Rangan Maitra

List of Publications by Year in descending order

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34 783 16 27
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all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Inhibition of NFÎ $^{\circ}$ B by the natural product Withaferin A in cellular models of Cystic Fibrosis inflammation. Journal of Inflammation, 2009, 6, 15.	3.4	92
2	Synthesis and Biological Evaluation of Bivalent Ligands for the Cannabinoid 1 Receptor. Journal of Medicinal Chemistry, 2010, 53, 7048-7060.	6.4	62
3	Design and Synthesis of Cannabinoid Receptor 1 Antagonists for Peripheral Selectivity. Journal of Medicinal Chemistry, 2012, 55, 2820-2834.	6.4	57
4	Peripherally Selective Cannabinoid 1 Receptor (CB1R) Agonists for the Treatment of Neuropathic Pain. Journal of Medicinal Chemistry, 2016, 59, 7525-7543.	6.4	53
5	Regulation of the Apelinergic System and Its Potential in Cardiovascular Disease: Peptides and Small Molecules as Tools for Discovery. Journal of Medicinal Chemistry, 2015, 58, 7913-7927.	6.4	40
6	Decreased Maternal Plasma Apelin Concentrations in Preeclampsia. Hypertension in Pregnancy, 2012, 31, 398-404.	1.1	35
7	Dual-Acting Cholinesterase–Human Cannabinoid Receptor 2 Ligands Show Pronounced Neuroprotection in Vitro and Overadditive and Disease-Modifying Neuroprotective Effects in Vivo. Journal of Medicinal Chemistry, 2019, 62, 9078-9102.	6.4	35
8	Diphenyl Purine Derivatives as Peripherally Selective Cannabinoid Receptor 1 Antagonists. Journal of Medicinal Chemistry, 2012, 55, 10022-10032.	6.4	31
9	Blocking Alcoholic Steatosis in Mice with a Peripherally Restricted Purine Antagonist of the Type 1 Cannabinoid Receptor. Journal of Medicinal Chemistry, 2018, 61, 4370-4385.	6.4	30
10	Towards rational design of cannabinoid receptor 1 (CB1) antagonists for peripheral selectivity. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 5711-5714.	2.2	29
11	Discovery of a Potent, Selective, and Brain-Penetrant Small Molecule that Activates the Orphan Receptor GPR88 and Reduces Alcohol Intake. Journal of Medicinal Chemistry, 2018, 61, 6748-6758.	6.4	28
12	Identifying structural determinants of potency for analogs of apelin-13: Integration of C-terminal truncation with structure–activity. Bioorganic and Medicinal Chemistry, 2014, 22, 2992-2997.	3.0	27
13	Discovery of a novel small molecule agonist scaffold for the APJ receptor. Bioorganic and Medicinal Chemistry, 2016, 24, 3758-3770.	3.0	26
14	Peripherally Selective Diphenyl Purine Antagonist of the CB1 Receptor. Journal of Medicinal Chemistry, 2013, 56, 8066-8072.	6.4	19
15	Pyrazole antagonists of the CB1 receptor with reduced brain penetration. Bioorganic and Medicinal Chemistry, 2016, 24, 1063-1070.	3.0	19
16	Evaluation of Amide Bioisosteres Leading to 1,2,3-Triazole Containing Compounds as GPR88 Agonists: Design, Synthesis, and Structure–Activity Relationship Studies. Journal of Medicinal Chemistry, 2021, 64, 12397-12413.	6.4	19
17	A patent update on cannabinoid receptor 1 antagonists (2015-2018). Expert Opinion on Therapeutic Patents, 2019, 29, 261-269.	5.0	18
18	"Photo-Rimonabant― Synthesis and Biological Evaluation of Novel Photoswitchable Molecules Derived from Rimonabant Lead to a Highly Selective and Nanomolar " <i>Cis</i> -On―CB ₁ R Antagonist. ACS Chemical Neuroscience, 2021, 12, 1632-1647.	3. 5	17

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19	Design, synthesis, and pharmacological evaluation of JDTic analogs to examine the significance of the 3- and 4-methyl substituents. Bioorganic and Medicinal Chemistry, 2015, 23, 6379-6388.	3.0	14
20	Behavioral assessment of rimonabant under acute and chronic conditions. Behavioural Brain Research, 2020, 390, 112697.	2.2	14
21	Development and validation of a high-throughput calcium mobilization assay for the orphan receptor GPR88. Journal of Biomedical Science, 2017, 24, 23.	7.0	13
22	Identification of potent pyrazole based APELIN receptor (APJ) agonists. Bioorganic and Medicinal Chemistry, 2020, 28, 115237.	3.0	13
23	Metabolic Profiling of CB1 Neutral Antagonists. Methods in Enzymology, 2017, 593, 199-215.	1.0	12
24	Structural analogs of pyrazole and sulfonamide cannabinoids: Effects on acute food intake in mice. European Journal of Pharmacology, 2012, 695, 62-70.	3.5	11
25	Altered Biogenesis of ΔF508-CFTR Following Treatment with Doxorubicin. Cellular Physiology and Biochemistry, 2007, 20, 465-472.	1.6	10
26	Functionalized 6-(Piperidin-1-yl)-8,9-Diphenyl Purines as Peripherally Restricted Inverse Agonists of the CB1 Receptor. Journal of Medicinal Chemistry, 2019, 62, 6330-6345.	6.4	10
27	A Disease-Relevant High-Content Screening Assay to Identify Anti-Inflammatory Compounds for Use in Cystic Fibrosis. Journal of Biomolecular Screening, 2010, 15, 1204-1210.	2.6	9
28	Design, Synthesis, and Structure–Activity Relationship Studies of (4-Alkoxyphenyl)glycinamides and Bioisosteric 1,3,4-Oxadiazoles as GPR88 Agonists. Journal of Medicinal Chemistry, 2020, 63, 14989-15012.	6.4	9
29	Peripherally Selective CB1 Receptor Antagonist Improves Symptoms of Metabolic Syndrome in Mice. ACS Pharmacology and Translational Science, 2021, 4, 757-764.	4.9	9
30	Pyrazole Agonist of the Apelin Receptor Improves Symptoms of Metabolic Syndrome in Mice. Journal of Medicinal Chemistry, 2021, 64, 3006-3025.	6.4	7
31	Synthesis and characterization of an orally bioavailable small molecule agonist of the apelin receptor. Bioorganic and Medicinal Chemistry, 2022, 66, 116789.	3.0	6
32	Synthesis and pharmacological characterization of functionalized 6-piperazin-1-yl-purines as cannabinoid receptor 1 (CB1) inverse agonists. Bioorganic and Medicinal Chemistry, 2018, 26, 4518-4531.	3.0	5
33	Functionalized 6-(piperidin-1-yl)-8,9-diphenyl purines as inverse agonists of the CB1 receptor – SAR efforts towards selectivity and peripheralization. Bioorganic and Medicinal Chemistry, 2019, 27, 3632-3649.	3.0	2
34	Aplnr knockout mice display sex-specific changes in conditioned fear. Behavioural Brain Research, 2021, 400, 113059.	2.2	2