Christian Nanoff

List of Publications by Year
in descending order

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43 papers	1,678 citations	304701 22 h-index	276858 41 g-index
43 all docs	43 docs citations	43 times ranked	1912 citing authors

#	Article	IF	Citations
1	Functional Impact of the G279S Substitution in the Adenosine A₁-Receptor (A₁R-G279S^{7.44}), a Mutation Associated with Parkinsonâ€™s Disease. Molecular Pharmacology, 2020, 98, 250-266.	2.3	2
2	Functional impact of the G279S substitution in the adenosine A1-receptor (A1R-G279S), a mutation associated with Parkinson's disease. Molecular Pharmacology, 2020, 98, MOLPHARM-AR-2020-000003.	2.3	12
3	The fallacy of small sample size - Comment on: Geier D.A., Kern J.K., Homme K.G., Geier M.R., 2018. The risk of neurodevelopmental disorders following thimerosal-containing Hib vaccine in comparison to thimerosal-free Hib vaccine administered from 1995 to 1999 in the United States. Int. J. Hyg. Environ. Health, 221: 677â€“683. International Journal of Hygiene and Environmental Health, 2019, 222, 307-308.	4.3	2
4	Hyponatremia and V2 vasopressin receptor upregulation: a result of HSP90 inhibition. Cancer Chemotherapy and Pharmacology, 2017, 80, 673-684.	2.3	3
5	The value of [11C]-acetate PET and [18F]-FDG PET in hepatocellular carcinoma before and after treatment with transarterial chemoembolization and bevacizumab. European Journal of Nuclear Medicine and Molecular Imaging, 2017, 44, 1732-1741.	6.4	20
6	Relax, Cool Down and Scaffold: How to Restore Surface Expression of Folding-Deficient Mutant GPCRs and SLC6 Transporters. International Journal of Molecular Sciences, 2017, 18, 2416.	4.1	5
7	Comparison of the Â-Adrenergic Receptor Antagonists Landiolol and Esmolol: Receptor Selectivity, Partial Agonism, and Pharmacochaperoning Actions. Journal of Pharmacology and Experimental Therapeutics, 2016, 359, 73-81.	2.5	27
8	Chaperoning of the A₁-Adenosine Receptor by Endogenous Adenosineâ€”An Extension of the Retaliatory Metabolite Concept. Molecular Pharmacology, 2015, 87, 39-51.	2.3	17
9	A Two-state Model for the Diffusion of the A2A Adenosine Receptor in Hippocampal Neurons. Journal of Biological Chemistry, 2014, 289, 9263-9274.	3.4	21
10	Recruitment of a Cytoplasmic Chaperone Relay by the A2A Adenosine Receptor. Journal of Biological Chemistry, 2013, 288, 28831-28844.	3.4	38
11	ER-Bound Steps in the Biosynthesis of G Protein-Coupled Receptors. Sub-Cellular Biochemistry, 2012, 63, 1-21.	2.4	11
12	Constitutive activity of the A2A adenosine receptor and compartmentalised cyclic AMP signalling fine-tune noradrenaline release. Purinergic Signalling, 2012, 8, 677-692.	2.2	12
13	The noradrenaline transporter as site of action for the anti-Parkinson drug amantadine. Neuropharmacology, 2012, 62, 1708-1716.	4.1	19
14	Pharmacochaperoning of the A₁Adenosine Receptor Is Contingent on the Endoplasmic Reticulum. Molecular Pharmacology, 2010, 77, 940-952.	2.3	37
15	Limitations in Adjuvant Breast Cancer Therapy: The Predictive Potential of Pharmacogenetics and Pharmacogenomics. Breast Care, 2008, 3, 401-406.	1.4	2
16	The Carboxyl Terminus of the GÎ±-Subunit Is the Latch for Triggered Activation of Heterotrimeric G Proteins. Molecular Pharmacology, 2006, 69, 397-405.	2.3	33
17	The Ubiquitin-Specific Protease Usp4 Regulates the Cell Surface Level of the A2a Receptor. Molecular Pharmacology, 2006, 69, 1083-1094.	2.3	122
18	The Human D2 Dopamine Receptor Synergizes with the A2A Adenosine Receptor to Stimulate Adenylyl Cyclase in PC12 Cells. Neuropsychopharmacology, 2003, 28, 1317-1327.	5.4	49

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19	Truncation of the A1 Adenosine Receptor Reveals Distinct Roles of the Membrane-proximal Carboxyl Terminus in Receptor Folding and G Protein Coupling. Journal of Biological Chemistry, 2003, 278, 30283-30293.	3.4	61
20	Development of Gs-Selective Inhibitory Compounds. Methods in Enzymology, 2002, 344, 469-480.	1.0	4
21	Beyond G proteins: The role of accessory proteins in G protein-coupled receptor signalling. Pharmacochemistry Library, 2002, , 161-173.	0.1	0
22	Adenosine receptors: G protein-mediated signalling and the role of accessory proteins. Cellular Signalling, 2002, 14, 99-108.	3.6	235
23	Biased inhibition by a suramin analogue of A 1 -adenosine receptor/G protein coupling in fused receptor/G protein tandems: the A 1 -adenosine receptor is predominantly coupled to Go Î± in human brain. Naunyn-Schmiedeberg's Archives of Pharmacology, 2002, 365, 8-16.	3.0	26
24	Removal of the carboxy terminus of the A 2A -adenosine receptor blunts constitutive activity: differential effect on cAMP accumulation and MAP kinase stimulation. Naunyn-Schmiedeberg's Archives of Pharmacology, 2002, 366, 287-298.	3.0	52
25	Suramin and the suramin analogue NF307 discriminate among calmodulin-binding sites. Biochemical Journal, 2001, 355, 827-833.	3.7	23
26	Mapping of Calmodulin and GÎ²Î³ Binding Domains within the C-terminal Region of the Metabotropic Glutamate Receptor 7A. Journal of Biological Chemistry, 2001, 276, 30662-30669.	3.4	60
27	Inhibition of Adenylyl and Guanylyl Cyclase Isoforms by the Antiviral Drug Foscarnet. Journal of Biological Chemistry, 2001, 276, 3010-3016.	3.4	28
28	G protein-dependent signalling and ageingâ˜†. Experimental Gerontology, 2000, 35, 133-143.	2.8	13
29	Binding of Calmodulin to the D2-Dopamine Receptor Reduces Receptor Signaling by Arresting the G Protein Activation Switch. Journal of Biological Chemistry, 2000, 275, 32672-32680.	3.4	122
30	Tight Association of the Human Mel_{1a}-Melatonin Receptor and G_i: Precoupling and Constitutive Activity. Molecular Pharmacology, 1999, 56, 1014-1024.	2.3	100
31	Kinetics of Ternary Complex Formation with Fusion Proteins Composed of the A1-Adenosine Receptor and G Protein Î±-Subunits. Journal of Biological Chemistry, 1999, 274, 30571-30579.	3.4	46
32	Calmodulin Dependence of Presynaptic Metabotropic Glutamate Receptor Signaling. Science, 1999, 286, 1180-1184.	12.6	153
33	G protein antagonists. Trends in Pharmacological Sciences, 1999, 20, 237-245.	8.7	96
34	Metal-dependent nucleotide binding to the Escherichia coli rotamase SlyD. Biochemical Journal, 1999, 342, 33-39.	3.7	23
35	Metal-dependent nucleotide binding to the Escherichia coli rotamase SlyD. Biochemical Journal, 1999, 342, 33.	3.7	10
36	The C2 Catalytic Domain of Adenylyl Cyclase Contains the Second Metal Ion (Mn2+) Binding Siteâ€. Biochemistry, 1998, 37, 16183-16191.	2.5	21

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37	G Protein coupling of the rat A1-adenosine receptorâ€"Partial purification of a protein which stabilizes the receptor-G protein association. Neuropharmacology, 1997, 36, 1211-1219.	4.1	26
38	Solubilization and Characterization of the A2-Adenosine Receptor. Journal of Receptors and Signal Transduction, 1993, 13, 961-973.	1.2	33
39	Sensitization of Dopamine-Stimulated Adenylyl Cyclase in the Striatum of 1-Methyl-4-Phenyl-1,2,3,6-Tetrahydropyridine-Treated Rhesus Monkeys and Patients with Idiopathic Parkinson's Disease. Journal of Neurochemistry, 1992, 58, 1997-2004.	3.9	52
40	Characterization of the Î²-Adrenoceptor Blocking Property of Diprafenone in Rats. Journal of Cardiovascular Pharmacology, 1991, 18, 837-842.	1.9	5
41	P₂â€, but not P₁â€purinoceptors mediate formation of 1,4,5â€inositol trisphosphate and its metabolites via a pertussis toxinâ€insensitive pathway in the rat renal cortex. British Journal of Pharmacology, 1990, 100, 63-68.	5.4	36
42	The fallacy of non-selectivity of radioactive ligands. Trends in Pharmacological Sciences, 1988, 9, 261-264.	8.7	8
43	Stimulation of adenylate cyclase activity via A2-adenosine receptors in isolated tubules of the rabbit renal cortex. European Journal of Pharmacology, 1987, 138, 137-140.	3.5	13