

William L Blalock

List of Publications by Year in descending order

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67
papers

4,105
citations

230014

27
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134545

62
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69
all docs

69
docs citations

69
times ranked

5999
citing authors

#	ARTICLE	IF	CITATIONS
1	Combined Treatment with PI3K Inhibitors BYL-719 and CAL-101 Is a Promising Antiproliferative Strategy in Human Rhabdomyosarcoma Cells. <i>Molecules</i> , 2022, 27, 2742.	1.7	3
2	Innate Immunity: A Balance between Disease and Adaption to Stress. <i>Biomolecules</i> , 2022, 12, 737.	1.8	6
3	Ectopic Expression of Ankr2 Affects Proliferation, Motility and Clonogenic Potential of Human Osteosarcoma Cells. <i>Cancers</i> , 2021, 13, 174.	1.7	6
4	Opposing forces fight over the same ground to regulate interferon signaling. <i>Biochemical Journal</i> , 2021, 478, 1853-1859.	1.7	4
5	Revisiting the Role of GSK3, A Modulator of Innate Immunity, in Idiopathic Inclusion Body Myositis. <i>Cells</i> , 2021, 10, 3255.	1.8	6
6	Expression of the double-stranded RNA-dependent kinase PKR influences osteosarcoma attachment independent growth, migration, and invasion. <i>Journal of Cellular Physiology</i> , 2020, 235, 1103-1119.	2.0	4
7	Cancer therapy and treatments during COVID-19 era. <i>Advances in Biological Regulation</i> , 2020, 77, 100739.	1.4	30
8	Glycogen synthase kinase (GSK)-3 and the double-strand RNA-dependent kinase, PKR: When two kinases for the common good turn bad. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2020, 1867, 118769.	1.9	10
9	Signal Transduction in Ribosome Biogenesis: A Recipe to Avoid Disaster. <i>International Journal of Molecular Sciences</i> , 2019, 20, 2718.	1.8	69
10	AKT-dependent phosphorylation of the adenosine deaminases ADAR1 and 2 inhibits deaminase activity. <i>FASEB Journal</i> , 2019, 33, 9044-9061.	0.2	20
11	Therapeutic potential of nvp620 in human osteosarcomas cells. <i>Journal of Cellular Physiology</i> , 2019, 234, 10907-10917.	2.0	16
12	MIRNA-210: A Current Overview. <i>Anticancer Research</i> , 2017, 37, 6511-6521.	0.5	159
13	Cell Cycle Arrest and Apoptosis Induced by Kinamycin F in Human Osteosarcoma Cells. <i>Anticancer Research</i> , 2017, 37, 4103-4109.	0.5	5
14	BMP2 Induced Expression of PLC1 That is a Positive Regulator of Osteoblast Differentiation. <i>Journal of Cellular Physiology</i> , 2016, 231, 623-629.	2.0	26
15	Intolerant contact lens wearers exhibit ocular surface impairment despite 3 months wear discontinuation. <i>Graefe's Archive for Clinical and Experimental Ophthalmology</i> , 2016, 254, 1825-1831.	1.0	15
16	PLC1b affects Akt activation, cyclin E expression, and caspase cleavage, promoting cell survival in pro-B lymphoblastic cells exposed to oxidative stress. <i>FASEB Journal</i> , 2015, 29, 1383-1394.	0.2	10
17	Prohibitin 2: At a communications crossroads. <i>IUBMB Life</i> , 2015, 67, 239-254.	1.5	136
18	PLC1a and PLC1b Selective Regulation and Cyclin D3 Modulation Reduced by Kinamycin F During K562 Cell Differentiation. <i>Journal of Cellular Physiology</i> , 2015, 230, 587-594.	2.0	11

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19	Prohibitin 2 represents a novel nuclear AKT substrate during all- <i>trans</i> retinoic acid-induced differentiation of acute promyelocytic leukemia cells. <i>FASEB Journal</i> , 2014, 28, 2009-2019.	0.2	28
20	Identification of the PKR Nuclear Interactome Reveals Roles in Ribosome Biogenesis, mRNA Processing and Cell Division. <i>Journal of Cellular Physiology</i> , 2014, 229, 1047-1060.	2.0	23
21	Nuclear PI-PLC β 1: An appraisal on targets and pathology. <i>Advances in Biological Regulation</i> , 2014, 54, 2-11.	1.4	32
22	PLC-beta 1 regulates the expression of miR-210 during mithramycin-mediated erythroid differentiation in K562 cells. <i>Oncotarget</i> , 2014, 5, 4222-4231.	0.8	19
23	Phosphoinositide-specific Phospholipase C β 1b (PI-PLC β 1b) Interactome: Affinity Purification-Mass Spectrometry Analysis of PI-PLC β 1b with Nuclear Protein. <i>Molecular and Cellular Proteomics</i> , 2013, 12, 2220-2235.	2.5	21
24	A role for PLC β 1 in myotonic dystrophies type 1 and 2. <i>FASEB Journal</i> , 2012, 26, 3042-3048.	0.2	24
25	A rapid standardized quantitative microfluidic system approach for evaluating human tear proteins. <i>Molecular Vision</i> , 2012, 18, 2526-37.	1.1	15
26	Multiple forms of PKR present in the nuclei of acute leukemia cells represent an active kinase that is responsive to stress. <i>Leukemia</i> , 2011, 25, 236-245.	3.3	40
27	A role for PKR in hematologic malignancies. <i>Journal of Cellular Physiology</i> , 2010, 223, 572-591.	2.0	21
28	Tear proteomics in evaporative dry eye disease. <i>Eye</i> , 2010, 24, 1396-1402.	1.1	130
29	eEF1A Phosphorylation in the Nucleus of Insulin-stimulated C2C12 Myoblasts. <i>Molecular and Cellular Proteomics</i> , 2010, 9, 2719-2728.	2.5	26
30	A novel human fibronectin cryptic sequence unmasked by the insertion of the angiogenesis-associated extra type III domain B. <i>International Journal of Cancer</i> , 2009, 125, 751-758.	2.3	27
31	PKR activity is required for acute leukemic cell maintenance and growth: A role for PKR-mediated phosphatase activity to regulate GSK β phosphorylation. <i>Journal of Cellular Physiology</i> , 2009, 221, 232-241.	2.0	29
32	Use of Uteroglobin for the Engineering of Polyvalent, Polyspecific Fusion Proteins. <i>Journal of Biological Chemistry</i> , 2009, 284, 26646-26654.	1.6	6
33	PKR is activated in MDS patients and its subcellular localization depends on disease severity. <i>Leukemia</i> , 2008, 22, 2267-2269.	3.3	22
34	PI-PLC β 1 and activated Akt levels are linked to azacitidine responsiveness in high-risk myelodysplastic syndromes. <i>Leukemia</i> , 2008, 22, 198-200.	3.3	39
35	RAX is required for fly neuronal development and mouse embryogenesis. <i>Mechanisms of Development</i> , 2008, 125, 777-785.	1.7	24
36	Proapoptotic Activity and Chemosensitizing Effect of the Novel Akt Inhibitor (2S)-1-(1H-Indol-3-yl)-3-[5-(3-methyl-2H-indazol-5-yl)pyridin-3-yl]oxypropan-2-amine (A443654) in T-Cell Acute Lymphoblastic Leukemia. <i>Molecular Pharmacology</i> , 2008, 74, 884-895.	1.0	33

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37	Targeting the Phosphatidylinositol 3-Kinase/Akt/Mammalian Target of Rapamycin Module for Acute Myelogenous Leukemia Therapy: From Bench to Bedside. <i>Current Medicinal Chemistry</i> , 2007, 14, 2009-2023.	1.2	116
38	The AKT Inhibitor, A443654, Induces Cell Cycle Arrest, Apoptosis and Synergizes with Chemotherapeutic Drugs in Multi-Drug Resistant T-Cell Acute Lymphoblastic Leukemia - A Novel Agent for Therapy of Drug Resistant ALL. <i>Blood</i> , 2007, 110, 3344-3344.	0.6	1
39	RAX, the PKR activator, sensitizes cells to inflammatory cytokines, serum withdrawal, chemotherapy, and viral infection. <i>Blood</i> , 2006, 108, 821-829.	0.6	74
40	Serine 18 Phosphorylation of RAX, the PKR Activator, Is Required for PKR Activation and Consequent Translation Inhibition. <i>Journal of Biological Chemistry</i> , 2004, 279, 42687-42693.	1.6	53
41	Ability of the Activated PI3K/Akt Oncoproteins to Synergize with MEK1 and Induce Cell Cycle Progression and Abrogate the Cytokine-Dependence of Hematopoietic Cells. <i>Cell Cycle</i> , 2004, 3, 501-510.	1.3	17
42	Ability of the activated PI3K/Akt oncoproteins to synergize with MEK1 and induce cell cycle progression and abrogate the cytokine-dependence of hematopoietic cells. <i>Cell Cycle</i> , 2004, 3, 503-12.	1.3	21
43	Elucidation of Signal Transduction Pathways by Retroviral Infection of Cells with Modified Oncogenes. , 2003, 218, 221-252.		0
44	Fibroblastic, Hematopoietic, and Hormone Responsive Epithelial Cell Lines and Culture Conditions for Elucidation of Signal Transduction and Drug Resistance Pathways by Gene Transfer. , 2003, 218, 185-202.		0
45	Elucidation of Signal Transduction Pathways by Transfection of Cells with Modified Oncogenes. , 2003, 218, 203-220.		0
46	Involvement of PI3K/Akt pathway in cell cycle progression, apoptosis, and neoplastic transformation: a target for cancer chemotherapy. <i>Leukemia</i> , 2003, 17, 590-603.	3.3	1,055
47	Requirement for the PI3K/Akt pathway in MEK1-mediated growth and prevention of apoptosis: identification of an Achilles heel in leukemia. <i>Leukemia</i> , 2003, 17, 1058-1067.	3.3	59
48	Signal transduction mediated by the Ras/Raf/MEK/ERK pathway from cytokine receptors to transcription factors: potential targeting for therapeutic intervention. <i>Leukemia</i> , 2003, 17, 1263-1293.	3.3	632
49	Differential effects of kinase cascade inhibitors on neoplastic and cytokine-mediated cell proliferation. <i>Leukemia</i> , 2003, 17, 1765-1782.	3.3	55
50	Effects of the RAF/MEK/ERK and PI3K/AKT signal transduction pathways on the abrogation of cytokine-dependence and prevention of apoptosis in hematopoietic cells. <i>Oncogene</i> , 2003, 22, 2478-2492.	2.6	95
51	Enhanced ability of the progenipoiectin-1 to suppress apoptosis in human hematopoietic cells. <i>International Journal of Molecular Medicine</i> , 2002, 10, 385.	1.8	2
52	Enhanced ability of the progenipoiectin-1 to suppress apoptosis in human hematopoietic cells. <i>International Journal of Molecular Medicine</i> , 2002, 10, 385-94.	1.8	1
53	Synergistic effects of pi3k/akt on abrogation of cytokine-dependency induced by oncogenic raf. <i>Advances in Enzyme Regulation</i> , 2001, 41, 289-323.	2.9	22
54	Effects of inducible MEK1 activation on the cytokine dependency of lymphoid cells. <i>Leukemia</i> , 2001, 15, 794-807.	3.3	41

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55	Enhanced ability of daniplestim and myelopoietin-1 to suppress apoptosis in human hematopoietic cells. <i>Leukemia</i> , 2001, 15, 1203-1216.	3.3	15
56	Ceramide Regulates Protein Synthesis by a Novel Mechanism Involving the Cellular PKR Activator RAX. <i>Journal of Biological Chemistry</i> , 2001, 276, 11754-11758.	1.6	74
57	Differential abilities of the Raf family of protein kinases to abrogate cytokine dependency and prevent apoptosis in murine hematopoietic cells by a MEK1-dependent mechanism. <i>Leukemia</i> , 2000, 14, 642-656.	3.3	98
58	Synergy between Raf and BCL2 in abrogating the cytokine dependency of hematopoietic cells. <i>Leukemia</i> , 2000, 14, 1060-1079.	3.3	47
59	Combined effects of aberrant MEK1 activity and BCL2 overexpression on relieving the cytokine dependency of human and murine hematopoietic cells. <i>Leukemia</i> , 2000, 14, 1080-1096.	3.3	48
60	Effects of deregulated Raf activation on integrin, cytokine-receptor expression and the induction of apoptosis in hematopoietic cells. <i>Leukemia</i> , 2000, 14, 1921-1938.	3.3	27
61	A conditionally-active form of MEK1 results in autocrine transformation of human and mouse hematopoietic cells. <i>Oncogene</i> , 2000, 19, 526-536.	2.6	76
62	Synergistic effects of akt on abrogation of cytokine-dependency induced by raf and mek. <i>Experimental Hematology</i> , 2000, 28, 38.	0.2	0
63	Enhanced ability of myelopoietins, dual receptor agonists for human IL-3 and g-csf receptors and the IL-3 receptor agonist, daniplestim, to suppress apoptosis and stimulate cytokine-inducible gene expression. <i>Experimental Hematology</i> , 2000, 28, 39.	0.2	0
64	The Raf signal transduction cascade as a target for chemotherapeutic intervention in growth factor-responsive tumors. , 2000, 88, 229-279.		86
65	Effects of deregulated RAF and MEK1 expression on the cytokine-dependency of hematopoietic cells. <i>Advances in Enzyme Regulation</i> , 2000, 40, 305-337.	2.9	24
66	Signal transduction, cell cycle regulatory, and anti-apoptotic pathways regulated by IL-3 in hematopoietic cells: possible sites for intervention with anti-neoplastic drugs. <i>Leukemia</i> , 1999, 13, 1109-1166.	3.3	161
67	Differential abilities of activated Raf oncoproteins to abrogate cytokine dependency, prevent apoptosis and induce autocrine growth factor synthesis in human hematopoietic cells. <i>Leukemia</i> , 1998, 12, 1903-1929.	3.3	109