

Dorota AÅ^{1/4}ewska

List of Publications by Year in descending order

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65
papers

1,205
citations

331670

21
h-index

454955

30
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66
all docs

66
docs citations

66
times ranked

1125
citing authors

#	ARTICLE	IF	CITATIONS
1	The histamine H3R antagonist DL77 attenuates autistic behaviors in a prenatal valproic acid-induced mouse model of autism. <i>Scientific Reports</i> , 2018, 8, 13077.	3.3	58
2	Anticonvulsant and procognitive properties of the non-imidazole histamine H3 receptor antagonist DL77 in male adult rats. <i>Neuropharmacology</i> , 2016, 106, 46-55.	4.1	57
3	Recent advances in histamine H ₃ receptor antagonists/inverse agonists. <i>Expert Opinion on Therapeutic Patents</i> , 2010, 20, 1147-1169.	5.0	51
4	Aryl-1,3,5-triazine derivatives as histamine H4 receptor ligands. <i>European Journal of Medicinal Chemistry</i> , 2014, 83, 534-546.	5.5	46
5	The dual-active histamine H3 receptor antagonist and acetylcholine esterase inhibitor E100 ameliorates stereotyped repetitive behavior and neuroinflammation in sodium valproate induced autism in mice. <i>Chemico-Biological Interactions</i> , 2019, 312, 108775.	4.0	44
6	New developments around histamine H ₃ receptor antagonists/inverse agonists: a patent review (2010 – present). <i>Expert Opinion on Therapeutic Patents</i> , 2014, 24, 89-111.	5.0	40
7	Antinociceptive effects of novel histamine H ₃ and H ₄ receptor antagonists and their influence on morphine analgesia of neuropathic pain in the mouse. <i>British Journal of Pharmacology</i> , 2018, 175, 2897-2910.	5.4	36
8	Ether derivatives of 3-piperidinopropan-1-ol as non-imidazole histamine H3 receptor antagonists. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 3522-3529.	3.0	35
9	The computer-aided discovery of novel family of the 5-HT6 serotonin receptor ligands among derivatives of 4-benzyl-1,3,5-triazine. <i>European Journal of Medicinal Chemistry</i> , 2017, 135, 117-124.	5.5	33
10	Anticonvulsive effect of nonimidazole histamine H3 receptor antagonists. <i>Behavioural Pharmacology</i> , 2014, 25, 245-252.	1.7	31
11	Chlorophenoxy aminoalkyl derivatives as histamine H3R ligands and antiseizure agents. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 53-72.	3.0	28
12	The Histamine H3 Receptor Antagonist E159 Reverses Memory Deficits Induced by Dizocilpine in Passive Avoidance and Novel Object Recognition Paradigm in Rats. <i>Frontiers in Pharmacology</i> , 2017, 8, 709.	3.5	27
13	Progress in the development of histamine H ₃ receptor antagonists/inverse agonists: a patent review (2013-2017). <i>Expert Opinion on Therapeutic Patents</i> , 2018, 28, 175-196.	5.0	27
14	Search for new multi-target compounds against Alzheimer's disease among histamine H3 receptor ligands. <i>European Journal of Medicinal Chemistry</i> , 2020, 185, 111785.	5.5	27
15	Aryl-1,3,5-triazine ligands of histamine H4 receptor attenuate inflammatory and nociceptive response to carrageen, zymosan and lipopolysaccharide. <i>Inflammation Research</i> , 2017, 66, 79-95.	4.0	26
16	Dual-Acting Diether Derivatives of Piperidine and Homopiperidine with Histamine H ₃ Receptor Antagonistic and Anticholinesterase Activity. <i>Archiv Der Pharmazie</i> , 2012, 345, 591-597.	4.1	25
17	The Histamine H3 Receptor Antagonist DL77 Ameliorates MK801-Induced Memory Deficits in Rats. <i>Frontiers in Neuroscience</i> , 2018, 12, 42.	2.8	25
18	The Dual-Active Histamine H3 Receptor Antagonist and Acetylcholine Esterase Inhibitor E100 Alleviates Autistic-Like Behaviors and Oxidative Stress in Valproic Acid Induced Autism in Mice. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3996.	4.1	25

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19	(2-Arylethenyl)-1,3,5-triazin-2-amines as a novel histamine H4 receptor ligands. <i>European Journal of Medicinal Chemistry</i> , 2015, 103, 238-251.	5.5	24
20	Novel naphthyloxy derivatives – Potent histamine H3 receptor ligands. Synthesis and pharmacological evaluation. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 2573-2585.	3.0	24
21	Structure-activity relationships of imidazothiazinones and analogs as antagonists of the cannabinoid-activated orphan G protein-coupled receptor GPR18. <i>European Journal of Medicinal Chemistry</i> , 2018, 155, 381-397.	5.5	22
22	Simultaneous Blockade of Histamine H3 Receptors and Inhibition of Acetylcholine Esterase Alleviate Autistic-Like Behaviors in BTBR T+ tf/J Mouse Model of Autism. <i>Biomolecules</i> , 2020, 10, 1251.	4.0	22
23	Antagonism of Histamine H3 receptors Alleviates Pentylentetrazole-Induced Kindling and Associated Memory Deficits by Mitigating Oxidative Stress, Central Neurotransmitters, and c-Fos Protein Expression in Rats. <i>Molecules</i> , 2020, 25, 1575.	3.8	21
24	Cholinesterase inhibitory activity of chlorophenoxy derivatives – Histamine H3 receptor ligands. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 4140-4145.	2.2	20
25	Imidazo[2,1-b]thiazoles, imidazo[2,1-b]imidazoles and Pyrrolo[1,2-c]imidazoles. synthesis, structure and evaluation of benzodiazepine receptor binding. <i>Journal of Heterocyclic Chemistry</i> , 2002, 39, 243-253.	2.6	19
26	Azines as histamine H4 receptor antagonists. <i>Frontiers in Bioscience - Scholar</i> , 2012, S4, 967-987.	2.1	19
27	The novel non-imidazole histamine H3 receptor antagonist DL77 reduces voluntary alcohol intake and ethanol-induced conditioned place preference in mice. <i>Physiology and Behavior</i> , 2015, 151, 189-197.	2.1	18
28	Computer-Aided Studies for Novel Arylhydantoin 1,3,5-Triazine Derivatives as 5-HT6 Serotonin Receptor Ligands with Antidepressive-Like, Anxiolytic and Antiobesity Action In Vivo. <i>Molecules</i> , 2018, 23, 2529.	3.8	18
29	Studies on Anticonvulsant Effects of Novel Histamine H3R Antagonists in Electrically and Chemically Induced Seizures in Rats. <i>International Journal of Molecular Sciences</i> , 2018, 19, 3386.	4.1	18
30	Synthesis and computer-aided SAR studies for derivatives of phenoxyalkyl-1,3,5-triazine as the new potent ligands for serotonin receptors 5-HT6. <i>European Journal of Medicinal Chemistry</i> , 2019, 178, 740-751.	5.5	18
31	Are the Hydantoin-1,3,5-triazine 5-HT6R Ligands a Hope to a Find New Procognitive and Anti-Obesity Drug? Considerations Based on Primary In Vivo Assays and ADME-Tox Profile In Vitro. <i>Molecules</i> , 2019, 24, 4472.	3.8	18
32	Histamine H3 and H4 receptor affinity of branched 3-(1H-imidazol-4-yl)propyl N-alkylcarbamates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009, 19, 6682-6685.	2.2	17
33	Rational design of new multitarget histamine H3 receptor ligands as potential candidates for treatment of Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112743.	5.5	17
34	Structural and Molecular Insight into Piperazine and Piperidine Derivatives as Histamine H ₃ and Sigma-1 Receptor Antagonists with Promising Antinociceptive Properties. <i>ACS Chemical Neuroscience</i> , 2022, 13, 1-15.	3.5	17
35	Piperidine variations in search for non-imidazole histamine H3 receptor ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2008, 16, 8729-8736.	3.0	16
36	Biphenyloxy-alkyl-piperidine and azepane derivatives as histamine H3 receptor ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 5341-5354.	3.0	16

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37	Role of Histamine H3 Receptor Antagonists on Intraocular Pressure Reduction in Rabbit Models of Transient Ocular Hypertension and Glaucoma. <i>International Journal of Molecular Sciences</i> , 2019, 20, 981.	4.1	16
38	<p>Histamine H3 receptor antagonist E177 attenuates amnesia induced by dizocilpine without modulation of anxiety-like behaviors in rats</p>. <i>Neuropsychiatric Disease and Treatment</i> , 2019, Volume 15, 531-542.	2.2	14
39	Histamine H3R Antagonists: From Scaffold Hopping to Clinical Candidates. <i>Receptors</i> , 2016, , 109-155.	0.2	14
40	Cunninghamella as a Microbiological Model for Metabolism of Histamine H3 Receptor Antagonist 1-[3-(4-tert-Butylphenoxy)propyl]piperidine. <i>Applied Biochemistry and Biotechnology</i> , 2012, 168, 1584-1593.	2.9	13
41	Synthesis and biological activity of novel tert -amylphenoxyalkyl (homo)piperidine derivatives as histamine H 3 R ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 2701-2712.	3.0	13
42	4-tert-Pentylphenoxyalkyl derivatives â€“ Histamine H3 receptor ligands and monoamine oxidase B inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 3596-3600.	2.2	13
43	Synthesis and computer-aided analysis of the role of linker for novel ligands of the 5-HT6 serotonin receptor among substituted 1,3,5-triazinylpiperazines. <i>Bioorganic Chemistry</i> , 2019, 84, 319-325.	4.1	13
44	The Neuroprotective Effects of Histamine H3 Receptor Antagonist E177 on Pilocarpine-Induced Status Epilepticus in Rats. <i>Molecules</i> , 2019, 24, 4106.	3.8	12
45	Diether derivatives of homo- or substituted piperidines as non-imidazole histamine H3 receptor ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2009, 17, 3037-3042.	3.0	11
46	The lipophilicity estimation of 5-arylidene derivatives of (2-thio)hydantoin with antimycobacterial activity. <i>Biomedical Chromatography</i> , 2007, 21, 291-298.	1.7	10
47	The Synthesis of 1,3,5â€“triazine Derivatives and JNJ7777120 Analogues with Histamine H₄ Receptor Affinity and Their Interaction with <i>PTEN</i> Promoter. <i>Chemical Biology and Drug Design</i> , 2016, 88, 254-263.	3.2	10
48	Alkyl derivatives of 1,3,5-triazine as histamine H4 receptor ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 1254-1262.	3.0	10
49	Dual Target Ligands with 4-tert-Butylphenoxy Scaffold as Histamine H3 Receptor Antagonists and Monoamine Oxidase B Inhibitors. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3411.	4.1	10
50	Chalcones as Potential Ligands for the Treatment of Parkinsonâ€™s Disease. <i>Pharmaceuticals</i> , 2022, 15, 847.	3.8	9
51	Anticonvulsant evaluation of novel non-imidazole histamine H3R antagonists in different convulsion models in rats. <i>Pharmacology Biochemistry and Behavior</i> , 2018, 170, 14-24.	2.9	8
52	Cyanobiphenyls: Novel H3 receptor ligands with cholinesterase and MAO B inhibitory activity as multitarget compounds for potential treatment of Alzheimerâ€™s disease. <i>Bioorganic Chemistry</i> , 2021, 114, 105129.	4.1	8
53	LCâ€“MSâ€“MS Method for the Analysis of New Non-Imidazole Histamine H3 Receptor Antagonist 1-[3-(4-tert-Butylphenoxy)propyl]piperidine in Rat Serumâ€”Application to Pharmacokinetic Studies. <i>Chromatographia</i> , 2011, 73, 913-919.	1.3	6
54	Eosinophils adhesion assay as a tool for phenotypic drug screening - The pharmacology of 1,3,5 â€“ Triazine and 1H-indole like derivatives against the human histamine H4 receptor. <i>European Journal of Pharmacology</i> , 2021, 890, 173611.	3.5	5

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55	Monocyclic and Fused Azines and Azoles as Histamine H4Receptor Ligands. <i>Current Medicinal Chemistry</i> , 2016, 23, 1870-1925.	2.4	5
56	The Search for Histamine H 4 Receptor Ligands with Anticancer Activity among Novel (Thio)urea Derivatives. <i>ChemistrySelect</i> , 2019, 4, 10943-10952.	1.5	4
57	Binding of 1-[3-(4-tert-butyl-phenoxy)propyl]piperidine, a new non imidazole histamine H3 receptor antagonist to bovine serum albumin. <i>Acta Poloniae Pharmaceutica</i> , 2012, 69, 1043-7.	0.1	4
58	Pharmacokinetics and tissue distribution of the new non-imidazole histamine H3 receptor antagonist 1-[3-(4-tert-butylphenoxy) propyl]piperidine in rats. <i>Xenobiotica</i> , 2015, 45, 912-920.	1.1	3
59	In silico and in vitro studies on interaction of novel non-imidazole histamine H3R antagonists with CYP3A4. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127147.	2.2	3
60	Biphenylalkoxyamine Derivativesâ€“Histamine H3 Receptor Ligands with Butyrylcholinesterase Inhibitory Activity. <i>Molecules</i> , 2021, 26, 3580.	3.8	3
61	Ameliorating effects of histamine H3 receptor antagonist E177 on acute pentylenetetrazole-induced memory impairments in rats. <i>Behavioural Brain Research</i> , 2021, 405, 113193.	2.2	2
62	Search for histamine H4 receptor ligands in the group of 4-methylpiperazino amide derivatives. <i>Inflammation Research</i> , 2010, 59, 243-245.	4.0	1
63	Diether (substituted) piperidine derivatives as novel, histamine H3 receptor ligands. <i>Inflammation Research</i> , 2009, 58, 47-48.	4.0	0
64	Convenient way of synthesis and crystal structure of 1-[(5-chloro-1H-indol-2-yl)carbonyl]-4-methylpiperazine, a histamine H4 receptor antagonist. <i>Heterocyclic Communications</i> , 2011, 17, .	1.2	0
65	Determination of in vitro metabolism of new non-imidazole histamine H3 receptor antagonist 1-[3-(4-tert-butylphenoxy)propyl]piperidine. <i>Acta Poloniae Pharmaceutica</i> , 2019, 76, 877-884.	0.1	0