

Nigel J Pyne

List of Publications by Year in descending order

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182
papers

9,629
citations

36303

51
h-index

45317

90
g-index

189
all docs

189
docs citations

189
times ranked

8458
citing authors

#	ARTICLE	IF	CITATIONS
1	Validation of highly selective sphingosine kinase 2 inhibitors SLM6031434 and HWG-35D as effective anti-fibrotic treatment options in a mouse model of tubulointerstitial fibrosis. <i>Cellular Signalling</i> , 2021, 79, 109881.	3.6	7
2	A new model for regulation of sphingosine kinase 1 translocation to the plasma membrane in breast cancer cells. <i>Journal of Biological Chemistry</i> , 2021, 296, 100674.	3.4	2
3	Dihydroceramide Desaturase Functions as an Inducer and Rectifier of Apoptosis: Effect of Retinol Derivatives, Antioxidants and Phenolic Compounds. <i>Cell Biochemistry and Biophysics</i> , 2021, 79, 461-475.	1.8	6
4	Interleukin-7 receptor $\hat{\pm}$ mutational activation can initiate precursor B-cell acute lymphoblastic leukemia. <i>Nature Communications</i> , 2021, 12, 7268.	12.8	24
5	Structure-function analysis of lipid substrates and inhibitors of sphingosine kinases. <i>Cellular Signalling</i> , 2020, 76, 109806.	3.6	10
6	A Novel Selective Sphingosine Kinase 2 Inhibitor, HWG-35D, Ameliorates the Severity of Imiquimod-Induced Psoriasis Model by Blocking Th17 Differentiation of Na ⁺ ve CD4 T Lymphocytes. <i>International Journal of Molecular Sciences</i> , 2020, 21, 8371.	4.1	12
7	Recent advances in the role of sphingosine 1-phosphate in cancer. <i>FEBS Letters</i> , 2020, 594, 3583-3601.	2.8	35
8	The regulation of p53, p38 MAPK, JNK and XBP-1s by sphingosine kinases in human embryonic kidney cells. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2020, 1865, 158631.	2.4	1
9	Sphingosine Kinase 1 Regulates the Survival of Breast Cancer Stem Cells and Non-stem Breast Cancer Cells by Suppression of STAT1. <i>Cells</i> , 2020, 9, 886.	4.1	23
10	THE CONCISE GUIDE TO PHARMACOLOGY 2019/20: Enzymes. <i>British Journal of Pharmacology</i> , 2019, 176, S297-S396.	5.4	423
11	Small-molecule allosteric activators of PDE4 long form cyclic AMP phosphodiesterases. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 13320-13329.	7.1	54
12	Topographical Mapping of Isoform-Selectivity Determinants for J-Channel-Binding Inhibitors of Sphingosine Kinases 1 and 2. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 3658-3676.	6.4	23
13	Ceramide and sphingosine 1-phosphate in adipose dysfunction. <i>Progress in Lipid Research</i> , 2019, 74, 145-159.	11.6	30
14	Short Periods of Hypoxia Upregulate Sphingosine Kinase 1 and Increase Vasodilation of Arteries to Sphingosine 1-Phosphate (S1P) via S1P ₃ . <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2019, 371, 63-74.	2.5	8
15	Requirement for sphingosine kinase 1 in mediating phase 1 of the hypotensive response to anandamide in the anaesthetised mouse. <i>European Journal of Pharmacology</i> , 2019, 842, 1-9.	3.5	9
16	Sphingosine Kinases as Druggable Targets. <i>Handbook of Experimental Pharmacology</i> , 2018, 259, 49-76.	1.8	12
17	Cellular Signalling – Special issue to celebrate 75th birthday of Prof. Robert J. Lefkowitz. <i>Cellular Signalling</i> , 2018, 41, 1.	3.6	1
18	Sphingosine 1-phosphate and cancer. <i>Advances in Biological Regulation</i> , 2018, 68, 97-106.	2.3	82

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19	Does the Sphingosine 1-Phosphate Receptor-1 Provide a Better or Worse Prognostic Outcome for Breast Cancer Patients?. <i>Frontiers in Oncology</i> , 2018, 8, 417.	2.8	0
20	Native and Polyubiquitinated Forms of Dihydroceramide Desaturase Are Differentially Linked to Human Embryonic Kidney Cell Survival. <i>Molecular and Cellular Biology</i> , 2018, 38, .	2.3	16
21	The sphingosine 1-phosphate receptor 2 is shed in exosomes from breast cancer cells and is N-terminally processed to a short constitutively active form that promotes extracellular signal regulated kinase activation and DNA synthesis in fibroblasts. <i>Oncotarget</i> , 2018, 9, 29453-29467.	1.8	27
22	Sphingosine Kinase 2 in Autoimmune/Inflammatory Disease and the Development of Sphingosine Kinase 2 Inhibitors. <i>Trends in Pharmacological Sciences</i> , 2017, 38, 581-591.	8.7	34
23	Sphingosine Kinase 1: A Potential Therapeutic Target in Pulmonary Arterial Hypertension?. <i>Trends in Molecular Medicine</i> , 2017, 23, 786-798.	6.7	23
24	Effect of sphingosine kinase modulators on interleukin-1 β release, sphingosine 1-phosphate receptor 1 expression and experimental autoimmune encephalomyelitis. <i>British Journal of Pharmacology</i> , 2017, 174, 210-222.	5.4	8
25	Sphingosine 1-Phosphate Receptor 1 Signaling in Mammalian Cells. <i>Molecules</i> , 2017, 22, 344.	3.8	64
26	Sphingosine kinase 2 and multiple myeloma. <i>Oncotarget</i> , 2017, 8, 43596-43597.	1.8	3
27	A reflection of the lasting contributions from Dr. Robert Bittman to sterol trafficking, sphingolipid and phospholipid research. <i>Progress in Lipid Research</i> , 2016, 61, 19-29.	11.6	0
28	Effect of the sphingosine kinase 1 selective inhibitor, PF-543 on arterial and cardiac remodelling in a hypoxic model of pulmonary arterial hypertension. <i>Cellular Signalling</i> , 2016, 28, 946-955.	3.6	37
29	Therapeutic potential of targeting sphingosine kinases and sphingosine 1-phosphate in hematological malignancies. <i>Leukemia</i> , 2016, 30, 2142-2151.	7.2	34
30	Sphingosine Kinases: Emerging Structure-Function Insights. <i>Trends in Biochemical Sciences</i> , 2016, 41, 395-409.	7.5	62
31	Sphingosine 1-phosphate and sphingosine kinases in health and disease: Recent advances. <i>Progress in Lipid Research</i> , 2016, 62, 93-106.	11.6	153
32	Effect of ether glycerol lipids on interleukin-1 β release and experimental autoimmune encephalomyelitis. <i>Chemistry and Physics of Lipids</i> , 2016, 194, 2-11.	3.2	4
33	Role of sphingosine 1-phosphate receptors, sphingosine kinases and sphingosine in cancer and inflammation. <i>Advances in Biological Regulation</i> , 2016, 60, 151-159.	2.3	119
34	Proteasomal degradation of sphingosine kinase 1 and inhibition of dihydroceramide desaturase by the sphingosine kinase inhibitors, SKI or ABC294640, induces growth arrest in androgen-independent LNCaP-AI prostate cancer cells. <i>Oncotarget</i> , 2016, 7, 16663-16675.	1.8	66
35	The life and work of Dr. Robert Bittman (1942-2014). <i>Biological Chemistry</i> , 2015, 396, 827-830.	2.5	0
36	Resveratrol and its oligomers: modulation of sphingolipid metabolism and signaling in disease. <i>Archives of Toxicology</i> , 2014, 88, 2213-2232.	4.2	16

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37	Sphingosine kinase 1 enables communication between melanoma cells and fibroblasts that provides a new link to metastasis. <i>Oncogene</i> , 2014, 33, 3361-3363.	5.9	9
38	The role of sphingosine 1-phosphate in inflammation and cancer. <i>Advances in Biological Regulation</i> , 2014, 54, 121-129.	2.3	44
39	Crystal Structure of Sphingosine Kinase 1 with PF-543. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 1329-1333.	2.8	90
40	Sphingosine kinase 2 prevents the nuclear translocation of sphingosine 1-phosphate receptor-2 and tyrosine 416 phosphorylated c-Src and increases estrogen receptor negative MDA-MB-231 breast cancer cell growth: The role of sphingosine 1-phosphate receptor-4. <i>Cellular Signalling</i> , 2014, 26, 1040-1047.	3.6	23
41	Assessment of the effect of sphingosine kinase inhibitors on apoptosis, unfolded protein response and autophagy of T-cell acute lymphoblastic leukemia cells; indications for novel therapeutics. <i>Oncotarget</i> , 2014, 5, 7886-7901.	1.8	36
42	Identification of novel functional and spatial associations between sphingosine kinase 1, sphingosine 1-phosphate receptors and other signaling proteins that affect prognostic outcome in estrogen receptor-positive breast cancer. <i>International Journal of Cancer</i> , 2013, 132, 605-616.	5.1	40
43	Sphingosine 1-Phosphate Is a Missing Link between Chronic Inflammation and Colon Cancer. <i>Cancer Cell</i> , 2013, 23, 5-7.	16.8	29
44	New Perspectives on the Role of Sphingosine 1-Phosphate in Cancer. <i>Handbook of Experimental Pharmacology</i> , 2013, , 55-71.	1.8	20
45	Structure-Activity Relationships and Molecular Modeling of Sphingosine Kinase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 9310-9327.	6.4	43
46	The sphingosine kinase inhibitor 2-(4-chlorophenyl)-4-hydroxy-5-oxo-1,4-dihydro-2H-pyran-3-ylthiothiazole reduces androgen receptor expression via an oxidative stress-dependent mechanism. <i>British Journal of Pharmacology</i> , 2013, 168, 1497-1505.	5.4	16
47	Novel sphingosine-containing analogues selectively inhibit sphingosine kinase (SK) isozymes, induce SK1 proteasomal degradation and reduce DNA synthesis in human pulmonary arterial smooth muscle cells. <i>MedChemComm</i> , 2013, 4, 1394.	3.4	53
48	The p.Arg86Gln change in GARP2 (glutamic acid-rich protein-2) is a common West African-related polymorphism. <i>Gene</i> , 2013, 515, 155-158.	2.2	11
49	Synthesis of selective inhibitors of sphingosine kinase 1. <i>Chemical Communications</i> , 2013, 49, 2136.	4.1	52
50	The roles of sphingosine kinases 1 and 2 in regulating the Warburg effect in prostate cancer cells. <i>Cellular Signalling</i> , 2013, 25, 1011-1017.	3.6	46
51	Synthesis of (S)-FTY720 vinylphosphonate analogues and evaluation of their potential as sphingosine kinase 1 inhibitors and activators. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 2503-2510.	3.0	23
52	Role of sphingosine 1-phosphate and lysophosphatidic acid in fibrosis. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2013, 1831, 228-238.	2.4	54
53	The Roles of Sphingosine Kinase 1 and 2 in Regulating the Metabolome and Survival of Prostate Cancer Cells. <i>Biomolecules</i> , 2013, 3, 316-333.	4.0	13
54	Regulation of cell survival by sphingosine-1-phosphate receptor S1P1 via reciprocal ERK-dependent suppression of Bim and PI-3-kinase/protein kinase C-mediated upregulation of Mcl-1. <i>Cell Death and Disease</i> , 2013, 4, e927-e927.	6.3	74

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55	Expression of sphingosine 1-phosphate receptor 4 and sphingosine kinase 1 is associated with outcome in oestrogen receptor-negative breast cancer. <i>British Journal of Cancer</i> , 2012, 106, 1453-1459.	6.4	59
56	Sphingosine 1-phosphate receptors and sphingosine kinase 1: novel biomarkers for clinical prognosis in breast, prostate, and hematological cancers. <i>Frontiers in Oncology</i> , 2012, 2, 168.	2.8	37
57	Resveratrol dimers are novel sphingosine kinase 1 inhibitors and affect sphingosine kinase 1 expression and cancer cell growth and survival. <i>British Journal of Pharmacology</i> , 2012, 166, 1605-1616.	5.4	54
58	Targeting sphingosine kinase 1 in cancer. <i>Advances in Biological Regulation</i> , 2012, 52, 31-38.	2.3	37
59	Inhibition kinetics and regulation of sphingosine kinase 1 expression in prostate cancer cells: Functional differences between sphingosine kinase 1a and 1b. <i>International Journal of Biochemistry and Cell Biology</i> , 2012, 44, 1457-1464.	2.8	36
60	Sphingosine 1-phosphate signalling in cancer. <i>Biochemical Society Transactions</i> , 2012, 40, 94-100.	3.4	109
61	Translational aspects of sphingosine 1-phosphate biology. <i>Trends in Molecular Medicine</i> , 2011, 17, 463-472.	6.7	66
62	Receptor tyrosine kinaseâ€“G-protein-coupled receptor signalling platforms: out of the shadow?. <i>Trends in Pharmacological Sciences</i> , 2011, 32, 443-450.	8.7	105
63	Selectivity and Specificity of Sphingosine 1-Phosphate Receptor Ligands: â€œOff-Targetsâ€•or Complex Pharmacology?. <i>Frontiers in Pharmacology</i> , 2011, 2, 26.	3.5	32
64	(R)-FTY720 methyl ether is a specific sphingosine kinase 2 inhibitor: Effect on sphingosine kinase 2 expression in HEK 293 cells and actin rearrangement and survival of MCF-7 breast cancer cells. <i>Cellular Signalling</i> , 2011, 23, 1590-1595.	3.6	95
65	Sphingosine Kinase Inhibitors and Cancer: Seeking the Golden Sword of Hercules. <i>Cancer Research</i> , 2011, 71, 6576-6582.	0.9	77
66	FTY720 Analogues as Sphingosine Kinase 1 Inhibitors. <i>Journal of Biological Chemistry</i> , 2011, 286, 18633-18640.	3.4	107
67	Intracellular S1P Generation Is Essential for S1P-Induced Motility of Human Lung Endothelial Cells: Role of Sphingosine Kinase 1 and S1P Lyase. <i>PLoS ONE</i> , 2011, 6, e16571.	2.5	49
68	Targeting Î²-cell cyclic 3â€²5â€²adenosine monophosphate for the development of novel drugs for treating type 2 diabetes mellitus. A review. <i>Journal of Pharmacy and Pharmacology</i> , 2010, 56, 1477-1492.	2.4	29
69	Cyclic AMP Signaling in Pancreatic Islets. <i>Advances in Experimental Medicine and Biology</i> , 2010, 654, 281-304.	1.6	89
70	FTY720 and (S)-FTY720 vinylphosphonate inhibit sphingosine kinase 1 and promote its proteasomal degradation in human pulmonary artery smooth muscle, breast cancer and androgen-independent prostate cancer cells. <i>Cellular Signalling</i> , 2010, 22, 1536-1542.	3.6	169
71	(S)-FTY720-Vinylphosphonate, an analogue of the immunosuppressive agent FTY720, is a pan-antagonist of sphingosine 1-phosphate GPCR signaling and inhibits autotaxin activity. <i>Cellular Signalling</i> , 2010, 22, 1543-1553.	3.6	50
72	The sphingosine kinase inhibitor N,N'-dimethylsphingosine inhibits neointimal hyperplasia. <i>British Journal of Pharmacology</i> , 2010, 159, 543-553.	5.4	12

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73	Interaction between anandamide and sphingosine-1-phosphate in mediating vasorelaxation in rat coronary artery. <i>British Journal of Pharmacology</i> , 2010, 161, 176-192.	5.4	30
74	Sphingosine Kinase 1 Induces Tolerance to Human Epidermal Growth Factor Receptor 2 and Prevents Formation of a Migratory Phenotype in Response to Sphingosine 1-Phosphate in Estrogen Receptor-Positive Breast Cancer Cells. <i>Molecular and Cellular Biology</i> , 2010, 30, 3827-3841.	2.3	94
75	Sphingosine 1-Phosphate Receptor 4 Uses HER2 (ERBB2) to Regulate Extracellular Signal Regulated Kinase-1/2 in MDA-MB-453 Breast Cancer Cells. <i>Journal of Biological Chemistry</i> , 2010, 285, 35957-35966.	3.4	72
76	Sphingosine 1-phosphate and cancer. <i>Nature Reviews Cancer</i> , 2010, 10, 489-503.	28.4	765
77	High Expression of Sphingosine 1-Phosphate Receptors, S1P1 and S1P3, Sphingosine Kinase 1, and Extracellular Signal-Regulated Kinase-1/2 Is Associated with Development of Tamoxifen Resistance in Estrogen Receptor-Positive Breast Cancer Patients. <i>American Journal of Pathology</i> , 2010, 177, 2205-2215.	3.8	156
78	The Sphingosine Kinase 1 Inhibitor 2-(p-Hydroxyanilino)-4-(p-chlorophenyl)thiazole Induces Proteasomal Degradation of Sphingosine Kinase 1 in Mammalian Cells*. <i>Journal of Biological Chemistry</i> , 2010, 285, 38841-38852.	3.4	106
79	Sphingosine 1-Phosphate Regulation of Extracellular Signal-Regulated Kinase-1/2 in Embryonic Stem Cells. <i>Stem Cells and Development</i> , 2009, 18, 1319-1330.	2.1	41
80	Role of sphingosine kinases and lipid phosphate phosphatases in regulating spatial sphingosine 1-phosphate signalling in health and disease. <i>Cellular Signalling</i> , 2009, 21, 14-21.	3.6	124
81	The sphingosine 1-phosphate receptor 5 and sphingosine kinases 1 and 2 are localised in centrosomes: Possible role in regulating cell division. <i>Cellular Signalling</i> , 2009, 21, 675-684.	3.6	30
82	The role of the PDE4D cAMP phosphodiesterase in the regulation of glucagon-like peptide-1 release. <i>British Journal of Pharmacology</i> , 2009, 157, 633-644.	5.4	50
83	New aspects of sphingosine 1-phosphate signaling in mammalian cells. <i>Advances in Enzyme Regulation</i> , 2009, 49, 214-221.	2.6	28
84	Targeting sphingosine-1-phosphate signalling for cardioprotection. <i>Current Opinion in Pharmacology</i> , 2009, 9, 194-201.	3.5	40
85	Sphingosine 1-phosphate, lysophosphatidic acid and growth factor signaling and termination. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2008, 1781, 467-476.	2.4	34
86	Protein Kinase C- μ Regulates Sphingosine 1-Phosphate-mediated Migration of Human Lung Endothelial Cells through Activation of Phospholipase D2, Protein Kinase C- η , and Rac1. <i>Journal of Biological Chemistry</i> , 2008, 283, 11794-11806.	3.4	51
87	Lipid phosphate phosphatases form homo- and hetero-oligomers: catalytic competency, subcellular distribution and function. <i>Biochemical Journal</i> , 2008, 411, 371-377.	3.7	23
88	Receptor tyrosine kinase-G-protein coupled receptor complex signaling in mammalian cells. <i>Advances in Enzyme Regulation</i> , 2007, 47, 271-280.	2.6	26
89	Lipid phosphate phosphatase-1 regulates lysophosphatidic acid- and platelet-derived-growth-factor-induced cell migration. <i>Biochemical Journal</i> , 2006, 394, 495-500.	3.7	29
90	Integrin signalling regulates the nuclear localization and function of the lysophosphatidic acid receptor-1 (LPA1) in mammalian cells. <i>Biochemical Journal</i> , 2006, 398, 55-62.	3.7	32

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91	Protean agonism of the lysophosphatidic acid receptor-1 with Ki16425 reduces nerve growth factor-induced neurite outgrowth in pheochromocytoma 12 cells. <i>Journal of Neurochemistry</i> , 2006, 98, 1920-1929.	3.9	24
92	The effect of RGS12 on PDGF β receptor signalling to p42/p44 mitogen activated protein kinase in mammalian cells. <i>Cellular Signalling</i> , 2006, 18, 971-981.	3.6	39
93	The effect of hypoxia on lipid phosphate receptor and sphingosine kinase expression and mitogen-activated protein kinase signaling in human pulmonary smooth muscle cells. <i>Prostaglandins and Other Lipid Mediators</i> , 2006, 79, 278-286.	1.9	47
94	The functional PDGF β receptor α 1 receptor signaling complex is involved in regulating migration of mouse embryonic fibroblasts in response to platelet derived growth factor. <i>Prostaglandins and Other Lipid Mediators</i> , 2006, 80, 74-80.	1.9	29
95	Cell migration activated by platelet-derived growth factor receptor is blocked by an inverse agonist of the sphingosine 1-phosphate receptor α 1. <i>FASEB Journal</i> , 2006, 20, 509-511.	0.5	77
96	Regulation of Lysophosphatidic Acid-induced Epidermal Growth Factor Receptor Transactivation and Interleukin-8 Secretion in Human Bronchial Epithelial Cells by Protein Kinase C δ , Lyn Kinase, and Matrix Metalloproteinases. <i>Journal of Biological Chemistry</i> , 2006, 281, 19501-19511.	3.4	91
97	Lipid phosphate phosphatases and lipid phosphate signalling. <i>Biochemical Society Transactions</i> , 2005, 33, 1370.	3.4	87
98	Regulation of cell survival by lipid phosphate phosphatases involves the modulation of intracellular phosphatidic acid and sphingosine 1-phosphate pools. <i>Biochemical Journal</i> , 2005, 391, 25-32.	3.7	68
99	c-Src is involved in regulating signal transmission from PDGF β receptor α 1-PCR(s) complexes in mammalian cells. <i>Cellular Signalling</i> , 2005, 17, 263-277.	3.6	77
100	Experimental Systems for Studying the Role of G-Protein-Coupled Receptors in Receptor Tyrosine Kinase Signal Transduction. <i>Methods in Enzymology</i> , 2004, 390, 451-475.	1.0	6
101	Ectopic Expression of Bovine Type 5 Phosphodiesterase Confers a Renal Phenotype in Drosophila. <i>Journal of Biological Chemistry</i> , 2004, 279, 8159-8168.	3.4	35
102	Nerve growth factor signaling involves interaction between the Trk A receptor and lysophosphatidate receptor 1 systems: nuclear translocation of the lysophosphatidate receptor 1 and Trk A receptors in pheochromocytoma 12 cells. <i>Cellular Signalling</i> , 2004, 16, 127-136.	3.6	75
103	The role of G-protein coupled receptors and associated proteins in receptor tyrosine kinase signal transduction. <i>Seminars in Cell and Developmental Biology</i> , 2004, 15, 309-323.	5.0	84
104	Cyclic nucleotide phosphodiesterases in pancreatic islets. <i>Diabetologia</i> , 2003, 46, 1179-1189.	6.3	75
105	Interaction of caspase-3 with the cyclic GMP binding cyclic GMP specific phosphodiesterase (PDE5a1). <i>FEBS Journal</i> , 2003, 270, 962-970.	0.2	11
106	An assessment of the role of the inhibitory gamma subunit of the retinal cyclic GMP phosphodiesterase and its effect on the p42/p44 mitogen-activated protein kinase pathway in animal and cellular models of pulmonary hypertension. <i>British Journal of Pharmacology</i> , 2003, 138, 1313-1319.	5.4	16
107	Sphingosine 1-Phosphate and Platelet-derived Growth Factor (PDGF) Act via PDGF β Receptor-Sphingosine 1-Phosphate Receptor Complexes in Airway Smooth Muscle Cells. <i>Journal of Biological Chemistry</i> , 2003, 278, 6282-6290.	3.4	131
108	Ryanodine receptors of pancreatic β cells mediate a distinct context-dependent signal for insulin secretion. <i>FASEB Journal</i> , 2003, 17, 301-303.	0.5	60

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109	The Inhibitory \hat{I}^3 Subunit of the Type 6 Retinal cGMP Phosphodiesterase Functions to Link c-Src and G-protein-coupled Receptor Kinase 2 in a Signaling Unit That Regulates p42/p44 Mitogen-activated Protein Kinase by Epidermal Growth Factor. <i>Journal of Biological Chemistry</i> , 2003, 278, 18658-18663.	3.4	32
110	Receptor tyrosine kinaseâ€“GPCR signal complexes. <i>Biochemical Society Transactions</i> , 2003, 31, 1220-1225.	3.4	53
111	The Identification of the Inhibitory \hat{I}^3 -Subunits of the Type 6 Retinal Cyclic Guanosine Monophosphate Phosphodiesterase in Non-retinal Tissues: Differential Processing of mRNA Transcripts. <i>Genomics</i> , 2002, 79, 582-586.	2.9	18
112	Sphingosine 1-phosphate signalling and termination at lipid phosphate receptors. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2002, 1582, 121-131.	2.4	81
113	Streptozotocin diabetes protects against arrhythmias in rat isolated hearts: role of hypothyroidism. <i>European Journal of Pharmacology</i> , 2002, 435, 269-276.	3.5	36
114	Increased expression of the cGMPâ€inhibited cAMPâ€specific (PDE3) and cGMP binding cGMPâ€specific (PDE5) phosphodiesterases in models of pulmonary hypertension. <i>British Journal of Pharmacology</i> , 2002, 137, 1187-1194.	5.4	118
115	Nerve Growth Factor Stimulation of p42/p44 Mitogen-Activated Protein Kinase in PC12 Cells: Role of G_{i/o}, G Protein-Coupled Receptor Kinase 2, \hat{I}^2 -Arrestin I, and Endocytic Processing. <i>Molecular Pharmacology</i> , 2001, 60, 63-70.	2.3	87
116	Attenuation of G-protein coupled receptor activated p42/p44 mitogen activated protein kinase by lipid phosphate phosphatases 1,1a and 2. <i>Biochemical Society Transactions</i> , 2001, 29, A118-A118.	3.4	0
117	Assessment of agonism at G-protein coupled receptors by phosphatidic acid and lysophosphatidic acid in human embryonic kidney 293 cells. <i>British Journal of Pharmacology</i> , 2001, 134, 6-9.	5.4	30
118	The \hat{I}^3 subunit of the rod photoreceptor cGMP phosphodiesterase can modulate the proteolysis of two cGMP binding cGMP-specific phosphodiesterases (PDE6 and PDE5) by caspase-3. <i>Cellular Signalling</i> , 2001, 13, 735-741.	3.6	11
119	The Inhibitory \hat{I}^3 Subunit of the Type 6 Retinal Cyclic Guanosine Monophosphate Phosphodiesterase Is a Novel Intermediate Regulating p42/p44 Mitogen-activated Protein Kinase Signaling in Human Embryonic Kidney 293 Cells. <i>Journal of Biological Chemistry</i> , 2001, 276, 37802-37808.	3.4	31
120	G-protein-coupled Receptor Stimulation of the p42/p44 Mitogen-activated Protein Kinase Pathway Is Attenuated by Lipid Phosphate Phosphatases 1, 1a, and 2 in Human Embryonic Kidney 293 Cells. <i>Journal of Biological Chemistry</i> , 2001, 276, 13452-13460.	3.4	88
121	Short-Term Local Delivery of an Inhibitor of Ras Farnesyltransferase Prevents Neointima Formation In Vivo After Porcine Coronary Balloon Angioplasty. <i>Circulation</i> , 2001, 104, 1538-1543.	1.6	43
122	Tethering of the Platelet-derived Growth Factor \hat{I}^2 Receptor to G-protein-coupled Receptors. <i>Journal of Biological Chemistry</i> , 2001, 276, 28578-28585.	3.4	147
123	Sphingosine 1-phosphate signalling in mammalian cells. <i>Biochemical Journal</i> , 2000, 349, 385.	3.7	464
124	Sphingosine 1-phosphate signalling in mammalian cells. <i>Biochemical Journal</i> , 2000, 349, 385-402.	3.7	637
125	Effect of type-selective inhibitors on cyclic nucleotide phosphodiesterase activity and insulin secretion in the clonal insulin secreting cell line BRIN-BD11. <i>British Journal of Pharmacology</i> , 2000, 129, 1228-1234.	5.4	33
126	The role of the cyclic GMP-inhibited cyclic AMP-specific phosphodiesterase (PDE3) in regulating clonal BRIN-BD11 insulin secreting cell survival. <i>Cellular Signalling</i> , 2000, 12, 541-548.	3.6	14

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127	Ceramide-dependent regulation of p42/p44 mitogen-activated protein kinase and c-Jun N-terminal-directed protein kinase in cultured airway smooth muscle cells. <i>Cellular Signalling</i> , 2000, 12, 737-743.	3.6	23
128	Sphingosine 1-phosphate signalling via the endothelial differentiation gene family of G-protein-coupled receptors. , 2000, 88, 115-131.		169
129	The cAMP-specific Phosphodiesterase PDE4A5 Is Cleaved Downstream of Its SH3 Interaction Domain by Caspase-3. <i>Journal of Biological Chemistry</i> , 2000, 275, 28063-28074.	3.4	45
130	The Platelet-Derived Growth Factor Receptor Stimulation of p42/p44 Mitogen-Activated Protein Kinase in Airway Smooth Muscle Involves a G-Protein-Mediated Tyrosine Phosphorylation of Gab1. <i>Molecular Pharmacology</i> , 2000, 58, 413-420.	2.3	43
131	Assessment of the Extracellular and Intracellular Actions of Sphingosine 1-phosphate by Using the p42/p44 Mitogen-Activated Protein Kinase Cascade as a Model. <i>Cellular Signalling</i> , 1999, 11, 349-354.	3.6	32
132	Sphingosine 1-phosphate stimulation of the p42/p44 mitogen-activated protein kinase pathway in airway smooth muscle. <i>Biochemical Journal</i> , 1999, 338, 643.	3.7	42
133	Extracellular actions of sphingosine 1-phosphate through endothelial differentiation gene products in mammalian cells: role in regulating proliferation and apoptosis. <i>Biochemical Society Transactions</i> , 1999, 27, 404-409.	3.4	14
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