the		21.10.53			
	Eurasian Economic Union member		21.10.53.120			
	states		21.10.54			
			21.10.54.110			
			21.10.54.120			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			

1	2	3	4	5	6	7 243 pages
236	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.1.0015.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Powders and granules.	21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.21.130 21.20.21.139 21.10.32 21.10.51.120 21.10.51.121 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.125 21.10.51.126 21.10.51.129 21.10.54.110 21.10.54.130 21.10.54.140	3004	Determination of fractional composition / particle size distribution/ particle size	7 45 μm – 11,2mm
237	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0002.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Powders and granules (microgranules, pellets): - for preparation (solution for injection, for oral use, drops); - for oral use; - for external use; - for local application; Lyophilizate; Capsules; Ointments; Tablets and dragee; Pharmaceutical substances	21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.1 21.10.5 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.125 21.10.51.125 21.10.51.129 21.10.51.129 21.10.52.110	3003 - 3004	Moisture content/Water content/water/relative humidity/humidity	0,01 – 100%

1	2	3	4	5	6	243 pages, p
			21.10.53.120			
			21.10.54			
			21.10.54.110			
			21.10.54.120			
			21.10.54.130			
			21.10.54.140			
			21.10.54.150			
			21.10.54.160			
			21.10.54.170			
			21.10.54.180			
			21.10.54.190			
			21.20.1			
			21.20.10			
			21.20.21.130			
			21.20.21.139			
238	STATE PHARMACOPOEIA, XIV	Drugs:	21.1	3004	Density	$700 - 1840 \text{ kg/m}^3$
	edition, GENERAL	Solutions:	21.10			$0.001 - 3.000 \text{ mg/cm}^3$
	PHARMACOPOEIA	- for injection;	21.10.1			$0.0001 - 3.000 \text{ mg/cm}^3$
	ARTICLE.1.2.1.0014.15 and other	- for oral use;	21.10.20.120			
	normative documents approved in the	- for external use;	21.10.32			
	established order, specifying the	- for intrauterine administration;	21.10.5			
	application of the research (testing)	- for intracisternal injection;	21.10.51.120			
	method, measurements, establishing	- for local application;	21.10.51.121			
	requirements for drugs registered in the	- for infusion;	21.10.51.122			
	established order and included in State	Suspensions and emulsions:	21.10.51.123			
	registers of drug for veterinary use of the	- for injection;	21.10.51.124			
	Eurasian Economic Union member	- for oral use;	21.10.51.125			
	states	- for external use;	21.10.51.126			
239	STATE PHARMACOPOEIA, XIV	- for intrauterine administration;	21.10.51.129		Color / colour / colour	-
	edition, GENERAL	- for intracisternal injection;	21.10.52.110		solution / coloration /	
	PHARMACOPOEIA	- for local application;	21.10.53		solution coloration	
	ARTICLE.1.2.1.0006.15 and other	Drops:	21.10.53.120		(description)	
	normative documents approved in the	- eye;	21.10.54			
	established order, specifying the	- Ear;	21.10.54.110			
	application of the research (testing)	- nasal;	21.10.54.120			
	method, measurements, establishing	- sublingual	21.10.54.130			
	requirements for drugs registered in the	- for local application;	21.10.54.140			
	established order and included in State	- for oral use.	21.10.54.150			
	registers of drug for veterinary use of the		21.10.54.160			
	Eurasian Economic Union member		21.10.54.170			
<u> </u>	states		21.10.54.180			
240	STATE PHARMACOPOEIA, XIV		21.10.54.190		Transparency and degree	-

1	2	3	4	5	6	243 pages, p
	edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0007.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140		of turbidity of liquids (description)	
241	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0002.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Nominal Volume / recoverable volume / filling volume / filling volume of bottle / drug volume in bottle	0,1 – 1000 ml (cm3; l; dm3); 80 -150 % of nominal; Pass test/ fail test
242	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0003.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
243	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0007.15 and other	Drugs: Drops: - eye; - Ear;			Weight (volume) of package contents	0,1 - 25000 ml (cm3; l; dm3); 0,1 - 10 kg; 0,001 - 500 g;

1	2	3	4	5	6	7 7
	normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	- nasal; - sublingual; - for local application; - for oral use; Suspensions and emulsions; Aerosols, sprays, foams; Tinctures and Extracts: - for oral use; - for external use; - for local application; Ointments; Syrups; Balsams				1,0 – 5000 mg; Pass test/ fail test
244	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0002.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Aerosols, sprays, foams			Weight (volume) of package contents; output of package contents (for aerosols) Dosage mass uniformity (mass uniformity) Dosage uniformity Stratification (delamination)	0,1 – 25000 ml (cm3; l; dm3); 0,1 – 10 kg; 0,001 – 500 g; 1,0 – 5000 mg; Pass test/ fail test 0,01 – 50 % of average mass; Pass test/ fail test Pass test/ fail test (if necessary, specify conditions) 0,5 – 120 minutes; Pass test/ fail test (if necessary, specify conditions)
245	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0006.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State	Drugs: Drops: - eye; - Ear; - nasal; - sublingual; - for local application; - for oral use; Aerosols and sprays; Cornea (eye)			Package hermeticity (for aerosols) Number of doses per package Dosage mass uniformity (mass uniformity)	Pass test/ fail test; Hermetic / non-hermetic 0-1000; Pass test/ fail test 0,01 - 50 % of average mass; Pass test/ fail test

1	2	3	4	5	6	243 pages, j
	registers of drug for veterinary use of the Eurasian Economic Union member states					
246	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0015.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions; Suspensions and emulsions; Drops (eye). Pharmaceutical substances			Viscosity	0,0001–100000 mm2/c; Ps; cPs; PAHs; MPAHs; m2/c; St; cSt;
247	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0011.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Pharmaceutical substances	21.1 21.10	3003	Melting Temperature	25 – 400 °C
248	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0005.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Pharmaceutical substances. Drugs: Extracts; Powder	21.1 21.10 21.20.1 21.20.10	3003 - 3004	Solubility	Pass test/ fail test (if necessary, specify conditions)

1	2	3	4	5	6	7 7	
249	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.2.2.0013.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Pharmaceutical substances Medicinal plant raw material and collections.	21.1 21.10 21.20.1 21.20.10 02.30.40.140	3003 - 3004		Total ash	0,001 – 10,000%
250	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.2.2.0014.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Sulfated ash	0,001 – 10,000%	
251	STATE PHARMACOPOEIA, XIV edition, GENERAL	Pharmaceutical substances.	21.1 21.10	3003	Solubility	Pass test/ fail test (if necessary, specify conditions)	
	PHARMACOPOEIA ARTICLE.1.1.0006.15 and other				Total ash	0,001 – 10,000%	
	normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Sulfated ash	0,001 – 10,000%	

1	2	3	4	5	6	7 243 pages, p																								
252	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0008.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tablets and dragee, briquettes, pastels; Capsules; Suppositories; Powders and granules (microgranules, pellets); Drops (eye); Ointments; Cornea (eye); Aerosols and sprays; System: - Intravaginal administration	21.10.5 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120	21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	20 21 22 23 24 25 26 29 10	Dosage uniformity	Pass test/ fail test (if necessary, specify conditions)									
253	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0003.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Drops (eye) Drugs: Cornea (eye)	21.10.54 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1 21.20.10 21.20.10		Dosage uniformity Sedimentary stability Size	Pass test/ fail test (if necessary, specify conditions) 0,5 - 120 minutes; Pass test/ fail test (if necessary, specify conditions) 10-20000 µm 0,01-2000 mm 0,001-200 cm																								
254	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0009.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tablets, dragee, briquettes, pastilles; Capsules; Suppositories; Powders and granules (microgranules, pellets); System: - Intravaginal administration	21.20.10.159 02.30.40.140		Average mass and mass uniformity	0,1 – 10 kg; 0,001 – 500 g; 1,0 – 5000 mg; 0,01 – 50 % of average mass; Pass test/ fail test																								
255	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA	Drugs: Tablets, dragee, briquettes, pastilles; Capsules;			Disintegration	0,5 -120 minutes; Pass test/ fail test (if necessary, specify conditions)																								

1	2	3	4	5	6	7 7
	ARTICLE.1.4.2.0013.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Powders and granules.				
256	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0012.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tablets; Suppositories.				
257	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0014.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tablets, dragee, briquettes, pastilles; Capsules; Suppositories; Powders and granules.			Dissolution	0,1 – 120% of declared; Pass test/fail test
258	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0005.18 and other normative documents approved in the established order, specifying the	Drugs: Capsules	21.10.51.120 21.10.54 21.10.54.180 21.20.1 21.20.10	3004	Size Capsules	000-5 0,01-160 mm 0,001-16 cm

1	2	3	4	5	6	7 7
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
259	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0007.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; Suspensions and emulsions: - for injection; Drops (eye)	21.10.51.120 21.10.54 21.10.54.180 21.20.1 21.20.10	3004	Authenticity; Quantitative determination of antimicrobial preservatives/ quantitative content / mass fraction / mass concentration	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-10000 mg/ml (cm3; l; dm3; 100 ml; pipette; syringe; fl.); 0,00001-20%, 0,00001-20% weight, 0,00001-20% volume, 1,0 - 200,0 % of declared; not found
260	state Pharmacopoeia, XIV edition, General Pharmacopoeia ARTICLE.1.4.1.0019.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; and other normative documents approved in the established order that specify the application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and included in the State registries of feeds and feed additives of the	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products. Feed additives	21.1 21.10 21.10.53 21.10.53.120 21.20.1 21.20.10 02.30.40.140 10.91.10.230	3003 – 3004; 2309	Mass fraction of dry residue / dry residue / Content of pharmacologically active substances or biological activity	0,001 – 10% 10 ⁻⁸ – 10,0 %; 10 ⁻⁸ – 0,1 %/tab(bottle); 10 ⁻⁸ – 10,0 mg/g (mg; cm3; ml; dm3; l); 10 ⁻³ – 100 mcg/g (mg; cm3; ml; dm3; l)

						243 pages,
1	2	3	4	5	6	7
	Eurasian Economic Union member	!				
	states					
	States					
261	STATE PHARMACOPOEIA, XIV					
201	edition, GENERAL					
	PHARMACOPOEIA					
	ARTICLE.1.4.1.0020.15 and other					
	normative documents approved in the					
	established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;					
	and other normative documents					
	approved in the established order that					
	specify the application of the research					
	(testing) method, measurements that					
	establish requirements for feed and feed					
	additives registered in the established					
	order and included in the State registries					
	of feeds and feed additives of the					
	Eurasian Economic Union member					
	states					
262	STATE PHARMACOPOEIA, XIV					
	edition, GENERAL					
	PHARMACOPOEIA					
	ARTICLE.1.5.1.0001.15 and other					
	normative documents approved in the					
	established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					

1	2	3	4	5	6	7 7
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; and other normative documents approved in the established order that specify the application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and included in the State registries of feeds and feed additives of the Eurasian Economic Union member states					
263	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.5.3.0007.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tinctures and Extracts; Syrups; Balsams. Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products.			Weight loss during drying /humidity determination/ dry residue	0,001 – 50%
264	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.5.3.0004.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Impurity content	0,1-20%; Pass test/ fail test; Presence/absence (if necessary, specify conditions)

1	2	3	4	5	6	7
265	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0017.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for oral use; - for injection; Suspensions and emulsions: - for oral use; - for injection; Powders and granules (microgranules, pellets): - for oral use. Pharmaceutical substances	21.1 21.10 21.10.51.120 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.20.1 21.20.10 21.20.21.130 21.20.21.139 02.30.40.140	3003-3004	Authenticity; Quantitative determination / quantitative content / mass fraction / mass concentration	Compliant/ Non compliant; Pass test/fail test (if necessary, specify conditions); 0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-1000 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,1 - 1000000000 IU/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 1000000000 Unit/kg (g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 1000000000 Unit/kg (g; 100g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 - 200,0 % of declared; not found
266	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.4.0012.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination / quantitative content / mass fraction / mass concentration	0,001-10000 mkg/kg (mg; g; 10 mg; 100 g; ml; cm3; ml; 10 ml; l; dm3; 100 ml; tablet.; capsule.; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,000001-10000 mg/kg (g; 10 g; 100 g; ml; cm3; l; dm3; 10 ml; 100 ml; tablet.; capsule.; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 10 g; 100g; ml; 10 ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle, plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 10 g; 100g; ml; 10 ml; cm3; l; dm3; 100ml; tablet; capsule; pipette;

						243 pages,
1	2	3	4	5	6	7
						syringe; bottle.; plate, suppository) 0,00001-150%
267	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.5.3.0009.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tinctures and Extracts; Medicinal plant raw material and collections. Pharmaceutical substances of vegetable products.			Quantitative determination arsenic/ quantitative content / mass fraction / mass concentration Quantitative determination / quantitative content / mass fraction / mass concentration of heavy metals (cadmium, lead, arsenic, Mercury)	(0,010 – 500) mg/kg, mg/dm ³ (0,002 – 500) mg/kg, mg/dm ³
268	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.2.2.0005.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination Mercury/ quantitative content / mass fraction / mass concentration	(0,010-20) mg/kg, mg/dm ³

1	2	3	4	5	6	7 7
269	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.5.2.0001.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for oral use; - for injection; Suspensions and emulsions: - for oral use; - for injection; Powders and granules (microgranules, pellets): - for oral use. Tinctures and Extracts. Medicinal plant raw material and collections. Pharmaceutical substances			Quantitative determination of aromatic compounds/ quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-1000 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,1 – 1000000000 IU/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
270	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0016.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for local application; Tinctures and Extracts: - for oral use; - for external use; - for local application; Syrups	21.10.51.120 21.10.52.110 21.10.54 21.10.54.180 21.20.1 21.20.10 02.30.40.140	3004	Quantitative determination of ethyl alcohol / quantitative content / mass fraction / mass concentration	0,01-96 % (mass, volume); g/l (dm3, cm3, ml); mg/ml (cm3)

1	2	3	4	5	6	7 243 pages, p
271	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0004.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions; Ointments; Drops (eye); Extracts	21.10.51.120 21.10.52.110 21.10.54 21.10.54.180 21.20.1 21.20.10 02.30.40.140	3004	Acid value	0,01 -10 mg KOH/ml (cm3;g)
272	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0007.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Peroxide value	0,01-50 mmol O2/kg
273	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.19.0002.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for local application; - for infusion; Drops: - eye; - Ear; - nasal; - for local application;	21.1 21.10 21.10.20.120 21.10.51.120 21.10.54 21.10.54.180 21.20.1 21.20.10 21.20.10.213 21.20.10.158 21.20.10.159 21.20.21.130 21.20.21.130	3004 from 3808	Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,000001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet.; capsule; pipette; syringe; bottle.; plate; suppository; stick; package, bag); 0,00001-10,0 g/kg (g; 100g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag) 0,01 -10 mg KOH/g (cm3;g); 0,0001-150%,

1	2	3	4	5	6	7
274	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0013.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member	- for oral use; Powders and granules (microgranules, pellets); Tablets; Ointments; Aerosols and sprays; Tinctures and Extracts; Syrups; Balsams; Drug checker; Cord. Pharmaceutical substances.	21.20.14 21.20.14.000 02.30.40.140			0,0001-150% weight, 0,0001-150% volume, not found
275	states STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0015.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
276	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.2.0001.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states					
277	STATE PHARMACOPOEIA, XIV edition, GENERAL	Drugs: Solutions:	21.10.52.110 21.20.1	3004	Quantitative determination /	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; package; bag);

1	2	3	4	5	6	7 243 pages,
	PHARMACOPOEIA ARTICLE.1.7.2.0033.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	- for injection; Suspensions and emulsions: - for injection; Powders and granules (microgranules, pellets): - for preparation (solution for injection, drops); Lyophilizate	21.20.10 21.20.10.213		quantitative content / mass fraction / mass concentration	0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml); 0,1 – 100000000 Unit/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml); not found
278	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0021.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Powders and granules: - for oral use; Tablets; Capsules; Pastes: - for oral use; Gels: - for oral use; Colloidal Solutions	21.20.1 21.20.10 02.30.40.140	3004	Adsorption activity	1-1000 mg/g (mg/Tab.; mg/caps.; mcg/ml; μmol/g; cm3/g)
279	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.4.0002.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for oral use; - for external use; - for intrauterine administration; - for local application; Suspensions and emulsions: - for oral use; - for external use; - for intrauterine administration; - for local application; Drops: - Ear; - nasal; - for local application; - for oral use; Ointments (Gels, creams, liniments, pastes);	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53 21.10.53.120 21.10.54	3003 – 3004 from 5102 from 3305	Microbiological purity: Total number of aerobic bacteria; Total number of fungi; Enterobacteria, etc gram-negative bacteria; E. coli; Pseudomonas aeruginosa; Staphylococcus aureus; Salmonella	Pass test/fail test (if necessary, specify conditions)

1	2	3	4	5	6	243 pages, 9
		- for external use;	21.10.54.110			
		- for local application;	21.10.54.120			
		Powders and granules	21.10.54.130			
		(microgranules, pellets):	21.10.54.140			
		- for preparation (solution for oral	21.10.54.150			
		use, drops);	21.10.54.160			
		- for oral use;	21.10.54.170			
		- for external use;	21.10.54.180			
		- for local application;	21.10.54.190			
		Aerosols and sprays;	21.20.1			
		Capsules;	21.20.10			
		Suppositories (sticks);	21.20.10.158			
		Tablets, dragee, briquettes, pastilles;	21.20.10.159			
		Tinctures and Extracts:	21.20.10.213			
		- for oral use;	21.20.21.130			
		- for external use;	21.20.21.139			
		- for local application;	02.30.40.140			
		Syrups;	01.49.28.000			
		Balsams;	20.4			
		System:				
		- Intravaginal administration.				
		Pharmaceutical substances.				
		Washing zoohygienic liquid				
		products for unproductive animals:				
		- shampoos;				
		- soft soap;				
		- foam;				
		- mousse;				
		- gel;				
		- lotion;				
		- tonic.				
280	STATE PHARMACOPOEIA, XIV	Drugs:			Sterility	Sterile/non-sterile; pass test/fail
	edition, GENERAL	Solutions:				test
	PHARMACOPOEIA	- for injection;				
	ARTICLE.1.2.4.0003.15 and other	- for external use (when applied to				
	normative documents approved in the	wounds);				
	established order, specifying the	- for intracisternal injection;				
	application of the research (testing)	- for infusion;				
	method, measurements, establishing	Suspensions and emulsions:				
	requirements for drugs registered in the	- for injection;				
	established order and included in State	- for intracisternal injection;				

1	2	3	4	5	6	7 7
281	registers of drug for veterinary use of the Eurasian Economic Union member states STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.4.0006.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drops (eye); Ointments (Gels, creams, liniments, pastes); - for external use; - for intracisternal injection; - eye; Powders and granules (microgranules, pellets): - for preparation solution for injection; - for external use (when applied to wounds). Conditions for semen dilution by farm animal manufacturers Drugs: Solutions: - for injection; - for infusion; Suspensions and emulsions: - for injection; Powders and granules: - for preparation solution for injection	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.52.110 21.10.53	3003 - 3004	Bacterial endotoxins	Pass test/fail test; Compliant/ Non compliant; 0,001-1000000000 Endotoxin units mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 EU/ ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit)
282	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.4.0010.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the	Drugs: Solutions: - for injection; - for oral use; - for external use; - for intrauterine administration; - for intracisternal injection; - for local application; Suspensions and emulsions: - for injection; - for oral use;	21.10.53.120 21.10.54 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190 21.20.1		Determination of antimicrobial activity of antibiotics Agar Diffusion Method/ Quantitative determination /quantitative content / mass fraction / mass concentration	0,001-1000000 mkg/kg (mg; g; 10 mg; 100g; ml; cm3; 10 ml; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle; plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (mg; g; 10 g; 100 g; ml; cm3; l; dm3; 10 ml; 100 ml; tabl; capsule; pipette; syringe; bottle; plate; suppository; stick; packing; bag); 0,00001-1000,0 g/kg (g; 10 g;

1	2	3	4	5	6	243 pages, j 7
	Eurasian Economic Union member states	- for external use; - for intrauterine administration; - for intracisternal injection; - for local application; Drops; Ointments; Powders and granules (microgranules, pellets): - for preparation (solution for injection, for oral use, drops); - for oral use; - for external use; - for local application; Lyophilizate; Tablets; Pharmaceutical substances.	21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140			100 g; ml; 10 ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle; plate; suppository; stick; package, bag) 0,1 – 1000000000 IU/kg (mg; g; 10 g; 100g; ml; 10 ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle, plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 10 g; 100g; ml; 10 ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository) 0,00001-150%; 1,0 – 200,0 % of declared; not found
283	state pharmacopoeia, XIV edition, General Pharmacopoeia ARTICLE.1.2.2.2.0009.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions	21.1 21.10 21.10.1	3003 - 3004	Mass fraction of chlorides	0,1-10000 mkg/kg (g; 100g; ml; cm3; l; dm3; 100 ml; bottle; package; bag); 0,000001-10000 mg/kg (g; 100g; ml; cm3; l; dm3; 100 ml; bottle; package; bag); 0,00001-10,0 g/kg (g; 100g; ml; cm3; l; dm3; 100 ml; bottle; package; bag); 0,0001-150%, 0,0001-150% weight, 0,0001-150% volume, not found
284	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.1.0001.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the	Drugs: Solutions; Drops; Aerosols and sprays; Suspensions and emulsions; Powders and granules; Suppositories (sticks) Tablets, dragee, briquettes, pastilles; Capsules; Ointments; Lyophilizate;	21.1 21.10 21.10.1 21.10.20.120 21.10.32 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.123	3003-3004	Authenticity; Quantitative determination / quantitative content / mass fraction / mass concentration	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml;

1	2	3	4	5	6	7 7
	Eurasian Economic Union member states	Pharmaceutical substances. Medicinal plant raw material and collections.	21.10.51.125 21.10.51.126 21.10.51.129			cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick;
285	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.1.0002.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.10.52.110 21.10.53 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.180			package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume,
286	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.1.1.0009.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states		21.10.54.190 21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140			1,0 – 200,0 % of declared; not found
287	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0012.15 (Method 1, Method 5, Method 7) and other	Drugs: Solutions; Lyophilizate.	21.20 21.20.1	3004	Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member				Quantitative determination of protein / quantitative content / mass fraction / mass concentration	0,2 – 2,0 mg/ml

	T					243 pages, ₁
1	2	3	4	5	6	7
	states					
288	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.4.1.0021.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tinctures; Extracts; Solutions	21.10.53 21.10.53.120 21.20.1 21.20.10 02.30.40.140	3003-3004	Mass fraction of dry residue / dry residue	0,001 – 10%
289	STATE PHARMACOPOEIA, XIV edition, GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0022.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions	21.20 21.20.1	3004	Determination of ammonia nitrogen	0,01 – 200 mg/ml (g)
290	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.1.0004.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State	Drugs: Powders and granules Suspensions	21.10.53 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.160	3004	Appearance (description) Authenticity pH Water Quantitative determination	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) from 0 to 14 0,01 – 100% 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet;

1	2	3	4	5	6	7
	registers of drug for veterinary use of the Eurasian Economic Union member states		21.10.54.170 21.10.54.180 21.10.54.190 21.20.10 21.20.21.130 21.20.21.139			capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
291	state pharmacopoeia, xiv edition, pharmacopoeia article 3.1.0005.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Tablets			Appearance (description) Authenticity Dissolution Water Dosage uniformity Quantitative determination	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1 – 120% of declared; Pass test/ fail test 0,01 – 100% Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g;

1		2	A	F	6	243 pages, 7
1	2	3	4	5	6	,
						ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150%, weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
292	STATE DHADMACODOEIA VIV	Deuge			Annagrance (description)	not found
292	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA	Drugs: Powder for preparation solution for			Appearance (description) Authenticity	Compliant/ Non compliant; Pass
	ARTICLE3.1.0006.18 and other normative documents approved in the	Powder for preparation solution for injection			Authenticity	test/ fail test (if necessary, specify conditions)
	established order, specifying the application of the research (testing)				Transparency solution (description)	-
	method, measurements, establishing requirements for drugs registered in the				Solution colour (description)	-
	established order and included in State				pH	from 0 to 14
	registers of drug for veterinary use of the Eurasian Economic Union member				Dissolution time	0,5 – 120 min; Pass test/ fail test
	states				Water	0,01 – 100%
					Quantitative	0,1-10000 mkg/kg (g; 100 g; ml;
					determination	cm3; 1; dm3; 100 ml; tablet;
						capsule; pipette; syringe; fl.
						plate; suppository; stick;
						package, bag); 0,00001-10000 mg/kg (g; 100 g;

1	2	3	4	5	6	7 243 pages,
						ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
293	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.1.0007.18 and other normative documents approved in the	Drugs: Tablets			Appearance (description) Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the application of the research (testing)				Dissolution	0,1 – 120% of declared; Pass test/fail test
	method, measurements, establishing				Water	0,01 – 100%
	requirements for drugs registered in the established order and included in State				Dosage uniformity	Pass test/ fail test (if necessary, specify conditions)
	registers of drugs for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick;

_							243 pages,
	1	2	3	4	5	6	7
							package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,0001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
	294	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.1.0009.18 and other normative documents approved in the established order, specifying the application of the research (testing)	, PHARMACOPOEIA CLE3.1.0009.18 and other ive documents approved in the shed order, specifying the		Appearance (description) Authenticity Dissolution	- Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1 – 120% of declared; Pass test/ fail test	
		method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Uniformity dosage Quantitative determination	50 -150 % of declared/ of average content; Pass test/ fail test 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet;

1	2	3	4	5	6	7 7
						capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
295	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.1.0021.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member	Drugs: Drops eye			Appearance (description) Authenticity pH Density Quantitative	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) from 0 to 14 700 – 1840 kg/m³ 0,001 – 3,000 mg/cm³ 0,0001 – 3,000 mg/cm³ 0,1-10000 mkg/kg (g; 100 g; ml;
	states				determination	cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag);

1	2	3	4	5	6	243 pages, 7
						0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%; 0,00001-150% weight; 0,00001-150% volume; 1,0 – 200,0 % of declared; not found
296	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-
	edition, PHARMACOPOEIA ARTICLE 3.1.0022.18 and other normative documents approved in the established	Tablets			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	order, specifying the application of the research (testing) method,				Dissolution	0,1 – 120% of declared; Pass test/ fail test
	measurements, establishing requirements for drugs registered in the established order and included in State registers of				Dosage uniformity	Pass test/ fail test (if necessary, specify conditions)
	drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; 1; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository);

1	2	3	4	5	6	7 7
						0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
297	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-
	edition, PHARMACOPOEIA ARTICLE3.1.0023.18 and other normative documents approved in the	Drops eye			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the				pН	from 0 to 14
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight,

						243 pages,
1	2	3	4	5	6	7
						1,0 – 200,0 % of declared; not found
					Transparency (description)	-
					Colour (description)	-
298	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-
	edition, PHARMACOPOEIA ARTICLE3.1.0024.18 and other normative documents approved in the	Solutions for injection			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the application of the research (testing)				Transparency (description)	-
	method, measurements, establishing				Colour (description)	-
	requirements for drugs registered in the				pH	from 0 to 14
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacterial endotoxins Ouantitative	Pass test/fail test; Compliant/ Non compliant; 0,001-1000000000 Endotoxin units mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-10000000000 EU/ ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-10000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit) 0,1-10000 mkg/kg (g; 100 g; ml;
					determination	cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; 1; dm3; 100ml; tablet;

1	2	3	4	5	6	243 pages, 7
						capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
299	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.1.0039.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member	Drugs: Solutions for injection			Appearance (description) Authenticity Transparency (description) Colour (description) pH Bacterial endotoxins	- Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) from 0 to 14 Pass test/fail test; Compliant/ Non compliant; 0,001-1000000000 Endotoxin units
	states				mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 EU/ ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit)	

1	2	3	4	5	6	7 245 pages,
					Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; table; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
300	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.1.0049.18 and other normative documents approved in the established order, specifying the	Drugs: Tablets			Appearance (description) Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the				Dissolution Dosage uniformity	0,1 – 120 % of declared; Pass test/ fail test Pass test/ fail test (if necessary, specify conditions)
	Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet;

1	2	3	4	5	6	243 pages, j 7
						capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
301	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.1.0067.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions for injection			Appearance (description) Authenticity Transparency (description) Colour (description) pH Bacterial endotoxins	- Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions from 0 to 14 Pass test/fail test; Compliant/ Non compliant; 0,001-1000000000 Endotoxin units mkg; mg; 10 mg; g; 10 g; 100 g; kg;

1	2	3	4	5	6	7
						powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 EU/ ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-10000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit)
					Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag);
						0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml;
						cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g;
						100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe;
						bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 - 200,0 % of
						declared; not found
302	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-

1	2	3	4	5	6	7
	edition, PHARMACOPOEIA ARTICLE3.1.0069.18 and other normative documents approved in the	Tablets			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the application of the research (testing)				Dissolution	0,1 – 120% of declared; Pass test/ fail test
	method, measurements, establishing requirements for drugs registered in the				Dosage uniformity	Pass test/ fail test (if necessary, specify conditions)
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; table; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of
						declared;
303	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-
	edition, PHARMACOPOEIA ARTICLE3.1.0082.18 and other normative documents approved in the	Drops: - eye; - Ear			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the				Transparency solution	-

1	2	3	4	5	6	7 7
	application of the research (testing)				(description)	
	method, measurements, establishing				pН	from 0 to 14
	requirements for drugs registered in the				Quantitative	0,1-10000 mkg/kg (g; 100 g; ml;
	established order and included in State				determination	cm3; l; dm3; 100 ml; tablet;
	registers of drug for veterinary use of the					capsule; pipette; syringe; fl.
	Eurasian Economic Union member					plate; suppository; stick;
	states					package, bag);
						0,00001-10000 mg/kg (g; 100 g;
						ml; cm3; l; dm3; 100 ml; tablet;
						capsule; pipette; syringe; bottle.;
						plate; suppository; stick;
						package; bag);
						0,00001-1000 g/kg (g; 100 g; ml;
						cm3; 1; dm3; 100ml; tablet;
						capsule; pipette; syringe; bottle.;
						plate; suppository; stick;
						package; bag); 0,1 – 100000000 IU/kg (mg; g;
						100g; ml; cm3; l; dm3; 100 ml;
						tablet; capsule; pipette; syringe;
						bottle.; plate, suppository);
						0,1 – 100000000 Unit/kg (mg; g;
						100 g; ml; cm3; l; dm3; 100 ml;
						tab.; caps.; pipette; syringe;
						bottle, plate, suppository);
						0,00001-150%,
						0,00001-150% weight,
						0,00001-150% volume,
						1,0 – 200,0 % of
						declared;
						not found
304	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-
	edition, PHARMACOPOEIA	Solutions			Authenticity	Compliant/ Non compliant; Pass
	ARTICLE3.1.0084.18 and other	- for infusion				test/ fail test (if necessary,
	normative documents approved in the					specify conditions)
	established order, specifying the				Transparency	-
	application of the research (testing)				(description)	
	method, measurements, establishing requirements for drugs registered in the				Colour (description)	
	established order and included in State				pH	from 0 to 14
	registers of drug for veterinary use of the				Bacterial endotoxins	Pass test/fail test; Compliant/ Non compliant;
	registers of drug for veterinary use of the					0,001-1000000000 Endotoxin units
	l	l				5,551 100000000 Endotoxin units

1	2	3	4	5	6	7 7
	Eurasian Economic Union member states				Quantitative determination	mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 EU/ ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-10000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 1000000000 IU/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150%, volume, 1,0 - 200,0 % of declared; not found

1	2	3	4	5	6	7
305	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.1.0085.18 and other normative documents approved in the	Drugs: Tablets			Appearance (description) Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary,
	established order, specifying the application of the research (testing)				Dissolution	specify conditions) 0,1 – 120% of declared; Pass test/ fail test
	method, measurements, establishing requirements for drugs registered in the				Water	0,01 – 100%
	established order and included in State				Uniformity	Uniform/non uniform; Pass test/ fail test
	registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; table; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
306	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-

1	2	3	4	5	6	7 7
	edition, PHARMACOPOEIA ARTICLE3.1.0101.18 and other normative documents approved in the	Tablets			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the application of the research (testing)				Dissolution	0,1 – 120% of declared; Pass test/ fail test
	method, measurements, establishing requirements for drugs registered in the				Dosage uniformity	Pass test/ fail test (if necessary, specify conditions)
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 100000000 IU/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 - 200,0 % of
307	STATE PHARMACOPOEIA, XIV	Druge			Appearance (description)	declared; not found
307	edition, PHARMACOPOEIA, AIV edition, PHARMACOPOEIA ARTICLE3.2.0002.18 and other normative documents approved in the established order, specifying the	Drugs: Solutions: - for local application; - for external use			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the				Acidity	Pass test/ fail test

1	2	3	4	5	6	7 7
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 - 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; table; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 - 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; table; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 - 200,0 % of declared; not found
308	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA	Drugs: Solutions:			Appearance (description)	-
	ARTICLE3.2.0003.18 and other normative documents approved in the established order, specifying the	for local application;for external use			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	application of the research (testing) method, measurements, establishing				Alcohol content	0,01-96 % (mass, volume); g/l (dm3, cm3, ml); mg/ml (cm3)
	requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g;

1	2	3	4	5	6	7 243 pages,
		3	7		J	ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150%, weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
309	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-
	edition, PHARMACOPOEIA ARTICLE3.2.0004.18 and other normative documents approved in the established order, specifying the	Solutions: - for external use			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the				Alcohol content	0,01-96 % (mass, volume); g/l (dm3, cm3, ml) ; mg/ml (cm3)
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml;

1	2	3	4	5	6	7 245 pages,
						cm3; 1; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; 1; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
310	O STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.2.0005.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing		Appearance (description) Authenticity Dissolution	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) 0,1 – 120% of declared; Pass test/ fail test		
	requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Osage uniformity Quantitative determination	Pass test/ fail test (if necessary, specify conditions) 0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag);

1	2	3	4	5	6	7 243 pages,					
						0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found					
311	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-					
	edition, PHARMACOPOEIA ARTICLE3.2.0006.18 and other normative documents approved in the Solutions: - for injection; Concentrate for preparation solution	Solutions: - for injection; Concentrate for preparation solution	Solutions: - for injection; Concentrate for preparation solution	Solutions: - for injection; Concentrate for preparation solution	Solutions: - for injection; Concentrate for preparation solution	Solutions: - for injection; Concentrate for preparation solution	Solutions: - for injection; ed in the Concentrate for preparation solution	on		Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the application of the research (testing)	for injection			Transparency (description)	-					
	method, measurements, establishing				Colour (description)	-					
	requirements for drugs registered in the established order and included in State				pН	from 0 to 14					
	registers of drug for veterinary use of the Eurasian Economic Union member states			Bacterial endotoxins	Pass test/fail test; Compliant/ Non compliant; 0,001-1000000000 Endotoxin units mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 EU/ ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit)						
					Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet;					
					uciëi iiiiiatioii	capsule; pipette; syringe; fl.					

1	2	3	4	5	6	7 7
						plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
312	STATE PHARMACOPOEIA, XIV edition, PHARMACOPOEIA ARTICLE3.2.0008.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs: Solutions: - for injection			Appearance (description) Authenticity Transparency (description) Colour (description) pH Manganese Bacterial endotoxins	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions) - from 0 to 14 Pass test/ fail test (if necessary, specify conditions) Pass test/fail test; Compliant/ Non compliant; 0,001-1000000000 Endotoxin units mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe;

amp.bottle, package, bag, Unit); 0,001-1000000000 EU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; kim3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit) Quantitative	1	2	3	4	5	6	243 pages,
determination cm3; l; dm3; 100 ml; tablet; capsule, pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository; 0,1 – 100000000 IU/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 IU/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; table; caps.; pipette; syringe; bottle, plate, suppository); 0,0001-150% weight, 0,00001-150% wolume, 1,0 – 200,0 % of declared; not found		_					0,001-1000000000 EU/ ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-10000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle,
							cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; 1; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; 1; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; 1; dm3; 100 ml; table; caps.; pipette; syringe; bottle, plate, suppository); 0,0001-150%, 0,00001-150%, 0,00001-150% volume, 1,0 – 200,0 % of declared;
	313	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	

1	2	3	4	5	6	7 7
	edition, PHARMACOPOEIA ARTICLE3.2.0012.18 and other normative documents approved in the	Solutions: - for injection - for infusion;			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	established order, specifying the application of the research (testing)	Solution for preparation dosage form for injection			Transparency (description)	-
	method, measurements, establishing	form for injection			Colour (description)	-
	requirements for drugs registered in the established order and included in State				pH Rectarial and stavins	from 0 to 14 Pass test/fail test; Compliant/ Non
	registers of drug for veterinary use of the Eurasian Economic Union member states				Bacterial endotoxins	compliant; 0,001-1000000000 Endotoxin units mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 EU/ ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit); 0,001-1000000000 IU/ml mkg; mg; 10 mg; g; 10 g; 100 g; kg; ml; cm3; 10 ml; l; dm3; 100 ml; powder; lyophilizate; syringe; amp.bottle, package, bag, Unit)
					Quantitative determination	0,1-10000 mkg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 100000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository);

			1	1	Ţ	243 pages,
1	2	3	4	5	6	7
						0,1 – 100000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume, 1,0 – 200,0 % of declared; not found
314	STATE PHARMACOPOEIA, XIV	Drugs:			Appearance (description)	-
	edition, PHARMACOPOEIA ARTICLE3.2.0015.18 and other normative documents approved in the established order, specifying the	Ointments			Authenticity	Compliant/ Non compliant; Pass test/ fail test (if necessary, specify conditions)
	application of the research (testing)				pН	from 0 to 14
	method, measurements, establishing				Quantitative	0,1-10000 mkg/kg (g; 100 g; ml;
	requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				determination	cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; fl. plate; suppository; stick; package, bag); 0,00001-10000 mg/kg (g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,00001-1000 g/kg (g; 100 g; ml; cm3; l; dm3; 100ml; tablet; capsule; pipette; syringe; bottle.; plate; suppository; stick; package; bag); 0,1 – 1000000000 IU/kg (mg; g; 100g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tablet; capsule; pipette; syringe; bottle.; plate, suppository); 0,1 – 1000000000 Unit/kg (mg; g; 100 g; ml; cm3; l; dm3; 100 ml; tab.; caps.; pipette; syringe; bottle, plate, suppository); 0,00001-150%, 0,00001-150% weight, 0,00001-150% volume,

1	2	3	4	5	6	7 243 pages,
						1,0 – 200,0 % of
						declared;
21.7	GT - TE DY - DY - GODOFY - YY	-	21 10 20 120	2004		not found
315	STATE PHARMACOPOEIA XI,	Drugs:	21.10.20.120	3004	Size	10-20000 μm
	STATE PHARMACOPOEIA XII,	Polymer tape (collar);	21.10.32	from 4201		0,01-2000 mm
	STATE PHARMACOPOEIA XIII,	Ear Tags;	21.10.5			0,001-200 cm
	STATE PHARMACOPOEIA XIV,	Plates	21.10.51.120		T	1.0. 120
	STATE PHARMACOPOEIA OF THE	Drugs:	21.10.51.121		Foam stability (foam	1,0 – 120 min;
	REPUBLIC OF BELARUS, USP, EP,	Suppositories	21.10.51.122		resistance)	Pass test/ fail test
	BP and other normative documents		21.10.51.123		Foam formation	50 – 7000 mm;
	approved in the established order,		21.10.51.124		(foam volume)	5 – 700 cm;
	specifying the application of the research		21.10.51.125			10 – 700 ml
	(testing) method, measurements,		21.10.51.126		Foaming time	1 – 30 min;
	establishing requirements for drugs		21.10.51.129 21.10.52.110			Pass test/ fail test (if necessary,
	registered in the established order and included in State registers of drug for		21.10.52.110			specify conditions)
	veterinary use of the Eurasian Economic	Drugs:	21.10.53		Stability of water	Pass test/ fail test (if necessary,
	Union member states	Emulsions;	21.10.53.120		Emulsions	specify conditions)
	Official member states	Solutions	21.10.54			
		Drugs:	21.10.54.110		Dissolution time	0,5 – 120 min; Pass test/ fail test
		Suppositories (sticks)	21.10.54.120			(if necessary, specify conditions)
			21.10.54.140		Uniformity	Uniform/non uniform; Pass test/
			21.10.54.140			fail test
		Solutions	21.10.54.160		Foaming capacity:	
			21.10.54.170		- foam value;	10 =00
			21.10.54.170		- foam resistance	10 – 700 mm;
			21.10.54.190			0.3 - 1
			21.20.1			
			21.20.10			
			21.20.10			
			21.20.10.159			
			21.20.10.213			
			21.20.21.130			
			21.20.21.139			
			02.30.40.140			
			15.12.11			
		Drugs:	21.10.20.120	3004	Weight (volume) of	0,1 – 25000 ml
		Drops:	21.10.32		package contents	(cm3; 1; dm3);
		- eye;	21.10.5			0.1 - 10 kg;
		- Ear;	21.10.51.120			0,001 – 500 g;
		- nasal;	21.10.51.121			1,0 – 5000 mg;
		- sublingual;	21.10.51.122			Compliant/ Non compliant; Pass

1 4
test
40 kg/m^3
$3,000 \text{ mg/cm}^3$
3,000 mg/cm ³
- , - ,
3,00

-	2		1 4	-		243 pages, j
1	2			5	6	1
1	2	Drops: - eye; - Ear; - nasal; - sublingual - for local application; - for oral use.	4 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.160 21.10.54.170 21.10.54.180 21.10.54.190	5	6	7
			21.20.1 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140	2004	S' - C - 1	000.5
		Drugs: Capsules	21.10.51.120 21.10.54 21.10.54.180 21.20.1 21.20.10	3004	Size Capsules	000-5 0,01-160 mm 0,001-16 cm
316	PCR Reagent Kit Instruction manual- SCHMALLENBERG FACTOR (manufacturing company - VET FAKTOR LLC)	Biological material	-	-	Schmallenberg virus RNA	positive (detected) / negative (not detected)
317	PCR Reagent Kit Instruction manual- NODULAR DERMATITIS CATTLE- FACTOR (manufacturing company - VET FAKTOR LLC)	Biological material	-	-	Nodular dermatitis virus DNA (Limpy skin disease virus, LSDV)	positive (detected) / negative (not detected)
318	Meteorological recommendations for identification and differentiation of bird metapneumovirus based on PCR with hybridization-fluorescent detection of amplification products; FGBU «VGNKI», approved 16.11.2016	Immunobiological drugs for veterinary use Cell culture	21.20.21.137	3002 30 000	Fragment of the genome of the (aMPV) virus subtype A Fragment of the genome of the (aMPV) virus subtype B	detected / not detected detected / not detected

1	2	3	4	5	6	243 pages, j
319	Methodological guidelines for detection of infectious anaemia virus in chickens based on PCR with hybridization-fluorescent detection of amplification products; FGBU «VGNKI», approved 16.11.2016	Immunobiological drugs for veterinary use Cell culture	21.20.21.137	3002 30 000	Genome fragment of Chicken anaemia virus (CAV)	detected / does not contain
320	Methodological guidelines for the detection of egg-drop syndrome virus PCR method in trials for Foreign matter in live vaccines for birds; FGBU «VGNKI», approved 16.11.2016	Immunobiological drugs for veterinary use Cell culture	21.20.21.137	3002 30 000	genome fragment of Egg drop syndrome (EDS)	contain / does not contain
321	Methodology for detection and differentiation of Mycoplasma bovigenitalium and Mycoplasma califormicum based on PCR with hybridization-fluorescent detection of	Semen Biological material	01.42.20.000	0511 10 000 0 0511 99 853	genome fragment Mycoplasma bovigenitalium (M. bovigenitalium)	detected / not detected detected / not detected
	amplification products, FGBU «VGNKI», approved 16.02.2018		24.42.20.000	0511 10 000	genome fragment Mycoplasma califormicum (M. califormicum	
322	Methodology DNA Mycoplasma bovis detection based on PCR with hybridization-fluorescent detection of amplification products; FGBU «VGNKI», approved 09.06.2018	Semen Biological material	01.42.20.000	0511 10 000 0 0511 99 853	genome fragment Mycoplasma bovis (M. bovis)	detected / not detected
323	Methodology DNA Ureaplasma diversum detection based on PCR with hybridization-fluorescent detection of amplification products; FGBU «VGNKI», approved 09.06.2018	Semen Biological material	01.42.20.000	0511 10 000 0 0511 99 853	genome fragment Ureaplasma diversum (U. diversum)	detected / not detected
324	Method of detection of genome virus of disease Schmallenberg based on PCR with hybridization-fluorescent detection of amplification products FGBU «VGNKI», approved 28.11.2018	Semen Biological material	01.42.20.000	0511 10 000 0 0511 99 853	RNA virus of diseaseSchmallenberg	detected / not detected
325	Method of detection of Nodular dermatitis virus DNA based on PCR with hybridization-fluorescent detection of amplification products; FGBU «VGNKI», approved 21.06.2018	Semen Biological material	01.42.20.000	0511 10 000 0 0511 99 853	genome fragment LSDV (Limpy skin disease virus)	detected / not detected

1	2	3	4	5	6	7 7
326	Marek's disease identification kit Method polymerase chain reaction; manufacturing company–FractalBio	Biological material	-	-	DNA virus of Marek's disease	detected / not detected
327	Instructions for using the kit to identify DNA porcine circovirus type 2 (manufacturing company– FractalBio);	Biological material	-	-	DNA porcine circovirus type 2	detected / not detected
328	Instructions for the kit "LSI VETMAX for detection of M. paratuberculosis Method Advanced" Method real-time PCR	Biological material	-	-	DNA M. paratuberculosis	positive (detected) / negative (not detected)
329	GOST ISO 21527-1	Food products (with water activity of more than 95%), feed for animals	10.110.8 10.91-10.92	02-05 07-11 14,15 17-21	Yeast Moulds	10 ¹ – 9,9·10 ⁹ CFU/g (cm3) 0 – 500 CFU/g (cm3)
330	GOST ISO 21527-2	Food products (with water activity of more than 95%), feed for animals	10.110.8 10.91-10.92	02-05 07-11 14,15 17-21	Yeast Moulds	10 ¹ – 9,9·10 ⁹ CFU/g (cm3) 0 – 500 CFU/g (cm3)
331	GOST 33566	Milk and dairy products Milk and dairy products	10.51, 10.52 10.51, 10.52	0401-0406 0401-0406	Yeast Moulds	$10^{1} - 9.9 \cdot 10^{9} \text{ CFU/g (cm3)}$ 0 - 500 CFU/g (cm3)
332	GOST 33951	Milk and dairy products	10.51, 10.52	0401-0406	Lactic acid microorganisms	10 ¹ – 9,9·10 ⁹ CFU/g (cm3)
333	GOST 7702.2.1 Coming into force since 01.01.2019 P. 7.1 P.8.1 P. 8.2	Poultry meat Offal, semifinished poultry meat	10.12.1, 10.12.2, 10.12.4, 10.12.50.200, 10.12.50.300	0207	Mesophilic aerobic and facultative anaerobic microorganisms	10 ¹ – 9,9·10 ⁹ CFU/g (cm3)
334	METHODOLOGICAL GUIDELINES 4.2.1890-04	Food products and flush from production facilities. Biological agents	-	-	Determination of antibiotic resistance of microorganisms Determination of the minimum inhibitory concentration	Sensitive/Resistant 0,015-512- μg / ml
335	Methodical guidelines «Identification of microorganisms with use of microflex MALDI Biotyper mass spectrometer in the study of food raw materials and food	Food Raw material and food products	10.1-10.8	02-05 07-11 14,15 17-21,23	Species identification of microorganisms	-

1	2	3	4	5	6	7 7
	products»					
336	GOST 34106	One-component food products and raw materials from animal meat, fish, caviar	03.11.12, 03.11.2, 03.12.12, 03.12.2, 03.21.12, 03.21.2, 03.21.5, 03.22.1, 03.22.2, 03.22.4, 10.20.1, 10.20.2, 10.11.1, 10.11.3, 10.13.11, 10.13.12	0301-0305; 1604, 0201-0205; 0208; 0210; 1602 41, 1602 50, 1602 90	Types of fish (description) Kind of mammal meat (description)	-
337	Methodological guidelines «Detection and identification of Corynebacterium glutamicum molecular genetic methods»	Feed additives containing amino acids obtained by microbiological synthesis	10.91.10.170, 10.91.10.180	-	DNA Corynebacterium glutamicum Identification of DNA Corynebacterium glutamicum	Detected/ Not detected
338	Methodological guidelines. «Identification of mutations associated with the most common hereditary pathologies of Holstein breed cattle, using molecular genetic methods»	Semen bovine native and frozen, venous blood	01.42.20	05 11 10 0000	Mutation carriage in the FANCI gene associated with brahispinal syndrome (BY) Mutation carriage in the gene APOB, associated with cholesterol deficiency (CDH) Mutation carriage in the gene FXI, associated with deficiency XI FACTOR blood coagulation (FXID)	Normal genotype (homozygous by normal allele)/ Mutation Carrier (heterozygote, present one mutant and one normal gene copy)/ Mutant genotype (homozygous by mutant allele)

1	2	3	4	5	6	7 243 pages,
339	Methodological guidelines. «Identification of mutations associated with the most common hereditary pathologies of brown svitz cattle by molecular genetic methods»	Semen bovine native and frozen, venous blood	01.42.20	05 11 10 0000	Mutation carriage in the gene SUOX, associated with arachnomelia and arthrogryposis syndrome (SAA)	Normal genotype (homozygous by normal allele)/ Mutation Carrier (heterozygote, present one mutant and one normal gene copy)/ Mutant genotype (homozygous by mutant allele)
340	Methodological guidelines. «Identification of mutations associated with the most common hereditary pathologies of aberdeen angus cattle breed, molecular genetic methods»	Semen bovine native and frozen, venous blood	01.42.20	05 11 10 0000	Mutation carriage in the gene SLC4A2, associated with osteopetrosis (OS)	Normal genotype (homozygous by normal allele)/ Mutation Carrier (heterozygote, present one mutant and one normal gene copy)/ Mutant genotype (homozygous by mutant allele)
					Mutation carriage in the gene MSTN, associated with muscle hypertrophy (DM)	by mutant anere)
341	Guidelines for the use of a set of reagents Amplisens® CamV-FL, Manufacturing company-Central Research Institute of Epidemiology of Rospotrebnadzor, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	DNA mosaic and cauliflower viruses (DNA CamV)	Detected/ not detected
342	Guidelines for the use of a set of reagents Amplisens® GM Soybean-lines-1-FL, Manufacturing company-Central Research Institute of Epidemiology of Rospotrebnadzor, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM maize lines 40-3-2, A5547-127, A2704-12, FG72, Syht0h2.	Detected/ not detected
343	Guidelines for the use of a set of reagents Amplisens® GM Soybean-lines-2-FL, Manufacturing company-Central Research Institute of Epidemiology of Rospotrebnadzor, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1,	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM maize lines CV127, MON 87701, MON 89788.	Detected/ not detected

1	2	3	4	5	6	7 243 pages, j
			10.2, 10.5, 10.6, 10.7, 10.8, 10.9			
344	Methodology for determining the species of animal ingredients. Method polymerase chain reaction	Food products, feed for animals, feed additives and raw materials for their production	10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	0201-0205; 0208; 0210; 1602 41, 1602 50, 1602 90	DNA horse (<i>Equus</i> caballus), DNA furbearing animals	Detected/ not detected
345	Method of detection of Plant DNA «Soybean/Raps/maize» Method multiplex polymerase chain reaction with real-time hybridization-fluorescent detection.	Food products, feed, feed additives, agricultural raw materials and seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of soybean, Raps, maize (screening Method: qualitative determination DNA soybean /maize, DNA Raps)	Detected/ not detected
346	-Guidelines for the use of a set of reagents to detect DNA of genetically modified plants in foods Method polymerase chain reaction (PCR) with hybridization-fluorescent detection «Amplisens®GM Plant-1-FL». Manufacturing company— Central Research Institute of Epidemiology of Rospotrebnadzor, Moscow;	Food products, feed for animals, raw materials, seeds.	10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9 01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005,1201, 2304000001, 1901-1902, 2103, 2104, 2106, 2301- 2304, 2308, 2309	Plant DNA DNA P-35S DNA T-NOS; DNA P-FMV (Detection of genetically modified organisms of vegetable products(screening))	Found /not found
347	METHODOLOGICAL GUIDELINES A - 1/047 Method identification and quantification GM potato line H92-527-1 Method Real-time PCR	Feed, feed additives, Food products and raw materials	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM potato line EH92-527-1	Detected/ not detected
348	METHODOLOGICAL GUIDELINES A- 1/046 Method identification and quantification GM rice line LL62	Feed, feed additives, Food products and raw materials	01.11, 01.12, 01.13.39, 01.13.49.110,	1005, 1201, 2304, 2103, 2301-2304,	Identification of GM rice line LL62	Detected/ not detected

						243 pages,
1	2	3	4	5	6	7
	Method Real-time PCR		01.13.51,	2308, 2309		
			01.13.7,			
			01.19.10, 10.1,			
			10.2, 10.5,			
			10.6, 10.7,			
			10.8, 10.9			
349	Method of detection of genetic	Feed for animals, feed additives and	01.11, 01.12,	1005, 1201,	Genetic constructions	Detected/ not detected
	constructions bar, cp4epsps, nptII, P-	raw materials for their production.	01.13.39,	2304, 2103,	bar, cp4epsps, nptII, P-	
	rice-Act1 and T-35S for screening		01.13.49.110,	2301-2304,	rice-Act1 and T-35S	
	studies for the presence of components		01.13.51,	2308, 2309	(Detection of genetically	
	GM in products of vegetable products		01.13.7,		modified organisms of	
			01.19.10, 10.1,		vegetable	
			10.2, 10.5,		products(screening)	
			10.6, 10.7,		F	
			10.8, 10.9			
350		Feed for animals, food products	01.11, 01.12,	1005, 1201,	Identification of GM	Detected/ not detected
000		Tood for minimus, 1000 products	01.13.39,	2304, 2103,	soybean lines BPS-	Bettetten, not accepted
			01.13.49.110,	2301-2304,	CV127-9	
	Instructions for using the test system/set		01.13.51,	2308, 2309	C 1 12 / 3	
	of reagents«Soybean BPS-CV127-9		01.13.7,			
	Identification of» Organization-manufacter		01.19.10, 10.1,			
	CJSC Syntol, Moscow		10.2, 10.5,			
			10.2, 10.3, 10.6, 10.7,			
			10.8, 10.9			
351		Earl for animals for domainst	01.11, 01.12,	1005, 1201,	Identification of GM	Detected/ not detected
331		Feed for animals, food products		2304, 2103,		Detected/ not detected
			01.13.39,	2301-2304,	soybean lines MON 89788	
	Instructions for using the test system/set		01.13.49.110,	2308, 2309		
	of reagentsSoybean MON 89788		01.13.51,	2000, 2000		
	Identification of, Organization-manufacter		01.13.7,			
	CJSC Syntol, Moscow		01.19.10, 10.1,			
	- '		10.2, 10.5,			
			10.6, 10.7,			
<u> </u>			10.8, 10.9			
352		Feed for animals, food products	01.11, 01.12,	1005, 1201,	Identification of GM	Detected/ not detected
			01.13.39,	2304, 2103,	soybean lines MON 87701	
	Instructions for using the test system/set		01.13.49.110,	2301-2304, 2308, 2309		
	of reagentsSoybean MON 87701		01.13.51,	2308, 2309		
	Identification of, Organization-manufacter		01.13.7,			
	CJSC Syntol, Moscow		01.19.10, 10.1,			
			10.2, 10.5,			
			10.6, 10.7,			

1	2	3	4	5	6	7 7
			10.8, 10.9			
353	Instructions for using the test system/set of reagentsMaize MON89034 Identification of, Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM soybean lines MON89034	Detected/ not detected
354	Instructions for using the test system/set of reagentsMaize 5307 Identification of, Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM line maize line 5307	Detected/ not detected
355	Instructions for using the test system/set of reagentsMaize MIR162 Identification of, Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM line maize line MIR162	Detected/ not detected
356	Instructions for using the test system/set of reagents"Soybean/GTS 40-3-2 amount" Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM Soybean line GTS 40-3-2	0,1-10 %
357	Instructions for using the test system/set of reagents "Maize MON810 Amount" Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51,	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM maize MON810	0,5-10%

			ı	1		243 pages,
1	2	3	4	5	6	7
			01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9			
358	Instructions for using the test system/set of reagents«Soybean A2704-12 amount» Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM Soybean A2704-12	0,1-10 %
359	Instructions for using the test system/set of reagents«Soybean A5547-127 amount» Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM Soybean line A5547-127	0,1-10 %
360	Instructions for using the test system/set of reagents«Maize 5307 amount» Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM- Maize 5307	0,1-10 %
361	Instructions for using the test system/set of reagents«Maize Bt11 amount» Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM- Maize line Bt11	0,098-4,89 %

_	,		1		,	243 pages,
1	2	3	4	5	6	7
362	Instructions for using the test system/set of reagents«Soybean SYHTOH2 amount» Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM Soybean line SYHTOH2	0,1-10 %
363	Instructions for using the test system/set of reagents«Soybean FG72 amount» Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM Soybean line FG72	0,1-10 %
364	Instructions for using the test system/set of reagents«Maize T25 amount» Organization-manufacter CJSC Syntol, Moscow	Feed for animals, food products	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Quantitative content GM- Maize line T25	0,1-10 %
365	Guidelines for the use of a set of reagents (diplex option) «Mon87708/Mon87769», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM Soybean line Mon87708, Mon87769	Detected/ not detected
366	Guidelines for the use of a set of reagents (triplex option) «Soybean/maize/Raps», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7,	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of DNA soybean, maize and Raps	Detected/ not detected

1	2	3	4	5	6	7 243 pages,
			01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9			
367	Guidelines for the use of a set of reagents (triplex option) «CTP2-cp4-epsps/tE9/Pisum sativum», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Genetic constructions CTP2- CP4-epsps, terminator tE9, DNA pea (Pisum sativum)	Detected/ not detected
368	Guidelines for the use of a set of reagents «GT73-amount», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of Quantitative determination GM-Raps line GT73	0,1-5 %
369	Guidelines for the use of a set of reagents (triplex option) «Mon87705/Mon87708/Mon87769», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM Soybean line Mon87705, Mon87708, Mon87769	Detected/ not detected
370	Guidelines for the use of a set of reagents «Mon89788-amount», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of quantitative GM Soybean line Mon89788	0,1-5 %
371	Guidelines for the use of a set of reagents	Food products, Feed and feed	01.11, 01.12,	1005, 1201, 2304, 2103,	Gen pat and promoter	Detected/ not detected

						243 pages,
1	2	3	4	5	6	7
	(diplex option) «pat/pSsuAra», manufacturing company FGBU «VGNKI», Moscow	additives, agricultural raw materials, seeds.	01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	2301-2304, 2308, 2309	pSSuAra	
372	Guidelines for the use of a set of reagents (diplex option) «CTP2-CP4-epsps/tE9», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Genetic construction CTP2-CP4-epsps and tE9 terminator.	Detected/ not detected
373	Guidelines for the use of a set of reagents (monoplex option) «FG72», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM Soybean line FG72.	Detected/ not detected
374	Guidelines for the use of a set of reagents (diplex option) «FG72/40-3-2», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM Soybean line FG72, 40-3-2.	Detected/ not detected
375	Guidelines for the use of a set of reagents (triplex option) «BPS-CV127-09/DP305423/DP356043», manufacturing company FGBU «VGNKI», Moscow	Food products, Feed and feed additives, agricultural raw materials, seeds.	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1,	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM Soybean line BPS-CV127-09, DP305423, DP356043.	Detected/ not detected

1	2	3	4	5	6	7 243 pages,
			10.2, 10.5, 10.6, 10.7, 10.8, 10.9			
376	Guidelines for the use of a set of reagents to detect DNA fur-bearing animals in feed Method polymerase chain reaction (PCR) with hybridization-fluorescent detection «PCR-DNA-CARNIVORE-1-FACTOR», Organization-manufacter - "VET FACTOR", Troitsk	Feed and feed additives	01.11, 01.12, 01.13.39, 01.13.49.110, 01.13.51, 01.13.7, 01.19.10, 10.1, 10.2, 10.5, 10.6, 10.7, 10.8, 10.9	0301-0305; 1604, 0201-0205; 0208; 0210; 1602 41, 1602 50, 1602 90	DNA fur-bearing animals Mustelidae family, <i>Mustelidae</i> .	Detected/ not detected
377	GOST 34104	Feed for animals and plant Raw material	01.11, 01.12, 01.13.39, 01.13.49.110,	1005, 1201, 2304, 2103, 2301-2304, 2308, 2309	Identification of GM soybean line 40-3-2; Identification of GM	Detected/ not detected Detected/ not detected
			01.13.51, 01.13.7, 01.19.10, 10.1,	2308, 2309	soybean line A2704-12; Identification of GM soybean line A5547-127;	Detected/ not detected
			10.2, 10.5, 10.6, 10.7,		Identification of GM soybean line FG 72;	Detected/ not detected
			10.8, 10.9		Identification of GM soybean line MON89788;	Detected/ not detected
					Identification of GM soybean line MON87701;	Detected/ not detected
					Identification of GM soybean line BPS-CV127-9;	Detected/ not detected
					Identification of GM soybean line SYHT0H2;	Detected/ not detected
					Identification of GM soybean line MON87705;	Detected/ not detected
					Identification of GM soybean line MON87708;	Detected/ not detected
					Identification of GM soybean line MON87769;	Detected/ not detected

1 2	3	4 5	6	7 243 pages,
			Identification of GM soybean line DP-305423;	Detected/ not detected
			Identification of GM soybean line DP-356043	Detected/ not detected
			Identification of GM soybean line DAS-44406	Detected/ not detected
			Identification of GM soybean line DAS-81419	Detected/ not detected
			Identification of GM soybean line DAS-68416	Detected/ not detected
			Identification of GM maize line MON810;	Detected/ not detected
			Identification of GM maize line NK 603;	Detected/ not detected
			Identification of GM maize line T 25;	Detected/ not detected
			Identification of GM maize line GA 21;	Detected/ not detected
			Identification of GM maize line MIR 604;	Detected/ not detected
			Identification of GM maize line MON 863;	Detected/ not detected
			Identification of GM maize line 3272;	Detected/ not detected
			Identification of GM maize line MON 88017;	Detected/ not detected
			Identification of GM maize line Bt 11;	Detected/ not detected
			Identification of GM maize line 5307;	Detected/ not detected
			Identification of GM maize line MON 89034;	Detected/ not detected
			Identification of GM maize line Bt176;	Detected/ not detected
			Identification of GM maize line MON 98140;	Detected/ not detected
			Identification of GM maize line MON 87460;	Detected/ not detected
			Identification of GM maize line TC1507;	Detected/ not detected

1	2	3	4	5	6	7 243 pages,
					Identification of GM maize line 59122;	Detected/ not detected
					Identification of GM maize line LY038;	Detected/ not detected
					Identification of GM maize line DAS40278;	Detected/ not detected
					Identification of GM maize line MIR 162	Detected/ not detected
					Identification of GM Raps line GT73;	Detected/ not detected
					Identification of GM Raps line MON88302;	Detected/ not detected
					Identification of GM Raps line MS1;	Detected/ not detected
					Identification of GM Raps line MS8;	Detected/ not detected
					Identification of GM Raps line T45;	Detected/ not detected
					Identification of GM Raps line RF1;	Detected/ not detected
					Identification of GM Raps line RF2;	Detected/ not detected
					Identification of GM Raps line RF3;	Detected/ not detected
					Identification of GM Raps line Topas;19/2	Detected/ not detected
					Quantitative content GM Soybean line 40-3-2;	0,1-5 %
					Quantitative content GM Soybean line A2704-12;	0,1-5 %
					Quantitative content GM Soybean line A5547-127;	0,1-5 %
					Quantitative content GM Soybean line MON89788;	0,1-5 %
					Quantitative content GM Soybean line MON87701;	0,1-5 %
					Quantitative content GM Soybean line BPS-CV-127-9;	0,1-5 %
					Quantitative content GM Soybean line FG 72;	0,1-5 %

1	2	3	4	5	6	243 pages, j
1	<u> </u>	<u> </u>	_	<u> </u>	Quantitative content GM	0,1-5 %
					Soybean line SYHT0H2;	0,1-3 /0
					Quantitative content GM	0,1-5 %
					Soybean line MON87705; Quantitative content GM	,
					Soybean line MON87708;	0,1-5 %
					Quantitative content GM Soybean line MON87769;	0,1-5 %
					Quantitative content GM Soybean line DP-305423;	0,1-5 %
					Quantitative content GM	
					Soybean line DP-356043;	0,1-5 %
					Quantitative content GM	0.1.5.0/
					Soybean GM soybean lines DAS-44406	0,1-5 %
					Quantitative content GM	
					Soybean GM soybean lines	0,1-5 %
					DAS-81419	, , ,
					Quantitative content GM	
					Soybean line GM soybean	0,1-5 %
					line DAS-68416 Quantitative content GM	
					maize line MON810;	0,1-5 %
					Quantitative content GM	0.1.7.0
					maize line NK 603;	0,1-5 %
					Quantitative content GM	0,1-5 %
					maize line T 25;	0,1 5 /0
					Quantitative content GM maize line GA 21;	0,1-5 %
					Quantitative content GM	
					maize line MIR 604;	0,1-5 %
					Quantitative content GM	0.1.5.0/
					maize line MON 863;	0,1-5 %
					Quantitative content GM	0,1-5 %
					maize line 3272;	5,1 5 /6
					Quantitative content GM maize line MON 88017;	0,1-5 %
					Quantitative content GM maize line Bt 11;	0,1-5 %
					Quantitative content GM maize line 5307;	0,1-5 %

						243 pages,
1	2	3	4	5	6	7
					Quantitative content GM maize line MON 89034;	0,1-5 %
					Quantitative content GM maize line Bt176;	0,1-5 %
					Quantitative content GM maize line 98140;	0,1-5 %
					Quantitative content GM maize line MON 87460;	0,1-5 %
					Quantitative content GM maize line TC1507;	0,1-5 %
					Quantitative content GM maize line MON 59122;	0,1-5 %
					Quantitative content GM maize line LY038;	0,1-5 %
					Quantitative content GM maize line DAS40278;	0,1-5 %
					Quantitative content GM maize line MIR 162	0,1-5 %
					Quantitative content of GM Raps line GT73	0,1-5 %
					Quantitative content of GM Raps line MON88302	0,1-5 %
					Quantitative content of GM Raps line MS1	0,1-5 %
					Quantitative content of GM Raps line MS8	0,1-5 %
					Quantitative content of GM Raps line T45	0,1-5 %
					Quantitative content of GM Raps line RF1	0,1-5 %
					Quantitative content of GM Raps line RF2	0,1-5 %
					Quantitative content of GM Raps line RF3	0,1-5 %
					Quantitative content Of GM Raps line Topas 19/2	0,1-5 %

Zvenigorodskoye shosse 5, 123022, Moscow, Russia

143511, Russia, Moscow region, Istra district, Manikhino laboratory facility 1, laboratory facility 2, laboratory facility 4, office building 2

1	2	3	4	5	6	243 pages, j
378	GOST P 57221 p. 20	Feed yeast, microbial synthesis protein feed products	-	2308 2309	Bacterial semination,	(0-10 ⁸) CFU/g (ml)-
	P.21		-	2308 2309	salmonella	Detected/Not found
379	GOST 23050 p.6.1	Virus vaccine against Aujeszky's disease	-	3002	Appearance and color. Presence of foreign matter, bottle and ampule crack, labelling violation	Homogeneous dry porous mass of white and yellow with a pink tint. Not allowed.
	p.6.3				Rehydration time	1-2 min
	p.6.4				Contamination of bacterial, fungal microflora	Not allowed
	p.6.5				Mycoplasma contamination	Not allowed
	p.6.6				Activity by infectivity titer in cell culture	Less 10 5,0 TCID 50/cm3
	p.6.7				Activity by infectivity titer for rabbits	Less 10 ^{3,0} LD50/cm3
	p.6.8				Harmlessness	Must be harmless for sheep aged between one and five years
	p.6.9				Immunogenic activity	Must be immunogenic for sheep aged between one and five years.
380	GOST P 55283 p.7.1	Inactivated rabies vaccines	-	3002	Appearance and color. Presence of foreign matter, violation of the consistency, integrity of the ampoules (of bottle)	Homogeneous cream colour dry porous mass
	p.7.2				PH	7,2±0,5
	p.7.3	7			Resuspension time	1-2 min
	p.7.5				Presence ampoule vacuum	Compliant/non compliant
	p.7.6				Contamination of bacterial, fungal microflora and mycoplasms	Compliant/non compliant
	p.7.7				Complete inactivation	-
	p.7.8				Immunogenic activity	Compliant/non compliant
381	METHODOLOGICAL RECOMMENDATIONS Method for microbiological detection of	Feed and feed additives	-	3501	Primary producer strain Corynebacterium glutamicum	Compliant/non compliant

1	2	3	4	5	6	7 7
	Corynebacterium glutamicum - amino acid producers in feed and feed additives, approved by FGBU VGNKI» 7.02.2018					
382	European pharmacopoeia in.8.0 art. 2.2.3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	Immunobiological drugs for veterinary use	-	3002 3003 3004	pН	0-14 Unit pH
	art. 2.6.12 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	Non-sterile Immunobiological drugs for veterinary use	-	3002 3003 3004	Total number of viable aerobic	Compliant/non compliant
	art. 2.6.13 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	Non-sterile Immunobiological drugs for veterinary use	-	3002	Presence of certain microorganisms	Compliant/non compliant
	Art. 2.6.7 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State	Immunobiological drugs for veterinary use	-	3002 3003 3004	Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant

[2	3	4	5	6	243 pages, 7
	registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art. 2.6.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Immunobiological drugs for veterinary use	-	3002 3003 3004	Sterility (contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	art. 2.6.24 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Immunobiological drugs	-	3002 3003 3004	Presence of foreign matter (contamination by foreign matter, contamination by foreign viruses (viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	art. 2.6.25 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Immunobiological drugs	-	3002 3003 3004	Presence of foreign matter (contamination by foreign matter, contamination by foreign viruses (viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	art.5.2.4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Presence of foreign matter (contamination by foreign matter, contamination by foreign viruses (viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	3 mon 1 mon				Activity (titer, virus titration, efficacy,	Compliant/non compliant

1	2	3	4	5	6	7 7
					antigenic component titer, viral component titration)	
	art. 04/2013:0870 and other normative documents approved in the established order, specifying the	Newcastle disease vaccine inactivated	-	3002	Authenticity (authenticity, Identificatio)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
					Activity (titer, virus titre, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
					Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant
	art. 04/2013:0450 and other normative documents approved in the established order, specifying the	Newcastle disease vaccine (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi,	Compliant/non compliant

1	2	3	4	5	6	7 7
	Eurasian Economic Union member states				bacterial and fungal sterility)	
					Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
					Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art. 04/2013:2038 and other normative documents approved in the established order, specifying the	Infectious anemia vaccine for chicken (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter,	Compliant/non compliant

1	2	3	4	5	6	7 7
					contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	
					Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art. 04/2013:0588 and other normative documents approved in the established order, specifying the	Avian infectious encephalomyelitis vaccine (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
					Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant

			5		
art. 04/2013:0587 and other normative documents approved in the established order, specifying the	Vaccine against avian infectious bursal disease (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
art. 04/2013:0960 and other normative documents approved in the established order, specifying the	Vaccine against avian infectious bursal disease (inactivated)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility) Residual live virus	Compliant/non compliant Compliant/non compliant

2	3	4	5	6	243 pages,
	3	1 7		(Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation) Specific foreign matters	,
				(Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
art. 04/2013:1202 and other normative documents approved in the established order, specifying the application of the research (testing)	Vaccine against egg-drop syndrome-76 (inactivated)	-	3002	Authenticity (authenticity, Identification) Bacteria and fungi	Compliant/non compliant
method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				(Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
				Residual live virus	Compliant/non compliant

1	2	3	4	5	6	7 243 pages,
					(Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
					Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art. 04/2013:1951 p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Adenovirus vaccine (live) for dogs	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-2 and other normative documents approved				Bacteria and fungi (Sterility, contamination	Compliant/non compliant

						243 pages,
1	2	3	4	5	6	7
	in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-3 and other normative documents approved in the established order, specifying the				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant

1	2	3	4	5	6	7 7
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art. 04/2013:1298 p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Adenovirus vaccine (inactivated) for dogs	-	3002	Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant
	p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing)				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component	Compliant/non compliant

1	2	3	4	5	6	243 pages,
1			4	<u> </u>	-	,
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				titration)	
	p.2-3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					Compliant/non compliant
	art.04/2013:0451 p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Rabies vaccine (inactivated) for veterinary use	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing				Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant

1	2	3	4	5	6	7 7
	requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art.04/2013:0744 p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Vaccine against Auezki's disease (inactivated) for pigs	-	3002	Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant
	p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal	Compliant/non compliant

1	2	3	4	5	6	7 7
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				sterility)	
	p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	art.04/2013:0745 p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Vaccine against Auezki's disease (live) for pigs for parenteral use	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant

1	2	3	4	5	6	243 pages,
1	registers of drug for veterinary use of the Eurasian Economic Union member		7	3		,
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Foreign matter (Specific foreign matters, Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State				Activity	Compliant/non compliant

1	2	3	4	5	6	7 7
	registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art.04/2013:0589 and other normative documents approved in the established order, specifying the	Vaccine for marek's disease (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
					Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art.04/2013:1177 p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State	Bovine respiratory syncytial virus vaccine (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant

1	2	3	4	5	6	243 pages,
	registers of drug for veterinary use of the Eurasian Economic Union member states			-		
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing)				Activity	Compliant/non compliant

1	2	3	4	5	6	7 7
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-3					
	and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					Compliant/non compliant
	art.04/2013:1952 p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Bovine viral diarrhea vaccine (inactivated)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing				Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant

1	2	3	4	5	6	243 pages, j
	requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art.04/2013:1315 and other normative documents approved in the established order, specifying the	Hepatitis virus vaccine I (live) for ducks	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility) Specific foreign matters	Compliant/non compliant Compliant/non compliant

1	2	3	4	5	6	7 7
					(Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign wiruses, foreign pathogens) Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art.04/2013:1613 p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Herpes virus vaccine (inactivated) for horses	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State				Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant

1	2	3	4	5	6	7 7
	registers of drug for veterinary use of the Eurasian Economic Union member states;					
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art. 04/2013:1955 p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Parainfluenza virus vaccine for dogs (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant

	, , , , , , , , , , , , , , , , , , ,		•	1		243 pages,
1	2	3	4	5	6	7
	Eurasian Economic Union member					
	states;					
	p.3-3				Mycoplasmas	
	and other normative documents approved				(Mycoplasma	
	in the established order, specifying the				contamination, Sterility	
	application of the research (testing)				for mycoplasma,	
	method, measurements, establishing				mycoplasma sterility)	Compliant/non compliant
	requirements for drugs registered in the					Compitant/non compitant
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;					
	p.3-4				Specific foreign matters	
	and other normative documents approved				(Presence of foreign	
	in the established order, specifying the				matter, contamination by	
	application of the research (testing)				foreign matter,	
	method, measurements, establishing				contamination by foreign	Compliant/non compliant
	requirements for drugs registered in the				viruses, viral purity,	Compitant non Compitant
	established order and included in State				contamination by foreign	
	registers of drug for veterinary use of the				matter, contamination by	
	Eurasian Economic Union member				foreign viruses, foreign	
	states;				pathogens)	
	p.3-5				Virus titre (activity, titer,	
	and other normative documents approved				virus titration, efficacy,	
	in the established order, specifying the				antigenic component	
ł	application of the research (testing)				titer, viral component	
	method, measurements, establishing				titration)	Compliant/non compliant
ł	requirements for drugs registered in the					Compilation Compilation
	established order and included in State					
ĺ	registers of drug for veterinary use of the					
ĺ	Eurasian Economic Union member					
ł	states;					
	p.3-6				Activity	Compliant/non compliant
	and other normative documents approved					
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					

				_	T	243 pages,
1	2	3	4	5	6	7
	states; p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member					
	states; art.04/2013:1206 p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Rhinotracheitis virus vaccine (live) for cats	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant

1	2	3	4	5	6	7 7
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens) Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity	Compliant/non compliant
	art. 04/2013:1207	Rhinotracheitis virus vaccine	=	3002	Authenticity	Compliant/non compliant

1	2	3	4	5	6	7 7
	p.3-1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State	(inactivated) for cats			(authenticity, Identification)	
	registers of drug for veterinary use of the Eurasian Economic Union member states p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	states; p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art. 04/2013:1176 p.3-1	Bovine parainfluenza vaccine (live)	-	-	Authenticity (authenticity,	Compliant/non compliant

1	2	2	4			243 pages,
1	2	3	4	5		1
1	and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing)	3	4	5	Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of	7
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	p.3-5 and other normative documents approved in the established order, specifying the				Virus titre (activity, titer, virus titration, efficacy, antigenic component	Compliant/non compliant

1	2	3	4	5	6	7 7
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				titer, viral component titration)	
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity	Compliant/non compliant
	art. 04/2013:2325 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Hemorrhagic disease vaccine (inactivated) for rabbits	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing)				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of	Compliant/non compliant

2	3	4	5	6	7
method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member				bacteria and fungi, bacterial and fungal sterility)	
states; p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non complia
p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-2 and other normative documents approved in the established order, specifying the				Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non complia
application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
art. 04/2013:0249 p.3-1 and other normative documents approved in the established order, specifying the	Influenza vaccine (inactivated) for horses	-	3002	Authenticity (authenticity, Identification)	Compliant/non complia

1		2	4	_		243 pages, j
1	2	3	4	5	6	1
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the					
	established order and included in State registers of drug for veterinary use of the					
	Eurasian Economic Union member states				Destruit and Consi	
	p.3-2 and other normative documents approved				Bacteria and fungi (Sterility, contamination	
	in the established order, specifying the				with bacterial and fungal microflora, presence of	
	application of the research (testing) method, measurements, establishing				bacteria and fungi,	
	requirements for drugs registered in the				bacterial and fungal	Compliant/non compliant
	established order and included in State				sterility)	
	registers of drug for veterinary use of the				•	
	Eurasian Economic Union member					
	states;					
	p.3-3				Residual live virus	
	and other normative documents approved in the established order, specifying the				(Complete inactivation, virus inactivation,	
	application of the research (testing)				avirulence, residual	
	method, measurements, establishing				virulence, residual virulence, inactivation)	
	requirements for drugs registered in the				,	Compliant/non compliant
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states; p.3-4					
	and other normative documents approved					
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing				Activity (virus titer, titer,	
	requirements for drugs registered in the				virus titration, efficacy,	
	established order and included in State				antigenic component	
	registers of drug for veterinary use of the				titer, viral component	Compliant/non compliant
	Eurasian Economic Union member states:				titration)	
	p.2-3-2					
	and other normative documents approved					
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					

1	2	3	4	5	6	7 7
	requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art.04/2013:0249 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Influenza vaccine (inactivated) for pigs	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-1 p.32 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity,	Compliant/non compliant

1	2	3	4	5	6	7 7
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.3-5				contamination by foreign matter, contamination by foreign viruses, foreign pathogens) Activity (virus titre, titer,	
	and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.2-3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					Соприанопон соприан
	art. 04/2013:0696 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Infectious bull rhinotracheitis vaccine (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State		-	-	Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant

1	2	3	4	5	6	243 pages, j
	registers of drug for veterinary use of the Eurasian Economic Union member states;					
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;		-	-	Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;		-	-	Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;		-	-	Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the		-	-	Activity	Compliant/non compliant

1	2	3	4	5	6	7 243 pages, j
	Eurasian Economic Union member					
	states;					
	p.2-3-4		-	-		
	and other normative documents approved					
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;					
	art. 04/2013:2461	Infectious rhinotracheitis vaccine	_	3002	Authenticity	
	and other normative documents approved	(live) for turkey		3002	(authenticity,	Compliant/non compliant
	in the established order, specifying the	(live) for turkey			Identification)	Compilant non Compilant
	application of the research (testing)				Bacteria and fungi	
	method, measurements, establishing				(Sterility, contamination	
	requirements for drugs registered in the				with bacterial and fungal	
	established order and included in State				microflora, presence of	Compliant/non compliant
	registers of drug for veterinary use of the				bacteria and fungi,	
	Eurasian Economic Union member states				bacterial and fungal	
					sterility)	
					Mycoplasmas	
					(Mycoplasma	
					contamination, Sterility	Compliant/non compliant
					for mycoplasma,	
					mycoplasma sterility)	
					Specific foreign matters	Compliant/non compliant
					(Presence of foreign	
					matter, contamination by	
					foreign matter, contamination by foreign	
					viruses, viral purity,	
					contamination by foreign	
					matter, contamination by	
					foreign viruses, foreign	
					pathogens)	
					Activity (virus titre, titer,	
					virus titration, efficacy,	
					antigenic component	Compliant/non compliant
					titer, viral component	

1	2	3	4	5	6	243 pages,
					titration)	
	art. 04/2013:0251 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Infectious enteritis vaccine (cat panleukopenia) (live) for cats	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant

1	2	3	4	5	6	7 7
	p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity	Compliant/non compliant
	art. 04/2013:1102 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Calicivirus vaccine (live) for cats	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2				Bacteria and fungi	Compliant/non compliant

					,	243 pages,
1	2	3	4	5	6	7
	and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				(Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	3-6 and other normative documents approved				Activity	Compliant/non compliant

1	2	3	4	5	6	7 7
	in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art.04/2013:1101 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Calicivirus vaccine (inactivated) for cats	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the				Residual live virus (Complete inactivation, virus inactivation,	Compliant/non compliant

1	2	3	4	5	6	7 7
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				avirulence, residual virulence. inactivation)	
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	states; art.2326 and other normative documents approved	Coccidiosis vaccine (live) for chickens	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma,	Compliant/non compliant

1	2	3	4	5	6	243 pages,
1	2	3	4	5	mycoplasma sterility) Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens) Number of sporulated	Compliant/non compliant Compliant/non compliant
					oocysts Activity (titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art. 04/2013:1953 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Coronavirus diarrhea vaccine(inactivated) for calf	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the				Residual live virus (Complete inactivation, virus inactivation,	Compliant/non compliant

1	2	3	4	5	6	7 243 pages,
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				avirulence, residual virulence. inactivation)	
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.2-3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					Compliant/non compliant
	art.04/2013:1321 and other normative documents approved in the established order, specifying the application of the research (testing)	Leukemia vaccine (inactivated) for cats	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant

1	2	2	1			243 pages,
1	2	3	4	5	6	1
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1					
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-2-2				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing					-

1	2	3	4	5	6	7 7
	requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art.04/2013:1942 and other normative documents approved in the established order, specifying the	Mycoplasma vaccine GALLISEPTICUM (inactivated)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	nents, establishing lrugs registered in the and included in State or veterinary use of the			Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Residual live mycoplasmas	Compliant/non compliant
					Activity (titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art.04/2013:1943 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Myxomatosis vaccine (live) for rabbits	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant

						243 pages,
1	2	3	4	5	6	7
	states;					
	p.3-3				Mycoplasmas	Compliant/non compliant
	and other normative documents approved				(Mycoplasma	
	in the established order, specifying the				contamination, Sterility	
	application of the research (testing)				for mycoplasma,	
	method, measurements, establishing				mycoplasma sterility)	
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;					
	p.3-4				Specific foreign matters	Compliant/non compliant
	and other normative documents approved				(Presence of foreign	
	in the established order, specifying the				matter, contamination by	
	application of the research (testing)				foreign matter,	
	method, measurements, establishing				contamination by foreign	
	requirements for drugs registered in the				viruses, viral purity,	
	established order and included in State				contamination by foreign	
	registers of drug for veterinary use of the				matter, contamination by	
	Eurasian Economic Union member				foreign viruses, foreign	
	states;				pathogens)	
	p.3-5				Virus titre (activity, titer,	Compliant/non compliant
	and other normative documents approved				virus titration, efficacy,	compliant non compliant
	in the established order, specifying the				antigenic component	
	application of the research (testing)				titer, viral component	
	method, measurements, establishing				titration)	
	requirements for drugs registered in the				titudion)	
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;					
	p.3-6				Activity	Compliant/non compliant
	and other normative documents approved				renvity	Compitant non compitant
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;					
	states,					

1	2	3	4	5	6	7 243 pages,
	p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art.04/2013:0649 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Smallpox vaccine (live) for poultry	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	p.3-4	1			Specific foreign matters	Compliant/non compliant

1	2	3	4	5	6	7 7
	and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				(Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens) Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-4-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity	Compliant/non compliant
	art.04/2013:0795 and other normative documents approved	Parvovirus vaccine (live) for dogs	-	3002	Authenticity (authenticity,	Compliant/non compliant

1		2	4	-		243 pages, j
1	2	3	4	3		/
	in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1 p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the	3	4	5	Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	Eurasian Economic Union member states; p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	p.3-5 and other normative documents approved in the established order, specifying the				Virus titre (activity, titer, virus titration, efficacy, antigenic component	Compliant/non compliant

1	2	3	4	5	6	7 7
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				titer, viral component titration)	
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity	Compliant/non compliant
	art.04/2013:0795 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1 p.3-2 and other normative documents approved	Parvovirus vaccine (inactivated) for dogs	-	3002	Authenticity (authenticity, Identification) Bacteria and fungi (Sterility, contamination	Compliant/non compliant Compliant/non compliant
	in the established order, specifying the application of the research (testing)				with bacterial and fungal microflora, presence of	

1	2	3	4	5	6	7
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				bacteria and fungi, bacterial and fungal sterility)	
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art.04/2013:0965 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Parvovirus vaccine (inactivated) for pigs	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing				Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant

1 2	3	4	5	6	7
requirements for drugs registered in the established order and included in State registers of drug for veterinary use of Eurasian Economic Union member states; p.3-4 and other normative documents approximate the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of Eurasian Economic Union member states; p.3-5 and other normative documents approximate the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the establishing requ	ne e the oved e the ov	4	5	Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens) Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant Compliant/non compliant
p.3-5 and other normative documents approin the established order, specifying the application of the research (testing)	ne e the the e the			Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component	Compliant/non compliant
registers of drug for veterinary use of Eurasian Economic Union member states; art.04/2013:1956 and other normative documents approin the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in t	Avian Viral Tenosynovitis Vaccine (live)	-	3002	Authenticity (authenticity, Identification) Bacteria and fungi (Sterility, contamination with bacterial and fungal	Compliant/non compliant Compliant/non compliant

1	2	3	4	5	6	243 pages, 7
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				microflora, presence of bacteria and fungi, bacterial and fungal sterility)	
					Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
					Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	art.04/2013:0442 and other normative documents approved in the established order, specifying the	Avian infectious bronchitis vaccine (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
					Specific foreign matters (Presence of foreign	Compliant/non compliant

1	2	3	4	5	6	7 243 pages,
					matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens) Activity (virus titre, titer, virus titration, efficacy, antigenic component titration)	Compliant/non compliant
	art.04/2013:0959 and other normative documents approved in the established order, specifying the	Avian infectious bronchitis vaccine (inactivated)	-	3002	titration) Authenticity (authenticity, Identification)	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
					Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
					Activity (virus titre, titer,	Compliant/non compliant

1	2	2	4	1 5		243 pages,	
I	2	3	4	5	6	1	
					virus titration, efficacy, antigenic component titer, viral component titration)		
	art.04/2013:1068 and other normative documents approved in the established order, specifying the	Avian Infectious Laryngotracheitis Vaccine (live)	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant	
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	g) ng in the State e of the			Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant	
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant	
					Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant	
	art.04/2013:1392 and other normative documents approved in the established order, specifying the	Avian Paramixovirus vaccine 3 (inactivated) for turkey	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant	
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of	Compliant/non compliant	

1	2	3	4	5	6	7 7
	registers of drug for veterinary use of the Eurasian Economic Union member states				bacteria and fungi, bacterial and fungal sterility)	
					Residual live virus (Complete inactivation, virus inactivation, avirulence, residual virulence. inactivation)	Compliant/non compliant
					Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
					Activity (virus titre, virus titration, efficiency, antigenic component titration, virus component titration)	Compliant/non compliant
	art.04/2013:1954 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Rotavirus diarrhea vaccine (inactivated) for calf	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant

					1	243 pages,
1	2	3	4	5	6	7
	Eurasian Economic Union member					
	states;					
	p.3-3				Residual live virus	Compliant/non compliant
	and other normative documents approved				(Complete inactivation,	
	in the established order, specifying the				virus inactivation,	
	application of the research (testing)				avirulence, residual	
	method, measurements, establishing				virulence. inactivation)	
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;					
	p.3-4				Specific foreign matters	Compliant/non compliant
	and other normative documents approved				(Presence of foreign	
	in the established order, specifying the				matter, contamination by	
	application of the research (testing)				foreign matter,	
	method, measurements, establishing				contamination by foreign	
	requirements for drugs registered in the				viruses, viral purity,	
	established order and included in State				contamination by foreign	
	registers of drug for veterinary use of the				matter, contamination by	
	Eurasian Economic Union member				foreign viruses, foreign	
	states;				pathogens)	
	p.3-5				Activity (virus titre, virus	Compliant/non compliant
	and other normative documents approved				titration, efficiency,	
	in the established order, specifying the				antigenic component	
	application of the research (testing)				titration, virus	
	method, measurements, establishing				component titration)	
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;					
	p.2-3-2				Activity (virus titre, virus	Compliant/non compliant
	and other normative documents approved				titration, efficiency,	_
	in the established order, specifying the				antigenic component	
	application of the research (testing)				titration, virus	
	method, measurements, establishing				component titration)	
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					

1	2	3	4	5	6	7 243 pages,
	states;					
	art.04/2013:0449 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Plague Vaccine (live) for Mustelids	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-1 p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Mycoplasmas (Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility)	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant

1	2	3	4	5	6	243 pages.
	p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states:				Virus titre (activity, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states:				Activity	Compliant/non compliant
	p.2-3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity	Compliant/non compliant
	art.04/2013:1938 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Plague Vaccine (live) for ducks	-	3002	Authenticity (authenticity, Identification) Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility) Mycoplasmas	Compliant/non compliant Compliant/non compliant Compliant/non compliant

1	2	3	4	5	6	7
1		3	4	5	(Mycoplasma contamination, Sterility for mycoplasma, mycoplasma sterility) Specific foreign matters (Presence of foreign matter, contamination by foreign matter, contamination by foreign wiruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign matter, contamination by foreign viruses, foreign pathogens) Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant Compliant/non compliant
	art.04/2013:0448 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1 p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing)	Plague Vaccine for dogs (live)	-	3002	Authenticity (authenticity, Identification) Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of	Compliant/non compliant
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.3-3 and other normative documents approved				bacteria and fungi, bacterial and fungal sterility) Mycoplasmas (Mycoplasma	Compliant/non compliant Compliant/non compliant

1	2	3	4	5	6	
1	in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states; p.3-4 and other normative documents approved in the established order, specifying the	3	4	5	contamination, Sterility for mycoplasma, mycoplasma sterility) Specific foreign matters (Presence of foreign matter, contamination by	7 243 pages.
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				foreign matter, contamination by foreign viruses, viral purity, contamination by foreign matter, contamination by foreign viruses, foreign pathogens)	Compliant/non compliant
	p.3-5 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	p.3-6 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity	Compliant/non compliant
	p.2-3-3 and other normative documents approved in the established order, specifying the					-

1	2	3	4	5	6	7 243 pages,
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art.04/2013:2448 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states p.3-1	Enzootic pneumonia vaccine (inactivated) for pigs	-	3002	Authenticity (authenticity, Identification)	Compliant/non compliant
	p.3-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi (Sterility, contamination with bacterial and fungal microflora, presence of bacteria and fungi, bacterial and fungal sterility)	Compliant/non compliant
	p.3-3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Residual live mycoplasmas	Compliant/non compliant
	p.3-4 and other normative documents approved in the established order, specifying the application of the research (testing)				Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component	Compliant/non compliant

1	2	3	4	5	6	7 243 pages,
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				titration)	
	p.2-2-2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Activity (virus titre, titer, virus titration, efficacy, antigenic component titer, viral component titration)	Compliant/non compliant
	Art. 2.2.8 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	Immunobiological drugs	-	3002	Viscosity	Compliant/non compliant
	Art. 2.2.9 Capillary Viscometry Method and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Viscosity	Compliant/non compliant
	Art. 2.2.10 Method of rotational viscometer and other normative documents approved in the established order, specifying the				Viscosity	Compliant/non compliant

1	2	3	4	5	6	7
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art. 2.2.1 (visual method) and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Immunobiological drugs	-	3002	Determination of the transparency and degree of turbidity of liquids	Compliant/non compliant
	art. 2.9.17 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Immunobiological drugs	-	3002	Volume	(0,00-500,00) ml
	Art. 04/2013:2525 p. 3.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	Dog Bordetellosis Vaccine (live)	-	3002	Authenticity	Compliant/non compliant
	p.3.2 and other normative documents approved in the established order, specifying the application of the research (testing)				Bacteria and fungi	Compliant/non compliant

1	2	3	4	5	6	7 7
	method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	p.3.3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Living Bacteria	(0-10 ¹²) CFU/g (ml)
	Art. 04/2013:1946 p. 3.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	Pasteurellosis vaccine haemolytica (inactivated) for sheep	-	3002	Authenticity	Compliant/non compliant
	p.3.2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi	Compliant/non compliant
	Art. 04/2013:2072 p. 3.1 and other normative documents approved in the established order, specifying the	Pasteurellosis vaccine trehalosi (inactivated) for sheep	-	3002	Authenticity	Compliant/non compliant

						243 pages,
1	2	3	4	5	6	7
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;			D. C. L. C. C.		
	p.3.2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi	Compliant/non compliant
	Art. 04/2013:1361 p. 3.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	Progressive Atrophic Rhinitis Vaccine (inactivated) for pigs		3002	Authenticity	Compliant/non compliant
	p.3.2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi	Compliant/non compliant
	p.3.3 and other normative documents approved in the established order, specifying the				Residual toxicity	Compliant/non compliant

1	2	3	4	5	6	7 243 pages,
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;					
	art. 04/2013:1944 p. 3.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	for cattle other normative documents approved e established order, specifying the ication of the research (testing) and, measurements, establishing irements for drugs registered in the olished order and included in State sters of drug for veterinary use of the sian Economic Union member s; other normative documents approved e established order, specifying the ication of the research (testing) and, measurements, establishing irements for drugs registered in the olished order and included in State others of drug for veterinary use of the sian Economic Union member	-	3002	Authenticity	Compliant/non compliant
	p.3.2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;				Bacteria and fungi	Compliant/non compliant
	art. 01/2013:0062 p. 3.1 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states;	Vaccines for veterinary use	-	3002	Authenticity	Compliant/non compliant
	p. 3.4 and other normative documents approved				Sterility	Compliant/non compliant

	2	3	4	5	6	7
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member					
	states;		24.20.24.422			
	0361	Clostridium shavo vaccine for	21.20.21.132	3002	Safety	Compliant/non complian
	p. 2-2-1.	veterinary use				
	p. 3-1.	4			Authenticity	Compliant/non complian
	p. 3-2.				Bacteria and fungi	Compliant/non complian
r	p. 3-3.	-			Activity	Compliant/non complian
	0364	Septicum clostridium vaccine for	21.20.21.132	3002	Safety	Compliant/non complian
	p. 2-2-1.	veterinary use				I was a second
	p. 3-1.	1			Authenticity	Compliant/non complian
	p. 3-2.	1			Bacteria and fungi	Compliant/non complian
	p. 3-3.	1			Residual toxicity;	Compliant/non compliant
	p. 3-4.				Activity	0- 10 IU/ml
	r					Compliant/non complian
	0362	Clostridium vaccine Novyi (type B)	21.20.21.132	3002	Authenticity	Compliant/non complian
ŀ	p. 3-1.	for veterinary use				r
	p. 3-1.		I		Bacteria and fungi	Compliant/non complian
		1			Dacteria and rungi	
	p. 3-2.					
	p. 3-2. p. 3-3.				Residual toxicity	Compliant/non complian
	p. 3-2.					Compliant/non complian 0 - 10 IU/ml
	p. 3-2. p. 3-3. p. 3-4.	Clostridium vaccine perfringens for	21.20.21.132	3002	Residual toxicity Activity	Compliant/non complian 0 - 10 IU/ml Compliant/non complian
	p. 3-2. p. 3-3.	Clostridium vaccine perfringens for veterinary use	21.20.21.132	3002	Residual toxicity	Compliant/non complian
	p. 3-2. p. 3-3. p. 3-4.		21.20.21.132	3002	Residual toxicity Activity	Compliant/non complian 0 - 10 IU/ml Compliant/non complian

2	3	4	5	6	7
p. 3-4.				Activity	Compliant/non complia
0360 p. 3-1.	Botulism vaccine for veterinary use	21.20.21.132	3002	Authenticity	Compliant/non complia
p. 3-2.				Bacteria and fungi	Compliant/non complia
p. 3-3.				Residual toxicity	Compliant/non complia
0697 p. 3-1.	Tetanus vaccine for veterinary use	21.20.21.132	3002	Authenticity	Compliant/non complia
p. 3-2.				Bacteria and fungi	Compliant/non complia
p. 3-3.				Residual toxicity	Compliant/non complia
0339	Antitoxic serum against alpha- toxin clostridium novyi for veterinary use	-	3002	Activity	Compliant/non complia
0340	Antitoxin serum against beta-toxin clostridium perfringens for veterinary use	-	3002	Activity	Compliant/non complia
20613 2.6.13 p 4-3	Immunobiological drugs for veterinary use	-	3002	Salmonella	Compliant/non complia
0341	Antitoxic Serum Against Epsilon - Clostridium Toxin perfringens for veterinary use	-	3002	Activity	Compliant/non complia
2520 p 2-2-1-1	Vaccine against salmonella Enteritidis (live, oral vaccine) for chickens	21.20.21.131	3002	Safety	Compliant/non complia
1947 p. 2-2-1	Vaccine against salmonella Enteritidis (inactivated) for chickens	21.20.21.132	3002	Safety	Compliant/non compliant
p.2-3-1				Activity	Compliant/non complia
2521 p. 2-2-1-1	Vaccine against salmonella Typhimurium (live, oral vaccine) for chickens	21.20.21.131	3002	Safety	Compliant/non complia
2361 p.2-2-1	Vaccine against salmonella Typhimurium (inactivated) for	21.20.21.132	3002	Safety	Compliant/non compliant
p.2-3-1	chickens			Activity	Compliant/non complia
0961 p.2-3-1.	Vaccine against colibacteriosis	21.20.21.132	3002	Activity	Compliant/non complia
p3-1.	(inactivated) for newborn ruminants			Authenticity	0 – 3,0 (OD 492) 0 - 3,5 RP Compliant/non complia
p.3-2.				Bacteria and fungi	Compliant/non compliant
0962 p.2-3-1. Vaccine against	Vaccine against colibacteriosis (inactivated) for newborn pigs	21.20.21.132	3002	Activity	0 – 3,5 RP (PR) 0 – 15 log2; 0 -100% ER25 – 70
p3-1.				Authenticity	0 – 3,5 RP (PR) 0 – 15 log2;

1	2	3	4	5	6	7 7
						0 -100% ER25 – 70
						Compliant/non compliant
	p.3-2.				Bacteria and fungi	Compliant/non compliant
	0447 p.2-3-1.	Vaccine against leptospirosis for	21.20.21.132	3002	Activity	Compliant/non compliant
	p3-1.	dogs (inactivated)			Authenticity	Compliant/non compliant
	p.3-2.				Bacteria and fungi	Compliant/non compliant
	0064 p.2-2-1.	Vaccine against Erysipelas	21.20.21.132	3002	Activity	0 - 10 METHODOLOGICAL
		(inactivated) for pigs				RECOMMENDATIONS
						0 - 200 IU/dose
						0 - 15 log 2 IE50%
						0 - 100 ppd (protective dose)
						0 - 5,0 ELISA unit
						0 – 3,5 RP (PR)
	p3-1.				Authenticity	Compliant/non compliant
	p.3-2.				Bacteria and fungi	Compliant/non compliant
383	P 4.2.2643,	Disinfectants	-	-	Fungicide activity	Effectively/ineffectively
	p. 5.3.2					
384	GOST P ISO 16256,	Pure yeast fungi culture	-	-	Minimum suppression	(0,03125 – 128) mg/ml
	p. 3				concentration	"sensitive" / "sensitive, dose
	European Committee on Antimicrobial					dependent" / "intermediate" /
	Susceptibility Testing (EUCAST)					"insensitive" /
	Antifungal Agents					resistant"
	Breakpoint tables for interpretation of					
	MICs					
	Version 9.0, valid since 2018-02-12					
385	GOST 54951	All kinds of animal feed	-	-	Mass fraction of	-
		The standard does not apply::			humidity (humidity,	
		a) Dairy products;			humidity)	
		b) mineral substances;c) mixtures containing a large amount				
		of dairy products or minerals (e.g. milk				
		substitutes);				
		d) feed for animals containing				
		humectants (e.g. propylene glycol);				
		e) animal and vegetable fats and oils,				
		oilseeds, oilcake, grain and cereal				
		products.				
386	GOST 34310	Biological drugs for veterinary use	-	3002	Phenol	(0-5)%
					26 11 1	(0,1-10000) mcg/ml
					Merthiolate	(0-0,1)%
						(0-1) mg/ml
						(0-1000) mcg/ml

1	2	3	4	5	6	7 243 pages,
					Formaldehyde	(0-1)%
						(0-500) mcg/ml
						(0-0,5) g/l
						(0-5) mg/ml
387		Biological, pathological material,	-	-	Specific antibodies to the	(0 - 1: 400)
	serological diagnosis of brucellosis in	blood serum from animals.			brucellosic Antigen	Presence of specific antibodies
	cattle and small cattle in an indirect	Immunobiological drugs for				to brucellosis Antigen / Absence
	hemagglutination reaction(Indirect	veterinary use.				of specific antibodies to
	hemagglutination test)	Diagnostic drugs.				brucellosis Antigen
						Positive reaction - Presence of
						agglutination of sensitized
						erythrocytes with test sera at a
						dilution of 1:100 with a score of
						Less than two crosses (++).
						Doubtful reaction - Presence of
						agglutination of sensitized
						erythrocytes with test sera at a dilution of 1:50 with a score of
						Less than two crosses (++) and
						1: 100 with a score of no higher
						than one cross (+).
						The reaction is negative -
						Presence of erythrocyte
						agglutination in a 1:50 dilution
						of sera with an assessment of
						one cross (+) and with its
						complete absence (-).
388	Instruction for use of the set of	Biological, pathological material,	_	_	Specific antibodies to	0 - 1: 400)
	preparations for diagnostics of infectious	blood serum from animals.			B.ovis	Presence of specific antibodies
	epididymitis of sheep in Indirect	Immunobiological drugs for				to B.ovis / Absence of specific
	hemagglutination test and Antibody	veterinary use.				antibodies to B.ovis
	Neutralization Reaction	Diagnostic drugs.				Positive reaction - Presence of
						agglutination of sensitized
						erythrocytes with test sera at a
						dilution of 1: 100 with a score
						of Less than two crosses (++).
						Doubtful reaction - Presence of
						agglutination of sensitized
						erythrocytes with test sera at a
						dilution of 1:50 with a score of
						Less than two crosses (++) and
						1: 100 with a score of no higher

1	2	3	4	5	6	243 pages,
1	<u> </u>	<u> </u>	<u> </u>	<u> </u>	<u> </u>	than one cross (+).
						The reaction is negative -
						Presence of erythrocyte
						agglutination in a 1:50 dilution
						of sera with an assessment of
						one cross (+) and with its
						complete absence (-).
389	GOST 30347	Milk and dairy products	10.51	0401	Staphylococcus aureus	(0-10 ⁸) CFU/g (ml)
369	GOS1 30347	Wink and dairy products	10.52	0401	Staphylococcus aureus	(0-10) C1·0/g (IIII)
390	GOST 33924		10.51	0401	Bifidobacteria	(0-10 ¹⁰) CFU/g (ml)
390	GOS1 33924		10.52	0401	Billidobacteria	(0-10) CFO/g (IIII)
391	GOST 33951		10.51	0401	Lactic acid	(0-10 ¹⁰) CFU/g (ml)
391	GOS1 33931		10.51	0401	microorganisms	(0-10 °) CFO/g (IIII)
202	GOST 33566		10.52	0401		(0-10 ¹⁰) CFU/g (ml)
392	GOS1 33300		10.51	0401	Yeast and mold fungi	(0-10°°) CFU/g (mi)
393	GOST 33568		10.52	0401	Salt-tolerant	(0-10 ⁸) CFU/g (ml)-
393	GOS1 33308		10.51	0401		(0-10°) CFU/g (mi)-
20.4	COST 20005	F 1 1		0401	microorganisms	(0.1012) CELL/ (1)
394	GOST 28805	Food products	10.11 – 10.89	0401 1601	Yeast and mold fungi	(0-10 ¹²) CFU/g (ml)
				1602		
				1901		
				Groups		
				21,22,23		
395	GOST ISO 21527-1		10.11 – 10.89	0401	Yeast and mold fungi	(0-10 ¹²) CFU/g (ml)
				1601		
				1602		
				1901		
				Groups 21,22,23		
396	GOST ISO 21527-2		10.11 – 10.89	0401	Yeast and mold fungi	(0-10 ¹²) CFU/g (ml)
390	GOST ISO 21327-2		10.11 – 10.89	1601	reast and mold fungi	(0-10) Cr0/g (IIII)
				1602		
				1901		
				Groups		
				21,22,23		
397	GOST 30706	Dairy products for child nutrition	10.86.10.100	0401 20 110	Yeast and mold fungi	(0-10 ¹²) CFU/g (ml)-
				1		
				0403 90 510		
				1 0406 10 500		
				1		
398		Drugs for veterinary use	-	-	Microbiological purity	Compliant/non compliant
	PHARMACOPOEIA XIV					

1	2	3	4	5	6	7 7
	GENERAL PHARMACOPOEIA ARTICLE.1.2.4.0002.18					
	and other normative documents approved					
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states					
	GENERAL PHARMACOPOEIA	Probiotic Drugs for veterinary use	-	-	Safety of test probiotics	
	ARTICLE.1.7.2.0001.15				in vivo	
	and other normative documents approved					
	in the established order, specifying the					
	application of the research (testing)					Compliant/non compliant
	method, measurements, establishing					Compilation Compilation
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states					
	GENERAL PHARMACOPOEIA	Probiotic Drugs for veterinary use	-	-	Specific activity of	
	ARTICLE. 1.7.2.0009.15				probiotics	
	and other normative documents approved					
	in the established order, specifying the					
	application of the research (testing)					Compliant/non compliant
	method, measurements, establishing					r r
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states				D : : : : : : : : : : : : : : : : : : :	
	GENERAL PHARMACOPOEIA	Colic Probiotic Drugs for veterinary		-	Determination of the	
	ARTICLE. 1.7.1.0005.15	use			number of viable bacteria	
	and other normative documents approved				in 1 dose of	Compliant/non compliant
	in the established order, specifying the				immunobiological drugs	•
	application of the research (testing)				(amount of live bacterial cells)	
	method, measurements, establishing requirements for drugs registered in the				,	
	established order and included in State				Description	Compliant/non compliant
	registers of drug for veterinary use of the				(Appearance)	r
	Eurasian Economic Union member states				Acid generation activity	Compliant/non compliant
	Lurasian Leonomic Omon member states					Compitant non compitant

-	2	2	4			243 pages,
1	2	3	4	5	6	·1
					Antagonistic activity	Compliant/non compliant
					Microbiological purity	Compliant/non compliant
					Authenticity	Compliant/non compliant
	GENERAL PHARMACOPOEIA ARTICLE. 1.7.1.0008.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Probiotic Drugs for veterinary use	-	-	Determination of the number of viable bacteria in 1 dose of immunobiological drugs (amount of live bacterial cells) Description (Appearance)	Compliant/non compliant
	GENERAL PHARMACOPOEIA ARTICLE. 1.7.1.0006.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing		-	-	Determination of the number of viable bacteria in 1 dose of immunobiological drugs (amount of live bacterial cells)	Compliant/non compliant
	requirements for drugs registered in the established order and included in State				Description (Appearance)	Compliant/non compliant
	registers of drug for veterinary use of the Eurasian Economic Union member states				Acid generation activity	Compliant/non compliant
	Larasian Leonomic Cinon member states				Antagonistic activity	Compliant/non compliant
					Microbiological purity	Compliant/non compliant
					Authenticity	Compliant/non compliant
	GENERAL PHARMACOPOEIA ARTICLE. 1.7.1.0003.15 and other normative documents approved in the established order, specifying the	Bifid Probiotic Drugs for veterinary use	-	-	Determination of the number of viable bacteria in 1 dose of immunobiological drugs	Compliant/non compliant
	application of the research (testing) method, measurements, establishing				Description (Appearance)	Compliant/non compliant
	requirements for drugs registered in the				Acid generation activity	Compliant/non compliant

1	2	3	4	5	6	7 7
	established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states				Antagonistic activity	Compliant/non compliant
	GENERAL PHARMACOPOEIA ARTICLE. 1.7.1.0009.15 and other normative documents approved in the established order, specifying the	Spore Probiotic Drugs for veterinary use	-	-	Determination of the number of viable bacteria in 1 dose of immunobiological drugs	Compliant/non compliant
	application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the				Description (Appearance)	Compliant/non compliant
	Eurasian Economic Union member states				Acid generation activity	Compliant/non compliant
					Antagonistic activity	Compliant/non compliant
					Microbiological purity	Compliant/non compliant
					Authenticity	Compliant/non compliant
	GENERAL PHARMACOPOEIA ARTICLE.1.7.2.0024.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Biological drugs for veterinary use	-	3002	Formaldehyde	(0-1)% (0-500) mcg/ml (0-0,5) g/l (0-5) mg/ml
	GENERAL PHARMACOPOEIA ARTICLE.1.7.2.0025.18 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the	Biological drugs for veterinary use	-	3002	Thimerosal	(0-0,1)% (0-1) mg/ml (0-1000) mcg/ml

1	2	3	4	5	6	7 243 pages,
	Eurasian Economic Union member states					
	GENERAL PHARMACOPOEIA	Biological drugs for veterinary use	-	3002	Phenol	(0-5)%
	ARTICLE.1.7.2.0028.18					(0,1-10000) mcg/ml
	and other normative documents approved					
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states					
	GENERAL PHARMACOPOEIA	Drugs, which, according to the	21.20.21.131	3002300000	Sterility	Compliant/non compliant
	ARTICLE.1.2.4.0003.15	regulatory documentation or	21.20.21.132	3002905000 3002909000		
	and other normative documents approved	pharmacopoeia articles, should be	21.20.21.133	3002909000		
	in the established order, specifying the	sterile.	21.20.21.134	3002120002		
	application of the research (testing)		21.20.21.139	3002190000		
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states	Y' '1 1 1'1	21 20 21 121	3002300000	Visible mechanical	G I' // I'
	GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0005.18	Liquid and solid parenteral dosage forms	21.20.21.131 21.20.21.132	3002300000	impurities	Compliant/non compliant
		TOTTIS	21.20.21.132	3002909000	impurities	
	and other normative documents approved in the established order, specifying the		21.20.21.133	3002120002		
	application of the research (testing)		21.20.21.134	3002150000		
	method, measurements, establishing		21.20.21.139	3002190000		
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states					
	GENERAL PHARMACOPOEIA	Immunobiological drugs for	21.20.21.131	3002300000	Solubility	Compliant/non compliant
	ARTICLE.1.2.1.0005.15	veterinary use	21.20.21.131	3002905000	Solubility	Compilate for compilate
	and other normative documents approved	, continuity use	21.20.21.132	3002909000		
	in the established order, specifying the		21.20.21.134	3002120002		
	application of the research (testing)		21.20.21.139	3002150000		
	method, measurements, establishing		=1.20.21.109	3002190000		
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states					

1	2	3	4	5	6	7 7
	GENERAL PHARMACOPOEIA ARTICLE.1.2.4.0004.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Substances of natural origin, drugs, derived from the blood, organs, tissues of humans or animals, plant raw materials, microorganisms and products of their vital functions in the production of ready-made forms, mainly for parenteral use	21.20.21.131 21.20.21.132 21.20.21.133 21.20.21.134 21.20.21.139 21.1 21.10 21.10.1 21.10.20.120 21.10.5 21.10.5 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.129 21.10.51.129 21.10.51.129	3002300000 3002905000 3002909000 3002120002 3002150000 3003 – 3004 from 4201 from 3808	Abnormal Toxicity/Toxicity in test dose	Compliant/non compliant
	GENERAL PHARMACOPOEIA ARTICLE.1.2.4.0005.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Injection Solutions and Pharmaceutical substances	21.10.53 21.10.53.120 21.10.54 21.10.54.110 21.10.54.120 21.10.54.130 21.10.54.140 21.10.54.150 21.10.54.160 21.10.54.170 21.10.54.190 21.20.1 21.20.10 21.20.10 21.20.10.158 21.20.10.159 21.20.10.213 21.20.21.130 21.20.21.139 02.30.40.140 15.12.11 20.20.14	3002	Pyrogenicity	Compliant/non compliant

1	2	3	4	5	6	243 pages, j
			20.20.14.000			
	GENERAL PHARMACOPOEIA	Immunobiological drugs, blood sera	21.20.21.131	3002300000	Immunogenic activity	Compliant/non compliant
	ARTICLE.1.7.2.0033.15	of target animals, including poultry,	21.20.21.132	3002905000	Activity	Compliant/non compliant
	and other normative documents approved	biological, pathological material	21.20.21.133	3002909000	Specificity	Compliant/non compliant
	in the established order, specifying the		21.20.21.134	3002120002 3002150000	Antigen specificity	Compliant/non compliant
	application of the research (testing)		21.20.21.139	3002130000	Antibodies	Compliant/non compliant
	method, measurements, establishing			2002170000	Antibody titer	Compliant/non compliant
	requirements for drugs registered in the				-	
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states		21 20 21 121	200220000		
	GENERAL PHARMACOPOEIA	Immunobiological drugs for	21.20.21.131	3002300000 3002905000	Immunogenic activity	Compliant/non compliant
	ARTICLE.1.7.2.0008.15	veterinary use	21.20.21.132	3002903000	Antigenic activity	Compliant/non compliant
	and other normative documents approved		21.20.21.133	3002120002	Live microbial cells /	Compliant/non compliant
	in the established order, specifying the application of the research (testing)		21.20.21.134 21.20.21.139	3002150000	Concentration, amount of	
	method, measurements, establishing		21.20.21.139	3002190000	microbial cells	
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states					
	GENERAL PHARMACOPOEIA	Drugs and immunobiological drugs	_	3002	Viscosity	(0,3 – 10000) mPa*s
	ARTICLE.1.2.1.0015.15	2 rugo unu mimumorrorogram urugo		5002	dynamic	$(0.6 - 300) \text{ mm}^2/\text{c}$
	and other normative documents approved				kinetic	(-,,
	in the established order, specifying the					
	application of the research (testing)					
	method, measurements, establishing					
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states					
	GENERAL PHARMACOPOEIA	Drugs and immunobiological drugs	-	3002	Mass fraction of	(0,00-25.0)%
	ARTICLE.1.2.1.0010.15 method 1				humidity (humidity,	
	and other normative documents approved				relative humidity,	
	in the established order, specifying the				Moisture content, weight	
	application of the research (testing)				loss during drying,	
	method, measurements, establishing				water)	
	requirements for drugs registered in the					
	established order and included in State					
	registers of drug for veterinary use of the					
	Eurasian Economic Union member states					

1	2	3	4	5	6	7 7
	GENERAL PHARMACOPOEIA ARTICLE.1.2.3.0012.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs, isolated from natural sources or obtained by biotechnological methods	-	3002	Determination of protein mass fraction	(0-50)% (0-10)mg/ml
	GENERAL PHARMACOPOEIA ARTICLE.1.7.2.0016.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Immunobiological drugs for veterinary use	-	3002	Determination of aluminium ions (mass fraction of aluminium, aluminium hydroxide) AL(OH)3	(0-10)mg/ml (0-10)%
	GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0006.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Immunobiological drugs for veterinary use	-	3002	Colour	Compliant/non compliant
	GENERAL PHARMACOPOEIA ARTICLE.1.4.2.0003.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states and other normative documents approved in the established order that specify the	Dosage forms for parenteral use	-	3002	Volume (extractable, nominal, extractable, fill volume) Volume control. Volume of vaccine in consumer package. Average Volume of Filling. Volume of primary package. Amount of drug in bottles).	(0,00-500,00) ml

1	2	3	4	5	6	7 243 pages,
	application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and included in the State registries of feeds and feed additives of the Eurasian Economic Union member states GENERAL PHARMACOPOEIA	Drugs	_	3002	nH (nH concentration	(0-14) un pH
	GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0004.15 p.2 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states and other normative documents approved in the established order that specify the application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and included in the State registries of feeds and feed additives of the Eurasian Economic Union member states p.3 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states and other normative documents approved in the established order that specify the application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and included order and included in State registered in the established order that specify the application of the research (testing) method, measurements that establish requirements for feed and feed additives registered in the established order and	Drugs	-	3002 3003 – 3004; 2308-2309 3501	pH (pH, concentration hydrogen ions, concentration hydrogen ions in 1% solution, active acidity, concentration hydrogen ions in 5% solution, concentration hydrogen ions in 10% solution, etc.)	(0-14) un. pH

1	2	3	4	5	6	7
	included in the State registries of feeds and feed additives of the Eurasian Economic Union member states					
	GENERAL PHARMACOPOEIA ARTICLE.1.2.1.0014.15 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	(Biological drugs for veterinary use (Veterinary serum allergens, blood products and obtained by genetic engineering used in veterinary medicine)	-	3002	Density	(0,7-1,840) g/cm3
399	GOST 32296 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances. Drugs: Solutions: - for injection; Suspensions and emulsions: - for injection; Drops (eye)	10.91.10.110 10.91.10.120 10.91.10.180 10.92.10.300 21.1 21.10 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.126 21.10.51.129 21.10.52.110 21.20.1 21.20.10 21.20.10.213 02.30.40.140	-	Determination of acute toxicity in the intragastric administration of the fixed-dose method	Grades 1 to 5 on a globally harmonized system for the classification and labeling of chemicals
400	GOST 32373 and other normative documents approved in the established order, specifying the application of the research (testing) method, measurements, establishing requirements for drugs registered in the established order and included in State registers of drug for veterinary use of the Eurasian Economic Union member states	Drugs. Medicinal plant raw material and collections. Pharmaceutical substances. Drugs: Solutions: - for injection; Suspensions and emulsions: - for injection; Drops (eye)	10.91.10.120 10.91.10.180 10.92.10.300 21.1 21.10 21.10.51.120 21.10.51.121 21.10.51.122 21.10.51.123 21.10.51.124 21.10.51.125 21.10.51.126 21.10.51.126	-	Determination of acute toxicity in skin intake	Determination of LD ₅₀

1	2	3	4	5	6	7
			21.10.52.110			
			21.20.1 21.20.10 21.20.10.213 02.30.40.140			
			21.20.10			
			21.20.10.213			
			02.30.40.140			

Director VGNKI		L. K. Kish
post of the authorized person	signature of the authrized person	initials, surname of the authorized person