## Konstantin Chegaev

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

56 1,037 30 20 h-index g-index citations papers 60 1,186 3.62 5.3 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
56	Sdox, a HS releasing anthracycline, with a safer profile than doxorubicin toward vasculature <i>Vascular Pharmacology</i> , <b>2022</b> , 143, 106969	5.9	1
55	A Comprehensive Evaluation of Sdox, a Promising HS-Releasing Doxorubicin for the Treatment of Chemoresistant Tumors <i>Frontiers in Pharmacology</i> , <b>2022</b> , 13, 831791	5.6	0
54	NO release regulated by doxorubicin as the green light-harvesting antenna. <i>Chemical Communications</i> , <b>2020</b> , 56, 6332-6335	5.8	2
53	In vitro vascular toxicity assessment of NitDOX, a novel NO-releasing doxorubicin. <i>European Journal of Pharmacology</i> , <b>2020</b> , 880, 173164	5.3	3
52	MRP5 nitration by NO-releasing gemcitabine encapsulated in liposomes confers sensitivity in chemoresistant pancreatic adenocarcinoma cells. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , <b>2020</b> , 1867, 118824	4.9	5
51	Paracetamol-Galactose Conjugate: A Novel Prodrug for an Old Analgesic Drug. <i>Molecular Pharmaceutics</i> , <b>2019</b> , 16, 4181-4189	5.6	5
50	Hyaluronated liposomes containing H2S-releasing doxorubicin are effective against P-glycoprotein-positive/doxorubicin-resistant osteosarcoma cells and xenografts. <i>Cancer Letters</i> , <b>2019</b> , 456, 29-39	9.9	26
49	In Vitro Assessment of NitDox Toxicity Toward Vasculature <b>2019</b> , 319-320		
48	Endoplasmic reticulum-targeting doxorubicin: a new tool effective against doxorubicin-resistant osteosarcoma. <i>Cellular and Molecular Life Sciences</i> , <b>2019</b> , 76, 609-625	10.3	32
47	New tetrahydroisoquinoline-based P-glycoprotein modulators: decoration of the biphenyl core gives selective ligands. <i>MedChemComm</i> , <b>2018</b> , 9, 862-869	5	9
46	New NO- and H2S-releasing doxorubicins as targeted therapy against chemoresistance in castration-resistant prostate cancer: in vitro and in vivo evaluations. <i>Investigational New Drugs</i> , <b>2018</b> , 36, 985-998	4.3	19
45	Mitochondrial Delivery of Phenol Substructure Triggers Mitochondrial Depolarization and Apoptosis of Cancer Cells. <i>Frontiers in Pharmacology</i> , <b>2018</b> , 9, 580	5.6	16
44	New Tetrahydroisoquinoline Derivatives Overcome Pgp Activity in Brain-Blood Barrier and Glioblastoma Multiforme in Vitro. <i>Molecules</i> , <b>2018</b> , 23,	4.8	9
43	Aceclofenac-Galactose Conjugate: Design, Synthesis, Characterization, and Pharmacological and Toxicological Evaluations. <i>Molecular Pharmaceutics</i> , <b>2018</b> , 15, 3101-3110	5.6	7
42	Structural and biological characterization of new hybrid drugs joining an HDAC inhibitor to different NO-donors. <i>European Journal of Medicinal Chemistry</i> , <b>2018</b> , 144, 612-625	6.8	13
41	Folate-targeted liposomal nitrooxy-doxorubicin: An effective tool against P-glycoprotein-positive and folate receptor-positive tumors. <i>Journal of Controlled Release</i> , <b>2018</b> , 270, 37-52	11.7	47
40	Galactosylated Pro-Drug of Ursodeoxycholic Acid: Design, Synthesis, Characterization, and Pharmacological Effects in a Rat Model of Estrogen-Induced Cholestasis. <i>Molecular Pharmaceutics</i> , <b>2018</b> , 15, 21-30	5.6	8

## (2014-2018)

Discovery of phenylsulfonylfuroxan derivatives as gamma globin inducers by histone acetylation. <i>European Journal of Medicinal Chemistry</i> , <b>2018</b> , 154, 341-353	6.8	7
Light-Regulated NO Release as a Novel Strategy To Overcome Doxorubicin Multidrug Resistance. <i>ACS Medicinal Chemistry Letters</i> , <b>2017</b> , 8, 361-365	4.3	35
Solid Lipid Nanoparticles Loaded with Antitumor Lipophilic Prodrugs Aimed to Glioblastoma Treatment: Preliminary Studies on Cultured Cells. <i>Journal of Nanoscience and Nanotechnology</i> , <b>2017</b> , 17, 3606-3614	1.3	5
New furoxan derivatives for the treatment of ocular hypertension. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2017</b> , 27, 479-483	2.9	5
Design, Synthesis, and Characterization of N-Oxide-Containing Heterocycles with in Vivo Sterilizing Antitubercular Activity. <i>Journal of Medicinal Chemistry</i> , <b>2017</b> , 60, 8647-8660	8.3	29
-Dinitroalkyl Benzenes: A Novel Class of IOP-Lowering Agents for the Treatment of Ocular Hypertension. <i>ACS Medicinal Chemistry Letters</i> , <b>2017</b> , 8, 1054-1059	4.3	3
Direct introduction of cyano group on furoxan ring. Mendeleev Communications, 2017, 27, 565-566	1.9	4
Synthesis and Biological Evaluation of N(2) -Substituted 2,4-Diamino-6-cyclohexylmethoxy-5-nitrosopyrimidines and Related 5-Cyano-NNO-azoxy Derivatives as Cyclin-Dependent Kinase 2 (CDK2) Inhibitors. <i>ChemMedChem</i> , <b>2016</b> , 11, 1705-8	3.7	5
Structure-Activity Relationship Studies on Tetrahydroisoquinoline Derivatives: [4V(6,7-Dimethoxy-3,4-dihydro-1H-isoquinolin-2-ylmethyl)biphenyl-4-ol] (MC70) Conjugated through Flexible Alkyl Chains with Furazan Moieties Gives Rise to Potent and Selective Ligands of	8.3	17
Overcoming multidrug resistance by targeting mitochondria with NO-donating doxorubicins.  Bioorganic and Medicinal Chemistry, <b>2016</b> , 24, 967-75	3.4	21
Solid lipid nanoparticles carrying lipophilic derivatives of doxorubicin: preparation, characterization, and in vitro cytotoxicity studies. <i>Journal of Microencapsulation</i> , <b>2016</b> , 33, 381-90	3.4	14
H2S-Donating Doxorubicins May Overcome Cardiotoxicity and Multidrug Resistance. <i>Journal of Medicinal Chemistry</i> , <b>2016</b> , 59, 4881-9	8.3	35
Synthesis and biological activity of furoxan derivatives against Mycobacterium tuberculosis. <i>European Journal of Medicinal Chemistry</i> , <b>2016</b> , 123, 523-531	6.8	48
A nitric oxide-donor furoxan moiety improves the efficacy of edaravone against early renal dysfunction and injury evoked by ischemia/reperfusion. <i>Oxidative Medicine and Cellular Longevity</i> , <b>2015</b> , 2015, 804659	6.7	18
NO-donor thiacarbocyanines as multifunctional agents for Alzheimer disease. <i>Bioorganic and Medicinal Chemistry</i> , <b>2015</b> , 23, 4688-4698	3.4	16
Furazan and furoxan sulfonamides are strong Etarbonic anhydrase inhibitors and potential antiglaucoma agents. <i>Bioorganic and Medicinal Chemistry</i> , <b>2014</b> , 22, 3913-21	3.4	25
Leishmanicidal activities of novel synthetic furoxan and benzofuroxan derivatives. <i>Antimicrobial Agents and Chemotherapy</i> , <b>2014</b> , 58, 4837-47	5.9	32
Liposomal nitrooxy-doxorubicin: one step over caelyx in drug-resistant human cancer cells.  Molecular Pharmaceutics, 2014, 11, 3068-79	5.6	27
	European Journal of Medicinal Chemistry, 2018, 154, 341-353  Light-Regulated NO Release as a Novel Strategy To Overcome Doxorubicin Multidrug Resistance.  ACS Medicinal Chemistry Letters, 2017, 8, 361-365  Solid Lipid Nanoparticles Loaded with Antitumor Lipophilic Prodrugs Aimed to Glioblastoma  Treatment: Preliminary Studies on Cultured Cells. Journal of Nanoscience and Nanotechnology, 2017, 17, 3606-3614  New Furoxan derivatives for the treatment of ocular hypertension. Bioorganic and Medicinal  Chemistry Letters, 2017, 27, 479-483  Design, Synthesis, and Characterization of N-Oxide-Containing Heterocycles with in Vivo Sterilizing  Antitubercular Activity. Journal of Medicinal Chemistry, 2017, 60, 8647-8660  -Dinitroalkyl Benzenes: A Novel Class of IOP-Lowering Agents for the Treatment of Ocular  Hypertension. ACS Medicinal Chemistry Letters, 2017, 8, 1054-1059  Direct introduction of cyano group on furoxan ring. Mendeleev Communications, 2017, 27, 565-566  Synthesis and Biological Evaluation of N(2) -Substituted  2,4-Diamino-6-cyclohexylmethoxy-5-nitrosopyrimidines and Related 5-Cyano-NNO-azoxy  Derivatives as Cyclin-Dependent Kinase (CORX) Inhibitors. ChemMed.hem., 2016, 11, 1705-8  Structure-Activity, Relationship Studies on Tetrahydroisoquinoline Derivatives:  [4V(6,7-Dimethoxy-3,4-dihydro-1H-isoquinolin-2-ylmethyl)biphenyl-4-ol] (MC70) Conjugated  through Flexible Alkyl Chains with Furazan Moieties Gives Rise to Potent and Selective Ligands of  P-clycoprotein. Journal of Medicinal Chemistry, 2016, 59, 6729-38  Overcoming multidrug resistance by targeting mitochondria with NO-donating doxorubicins.  Bioorganic and Medicinal Chemistry, 2016, 24, 967-75  Solid lipid nanoparticles carrying lipophilic derivatives of doxorubicin: preparation, characterization,  and in vitro cytotoxicity studies. Journal of Microencapsulation, 2016, 33, 381-90  H2S-Donating Doxorubicins May Overcome Cardiotoxicity and Multidrug Resistance. Journal of Medicinal Chemistry, 2016, 123, 523-531  A Nitric oxide-donor furoxan moiety i	Light-Regulated NO Release as a Novel Strategy To Overcome Doxorubicin Multidrug Resistance.  ACS Medicinal Chemistry Letters, 2017, 8, 361-365  Solid Lipid Nanoparticles Loaded with Antitumor Lipophilic Prodrugs Aimed to Clioblastoma Treatment: Preliminary Studies on Cultured Cells. Journal of Nanoscience and Nanotechnology, 2017, 17, 3606-3614  New furoxan derivatives for the treatment of ocular hypertension. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 479-483  Design, Synthesis, and Characterization of N-Oxide-Containing Heterocycles with in Vivo Sterilizing Antitubercular Activity. Journal of Medicinal Chemistry, 2017, 60, 8647-8660  Dinitroalkyl Benzenes: A Novel Class of IOP-Lowering Agents for the Treatment of Ocular Hypertension. ACS Medicinal Chemistry Letters, 2017, 8, 1054-1059  Direct introduction of cyano group on furoxan ring. Mendeleev Communications, 2017, 27, 565-566  Synthesis and Biological Evaluation of N(2) - Substituted 2,4-Diamino-6-cyclohexylmethoxy-5-introsopyrimidines and Related 5-Cyano-NNO-azoxy Derivatives as Cyclin-Dependent Kinase 2 (CDK2) Inhibitors. ChemMedChem, 2016, 11, 1705-8  Structure-Activity Relationship Studies on Tetrahydroisoquinoline Derivatives: [4V(6):7-Dimethoxy-3,4-d'hydro-IH-isoquinolin-2yimethyl)biphenyl-4-0] (MC70) Conjugated through Flexible Alkyl Chains with Furazan Moleties Gives Rise to Potent and Selective Ligands of P-dycoprotein. Journal of Medicinal Chemistry, 2016, 89, 892-9-88  Overcoming multidrug resistance by targeting mitochondria with NO-donating doxorubicins. Bioorganic and Medicinal Chemistry, 2016, 24, 967-75  Solid lipid nanoparticles carrying lipophilic derivatives of doxorubicin; preparation, characterization, and in vitro cytotoxicity studies. Journal of Microencapsulation, 2016, 33, 381-90  Synthesis and biological activity of furoxan derivatives against Mycobacterium tuberculosis. European Journal of Medicinal Chemistry, 2016, 29, 967-75  Solid lipid nanoparticles of movel synthetic furoxan and benzofuroxan derivatives. Antimic

21	Doxorubicin-antioxidant co-drugs. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2013</b> , 23, 5307-10	2.9	19
20	Mitochondrial-targeting nitrooxy-doxorubicin: a new approach to overcome drug resistance. <i>Molecular Pharmaceutics</i> , <b>2013</b> , 10, 161-74	5.6	52
19	Synthesis physicochemical profile and PAMPA study of new NO-donor edaravone co-drugs. <i>Bioorganic and Medicinal Chemistry</i> , <b>2012</b> , 20, 841-50	3.4	5
18	New nitric oxide or hydrogen sulfide releasing aspirins. <i>Journal of Medicinal Chemistry</i> , <b>2011</b> , 54, 5478-8	48.3	28
17	Nitric oxide donor doxorubicins accumulate into Doxorubicin-resistant human colon cancer cells inducing cytotoxicity. <i>ACS Medicinal Chemistry Letters</i> , <b>2011</b> , 2, 494-7	4.3	58
16	Phenylsulfonylfuroxans as modulators of multidrug-resistance-associated protein-1 and P-glycoprotein. <i>Journal of Medicinal Chemistry</i> , <b>2010</b> , 53, 5467-75	8.3	49
15	Effects of nitric oxide donor antioxidants containing the phenol vitamin E substructure and a furoxan moiety on ischemia/reperfusion injury. <i>Arzneimittelforschung</i> , <b>2009</b> , 59, 111-6		1
14	Unsymmetrically substituted furoxans. Part 19. Methyl and phenylfuroxansulfonic acids and related sulfonamides. <i>Journal of Heterocyclic Chemistry</i> , <b>2009</b> , 46, 866-872	1.9	10
13	Edaravone derivatives containing NO-donor functions. <i>Journal of Medicinal Chemistry</i> , <b>2009</b> , 52, 574-8	8.3	31
12	(Nitrooxyacyloxy)methyl esters of aspirin as novel nitric oxide releasing aspirins. <i>Journal of Medicinal Chemistry</i> , <b>2009</b> , 52, 5058-68	8.3	30
11	Synthesis of some novel organic nitrates and comparative in vitro study of their vasodilator profile. Journal of Medicinal Chemistry, <b>2009</b> , 52, 4020-5	8.3	4
10	Multitarget drugs: Focus on the NO-donor hybrid drugs. <i>Pure and Applied Chemistry</i> , <b>2008</b> , 80, 1693-170	12.1	17
9	Novel antioxidant agents deriving from molecular combination of Vitamin C and NO-donor moieties. <i>Bioorganic and Medicinal Chemistry</i> , <b>2008</b> , 16, 5199-206	3.4	15
8	NO-donor COX-2 inhibitors. New nitrooxy-substituted 1,5-diarylimidazoles endowed with COX-2 inhibitory and vasodilator properties. <i>Journal of Medicinal Chemistry</i> , <b>2007</b> , 50, 1449-57	8.3	52
7	Amphiphilic NO-donor antioxidants. <i>ChemMedChem</i> , <b>2007</b> , 2, 234-40	3.7	2
6	NO-donor melatonin derivatives: synthesis and in vitro pharmacological characterization. <i>Journal of Pineal Research</i> , <b>2007</b> , 42, 371-85	10.4	12
5	Synthesis, chiral HPLC resolution and configuration assignment of 1-phenylglyceryl trinitrate stereomers. <i>Chirality</i> , <b>2006</b> , 18, 430-6	2.1	6
4	NO-donor phenols: a new class of products endowed with antioxidant and vasodilator properties. Journal of Medicinal Chemistry, <b>2006</b> , 49, 2886-97	8.3	43

## LIST OF PUBLICATIONS

3	Development of a new class of potential antiatherosclerosis agents: NO-donor antioxidants. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2004</b> , 14, 5971-4	2.9	23
2	Synthesis of 2-monofunctionalized 2,4,6,8-tetraazabicyclo[3.3.0]octane-3,7-diones. <i>Russian Chemical Bulletin</i> , <b>2003</b> , 52, 192-197	1.7	14
1	New functional glycoluril derivatives. <i>Mendeleev Communications</i> , <b>2001</b> , 11, 32-33	1.9	16