

Marek Bednarski

List of Publications by Citations

Source: <https://exaly.com/author-pdf/4616943/marek-bednarski-publications-by-citations.pdf>

Version: 2024-04-24

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

57
papers

546
citations

14
h-index

21
g-index

61
ext. papers

678
ext. citations

4.4
avg, IF

3.18
L-index

#	Paper	IF	Citations
57	The influence of the route of administration of gold nanoparticles on their tissue distribution and basic biochemical parameters: In vivo studies. <i>Pharmacological Reports</i> , 2015 , 67, 405-9	3.9	59
56	Synthesis and evaluation of in vivo activity of diphenylhydantoin basic derivatives. <i>European Journal of Medicinal Chemistry</i> , 2004 , 39, 1013-27	6.8	39
55	Novel multi-target azinesulfonamides of cyclic amine derivatives as potential antipsychotics with pro-social and pro-cognitive effects. <i>European Journal of Medicinal Chemistry</i> , 2018 , 145, 790-804	6.8	28
54	Alpha lipoic acid protects the heart against myocardial post ischemia-reperfusion arrhythmias via KATP channel activation in isolated rat hearts. <i>Pharmacological Reports</i> , 2014 , 66, 499-504	3.9	26
53	Antiarrhythmic properties of phenylpiperazine derivatives of phenytoin with β -adrenoceptor affinities. <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 2290-303	3.4	25
52	The role of lipoic acid in prevention of nitroglycerin tolerance. <i>European Journal of Pharmacology</i> , 2008 , 591, 203-10	5.3	22
51	Synthesis and evaluation of some xanthone derivatives for anti-arrhythmic, hypotensive properties and their affinity for adrenergic receptors. <i>Archiv Der Pharmazie</i> , 2008 , 341, 90-8	4.3	20
50	A Comparison of the Anorectic Effect and Safety of the Alpha2-Adrenoceptor Ligands Guanfacine and Yohimbine in Rats with Diet-Induced Obesity. <i>PLoS ONE</i> , 2015 , 10, e0141327	3.7	19
49	Synthesis and adrenolytic activity of 1-(1H-indol-4-yloxy)-3-[[2-(2-methoxyphenoxy)ethyl]amino]propan-2-ol and its enantiomers. Part 1. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 809-17	6.8	18
48	H3 histamine receptor antagonist pitolisant reverses some subchronic disturbances induced by olanzapine in mice. <i>Metabolic Brain Disease</i> , 2016 , 31, 1023-9	3.9	18
47	Are anti-inflammatory properties of lipoic acid associated with the formation of hydrogen sulfide?. <i>Pharmacological Reports</i> , 2013 , 65, 1018-24	3.9	16
46	Tissue distribution of gold nanoparticles after single intravenous administration in mice. <i>Pharmacological Reports</i> , 2013 , 65, 1033-8	3.9	15
45	Design, synthesis, anticonvulsant, and antiarrhythmic properties of novel N-Mannich base and amide derivatives of Etetralinohydantoin. <i>Pharmacological Reports</i> , 2016 , 68, 886-93	3.9	14
44	Antidepressant-like activity of aroxyalkyl derivatives of 2-methoxyphenylpiperazine and evidence for the involvement of serotonin receptor subtypes in their mechanism of action. <i>Pharmacology Biochemistry and Behavior</i> , 2016 , 141, 28-41	3.9	14
43	Investigations on the synthesis and pharmacological properties of N-substituted derivatives of 4-alkoxy-6-methyl-1H-pyrrolo[3,4-c]pyridine-1,3(2H)-diones. <i>Il Farmaco</i> , 2005 , 60, 53-9		14
42	In vivo anti-inflammatory activity of lipoic acid derivatives in mice. <i>Postepy Higieny I Medycyny Doswiadczalnej</i> , 2013 , 67, 331-8	0.3	13
41	Pyrrolidin-2-one derivatives may reduce body weight in rats with diet-induced obesity. <i>European Journal of Pharmacology</i> , 2016 , 776, 146-55	5.3	12

40	Studies on novel pyridine and 2-pyridone derivatives of N-arylpiperazine as β -adrenoceptor ligands. <i>Medicinal Chemistry</i> , 2014 , 10, 144-53	1.8	12
39	Synthesis and adrenergic activity of 1-(1H-indol-4-yloxy)-3-(2-(2-methoxyphenoxy)ethylamino)propan-2-ol analogs and its enantiomers. Part 2. <i>European Journal of Medicinal Chemistry</i> , 2009 , 44, 5103-11	6.8	11
38	PSB 603 - a known selective adenosine A2B receptor antagonist - has anti-inflammatory activity in mice. <i>Biomedicine and Pharmacotherapy</i> , 2021 , 135, 111164	7.5	10
37	Structural modifications and in vitro pharmacological evaluation of 4-pyridyl-piperazine derivatives as an active and selective histamine H receptor ligands. <i>Bioorganic Chemistry</i> , 2019 , 91, 103071	5.1	9
36	Anti-Alzheimer's multitarget-directed ligands with serotonin 5-HT antagonist, butyrylcholinesterase inhibitory, and antioxidant activity. <i>Archiv Der Pharmazie</i> , 2019 , 352, e1900041	4.3	9
35	KSK19 - Novel histamine H3 receptor ligand reduces body weight in diet induced obese mice. <i>Biochemical Pharmacology</i> , 2019 , 168, 193-203	6	9
34	Novel mannich bases, 5-arylimidazolidine-2,4-dione derivatives with dual 5-HT(1A) receptor and serotonin transporter affinity. <i>Archiv Der Pharmazie</i> , 2013 , 346, 98-109	4.3	9
33	Evaluation of anticonvulsant activity of novel pyrrolidin-2-one derivatives. <i>Pharmacological Reports</i> , 2014 , 66, 708-11	3.9	8
32	Ergotamine and nicergoline - facts and myths. <i>Pharmacological Reports</i> , 2015 , 67, 360-3	3.9	8
31	Bioactivation of nitroglycerin to nitric oxide (NO) and S-nitrosothiols in the rat liver and evaluation of the coexisting hypotensive effect. <i>Fundamental and Clinical Pharmacology</i> , 2004 , 18, 449-56	3.1	8
30	Synthesis and biological evaluation of N-arylpiperazine derivatives of 4,4-dimethylisoquinoline-1,3(2H,4H)-dione as potential antiplatelet agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 536-545	5.6	7
29	Antiarrhythmic activity of some xanthone derivatives with β -adrenoceptor affinities in rats. <i>European Journal of Pharmacology</i> , 2014 , 738, 14-21	5.3	7
28	β -Adrenoceptor antagonistic and hypotensive properties of novel arylpiperazine derivatives of pyrrolidin-2-one. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 2104-11	3.4	6
27	Metabolic benefits of 1-(3-(4-(o-tolyl)piperazin-1-yl)propyl)pyrrolidin-2-one: a non-selective β -adrenoceptor antagonist. <i>Journal of Endocrinological Investigation</i> , 2018 , 41, 609-619	5.2	6
26	Synthesis and Pharmacological Activity of a New Series of 1-(1H-Indol-4-yloxy)-3-(2-(2-methoxyphenoxy)ethylamino)propan-2-ol Analogs. <i>Archiv Der Pharmazie</i> , 2016 , 349, 211-23	4.3	5
25	Characteristics of metabolic stability and the cell permeability of 2-pyrimidinyl-piperazinyl-alkyl derivatives of 1H-imidazo[2,1-f]purine-2,4(3H,8H)-dione with antidepressant- and anxiolytic-like activities. <i>Chemical Biology and Drug Design</i> , 2019 , 93, 511-521	2.9	5
24	KD-64-A new selective A2A adenosine receptor antagonist has anti-inflammatory activity but contrary to the non-selective antagonist-Caffeine does not reduce diet-induced obesity in mice. <i>PLoS ONE</i> , 2020 , 15, e0229806	3.7	4
23	The effect of nitroglycerin tolerance on oxidative stress and anaerobic sulfur metabolism in rat tissues. <i>Fundamental and Clinical Pharmacology</i> , 2010 , 24, 47-53	3.1	4

22	Isolation of 1-(3,4,5-trihydroxyphenyl)-3-(2,4,6-trihydroxyphenyl)-propan-2-ol from Grape Seed Extract and Evaluation of its Antioxidant and Antispasmodic Potential. <i>Molecules</i> , 2019 , 24,	4.8	3
21	Antiarrhythmic and β -Adrenoceptor Antagonistic Properties of Novel Arylpiperazine Derivatives of Pyrrolidin-2-one. <i>Archiv Der Pharmazie</i> , 2015 , 348, 861-7	4.3	3
20	Application of liquid chromatography-tandem mass spectrometry method for the analysis of new nonselective beta-adrenergic blocker 1-(1-H-indol-4-yloxy)-3-[[2-(2-methoxyphenoxy)ethyl]amino]propan-2-ol (2F109) in rat plasma. <i>Chirality</i> , 2007 , 19, 536-41	2.1	3
19	Arylsulfonamide derivatives of (aryloxy)ethyl pyrrolidines and piperidines as β -adrenergic receptor antagonist with uro-selective activity. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 5582-5591	3.4	3
18	Structural modifications in the distal, regulatory region of histamine H receptor antagonists leading to the identification of a potent anti-obesity agent. <i>European Journal of Medicinal Chemistry</i> , 2021 , 213, 113041	6.8	3
17	Synthesis and Analgesic Activity of Annelated Xanthine Derivatives in Experimental Models in Rodents. <i>Archiv Der Pharmazie</i> , 2015 , 348, 704-14	4.3	2
16	The nitric oxide/soluble cyclic guanylate/cyclic guanosine monophosphate pathway is involved in the cardiovascular effects of a novel α - and β -adrenoceptor antagonist. <i>Pharmacology</i> , 2014 , 94, 287-95	2.3	2
15	Synthesis and adrenergic activity of new propanolamines. <i>Molecules</i> , 2010 , 15, 3887-904	4.8	2
14	Antidepressant-like activity and safety profile evaluation of 1H-imidazo[2,1-f]purine-2,4(3H,8H)-dione derivatives as 5-HT _{1A} receptor partial agonists. <i>PLoS ONE</i> , 2020 , 15, e0237196	3.7	2
13	MH-76, a Novel Non-Quinazoline β -Adrenoceptor Antagonist, but Not Prazosin Reduces Inflammation and Improves Insulin Signaling in Adipose Tissue of Fructose-Fed Rats. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	2
12	Design, Sustainable Synthesis and Biological Evaluation of a Novel Dual α A/5-HT ₇ Receptor Antagonist with Antidepressant-Like Properties. <i>Molecules</i> , 2021 , 26,	4.8	2
11	Synthesis and Pharmacological Evaluation of Novel Silodosin-Based Arylsulfonamide Derivatives as β -Adrenergic Receptor Antagonist with Potential Uroselective Profile. <i>Molecules</i> , 2018 , 23,	4.8	2
10	The GPR18 Agonist PSB-KD-107 Exerts Endothelium-Dependent Vasorelaxant Effects. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	2
9	KD-64 is a new selective A _{2A} adenosine receptor antagonist has anti-inflammatory activity but contrary to the non-selective antagonist caffeine does not reduce diet-induced obesity in mice		1
8	The Structural Determinants for β -Adrenergic/Serotonin Receptors Activity among Phenylpiperazine-Hydantoin Derivatives. <i>Molecules</i> , 2021 , 26,	4.8	1
7	The antidepressant-like activity of chiral xanthone derivatives may be mediated by 5-HT _{1A} receptor and β arrestin signalling. <i>Journal of Psychopharmacology</i> , 2020 , 34, 1431-1442	4.6	1
6	Metabolic benefits of novel histamine H receptor ligands in the model of excessive eating: The importance of intrinsic activity and pharmacokinetic properties. <i>Biomedicine and Pharmacotherapy</i> , 2021 , 142, 111952	7.5	1
5	Antiarrhythmic activity in occlusion-reperfusion model of 1-(1H-indol-4-yloxy)-3-[[2-(2-methoxyphenoxy)ethyl]amino]propan-2-ol and its enantiomers. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2016 , 43, 81-7	3	

4 KD-64A new selective A2A adenosine receptor antagonist has anti-inflammatory activity but contrary to the non-selective antagonist caffeine does not reduce diet-induced obesity in mice **2020**, 15, e0229806

3 KD-64A new selective A2A adenosine receptor antagonist has anti-inflammatory activity but contrary to the non-selective antagonist caffeine does not reduce diet-induced obesity in mice **2020**, 15, e0229806

2 KD-64A new selective A2A adenosine receptor antagonist has anti-inflammatory activity but contrary to the non-selective antagonist caffeine does not reduce diet-induced obesity in mice **2020**, 15, e0229806

1 KD-64A new selective A2A adenosine receptor antagonist has anti-inflammatory activity but contrary to the non-selective antagonist caffeine does not reduce diet-induced obesity in mice **2020**, 15, e0229806