

# Rangan Maitra

## List of Publications by Year in descending order

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34  
papers

783  
citations

516710

16  
h-index

526287

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g-index

34  
all docs

34  
docs citations

34  
times ranked

1167  
citing authors

#	ARTICLE	IF	CITATIONS
1	Synthesis and characterization of an orally bioavailable small molecule agonist of the apelin receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2022, 66, 116789.	3.0	6
2	Aplnr knockout mice display sex-specific changes in conditioned fear. <i>Behavioural Brain Research</i> , 2021, 400, 113059.	2.2	2
3	Peripherally Selective CB1 Receptor Antagonist Improves Symptoms of Metabolic Syndrome in Mice. <i>ACS Pharmacology and Translational Science</i> , 2021, 4, 757-764.	4.9	9
4	Pyrazole Agonist of the Apelin Receptor Improves Symptoms of Metabolic Syndrome in Mice. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 3006-3025.	6.4	7
5	Photo-Rimonabant: Synthesis and Biological Evaluation of Novel Photoswitchable Molecules Derived from Rimonabant Lead to a Highly Selective and Nanomolar Cis-On-CB <sub>1</sub> R Antagonist. <i>ACS Chemical Neuroscience</i> , 2021, 12, 1632-1647.	3.5	17
6	Evaluation of Amide Bioisosteres Leading to 1,2,3-Triazole Containing Compounds as GPR88 Agonists: Design, Synthesis, and Structure-Activity Relationship Studies. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 12397-12413.	6.4	19
7	Identification of potent pyrazole based APELIN receptor (APJ) agonists. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115237.	3.0	13
8	Design, Synthesis, and Structure-Activity Relationship Studies of (4-Alkoxyphenyl)glycinamides and Bioisosteric 1,3,4-Oxadiazoles as GPR88 Agonists. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 14989-15012.	6.4	9
9	Behavioral assessment of rimonabant under acute and chronic conditions. <i>Behavioural Brain Research</i> , 2020, 390, 112697.	2.2	14
10	Functionalized 6-(piperidin-1-yl)-8,9-diphenyl purines as inverse agonists of the CB1 receptor: SAR efforts towards selectivity and peripheralization. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 3632-3649.	3.0	2
11	Dual-Acting Cholinesterase-Human Cannabinoid Receptor 2 Ligands Show Pronounced Neuroprotection in Vitro and Overadditive and Disease-Modifying Neuroprotective Effects in Vivo. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 9078-9102.	6.4	35
12	Functionalized 6-(Piperidin-1-yl)-8,9-Diphenyl Purines as Peripherally Restricted Inverse Agonists of the CB1 Receptor. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 6330-6345.	6.4	10
13	A patent update on cannabinoid receptor 1 antagonists (2015-2018). <i>Expert Opinion on Therapeutic Patents</i> , 2019, 29, 261-269.	5.0	18
14	Blocking Alcoholic Steatosis in Mice with a Peripherally Restricted Purine Antagonist of the Type 1 Cannabinoid Receptor. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 4370-4385.	6.4	30
15	Discovery of a Potent, Selective, and Brain-Penetrant Small Molecule that Activates the Orphan Receptor GPR88 and Reduces Alcohol Intake. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 6748-6758.	6.4	28
16	Synthesis and pharmacological characterization of functionalized 6-piperazin-1-yl-purines as cannabinoid receptor 1 (CB1) inverse agonists. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 4518-4531.	3.0	5
17	Development and validation of a high-throughput calcium mobilization assay for the orphan receptor GPR88. <i>Journal of Biomedical Science</i> , 2017, 24, 23.	7.0	13
18	Metabolic Profiling of CB1 Neutral Antagonists. <i>Methods in Enzymology</i> , 2017, 593, 199-215.	1.0	12

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19	Peripherally Selective Cannabinoid 1 Receptor (CB1R) Agonists for the Treatment of Neuropathic Pain. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 7525-7543.	6.4	53
20	Discovery of a novel small molecule agonist scaffold for the APJ receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3758-3770.	3.0	26
21	Pyrazole antagonists of the CB1 receptor with reduced brain penetration. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 1063-1070.	3.0	19
22	Regulation of the Apelinergic System and Its Potential in Cardiovascular Disease: Peptides and Small Molecules as Tools for Discovery. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 7913-7927.	6.4	40
23	Design, synthesis, and pharmacological evaluation of JDTic analogs to examine the significance of the 3- and 4-methyl substituents. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 6379-6388.	3.0	14
24	Identifying structural determinants of potency for analogs of apelin-13: Integration of C-terminal truncation with structure-activity. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 2992-2997.	3.0	27
25	Peripherally Selective Diphenyl Purine Antagonist of the CB1 Receptor. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 8066-8072.	6.4	19
26	Decreased Maternal Plasma Apelin Concentrations in Preeclampsia. <i>Hypertension in Pregnancy</i> , 2012, 31, 398-404.	1.1	35
27	Diphenyl Purine Derivatives as Peripherally Selective Cannabinoid Receptor 1 Antagonists. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 10022-10032.	6.4	31
28	Structural analogs of pyrazole and sulfonamide cannabinoids: Effects on acute food intake in mice. <i>European Journal of Pharmacology</i> , 2012, 695, 62-70.	3.5	11
29	Design and Synthesis of Cannabinoid Receptor 1 Antagonists for Peripheral Selectivity. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 2820-2834.	6.4	57
30	Towards rational design of cannabinoid receptor 1 (CB1) antagonists for peripheral selectivity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 5711-5714.	2.2	29
31	A Disease-Relevant High-Content Screening Assay to Identify Anti-Inflammatory Compounds for Use in Cystic Fibrosis. <i>Journal of Biomolecular Screening</i> , 2010, 15, 1204-1210.	2.6	9
32	Synthesis and Biological Evaluation of Bivalent Ligands for the Cannabinoid 1 Receptor. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 7048-7060.	6.4	62
33	Inhibition of NF $\kappa$ B by the natural product Withaferin A in cellular models of Cystic Fibrosis inflammation. <i>Journal of Inflammation</i> , 2009, 6, 15.	3.4	92
34	Altered Biogenesis of $\text{F}^{508}$ -CFTR Following Treatment with Doxorubicin. <i>Cellular Physiology and Biochemistry</i> , 2007, 20, 465-472.	1.6	10