Qiong Xie

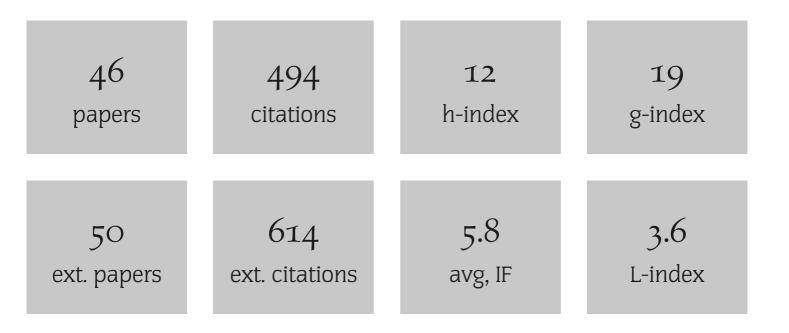
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#	Paper	IF	Citations
46	Bis-(-)-nor-meptazinols as novel nanomolar cholinesterase inhibitors with high inhibitory potency on amyloid-beta aggregation. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 2027-36	8.3	69
45	Cadmium(II)-Triazole Framework as a Luminescent Probe for Ca(2+) and Cyano Complexes. <i>Chemistry - A European Journal</i> , 2016 , 22, 10459-74	4.8	49
44	Microwave-Assisted Synthesis of Phenanthridines by Radical Insertion/Cyclization of Biphenyl Isocyanides. <i>Journal of Organic Chemistry</i> , 2016 , 81, 8426-35	4.2	34
43	Solvent-Induced Single Crystal-Single Crystal Transformation of an Interpenetrated Three-Dimensional Copper Triazole Catalytic Framework. <i>Inorganic Chemistry</i> , 2016 , 55, 4069-71	5.1	25
42	The crystal structure of a complex of acetylcholinesterase with a bis-(-)-nor-meptazinol derivative reveals disruption of the catalytic triad. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 2543-9	8.3	22
41	Adenosine A Receptor Antagonists for Cancer Immunotherapy. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 12196-12212	8.3	21
40	Bis(9)-(-)-nor-meptazinol as a novel dual-binding AChEI potently ameliorates scopolamine-induced cognitive deficits in mice. <i>Pharmacology Biochemistry and Behavior</i> , 2013 , 104, 138-43	3.9	18
39	Investigation of the binding mode of (-)-meptazinol and bis-meptazinol derivatives on acetylcholinesterase using a molecular docking method. <i>Journal of Molecular Modeling</i> , 2006 , 12, 390-7	2	18
38	Discovery of carbazole carboxamides as novel RORE inverse agonists. <i>European Journal of Medicinal Chemistry</i> , 2018 , 148, 465-476	6.8	17
37	(-)-Meptazinol-melatonin hybrids as novel dual inhibitors of cholinesterases and amyloid-□ aggregation with high antioxidant potency for Alzheimer therapy. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 3110-8	3.4	16
36	Microwave-assisted synthesis of hydroxyl-containing isoquinolines by metal-free radical cyclization of vinyl isocyanides with alcohols. <i>Organic and Biomolecular Chemistry</i> , 2017 , 15, 10044-10052	3.9	13
35	Novel bis-(-)-nor-meptazinol derivatives act as dual binding site AChE inhibitors with metal-complexing property. <i>Toxicology and Applied Pharmacology</i> , 2012 , 264, 65-72	4.6	13
34	Discovery of N-indanyl benzamides as potent RORE inverse agonists. <i>European Journal of Medicinal Chemistry</i> , 2019 , 167, 37-48	6.8	11
33	Synthetic access to isoxazoline-functionalized isoquinolines via microwave-assisted iminoxyl radical-participated cascade cyclization of vinyl isocyanides. <i>Organic and Biomolecular Chemistry</i> , 2018 , 16, 4996-5005	3.9	11
32	Meserine, a novel carbamate AChE inhibitor, ameliorates scopolamine-induced dementia and alleviates amyloidogenesis of APP/PS1 transgenic mice. <i>CNS Neuroscience and Therapeutics</i> , 2014 , 20, 165-71	6.8	11
31	On-Water Silver(I)-Catalyzed Cycloisomerization of Acetylenic Free Amines/Amides towards 7-Azaindole/Indole/Isoquinolone Derivatives. <i>Synthesis</i> , 2017 , 49, 4845-4852	2.9	11
30	Bis(9)-(-)-Meptazinol, a novel dual-binding AChE inhibitor, rescues cognitive deficits and pathological changes in APP/PS1 transgenic mice. <i>Translational Neurodegeneration</i> , 2018 , 7, 21	10.3	10

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29	Effects of BIS-MEP on Reversing Amyloid Plaque Deposition and Spatial Learning and Memory Impairments in a Mouse Model of EAmyloid Peptide- and Ibotenic Acid-Induced Alzheimer Disease. Frontiers in Aging Neuroscience, 2019, 11, 3	5.3	9
28	Discovery of novel BTK PROTACs for B-Cell lymphomas. <i>European Journal of Medicinal Chemistry</i> , 2021 , 225, 113820	6.8	9
27	The Pharmacological Heterogeneity of Nepenthone Analogs in Conferring Highly Selective and Potent Expioid Agonistic Activities. <i>ACS Chemical Neuroscience</i> , 2017 , 8, 766-776	5.7	8
26	Pharmacophore-based design and discovery of (-)-meptazinol carbamates as dual modulators of cholinesterase and amyloidogenesis. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 65	9 - 671	7
25	Design, synthesis, and bioavailability evaluation of coumarin-based prodrug of meptazinol. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 4953-6	2.9	7
24	Agonist Lock Touched and Untouched Retinoic Acid Receptor-Related Orphan Receptor-E (RORE) Inverse Agonists: Classification Based on the Molecular Mechanisms of Action. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 10519-10536	8.3	7
23	Discovery of a Highly Selective and Potent Dipioid Receptor Agonist from -Cyclopropylmethyl-7phenyl-6,14-endoethanotetrahydronorthebaines with Reduced Central Nervous System (CNS) Side Effects Navigated by the Message-Address Concept. Journal of	8.3	7
22	Medicinal Chemistry, 2019 , 62, 11054-11070 7EMethyl substituent is a structural locus associated with activity cliff for nepenthone analogues. Bioorganic and Medicinal Chemistry, 2018 , 26, 4254-4263	3.4	6
21	Discovery of novel 20S proteasome inhibitors by rational topology-based scaffold hopping of bortezomib. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018 , 28, 2148-2152	2.9	6
20	Discovery of aryl-substituted indole and indoline derivatives as RORE agonists. <i>European Journal of Medicinal Chemistry</i> , 2019 , 182, 111589	6.8	6
19	Synthesis and evaluation of Eppioid receptor agonistic activity and antinociceptive effect of novel morphine analogues, 7 phenyl-6 Amendo-etheno-tetrahydrothebaine with substituted o-, mand p-amino group. <i>Medicinal Chemistry Research</i> , 2011 , 20, 1364-1370	2.2	6
18	Discovery, synthesis, biological evaluation and structure-based optimization of novel piperidine derivatives as acetylcholine-binding protein ligands. <i>Acta Pharmacologica Sinica</i> , 2017 , 38, 146-155	8	5
17	Discovery of carboxyl-containing biaryl ureas as potent RORE inverse agonists. <i>European Journal of Medicinal Chemistry</i> , 2020 , 202, 112536	6.8	4
16	Microwave-Assisted Radical Insertion/Cyclization of Vinyl Isocyanides for the Synthesis of Multi-Substituted Isoquinolines. <i>ChemistrySelect</i> , 2017 , 2, 8033-8038	1.8	4
15	Discovery of tetrahydroquinolines and benzomorpholines as novel potent RORE agonists. <i>European Journal of Medicinal Chemistry</i> , 2021 , 211, 113013	6.8	4
14	Palladium-Catalyzed Cascade SaegusaHeck Reaction: Synthesis of Diarylacroleins from Arylpropanals and Aryl Iodides. <i>European Journal of Organic Chemistry</i> , 2017 , 2017, 5880-5883	3.2	3
13	Conformational re-analysis of (+)-meptazinol: an opioid with mixed analgesic pharmacophores. <i>Acta Pharmacologica Sinica</i> , 2006 , 27, 1247-52	8	3
12	Crystallography-guided discovery of carbazole-based retinoic acid-related orphan receptor gamma-t (RORE) modulators: insights into different protein behaviors with "short" and "long" inverse agonists. <i>Acta Pharmacologica Sinica</i> , 2021 , 42, 1524-1534	8	3

11	Exploration of the SAR Connection between Morphinan- and Arylacetamide-Based ©pioid Receptor (DR) Agonists Using the Strategy of Bridging. <i>ACS Chemical Neuroscience</i> , 2021 , 12, 1018-103	o ^{5.7}	3
10	Palladium-Catalyzed Synthesis of Diaryl Disaturated Ketones. <i>Synthesis</i> , 2019 , 51, 1455-1465	2.9	3
9	Determination of a novel carbamate AChE inhibitor meserine in mouse plasma, brain and rat plasma by LC-MS/MS: application to pharmacokinetic study after intravenous and subcutaneous administration. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2014 , 96, 156-61	3.5	2
8	Theoretical and NMR investigations on the conformations of (I)-meptazinol hydrochloride in solution. <i>Molecular Simulation</i> , 2013 , 39, 1065-1069	2	2
7	Determination of Bis(9)-(-)-Meptazinol, a bis-ligand for Alzheimer's disease, in rat plasma by liquid chromatography-tandem mass spectrometry: application to pharmacokinetics study. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2012 , 881-882, 126-30	3.2	2
6	RORE agonist enhances anti-PD-1 therapy by promoting monocyte-derived dendritic cells through CXCL10 in cancers <i>Journal of Experimental and Clinical Cancer Research</i> , 2022 , 41, 155	12.8	2
5	Discovery of an -Substituted -Cyclopropylmethyl-7phenyl-6,14-endoethanotetrahydronorthebaine as a Selective, Potent, and Orally Active Expioid Receptor Agonist with an Improved Central Nervous System Safety Profile.	8.3	1
4	Journal of Medicinal Chemistry, 2021 , 64, 12414-12433 RORE agonist synergizes with CTLA-4 antibody to inhibit tumor growth through inhibition of Treg cells via TGF-Bignaling in cancer. <i>Pharmacological Research</i> , 2021 , 172, 105793	10.2	1
3	Discovery of tert-amine-based RORE agonists. European Journal of Medicinal Chemistry, 2021, 224, 1137	704 8	1
2	1-Ethyl-4-hydr-oxy-9-aza-tricyclo-[7.4.1.0]tetra-deca-2,4,6-trien-8-one. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2009 , 65, o3008		
1	Discovery, cocrystallization and biological evaluation of novel piperidine derivatives as high affinity Ls-AChBP ligands possessing # nAChR activities. <i>European Journal of Medicinal Chemistry</i> , 2018 , 160, 37-48	6.8	