

Aaron M Bender

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.

24
papers

302
citations

10
h-index

17
g-index

24
ext. papers

359
ext. citations

5.1
avg, IF

3.36
L-index

#	Paper	IF	Citations
24	Synthesis and characterization of chiral 6-azaspiro[2.5]octanes as potent and selective antagonists of the M muscarinic acetylcholine receptor. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021 , 56, 128479 ^{2,9}		
23	Discovery of the First Selective M Muscarinic Acetylcholine Receptor Antagonists with Antiparkinsonian and Antidystonic Efficacy. <i>ACS Pharmacology and Translational Science</i> , 2021 , 4, 1306-1321	5.9	5
22	Discovery of VU6028418: A Highly Selective and Orally Bioavailable M Muscarinic Acetylcholine Receptor Antagonist. <i>ACS Medicinal Chemistry Letters</i> , 2021 , 12, 1342-1349	4.3	2
21	Discovery of a novel class of heteroaryl-pyrrolidinones as positive allosteric modulators of the muscarinic acetylcholine receptor M. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021 , 47, 128193	2.9	0
20	Synthesis of Substituted 6,7-Dihydro-5-pyrrolo[2,3-]pyridazines/pyrazines via Catalyst-Free Tandem Hydroamination-Aromatic Substitution. <i>Journal of Organic Chemistry</i> , 2020 , 85, 6123-6130	4.2	4
19	DARK Classics in Chemical Neuroscience: Carfentanil. <i>ACS Chemical Neuroscience</i> , 2020 ,	5.7	7
18	Dual Pharmacophores Explored via Structure-Activity Relationship (SAR) Matrix: Insights into Potent, Bifunctional Opioid Ligand Design. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 4193-4203	8.3	5
17	Discovery of Tricyclic Triazolo- and Imidazopyridine Lactams as M Positive Allosteric Modulators. <i>ACS Chemical Neuroscience</i> , 2019 , 10, 1035-1042	5.7	5
16	Biased M receptor-positive allosteric modulators reveal role of phospholipase D in M-dependent rodent cortical plasticity. <i>Science Signaling</i> , 2019 , 12,	8.8	7
15	The discovery of VU0652957 (VU2957, Valiglurax): SAR and DMPK challenges en route to an mGlu PAM development candidate. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019 , 29, 342-346	2.9	6
14	The Muscarinic Acetylcholine Receptor M: Therapeutic Implications and Allosteric Modulation. <i>ACS Chemical Neuroscience</i> , 2019 , 10, 1025-1034	5.7	12
13	Discovery and Optimization of Potent and CNS Penetrant M-Preferring Positive Allosteric Modulators Derived from a Novel, Chiral N-(Indanyl)piperidine Amide Scaffold. <i>ACS Chemical Neuroscience</i> , 2018 , 9, 1572-1581	5.7	7
12	Structure-Activity Relationships of Pan-G α Coupled Muscarinic Acetylcholine Receptor Positive Allosteric Modulators. <i>ACS Chemical Neuroscience</i> , 2018 , 9, 1818-1828	5.7	5
11	Discovery, Characterization, and Effects on Renal Fluid and Electrolyte Excretion of the Kir4.1 Potassium Channel Pore Blocker, VU0134992. <i>Molecular Pharmacology</i> , 2018 , 94, 926-937	4.3	24
10	Classics in Chemical Neuroscience: Xanomeline. <i>ACS Chemical Neuroscience</i> , 2017 , 8, 435-443	5.7	26
9	Synthesis and evaluation of 4,6-disubstituted pyrimidines as CNS penetrant pan-muscarinic antagonists with a novel chemotype. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017 , 27, 2479-2483	2.9	2
8	Discovery and optimization of 3-(4-aryl/heteroarylsulfonyl)piperazin-1-yl)-6-(piperidin-1-yl)pyridazines as novel, CNS penetrant pan-muscarinic antagonists. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017 , 27, 3576-3581	2.9	7

7	Preparation of Unsymmetrical 1,2,4,5-Tetrazines via a Mild Suzuki Cross-Coupling Reaction. <i>Organic Letters</i> , 2017 , 19, 5693-5696	6.2	22
6	Classics in Chemical Neuroscience: Memantine. <i>ACS Chemical Neuroscience</i> , 2017 , 8, 1823-1829	5.7	38
5	Effects of N-Substitutions on the Tetrahydroquinoline (THQ) Core of Mixed-Efficacy μ Opioid Receptor (MOR)/ δ Opioid Receptor (DOR) Ligands. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 4985-98	8.3	30
4	Rapid Synthesis of Boc-2 β -Dimethyl-L-tyrosine and Derivatives and Incorporation into Opioid Peptidomimetics. <i>ACS Medicinal Chemistry Letters</i> , 2015 , 6, 1199-203	4.3	13
3	Asymmetric synthesis and in vitro and in vivo activity of tetrahydroquinolines featuring a diverse set of polar substitutions at the 6 position as mixed-efficacy μ opioid receptor/ δ opioid receptor ligands. <i>ACS Chemical Neuroscience</i> , 2015 , 6, 1428-35	5.7	22
2	Synthesis and evaluation of 4-substituted piperidines and piperazines as balanced affinity μ opioid receptor (MOR) agonist/ δ opioid receptor (DOR) antagonist ligands. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 548-51	2.9	12
1	Opioid peptidomimetics: leads for the design of bioavailable mixed efficacy μ opioid receptor (MOR) agonist/ δ opioid receptor (DOR) antagonist ligands. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 2139-49	8.3	41