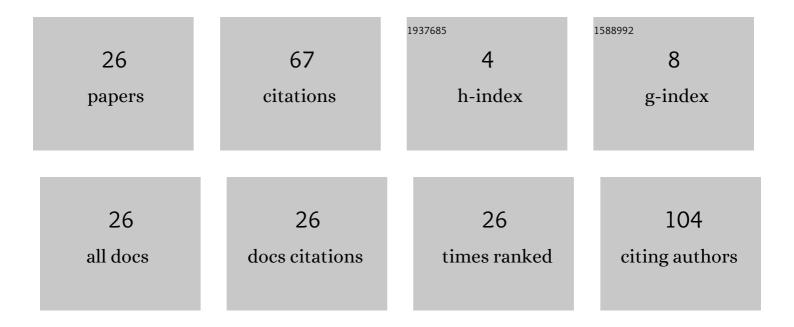
Natalia Sadchikova

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4203063/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Development and Validation of Pomalidomide Determination in Human Plasma by HPLC-MS/MS Method. Drug Development and Registration, 2020, 9, 146-154.	0.6	2
2	A facile synthesis and microtubule-destabilizing properties of 4-(1H-benzo[d]imidazol-2-yl)-furazan-3-amines. European Journal of Medicinal Chemistry, 2015, 94, 237-251.	5.5	26
3	Synthesis and antimicrobial activity of geminal bis-hydroperoxides. Pharmaceutical Chemistry Journal, 2010, 44, 248-250.	0.8	11
4	Evaluating the pharmaceutical equivalence of drugs in the registration stage. Pharmaceutical Chemistry Journal, 2009, 43, 171-175.	0.8	1
5	Use of the "dissolution―test for evaluation of the pharmaceutical equivalence of tablet formulations of phenazepam. Pharmaceutical Chemistry Journal, 2008, 42, 48-50.	0.8	3
6	Quantitative estimation of components of combined hormonal contraceptives by HPLC. Pharmaceutical Chemistry Journal, 2008, 42, 291-293.	0.8	5
7	Current state of IR spectroscopy applied to pharmaceutical analysis. Pharmaceutical Chemistry Journal, 2008, 42, 466-470.	0.8	2
8	Harmonization of quality indicators of the domestic parent substance fentanyl. Pharmaceutical Chemistry Journal, 2008, 42, 550-552.	0.8	0
9	Drug particle shape and size control: A necessary factor for high-quality drug production. Pharmaceutical Chemistry Journal, 2007, 41, 40-49.	0.8	5
10	Direct molding technology for the production of zolpidem tablets. Pharmaceutical Chemistry Journal, 2007, 41, 659-661.	0.8	1
11	Direct molding of tablets with prolonged drug release. Pharmaceutical Chemistry Journal, 2006, 40, 448-451.	0.8	0
12	Optimization of the Composition and Technology of Diazoline Tablets. Pharmaceutical Chemistry Journal, 2005, 39, 437-440.	0.8	0
13	Studying Prothionamide Release in vitro from Tablets. Pharmaceutical Chemistry Journal, 2004, 38, 327-329.	0.8	0
14	Quality Evaluation and Standardization of Fentanyl and the Related Injection Preparation. Pharmaceutical Chemistry Journal, 2004, 38, 336-338.	0.8	2
15	Composition and technology of tinidazole core tablets. Pharmaceutical Chemistry Journal, 2004, 38, 628-631.	0.8	2
16	A Comparative Analysis of Requirements to the Pharmacopoeial Drug Dissolution Test. Pharmaceutical Chemistry Journal, 2003, 37, 37-43.	0.8	0
17	Optimization of the Composition and Technology of Pentoxifyllin Core Tablets. Pharmaceutical Chemistry Journal, 2003, 37, 536-539.	0.8	0
18	The Problem of Device Calibration for the Pharmacopoeial Drug Dissolution Test. Pharmaceutical Chemistry Journal, 2003, 37, 550-555.	0.8	0

NATALIA SADCHIKOVA

#	Article	IF	CITATIONS
19	Development of the Composition and Production Technology of Carvedilol Tablets. Pharmaceutical Chemistry Journal, 2003, 37, 594-598.	0.8	2
20	Title is missing!. Pharmaceutical Chemistry Journal, 2001, 35, 453-457.	0.8	2
21	Principal aspects in the development of pharmacopoeial analysis. Pharmaceutical Chemistry Journal, 2000, 34, 271-272.	0.8	0
22	Mass-spectrometric identification of impurities in Ditilin. Pharmaceutical Chemistry Journal, 1999, 33, 665-670.	0.8	0
23	Evaluation of the quality of ditilin (suxamethonium iodide) preparations with respect to host impurities. Pharmaceutical Chemistry Journal, 1999, 33, 616-618.	0.8	1
24	Identification of impurities in commercial benzonal preparations by coupled chromatography—Mass spectrometry method. Pharmaceutical Chemistry Journal, 1999, 33, 568-572.	0.8	1
25	Spectrophotometric analysis of benzonal in tablets. Pharmaceutical Chemistry Journal, 1997, 31, 212-214.	0.8	0
26	Use of UV spectra for the identification of sulfanilamide preparations. Pharmaceutical Chemistry Journal, 1981, 15, 681-686.	0.8	1