

Constantine Mitsiades

List of Publications by Year in Descending Order

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Version: 2024-04-24

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

75
papers

9,715
citations

45
h-index

77
g-index

77
ext. papers

10,326
ext. citations

5.3
avg, IF

5.14
L-index

#	Paper	IF	Citations
75	A biphenyl inhibitor of eIF4E targeting an internal binding site enables the design of cell-permeable PROTAC-degraders. <i>European Journal of Medicinal Chemistry</i> , 2021 , 219, 113435	6.8	2
74	The power of proteasome inhibition in multiple myeloma. <i>Expert Review of Proteomics</i> , 2018 , 15, 1033-1052	10.52	25
73	Inference and Analysis of Gene Regulatory Networks in R 2016 , 289-306		0
72	Identification of Wee1 as a novel therapeutic target for mutant RAS-driven acute leukemia and other malignancies. <i>Leukemia</i> , 2015 , 29, 27-37	10.7	43
71	Upregulation of IGF1R by mutant RAS in leukemia and potentiation of RAS signaling inhibitors by small-molecule inhibition of IGF1R. <i>Clinical Cancer Research</i> , 2014 , 20, 5483-95	12.9	12
70	Pomalidomide for the treatment of relapsed and refractory multiple myeloma. <i>Expert Opinion on Orphan Drugs</i> , 2014 , 2, 1089-1108	1.1	
69	Novel Agents in Multiple Myeloma 2013 , 215-228		
68	New proteasome inhibitors in myeloma. <i>Current Hematologic Malignancy Reports</i> , 2012 , 7, 258-66	4.4	74
67	Lenalidomide for the treatment of relapsed and refractory multiple myeloma. <i>Cancer Management and Research</i> , 2012 , 4, 253-68	3.6	26
66	A novel panel of protein biomarkers for predicting response to thalidomide-based therapy in newly diagnosed multiple myeloma patients. <i>Proteomics</i> , 2011 , 11, 1391-402	4.8	26
65	A proto-oncogene BCL6 is up-regulated in the bone marrow microenvironment in multiple myeloma cells. <i>Blood</i> , 2010 , 115, 3772-5	2.2	52
64	Complications of multiple myeloma therapy, part 1: risk reduction and management of peripheral neuropathy and asthenia. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2010 , 8 Suppl 1, S4-S12	7.3	26
63	Lenalidomide in multiple myeloma: an evidence-based review of its role in therapy. <i>Core Evidence</i> , 2010 , 4, 215-45	4.9	16
62	Single-agent bortezomib in previously untreated multiple myeloma: efficacy, characterization of peripheral neuropathy, and molecular correlations with response and neuropathy. <i>Journal of Clinical Oncology</i> , 2009 , 27, 3518-25	2.2	213
61	Bortezomib induces canonical nuclear factor-kappaB activation in multiple myeloma cells. <i>Blood</i> , 2009 , 114, 1046-52	2.2	285
60	Functional interaction of plasmacytoid dendritic cells with multiple myeloma cells: a therapeutic target. <i>Cancer Cell</i> , 2009 , 16, 309-23	24.3	197
59	Novel Combination Therapies for the Treatment of Relapsed/Refractory Multiple Myeloma: Current Phase I/II Combinations. <i>Clinical Lymphoma and Myeloma</i> , 2009 , 9, S40-S42		

58	Anti-DKK1 mAb (BHQ880) as a potential therapeutic agent for multiple myeloma. <i>Blood</i> , 2009 , 114, 371-2.	2.2	331
57	Bortezomib in the front-line treatment of multiple myeloma. <i>Expert Review of Anticancer Therapy</i> , 2008 , 8, 1053-72	3.5	73
56	Phase I trial of oral vorinostat (suberoylanilide hydroxamic acid, SAHA) in patients with advanced multiple myeloma. <i>Leukemia and Lymphoma</i> , 2008 , 49, 502-7	1.9	174
55	Lenalidomide plus dexamethasone is efficacious in patients with relapsed or refractory multiple myeloma. <i>Nature Clinical Practice Oncology</i> , 2008 , 5, 374-5		1
54	TH17 Pathway Promotes Tumor Cell Growth and Suppresses Immune Function in Myeloma: Potential for Therapeutic Application. <i>Blood</i> , 2008 , 112, 2737-2737	2.2	
53	Understanding multiple myeloma pathogenesis in the bone marrow to identify new therapeutic targets. <i>Nature Reviews Cancer</i> , 2007 , 7, 585-98	31.3	699
52	Inhibition of Akt induces significant downregulation of survivin and cytotoxicity in human multiple myeloma cells. <i>British Journal of Haematology</i> , 2007 , 138, 783-91	4.5	94
51	Targeting the beta-catenin/TCF transcriptional complex in the treatment of multiple myeloma. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007 , 104, 7516-21	11.5	173
50	Emerging drugs in multiple myeloma. <i>Expert Opinion on Emerging Drugs</i> , 2007 , 12, 155-63	3.7	9
49	Potential of antileukemic therapies by Smac mimetic, LBW242: effects on mutant FLT3-expressing cells. <i>Molecular Cancer Therapeutics</i> , 2007 , 6, 1951-61	6.1	71
48	New drugs for myeloma. <i>Oncologist</i> , 2007 , 12, 664-89	5.7	138
47	The treatment of relapsed and refractory multiple myeloma. <i>Hematology American Society of Hematology Education Program</i> , 2007 , 2007, 317-23	3.1	45
46	Targeting mitochondrial factor Smac/DIABLO as therapy for multiple myeloma (MM). <i>Blood</i> , 2007 , 109, 1220-7	2.2	140
45	Gene expression profiling and correlation with outcome in clinical trials of the proteasome inhibitor bortezomib. <i>Blood</i> , 2007 , 109, 3177-88	2.2	314
44	The emerging role of novel therapies for the treatment of relapsed myeloma. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2007 , 5, 149-62	7.3	56
43	MLN120B, a novel I κ B kinase beta inhibitor, blocks multiple myeloma cell growth in vitro and in vivo. <i>Clinical Cancer Research</i> , 2006 , 12, 5887-94	12.9	118
42	Lenalidomide in multiple myeloma. <i>Expert Review of Anticancer Therapy</i> , 2006 , 6, 1165-73	3.5	42
41	Bortezomib: proteasome inhibition as an effective anticancer therapy. <i>Annual Review of Medicine</i> , 2006 , 57, 33-47	17.4	289

40	Perifosine, an oral bioactive novel alkylphospholipid, inhibits Akt and induces in vitro and in vivo cytotoxicity in human multiple myeloma cells. <i>Blood</i> , 2006 , 107, 4053-62	2.2	360
39	Beyond single-agent bortezomib: combination regimens in relapsed multiple myeloma. <i>Current Opinion in Oncology</i> , 2006 , 18, 598-608	4.2	29
38	Proteasome inhibition as a new therapeutic principle in hematological malignancies. <i>Current Drug Targets</i> , 2006 , 7, 1341-7	3	38
37	Gene expression analysis of B-lymphoma cells resistant and sensitive to bortezomib. <i>British Journal of Haematology</i> , 2006 , 134, 145-56	4.5	90
36	Bortezomib Induces Proliferation of Mesenchymal Progenitor Cells and Promotes Differentiation towards Osteoblastic Lineage.. <i>Blood</i> , 2006 , 108, 88-88	2.2	4
35	Proteasome Inhibition in the Treatment of Cancer. <i>Cell Cycle</i> , 2005 , 4, 289-295	4.7	77
34	Molecular characterization of PS-341 (bortezomib) resistance: implications for overcoming resistance using lysophosphatidic acid acyltransferase (LPAAT)-beta inhibitors. <i>Oncogene</i> , 2005 , 24, 3121-9	4.3	39
33	A novel orally active proteasome inhibitor induces apoptosis in multiple myeloma cells with mechanisms distinct from Bortezomib. <i>Cancer Cell</i> , 2005 , 8, 407-19	24.3	611
32	Proteasome inhibitor therapy in multiple myeloma. <i>Molecular Cancer Therapeutics</i> , 2005 , 4, 686-92	6.1	142
31	Bortezomib: proteasome inhibition as an effective anticancer therapy. <i>Future Oncology</i> , 2005 , 1, 161-71	3.6	30
30	Targeting Bcl-2 as Therapy for Multiple Myeloma.. <i>Blood</i> , 2005 , 106, 109-109	2.2	5
29	Requirement of Caspase-8 Versus Caspase-9 during Apoptosis in Multiple Myeloma Cells Induced by Bortezomib- or a Novel Proteasome Inhibitor NPI-0052.. <i>Blood</i> , 2005 , 106, 3378-3378	2.2	1
28	Proteasome inhibition in the treatment of cancer. <i>Cell Cycle</i> , 2005 , 4, 290-6	4.7	40
27	Blockade of ubiquitin-conjugating enzyme CDC34 enhances anti-myeloma activity of Bortezomib/Proteasome inhibitor PS-341. <i>Oncogene</i> , 2004 , 23, 3597-602	9.2	49
26	p38 MAPK inhibition enhances PS-341 (bortezomib)-induced cytotoxicity against multiple myeloma cells. <i>Oncogene</i> , 2004 , 23, 8766-76	9.2	112
25	Proteasome inhibition in hematologic malignancies. <i>Annals of Medicine</i> , 2004 , 36, 304-14	1.5	53
24	Thalidomide for patients with relapsed multiple myeloma after high-dose chemotherapy and stem cell transplantation: results of an open-label multicenter phase 2 study of efficacy, toxicity, and biological activity. <i>Mayo Clinic Proceedings</i> , 2004 , 79, 875-82	6.4	106
23	Identification of genes modulated in multiple myeloma using genetically identical twin samples. <i>Blood</i> , 2004 , 103, 1799-806	2.2	120

22	Immunomodulatory drug costimulates T cells via the B7-CD28 pathway. <i>Blood</i> , 2004 , 103, 1787-90	2.2	230
21	The bortezomib/proteasome inhibitor PS-341 and triterpenoid CDDO-Im induce synergistic anti-multiple myeloma (MM) activity and overcome bortezomib resistance. <i>Blood</i> , 2004 , 103, 3158-66	2.2	106
20	Targeting mitochondria to overcome conventional and bortezomib/proteasome inhibitor PS-341 resistance in multiple myeloma (MM) cells. <i>Blood</i> , 2004 , 104, 2458-66	2.2	70
19	Combination of the mTOR inhibitor rapamycin and CC-5013 has synergistic activity in multiple myeloma. <i>Blood</i> , 2004 , 104, 4188-93	2.2	167
18	A Novel Orally Available Proteasome Inhibitor NPI-0052 Induces Killing in Multiple Myeloma (MM) Cells Resistant to Conventional and Bortezomib Therapies.. <i>Blood</i> , 2004 , 104, 2405-2405	2.2	2
17	Comprehensive Genome-Wide Profile of Regional Gains and Losses in Multiple Myeloma Using Array-CGH: The 1q21 Amplification and Potential Role of the BCL-9 Gene in Multiple Myeloma Pathogenesis.. <i>Blood</i> , 2004 , 104, 785-785	2.2	4
16	The Akt pathway: molecular targets for anti-cancer drug development. <i>Current Cancer Drug Targets</i> , 2004 , 4, 235-56	2.8	215
15	Atiprimod (N-N-diethyl-8,8-dipropyl-2-azaspiro [4.5] decane-2-propanamine) Inhibits Myeloma in Vivo.. <i>Blood</i> , 2004 , 104, 2401-2401	2.2	
14	Increased TCF-4 Expression Correlates with Reduced Caspase-3 Induction and Confers Resistance to Bortezomib.. <i>Blood</i> , 2004 , 104, 285-285	2.2	1
13	Targeting Mitochondrial Factor Smac/DIABLO as Therapy for Multiple Myeloma (MM).. <i>Blood</i> , 2004 , 104, 764-764	2.2	
12	JNK-dependent release of mitochondrial protein, Smac, during apoptosis in multiple myeloma (MM) cells. <i>Journal of Biological Chemistry</i> , 2003 , 278, 17593-6	5.4	169
11	Molecular mechanisms mediating antimyeloma activity of proteasome inhibitor PS-341. <i>Blood</i> , 2003 , 101, 1530-4	2.2	488
10	Identification of genes regulated by 2-methoxyestradiol (2ME2) in multiple myeloma cells using oligonucleotide arrays. <i>Blood</i> , 2003 , 101, 3606-14	2.2	65
9	Hsp27 inhibits release of mitochondrial protein Smac in multiple myeloma cells and confers dexamethasone resistance. <i>Blood</i> , 2003 , 102, 3379-86	2.2	131
8	CD40 induces human multiple myeloma cell migration via phosphatidylinositol 3-kinase/AKT/NF-kappa B signaling. <i>Blood</i> , 2003 , 101, 2762-9	2.2	98
7	Proteasome inhibitor PS-341 abrogates IL-6 triggered signaling cascades via caspase-dependent downregulation of gp130 in multiple myeloma. <i>Oncogene</i> , 2003 , 22, 8386-93	9.2	151
6	Insights into the multistep transformation of MGUS to myeloma using microarray expression analysis. <i>Blood</i> , 2003 , 102, 4504-11	2.2	194
5	Insulin-like growth factor-1 induces adhesion and migration in human multiple myeloma cells via activation of beta1-integrin and phosphatidylinositol 3-kinase/AKT signaling. <i>Cancer Research</i> , 2003 , 63, 5850-8	10.1	138

4	Antitumor activity of lysophosphatidic acid acyltransferase-beta inhibitors, a novel class of agents, in multiple myeloma. <i>Cancer Research</i> , 2003 , 63, 8428-36	10.1	42
3	Immunomodulatory drug CC-5013 overcomes drug resistance and is well tolerated in patients with relapsed multiple myeloma. <i>Blood</i> , 2002 , 100, 3063-7	2.2	693
2	NF-kappa B as a therapeutic target in multiple myeloma. <i>Journal of Biological Chemistry</i> , 2002 , 277, 16639-47	9.4	723
1	Vascular endothelial growth factor triggers signaling cascades mediating multiple myeloma cell growth and migration. <i>Blood</i> , 2001 , 98, 428-35	2.2	358