Elena Kovaleva

List of Publications by Year
in descending order

25	46	1937685 4	1872680 6
papers	citations	h-index	g-index

25	25	25	46
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Modern Approaches to Estimating the Content of Genotoxic Impurities in Drugs (a Review). Pharmaceutical Chemistry Journal, 2016, 49, 765-770.	0.8	8
2	CURRENT REQUIREMENTS FOR THE QUALITY OF HERBAL MEDICINAL PRODUCTS. The Bulletin of the Scientific Centre for Expert Evaluation of Medicinal Products, 2018, 8, 170-178.	0.2	8
3	Comparative Analysis of Quality Assessment Requirements for Gelatin Used in Drug Production (Review). Pharmaceutical Chemistry Journal, 2017, 50, 820-825.	0.8	7
4	Developing methodological approaches to standardization of pharmaceutical substances. Pharmaceutical Chemistry Journal, 2010, 44, 33-39.	0.8	4
5	From Implementation of the State Standard for Drug Quality (OST 91.500.05.001-00) to Creation of the New Russian State Pharmacopoeia. Pharmaceutical Chemistry Journal, 2003, 37, 602-604.	0.8	3
6	Comparison of the Nomenclatures of Herbal Substances Used in the Russian and Foreign Pharmacopoeial Texts. The Bulletin of the Scientific Centre for Expert Evaluation of Medicinal Products, 2020, 10, 29-40.	0.2	3
7	Manufacturing Requirements for Excipients Used in Medicines (Review). Pharmaceutical Chemistry Journal, 2015, 49, 340-343.	0.8	2
8	General Principles for Composing Dosage Form Names. Pharmaceutical Chemistry Journal, 2015, 48, 683-686.	0.8	2
9	Current Requirements for Assessment of Elemental Impurities (Heavy Metals) in Medicines. Pharmaceutical Chemistry Journal, 2018, 52, 84-89.	0.8	2
10	Title is missing!. Pharmaceutical Chemistry Journal, 2001, 35, 680-682.	0.8	1
11	Current Problems of Drug Expertise and Standardization. Pharmaceutical Chemistry Journal, 2005, 39, 333-335.	0.8	1
12	Preparing a pharmacopoeial article on pharmaceutical substances. Pharmaceutical Chemistry Journal, 2007, 41, 34-36.	0.8	1
13	Improvement of methodological approaches to the standardization of medicines in the pharmaceutical dosage form of tablets. Pharmaceutical Chemistry Journal, 2009, 43, 668-676.	0.8	1
14	Use of Hydrophilic Interaction Liquid Chromatography to Separate Butylhydroxyanisole Isomers. Pharmaceutical Chemistry Journal, 2015, 49, 203-205.	0.8	1
15	Comparison of Approaches to Stability Testing of Medicines in the Russian Federation and the Eurasian Economic Union. The Bulletin of the Scientific Centre for Expert Evaluation of Medicinal Products, 2021, 11, 16-23.	0.2	1
16	Quality Control of Ethyl Alcohol Used as a Medicinal Product. The Bulletin of the Scientific Centre for Expert Evaluation of Medicinal Products, 2020, 10, 6-18.	0.2	1
17	The Standardization of Therapeutic Agents: Current Status. Pharmaceutical Chemistry Journal, 2000, 34, 617-618.	0.8	0
18	Title is missing!. Pharmaceutical Chemistry Journal, 2002, 36, 43-44.	0.8	0

#	Article	IF	Citations
19	Harmonization of approaches to evaluation of the disintegration characteristics of tablets and capsules. Pharmaceutical Chemistry Journal, 2007, 41, 232-234.	0.8	0
20	Developing approaches to dosage uniformity evaluation in the Russian State Pharmacopoeia. Pharmaceutical Chemistry Journal, 2012, 45, 640-645.	0.8	0
21	Quality Assessment of Excipients at the Drug Registration Stage. Pharmaceutical Chemistry Journal, 2015, 49, 393-397.	0.8	0
22	Determination of Organic Impurities in Combination Drugs. Pharmaceutical Chemistry Journal, 2017, 51, 126-129.	0.8	0
23	Development of an Approach to the Assessment of Changes to Approved Biological Products. BIOpreparations Prevention Diagnosis Treatment, 2019, 19, 109-117.	0.5	0
24	Nonspecific Impurities in Pharmaceutical Substances: Characteristics of Test Methods. The Bulletin of the Scientific Centre for Expert Evaluation of Medicinal Products, 2019, 9, 153-161.	0.2	0
25	Current Requirements for the Degree of Fineness of Herbal Substances and Herbal Medicinal Products. The Bulletin of the Scientific Centre for Expert Evaluation of Medicinal Products, 2020, 10, 218-227.	0.2	0