

Manuel Grundmann

List of Publications by Year in descending order

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Version: 2024-02-01

31
papers

1,789
citations

331259

21
h-index

454577

30
g-index

31
all docs

31
docs citations

31
times ranked

2532
citing authors

#	ARTICLE	IF	CITATIONS
1	Pharmacological inhibition of Vanin-1 is not protective in models of acute and chronic kidney disease. <i>American Journal of Physiology - Renal Physiology</i> , 2021, 320, F61-F73.	1.3	5
2	Allosteric targeting of the FFA2 receptor (GPR43) restores responsiveness of desensitized human neutrophils. <i>Journal of Leukocyte Biology</i> , 2021, 109, 741-751.	1.5	9
3	Direct Blood Pressure-Independent Anti-Fibrotic Effects by the Selective Nonsteroidal Mineralocorticoid Receptor Antagonist Finerenone in Progressive Models of Kidney Fibrosis. <i>American Journal of Nephrology</i> , 2021, 52, 588-601.	1.4	31
4	Pharmacology of Free Fatty Acid Receptors and Their Allosteric Modulators. <i>International Journal of Molecular Sciences</i> , 2021, 22, 1763.	1.8	55
5	A flow cytometry approach reveals heterogeneity in conventional subsets of murine renal mononuclear phagocytes. <i>Scientific Reports</i> , 2021, 11, 13251.	1.6	8
6	Establishment of a novel, cell-based autotaxin assay. <i>Analytical Biochemistry</i> , 2021, 630, 114322.	1.1	0
7	Lack of beta-arrestin signaling in the absence of active G proteins. <i>Nature Communications</i> , 2018, 9, 341.	5.8	297
8	Label-Free Whole Cell Biosensing for High-Throughput Discovery of Activators and Inhibitors Targeting G Protein-Activated Inwardly Rectifying Potassium Channels. <i>ACS Omega</i> , 2018, 3, 14814-14823.	1.6	10
9	Pro-Angiogenic Effects of Latent Heparanase and Thrombin Receptor-Mediated Pathways—Do They Share a Common Ground in Melanoma Cells?. <i>Thrombosis and Haemostasis</i> , 2018, 118, 1803-1814.	1.8	8
10	Temporal Bias: Time-Encoded Dynamic GPCR Signaling. <i>Trends in Pharmacological Sciences</i> , 2017, 38, 1110-1124.	4.0	68
11	Label-Free Dynamic Mass Redistribution and Bio-Impedance Methods for Drug Discovery. <i>Current Protocols in Pharmacology</i> , 2017, 77, 9.24.1-9.24.21.	4.0	8
12	The molecular mechanism by which saturated lysophosphatidylcholine attenuates the metastatic capacity of melanoma cells. <i>FEBS Open Bio</i> , 2016, 6, 1297-1309.	1.0	19
13	WNT Stimulation Dissociates a Frizzled 4 Inactive-State Complex with G $\alpha_{12/13}$. <i>Molecular Pharmacology</i> , 2016, 90, 447-459.	1.0	33
14	A Molecular Mechanism for Sequential Activation of a G Protein-Coupled Receptor. <i>Cell Chemical Biology</i> , 2016, 23, 392-403.	2.5	30
15	<i>N</i> -Benzylbenzamides: A Novel Merged Scaffold for Orally Available Dual Soluble Epoxide Hydrolase/Peroxisome Proliferator-Activated Receptor β Modulators. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 61-81.	2.9	44
16	Holistic Methods for the Analysis of cNMP Effects. <i>Handbook of Experimental Pharmacology</i> , 2015, 238, 339-357.	0.9	7
17	The experimental power of FR900359 to study Gq-regulated biological processes. <i>Nature Communications</i> , 2015, 6, 10156.	5.8	282
18	Activity of dietary fatty acids on FFA1 and FFA4 and characterisation of pinolenic acid as a dual FFA1/FFA4 agonist with potential effect against metabolic diseases. <i>British Journal of Nutrition</i> , 2015, 113, 1677-1688.	1.2	93

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19	Label-Free Biosensor Assays in GPCR Screening. <i>Methods in Molecular Biology</i> , 2015, 1272, 199-213.	0.4	29
20	Detection of free fatty acid receptor 1 expression: the critical role of negative and positive controls. <i>Diabetologia</i> , 2014, 57, 776-780.	2.9	8
21	cNMP-AMs mimic and dissect bacterial nucleotidyl cyclase toxin effects. <i>Biochemical and Biophysical Research Communications</i> , 2014, 451, 497-502.	1.0	28
22	A Cell-Permeable Inhibitor to Trap G β q Proteins in the Empty Pocket Conformation. <i>Chemistry and Biology</i> , 2014, 21, 890-902.	6.2	47
23	Discovery of a Potent and Selective Free Fatty Acid Receptor 1 Agonist with Low Lipophilicity and High Oral Bioavailability. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 982-992.	2.9	52
24	Discovery of TUG-770: A Highly Potent Free Fatty Acid Receptor 1 (FFA1/GPR40) Agonist for Treatment of Type 2 Diabetes. <i>ACS Medicinal Chemistry Letters</i> , 2013, 4, 441-445.	1.3	58
25	Reevaluation of Fatty Acid Receptor 1 as a Drug Target for the Stimulation of Insulin Secretion in Humans. <i>Diabetes</i> , 2013, 62, 2106-2111.	0.3	64
26	Chemically engineering ligand selectivity at the free fatty acid receptor 2 based on pharmacological variation between species orthologs. <i>FASEB Journal</i> , 2012, 26, 4951-4965.	0.2	75
27	Free Fatty Acid Receptor 1 (FFA1/GPR40) Agonists: Mesylpropoxy Appendage Lowers Lipophilicity and Improves ADME Properties. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 6624-6628.	2.9	50
28	Applying label-free dynamic mass redistribution technology to frame signaling of G protein-coupled receptors noninvasively in living cells. <i>Nature Protocols</i> , 2011, 6, 1748-1760.	5.5	154
29	Identification of a Potent and Selective Free Fatty Acid Receptor 1 (FFA1/GPR40) Agonist with Favorable Physicochemical and in Vitro ADME Properties. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 6691-6703.	2.9	65
30	Conjugated Linoleic Acids Mediate Insulin Release through Islet G Protein-coupled Receptor FFA1/GPR40. <i>Journal of Biological Chemistry</i> , 2011, 286, 11890-11894.	1.6	51
31	Selective Orthosteric Free Fatty Acid Receptor 2 (FFA2) Agonists. <i>Journal of Biological Chemistry</i> , 2011, 286, 10628-10640.	1.6	101