

Richard Neubig

List of Publications by Year in descending order

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Version: 2024-02-01

265
papers

13,276
citations

22153

59
h-index

30087

103
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416
all docs

416
docs citations

416
times ranked

12013
citing authors

#	ARTICLE	IF	CITATIONS
1	Ibrutinib Blocks YAP1 Activation and Reverses BRAF Inhibitor Resistance in Melanoma Cells. <i>Molecular Pharmacology</i> , 2022, 101, 1-12.	2.3	5
2	Mice with monoallelic <i>GNAO1</i> loss exhibit reduced inhibitory synaptic input to cerebellar Purkinje cells. <i>Journal of Neurophysiology</i> , 2022, 127, 607-622.	1.8	5
3	BRAF Inhibitor Resistance Confers Increased Sensitivity to Mitotic Inhibitors. <i>Frontiers in Oncology</i> , 2022, 12, 766794.	2.8	2
4	A Glowing Opportunity to Target YAP in Lung Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2022, 67, 1-2.	2.9	1
5	Mice With Monoallelic <i>GNAO1</i> Loss Exhibit Reduced Inhibitory Synaptic Input to Cerebellar Purkinje Cells. <i>FASEB Journal</i> , 2022, 36, .	0.5	0
6	Mice with an RGS-insensitive G_{i2} protein show growth hormone axis dysfunction. <i>Molecular and Cellular Endocrinology</i> , 2021, 521, 111098.	3.2	3
7	COVID-19 – A Theory of Autoimmunity Against ACE-2 Explained. <i>Frontiers in Immunology</i> , 2021, 12, 582166.	4.8	41
8	Inhibition of the Myocardin-Related Transcription Factor Pathway Increases Efficacy of Trametinib in NRAS-Mutant Melanoma Cell Lines. <i>Cancers</i> , 2021, 13, 2012.	3.7	6
9	Transforming Growth Factor β_1 Increases Expression of Contractile Genes in Human Pulmonary Arterial Smooth Muscle Cells by Potentiating Sphingosine-1-Phosphate Signaling. <i>Molecular Pharmacology</i> , 2021, 100, 83-90.	2.3	7
10	Convergent olfactory trace amine-associated receptors detect biogenic polyamines with distinct motifs via a conserved binding site. <i>Journal of Biological Chemistry</i> , 2021, 297, 101268.	3.4	6
11	Class A Orphans in GtoPdb v.2021.3. <i>IUPHAR/BPS Guide To Pharmacology CITE</i> , 2021, 2021, .	0.2	3
12	G protein-coupled estrogen receptor in GtoPdb v.2021.3. <i>IUPHAR/BPS Guide To Pharmacology CITE</i> , 2021, 2021, .	0.2	0
13	RGS4 negatively modulates Nociceptin/Orphanin FQ opioid receptor signaling: implication for L-Dopa-induced dyskinesia.. <i>British Journal of Pharmacology</i> , 2021, .	5.4	1
14	Rho-mediated signaling promotes BRAF inhibitor resistance in de-differentiated melanoma cells. <i>Oncogene</i> , 2020, 39, 1466-1483.	5.9	40
15	Two highly related odorant receptors specifically detect β -bile acid pheromones in sea lamprey (<i>Petromyzon marinus</i>). <i>Journal of Biological Chemistry</i> , 2020, 295, 12153-12166.	3.4	6
16	Mice with GNAO1 R209H Movement Disorder Variant Display Hyperlocomotion Alleviated by Risperidone. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2020, 373, jpet.119.262733.	2.5	14
17	Class A Orphans (version 2020.5) in the IUPHAR/BPS Guide to Pharmacology Database. <i>IUPHAR/BPS Guide To Pharmacology CITE</i> , 2020, 2020, .	0.2	7
18	Spermine in semen of male sea lamprey acts as a sex pheromone. <i>PLoS Biology</i> , 2019, 17, e3000332.	5.6	37

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19	Identifying chemopreventive agents for obesity-associated cancers using an efficient, 3D high-throughput transformation assay. <i>Scientific Reports</i> , 2019, 9, 10278.	3.3	4
20	RAC1P29S Induces a Mesenchymal Phenotypic Switch via Serum Response Factor to Promote Melanoma Development and Therapy Resistance. <i>Cancer Cell</i> , 2019, 36, 68-83.e9.	16.8	104
21	Mouse models of GNAO1-associated movement disorder: Allele- and sex-specific differences in phenotypes. <i>PLoS ONE</i> , 2019, 14, e0211066.	2.5	18
22	The Rho/MRTF pathway inhibitor CCG-222740 reduces stellate cell activation and modulates immune cell populations in <i>Kras</i> G12D; <i>Pdx1-Cre</i> (KC) mice. <i>Scientific Reports</i> , 2019, 9, 7072.	3.3	17
23	Cover Image, Volume 87, Issue 2. <i>Proteins: Structure, Function and Bioinformatics</i> , 2019, 87, C1.	2.6	0
24	Identification of Pirin as a Molecular Target of the CCG-1423/CCG-203971 Series of Antifibrotic and Antimetastatic Compounds. <i>ACS Pharmacology and Translational Science</i> , 2019, 2, 92-100.	4.9	28
25	5-Aryl-1,3,4-oxadiazol-2-ylthioalkanoic Acids: A Highly Potent New Class of Inhibitors of Rho/Myocardin-Related Transcription Factor (MRTF)/Serum Response Factor (SRF)-Mediated Gene Transcription as Potential Antifibrotic Agents for Scleroderma. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 4350-4369.	6.4	34
26	A Two-Pulse Cellular Stimulation Test Elucidates Variability and Mechanisms in Signaling Pathways. <i>Biophysical Journal</i> , 2019, 116, 962-973.	0.5	11
27	An Interhelical Salt Bridge Controls Flexibility and Inhibitor Potency for Regulators of G-protein Signaling Proteins 4, 8, and 19. <i>Molecular Pharmacology</i> , 2019, 96, 683-691.	2.3	8
28	Loss-of-Function Mutations in Human Regulator of G Protein Signaling RGS2 Differentially Regulate Pharmacological Reactivity of Resistance Vasculature. <i>Molecular Pharmacology</i> , 2019, 96, 826-834.	2.3	6
29	Interplay of cysteine exposure and global protein dynamics in small-molecule recognition by a regulator of G-protein signaling protein. <i>Proteins: Structure, Function and Bioinformatics</i> , 2019, 87, 146-156.	2.6	13
30	Class A Orphans (version 2019.5) in the IUPHAR/BPS Guide to Pharmacology Database. <i>IUPHAR/BPS Guide To Pharmacology CITE</i> , 2019, 2019, .	0.2	8
31	G protein-coupled estrogen receptor (version 2019.4) in the IUPHAR/BPS Guide to Pharmacology Database. <i>IUPHAR/BPS Guide To Pharmacology CITE</i> , 2019, 2019, .	0.2	1
32	A Salt Bridge between $\hat{I}\pm 4$ and $\hat{I}\pm 5$ Helices Drives Differences in Flexibility and Potency of Inhibition among Regulator of G-protein Signaling (RGS) Proteins. <i>FASEB Journal</i> , 2019, 33, 784.16.	0.5	0
33	Mice with <i>Gnao1</i> G203R Gain-of-Function (GOF) Mutation Phenocopy Combined Movement Disorder and Seizures of Patients. <i>FASEB Journal</i> , 2019, 33, 667.4.	0.5	0
34	Class A Orphans (version 2019.4) in the IUPHAR/BPS Guide to Pharmacology Database. <i>IUPHAR/BPS Guide To Pharmacology CITE</i> , 2019, 2019, .	0.2	0
35	Differential Protein Dynamics of Regulators of G-Protein Signaling: Role in Specificity of Small-Molecule Inhibitors. <i>Journal of the American Chemical Society</i> , 2018, 140, 3454-3460.	13.7	21
36	Role of signalling molecules in behaviours mediated by the \hat{I} opioid receptor agonist SNC80. <i>British Journal of Pharmacology</i> , 2018, 175, 891-901.	5.4	31

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37	In vitro and in vivo delivery of a sustained release nanocarrier-based formulation of an MRTF/SRF inhibitor in conjunctival fibrosis. <i>Journal of Nanobiotechnology</i> , 2018, 16, 97.	9.1	20
38	Role of hippocampal 5-HT1A receptors in the antidepressant-like phenotype of mice expressing RGS-insensitive Gi α 2 protein. <i>Neuropharmacology</i> , 2018, 141, 296-304.	4.1	2
39	Interpreting Hydrogen/Deuterium Exchange Events in Proteins Using Atomistic Simulations: Case Studies on Regulators of G-Protein Signaling Proteins. <i>Journal of Physical Chemistry B</i> , 2018, 122, 9314-9323.	2.6	30
40	A mechanistic review on GNAO1-associated movement disorder. <i>Neurobiology of Disease</i> , 2018, 116, 131-141.	4.4	62
41	Novel antifibrotic target related to RhoA-induced MRTF activation in fibrotic diseases. Proceedings for Annual Meeting of the Japanese Pharmacological Society, 2018, WCP2018, SY84-4.	0.0	0
42	Loss of function RGS2 mutations augment vascular contractility ex vivo. <i>FASEB Journal</i> , 2018, 32, 699.2.	0.5	0
43	Role of Protein Dynamics in Selectivity of Thiadiazolidinone Inhibition of RGS Proteins. <i>FASEB Journal</i> , 2018, 32, 557.9.	0.5	0
44	Chemerin-induced arterial contraction is Gi- and calcium-dependent. <i>Vascular Pharmacology</i> , 2017, 88, 30-41.	2.1	33
45	Pharmacokinetic optimization of CCG-203971: Novel inhibitors of the Rho/MRTF/SRF transcriptional pathway as potential antifibrotic therapeutics for systemic scleroderma. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 1744-1749.	2.2	42
46	Local delivery of novel MRTF/SRF inhibitors prevents scar tissue formation in a preclinical model of fibrosis. <i>Scientific Reports</i> , 2017, 7, 518.	3.3	52
47	The DRY motif and the four corners of the cubic ternary complex model. <i>Cellular Signalling</i> , 2017, 35, 16-23.	3.6	14
48	Regulator of G Protein Signaling 6 Protects the Heart from Ischemic Injury. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2017, 360, 409-416.	2.5	15
49	Human Missense Mutations in Regulator of G Protein Signaling 2 Affect the Protein Function Through Multiple Mechanisms. <i>Molecular Pharmacology</i> , 2017, 92, 451-458.	2.3	12
50	Movement disorder in GNAO1 encephalopathy associated with gain-of-function mutations. <i>Neurology</i> , 2017, 89, 762-770.	1.1	73
51	The role of regulator of G protein signaling 4 in delta-opioid receptor-mediated behaviors. <i>Psychopharmacology</i> , 2017, 234, 29-39.	3.1	19
52	Pharmacological Inhibition of Myocardin-related Transcription Factor Pathway Blocks Lung Metastases of RhoC-Overexpressing Melanoma. <i>Molecular Cancer Therapeutics</i> , 2017, 16, 193-204.	4.1	35
53	A central role for R7bp in the regulation of itch sensation. <i>Pain</i> , 2017, 158, 931-944.	4.2	11
54	Investigating Regulator of G-protein Signaling (RGS) Protein Dynamics by Hydrogen/Deuterium Exchange. <i>FASEB Journal</i> , 2017, 31, 665.8.	0.5	0

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55	Small-Molecule Inhibition of Rho/MKL/SRF Transcription in Prostate Cancer Cells: Modulation of Cell Cycle, ER Stress, and Metastasis Gene Networks. <i>Microarrays (Basel, Switzerland)</i> , 2016, 5, 13.	1.4	21
56	Optimisation of Intestinal Fibrosis and Survival in the Mouse. Typhimurium Model for Anti-fibrotic Drug Discovery and Preclinical Applications. <i>Journal of Crohn's and Colitis</i> , 2016, 11, jjw210.	1.3	6
57	RGS Proteins and G β 12 Modulate Sleep, Wakefulness, and Disruption of Sleep/ Wake States after Isoflurane and Sevoflurane Anesthesia. <i>Sleep</i> , 2016, 39, 393-404.	1.1	7
58	Digoxin-Mediated Upregulation of RGS2 Protein Protects against Cardiac Injury. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2016, 357, 311-319.	2.5	20
59	FBXO44-Mediated Degradation of RGS2 Protein Uniquely Depends on a Cullin 4B/DDB1 Complex. <i>PLoS ONE</i> , 2015, 10, e0123581.	2.5	21
60	Band-pass processing in a GPCR signaling pathway selects for NFAT transcription factor activation. <i>Integrative Biology (United Kingdom)</i> , 2015, 7, 1378-1386.	1.3	20
61	Inhibition of Myocardin-Related Transcription Factor/Serum Response Factor Signaling Decreases Lung Fibrosis and Promotes Mesenchymal Cell Apoptosis. <i>American Journal of Pathology</i> , 2015, 185, 969-986.	3.8	138
62	Selectivity and Anti-Parkinson's Potential of Thiadiazolidinone RGS4 Inhibitors. <i>ACS Chemical Neuroscience</i> , 2015, 6, 911-919.	3.5	41
63	RGS-Insensitive G Proteins as In Vivo Probes of RGS Function. <i>Progress in Molecular Biology and Translational Science</i> , 2015, 133, 13-30.	1.7	20
64	M4 Muscarinic Receptor Signaling Ameliorates Striatal Plasticity Deficits in Models of L-DOPA-Induced Dyskinesia. <i>Neuron</i> , 2015, 88, 762-773.	8.1	183
65	Regulator of G Protein Signaling Protein 6 (RGS6) Protects the Heart from Ischemic Injury. <i>FASEB Journal</i> , 2015, 29, 1026.8.	0.5	0
66	RGS4 Differentially Regulates Antidepressant and Locomotor Behaviors In Vivo. <i>FASEB Journal</i> , 2015, 29, 618.11.	0.5	0
67	RGS2 Protein Degradation is Mediated by a Novel Cullin 4B/F-box 44 E3 Ligase Complex. <i>FASEB Journal</i> , 2015, 29, 618.15.	0.5	0
68	Regulation of Protease-activated Receptor 1 Signaling by the Adaptor Protein Complex 2 and R4 Subfamily of Regulator of G Protein Signaling Proteins. <i>Journal of Biological Chemistry</i> , 2014, 289, 1580-1591.	3.4	13
69	Identification of Protein Kinase C Activation as a Novel Mechanism for RGS2 Protein Upregulation through Phenotypic Screening of Natural Product Extracts. <i>Molecular Pharmacology</i> , 2014, 86, 406-416.	2.3	15
70	Novel Rho/MRTF/SRF Inhibitors Block Matrix-stiffness and TGF- β -Induced Fibrogenesis in Human Colonic Myofibroblasts. <i>Inflammatory Bowel Diseases</i> , 2014, 20, 154-165.	1.9	155
71	Targeting the Myofibroblast Genetic Switch: Inhibitors of Myocardin-Related Transcription Factor/Serum Response Factor-Regulated Gene Transcription Prevent Fibrosis in a Murine Model of Skin Injury. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2014, 349, 480-486.	2.5	92
72	Redox Modification of Nuclear Actin by MICAL-2 Regulates SRF Signaling. <i>Cell</i> , 2014, 156, 563-576.	28.9	142

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73	Cellular Mechanisms of Tissue Fibrosis. 8. Current and future drug targets in fibrosis: focus on Rho GTPase-regulated gene transcription. American Journal of Physiology - Cell Physiology, 2014, 307, C2-C13.	4.6	71
74	Induction of the matricellular protein CCN1 through RhoA and MRTF-A contributes to ischemic cardioprotection. Journal of Molecular and Cellular Cardiology, 2014, 75, 152-161.	1.9	29
75	Gain-of-function mutation in Gnao1: A murine model of epileptiform encephalopathy (EIEE17)?. Mammalian Genome, 2014, 25, 202-210.	2.2	34
76	International Union of Basic and Clinical Pharmacology. XC. Multisite Pharmacology: Recommendations for the Nomenclature of Receptor Allosterism and Allosteric Ligands. Pharmacological Reviews, 2014, 66, 918-947.	16.0	189
77	Conditional disruption of interactions between G α 12 and regulator of G protein signaling (RGS) proteins protects the heart from ischemic injury. BMC Pharmacology & Toxicology, 2014, 15, 29.	2.4	9
78	Optimization of novel nipecotic bis(amide) inhibitors of the Rho/MKL1/SRF transcriptional pathway as potential anti-metastasis agents. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 3826-3832.	2.2	61
79	Conformational Dynamics of a Regulator of G-Protein Signaling Protein Reveals a Mechanism of Allosteric Inhibition by a Small Molecule. ACS Chemical Biology, 2013, 8, 2778-2784.	3.4	33
80	NMR Methods for Detection of Small Molecule Binding to RGS4. Methods in Enzymology, 2013, 522, 133-152.	1.0	5
81	Reversible inhibitors of regulators of G-protein signaling identified in a high-throughput cell-based calcium signaling assay. Cellular Signalling, 2013, 25, 2848-2855.	3.6	21
82	Microfluidic interrogation and mathematical modeling of multi-regime calcium signaling dynamics. Integrative Biology (United Kingdom), 2013, 5, 932.	1.3	11
83	Regulation of G protein signaling by the 70kDa heat shock protein. Cellular Signalling, 2013, 25, 389-396.	3.6	8
84	Differential Control of Opioid Antinociception to Thermal Stimuli in a Knock-In Mouse Expressing Regulator of G-Protein Signaling-Insensitive G α o Protein. Journal of Neuroscience, 2013, 33, 4369-4377.	3.6	29
85	International Union of Basic and Clinical Pharmacology. LXXXVIII. G Protein-Coupled Receptor List: Recommendations for New Pairings with Cognate Ligands. Pharmacological Reviews, 2013, 65, 967-986.	16.0	250
86	Detection of G Protein-selective G Protein-coupled Receptor (GPCR) Conformations in Live Cells. Journal of Biological Chemistry, 2013, 288, 17167-17178.	3.4	60
87	Design and synthesis of tag-free photoprobes for the identification of the molecular target for CCG-1423, a novel inhibitor of the Rho/MKL1/SRF signaling pathway. Beilstein Journal of Organic Chemistry, 2013, 9, 966-973.	2.2	22
88	RGS 2 and RGS 4 Differentially Modulate G Protein Coupled Receptor Signaling in the Mouse Aorta. FASEB Journal, 2013, 27, 1095.3.	0.5	0
89	Dynamic control of Allosteric Inhibitor Specificity for RGS4. FASEB Journal, 2013, 27, 1095.11.	0.5	0
90	Increased Go activity in C57Bl/6j mice enhances sensitivity to a model of epilepsy. FASEB Journal, 2013, 27, 660.1.	0.5	0

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91	Cardiotonic Steroids Stabilize Regulator of G Protein Signaling 2 Protein Levels. <i>Molecular Pharmacology</i> , 2012, 82, 500-509.	2.3	23
92	MScreen: An Integrated Compound Management and High-Throughput Screening Data Storage and Analysis System. <i>Journal of Biomolecular Screening</i> , 2012, 17, 1080-1087.	2.6	44
93	The Loss of RGS Protein-G α Interactions Results in Markedly Impaired Mouse Neutrophil Trafficking to Inflammatory Sites. <i>Molecular and Cellular Biology</i> , 2012, 32, 4561-4571.	2.3	32
94	A newly identified complex of spinophilin and the tyrosine phosphatase, SHP-1, modulates platelet activation by regulating G protein-dependent signaling. <i>Blood</i> , 2012, 119, 1935-1945.	1.4	57
95	Toll-like Receptor-Induced Inflammatory Cytokines are Suppressed by Gain of Function or Overexpression of G α i2 Protein. <i>Inflammation</i> , 2012, 35, 1611-1617.	3.8	16
96	G α i2 signaling: friend or foe in cardiac injury and heart failure?. <i>Naunyn-Schmiedeberg's Archives of Pharmacology</i> , 2012, 385, 443-453.	3.0	15
97	Small Molecule Inhibitors of Regulators of G Protein Signaling (RGS) Proteins. <i>ACS Medicinal Chemistry Letters</i> , 2012, 3, 146-150.	2.8	41
98	Increased CD39 Nucleotidase Activity on Microparticles from Patients with Idiopathic Pulmonary Arterial Hypertension. <i>PLoS ONE</i> , 2012, 7, e40829.	2.5	43
99	Targeting degradation pathways of RGS2 using high-throughput siRNA screening. <i>FASEB Journal</i> , 2012, 26, 838.9.	0.5	0
100	S1P induces CCN1 expression through RhoA/MRTF α activation and protects cardiomyocytes against cell death. <i>FASEB Journal</i> , 2012, 26, 1060.4.	0.5	0
101	Generation of G α i2 G184S conditional mutant mice to study regulator of G protein signaling (RGS) proteins. <i>FASEB Journal</i> , 2012, 26, 1114.10.	0.5	0
102	Hi-Fi transmission of periodic signals amid cell-to-cell variability. <i>Molecular BioSystems</i> , 2011, 7, 2238.	2.9	16
103	A Nanomolar-Potency Small Molecule Inhibitor of Regulator of G-Protein Signaling Proteins. <i>Biochemistry</i> , 2011, 50, 3181-3192.	2.5	55
104	RGS-Insensitive G α Subunits: Probes of G α Subtype-Selective Signaling and Physiological Functions of RGS Proteins. <i>Methods in Molecular Biology</i> , 2011, 756, 75-98.	0.9	14
105	Complementary Cell-Based High-Throughput Screens Identify Novel Modulators of the Unfolded Protein Response. <i>Journal of Biomolecular Screening</i> , 2011, 16, 825-835.	2.6	44
106	G α i2-mediated protection from ischaemic injury is modulated by endogenous RGS proteins in the mouse heart. <i>Cardiovascular Research</i> , 2011, 91, 45-52.	3.8	17
107	Glossary of terms used in biomolecular screening (IUPAC Recommendations 2011). <i>Pure and Applied Chemistry</i> , 2011, 83, 1129-1158.	1.9	8
108	Differential effects of G α o and G α i2 on seizure threshold. <i>FASEB Journal</i> , 2011, 25, 1010.2.	0.5	0

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109	RGS/Gi2 interactions modulate platelet accumulation and thrombus formation at sites of vascular injury. <i>Blood</i> , 2010, 116, 6092-6100.	1.4	52
110	Design, synthesis and prostate cancer cell-based studies of analogs of the Rho/MKL1 transcriptional pathway inhibitor, CCG-1423. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 665-672.	2.2	60
111	G α Subunit Coordinates with Ephrin-B to Balance Self-Renewal and Differentiation in Neural Progenitor Cells. <i>Stem Cells</i> , 2010, 28, 1581-1589.	3.2	17
112	Differential modulation of μ -opioid receptor signaling to adenylyl cyclase by regulators of G protein signaling proteins 4 or 8 and 7 in permeabilised C6 cells is G α subtype dependent. <i>Journal of Neurochemistry</i> , 2010, 112, 1026-1034.	3.9	29
113	Mind Your Salts: When the Inactive Constituent Isn't: Fig. 1.. <i>Molecular Pharmacology</i> , 2010, 78, 558-559.	2.3	18
114	Reversible, Allosteric Small-Molecule Inhibitors of Regulator of G Protein Signaling Proteins. <i>Molecular Pharmacology</i> , 2010, 78, 524-533.	2.3	70
115	Allosteric Inhibition of the Regulator of G Protein Signaling α -G α Protein α -Protein Interaction by CCG-4986. <i>Molecular Pharmacology</i> , 2010, 78, 360-365.	2.3	39
116	GPCR-OKB: the G Protein Coupled Receptor Oligomer Knowledge Base. <i>Bioinformatics</i> , 2010, 26, 1804-1805.	4.1	74
117	RGS inhibition at G α selectively potentiates 5-HT1A-mediated antidepressant effects. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 11086-11091.	7.1	60
118	Phase-Locked Signals Elucidate Circuit Architecture of an Oscillatory Pathway. <i>PLoS Computational Biology</i> , 2010, 6, e1001040.	3.2	40
119	Regulators of G Protein Signaling Proteins as Targets for Drug Discovery. <i>Progress in Molecular Biology and Translational Science</i> , 2010, 91, 81-119.	1.7	84
120	Analyzing Binding Data. <i>Current Protocols in Neuroscience</i> , 2010, 52, Unit 7.5.	2.6	41
121	Thinking Outside of the "RGS Box": New Approaches to Therapeutic Targeting of Regulators of G Protein Signaling: Fig. 1.. <i>Molecular Pharmacology</i> , 2010, 78, 550-557.	2.3	67
122	Use of Flow Cytometric Methods to Quantify Protein-Protein Interactions. <i>Current Protocols in Cytometry</i> , 2010, 51, Unit 13.11.1-15.	3.7	27
123	RGS7 Protein Suppression of G α -Mediated β -Adrenergic Receptor Inhibition of Mouse Hippocampal CA3 Epileptiform Activity. <i>FASEB Journal</i> , 2010, 24, 587.3.	0.5	0
124	International Union of Pharmacology. LXXII. Recommendations for Trace Amine Receptor Nomenclature. <i>Pharmacological Reviews</i> , 2009, 61, 1-8.	16.0	49
125	Polyplexed Flow Cytometry Protein Interaction Assay: A Novel High-Throughput Screening Paradigm for RGS Protein Inhibitors. <i>Journal of Biomolecular Screening</i> , 2009, 14, 610-619.	2.6	38
126	Regulator of G Protein Signaling Protein Suppression of G α Protein-Mediated β -Adrenergic Receptor Inhibition of Mouse Hippocampal CA3 Epileptiform Activity. <i>Molecular Pharmacology</i> , 2009, 75, 1222-1230.	2.3	26

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127	High-Throughput Screening for Small-Molecule Inhibitors of LARG-Stimulated RhoA Nucleotide Binding via a Novel Fluorescence Polarization Assay. <i>Journal of Biomolecular Screening</i> , 2009, 14, 161-172.	2.6	42
128	IUPHAR-DB: the IUPHAR database of G protein-coupled receptors and ion channels. <i>Nucleic Acids Research</i> , 2009, 37, D680-D685.	14.5	199
129	A Juxtamembrane Mutation in the N Terminus of the Dopamine Transporter Induces Preference for an Inward-Facing Conformation. <i>Molecular Pharmacology</i> , 2009, 75, 514-524.	2.3	61
130	A conserved hydrophobic surface of the LARG pleckstrin homology domain is critical for RhoA activation in cells. <i>Cellular Signalling</i> , 2009, 21, 1569-1578.	3.6	22
131	A covalent peptide inhibitor of RGS4 identified in a focused one-bead, one compound library screen. <i>BMC Pharmacology</i> , 2009, 9, 9.	0.4	18
132	GNAI2 and regulators of G protein signaling as a potential Noonan syndrome mechanism. <i>Medical Hypotheses</i> , 2009, 73, 56-59.	1.5	4
133	Small Molecule Protein-Protein Interaction Inhibitors as CNS Therapeutic Agents: Current Progress and Future Hurdles. <i>Neuropsychopharmacology</i> , 2009, 34, 126-141.	5.4	164
134	Isoflurane-Induced Changes in Righting Response and Breathing Are Modulated by RGS Proteins. <i>Anesthesia and Analgesia</i> , 2009, 109, 1500-1505.	2.2	16
135	Functional Selectivity at Adrenergic Receptors. , 2009, , 107-124.		0
136	In vitro protein kinase activity measurement by flow cytometry. <i>Analytical Biochemistry</i> , 2008, 383, 180-185.	2.4	7
137	Novel Peptide Ligands of RGS4 from a Focused One-Bead, One-Compound Library. <i>Chemical Biology and Drug Design</i> , 2008, 72, 111-119.	3.2	25
138	Microfabricated Channel Array Electrophoresis for Characterization and Screening of Enzymes Using RGS-G Protein Interactions as a Model System. <i>Analytical Chemistry</i> , 2008, 80, 5225-5231.	6.5	18
139	And the Winner Is RGS4!. <i>Circulation Research</i> , 2008, 103, 444-446.	4.5	8
140	Assembly of High Order G β q-Effector Complexes with RGS Proteins. <i>Journal of Biological Chemistry</i> , 2008, 283, 34923-34934.	3.4	46
141	Resistance to Diet-Induced Obesity and Improved Insulin Sensitivity in Mice With a Regulator of G Protein Signaling-Insensitive G184S Gnai2 Allele. <i>Diabetes</i> , 2008, 57, 77-85.	0.6	50
142	β 2A adrenergic receptor mediated antiepileptic effects via G β o proteins. <i>FASEB Journal</i> , 2008, 22, 729.3.	0.5	0
143	Genetic deletion of Regulators of G protein Signaling (RGS) protein activity enhances buprenorphine antinociception while limiting withdrawal behaviors associated with chronic administration. <i>FASEB Journal</i> , 2008, 22, 907.7.	0.5	0
144	Mice lacking RGS protein activity at G β i2 exhibit a 5HT1A receptor-mediated antidepressant-like phenotype. <i>FASEB Journal</i> , 2008, 22, 907.8.	0.5	0

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