

David C Huang

List of Publications by Year in descending order

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Version: 2024-02-01

234
papers

38,707
citations

3149

92
h-index

2736

192
g-index

241
all docs

241
docs citations

241
times ranked

33202
citing authors

#	ARTICLE	IF	CITATIONS
1	Clonal hematopoiesis, myeloid disorders and <i>BAX</i> -mutated myelopoiesis in patients receiving venetoclax for CLL. <i>Blood</i> , 2022, 139, 1198-1207.	0.6	34
2	Pharmacologic Reduction of Mitochondrial Iron Triggers a Noncanonical BAX/BAK-Dependent Cell Death. <i>Cancer Discovery</i> , 2022, 12, 774-791.	7.7	18
3	The Lck inhibitor, AMG-47a, blocks necroptosis and implicates RIPK1 in signalling downstream of MLKL. <i>Cell Death and Disease</i> , 2022, 13, 291.	2.7	10
4	Single-cell multiomics reveal the scale of multilayered adaptations enabling CLL relapse during venetoclax therapy. <i>Blood</i> , 2022, 140, 2127-2141.	0.6	28
5	The transcription factor IRF4 represses proapoptotic BMF and BIM to licence multiple myeloma survival. <i>Leukemia</i> , 2021, 35, 2114-2118.	3.3	18
6	Structure-Guided Development of Potent Benzoylurea Inhibitors of BCL-X _L and BCL-2. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 5447-5469.	2.9	5
7	PRMT1-mediated H4R3me2a recruits SMARCA4 to promote colorectal cancer progression by enhancing EGFR signaling. <i>Genome Medicine</i> , 2021, 13, 58.	3.6	62
8	Intact TP-53 function is essential for sustaining durable responses to BH3-mimetic drugs in leukemias. <i>Blood</i> , 2021, 137, 2721-2735.	0.6	75
9	TCF3 is epigenetically silenced by EZH2 and DNMT3B and functions as a tumor suppressor in endometrial cancer. <i>Cell Death and Differentiation</i> , 2021, 28, 3316-3328.	5.0	25
10	BCL2 and MCL1 inhibitors for hematologic malignancies. <i>Blood</i> , 2021, 138, 1120-1136.	0.6	78
11	Transcriptional silencing of fetal hemoglobin expression by NonO. <i>Nucleic Acids Research</i> , 2021, 49, 9711-9723.	6.5	7
12	Outcomes of patients with CLL sequentially resistant to both BCL2 and BTK inhibition. <i>Blood Advances</i> , 2021, 5, 4054-4058.	2.5	39
13	TNK1 is a ubiquitin-binding and 14-3-3-regulated kinase that can be targeted to block tumor growth. <i>Nature Communications</i> , 2021, 12, 5337.	5.8	14
14	Comprehensive characterization of single-cell full-length isoforms in human and mouse with long-read sequencing. <i>Genome Biology</i> , 2021, 22, 310.	3.8	83
15	Mesenchymal stromal cell apoptosis is required for their therapeutic function. <i>Nature Communications</i> , 2021, 12, 6495.	5.8	91
16	EBV BCL-2 homologue BHRF1 drives chemoresistance and lymphomagenesis by inhibiting multiple cellular pro-apoptotic proteins. <i>Cell Death and Differentiation</i> , 2020, 27, 1554-1568.	5.0	35
17	BH3 Mimetics for the Treatment of B-Cell Malignancies—Insights and Lessons from the Clinic. <i>Cancers</i> , 2020, 12, 3353.	1.7	12
18	Defining the susceptibility of colorectal cancers to BH3-mimetic compounds. <i>Cell Death and Disease</i> , 2020, 11, 735.	2.7	10

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19	Potent Inhibition of Necroptosis by Simultaneously Targeting Multiple Effectors of the Pathway. ACS Chemical Biology, 2020, 15, 2702-2713.	1.6	22
20	Cotargeting BCL-2 and MCL-1 in high-risk B-ALL. Blood Advances, 2020, 4, 2762-2767.	2.5	28
21	Loss of RIPK3 does not impact MYC-driven lymphomagenesis or chemotherapeutic drug-induced killing of malignant lymphoma cells. Cell Death and Differentiation, 2020, 27, 2531-2533.	5.0	6
22	MARCH5 requires MTCH2 to coordinate proteasomal turnover of the MCL1:NOXA complex. Cell Death and Differentiation, 2020, 27, 2484-2499.	5.0	33
23	Deep profiling of apoptotic pathways with mass cytometry identifies a synergistic drug combination for killing myeloma cells. Cell Death and Differentiation, 2020, 27, 2217-2233.	5.0	29
24	Molecular patterns of response and treatment failure after frontline venetoclax combinations in older patients with AML. Blood, 2020, 135, 791-803.	0.6	412
25	Multiple BCL2 mutations cooccurring with Gly101Val emerge in chronic lymphocytic leukemia progression on venetoclax. Blood, 2020, 135, 773-777.	0.6	115
26	Acquired Mutations in BAX Confer Resistance to BH3 Mimetics in Acute Myeloid Leukemia. Blood, 2020, 136, 7-8.	0.6	13
27	BAX-Mutated Clonal Hematopoiesis in Patients on Long-Term Venetoclax for Relapsed/Refractory Chronic Lymphocytic Leukemia. Blood, 2020, 136, 9-10.	0.6	4
28	Replication stress induces mitotic death through parallel pathways regulated by WAPL and telomere deprotection. Nature Communications, 2019, 10, 4224.	5.8	38
29	A small molecule interacts with VDAC2 to block mouse BAK-driven apoptosis. Nature Chemical Biology, 2019, 15, 1057-1066.	3.9	30
30	Characterization of a novel venetoclax resistance mutation (BCL2 Phe104Ile) observed in follicular lymphoma. British Journal of Haematology, 2019, 186, e188-e191.	1.2	37
31	Structures of BCL-2 in complex with venetoclax reveal the molecular basis of resistance mutations. Nature Communications, 2019, 10, 2385.	5.8	139
32	Venetoclax for the treatment of mantle cell lymphoma. Annals of Lymphoma, 2019, 3, 4-4.	4.5	1
33	Multiple myeloma with 1q21 amplification is highly sensitive to MCL-1 targeting. Blood Advances, 2019, 3, 4202-4214.	2.5	60
34	Combining BH3-mimetics to target both BCL-2 and MCL1 has potent activity in pre-clinical models of acute myeloid leukemia. Leukemia, 2019, 33, 905-917.	3.3	126
35	Dynamic molecular monitoring reveals that SWI65-SNF mutations mediate resistance to ibrutinib plus venetoclax in mantle cell lymphoma. Nature Medicine, 2019, 25, 119-129.	15.2	147
36	Recipient BCL2 inhibition and NK cell ablation form part of a reduced intensity conditioning regime that improves allo-bone marrow transplantation outcomes. Cell Death and Differentiation, 2019, 26, 1516-1530.	5.0	10

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37	KRAS-enhanced macropinocytosis and reduced FcRn-mediated recycling sensitize pancreatic cancer to albumin-conjugated drugs. <i>Journal of Controlled Release</i> , 2019, 296, 40-53.	4.8	39
38	Acquisition of the Recurrent Gly101Val Mutation in BCL2 Confers Resistance to Venetoclax in Patients with Progressive Chronic Lymphocytic Leukemia. <i>Cancer Discovery</i> , 2019, 9, 342-353.	7.7	306
39	Detection of Multiple Recurrent Novel BCL2 Mutations Co-Occurring with BCL2 Gly101Val in Patients with Chronic Lymphocytic Leukemia on Long Term Venetoclax. <i>Blood</i> , 2019, 134, 171-171.	0.6	5
40	BTK Leu528Trp - a Potential Secondary Resistance Mechanism Specific for Patients with Chronic Lymphocytic Leukemia Treated with the Next Generation BTK Inhibitor Zanubrutinib. <i>Blood</i> , 2019, 134, 170-170.	0.6	33
41	Loss of IRF4 Results in Multiple Myeloma Cell Apoptosis through the Transcriptional up-Regulation of the BH3-Only Proteins Bim and BIM. <i>Blood</i> , 2019, 134, 3103-3103.	0.6	2
42	BAK/BAX-Mediated Apoptosis Is a Myc-Induced Roadblock to Reprogramming. <i>Stem Cell Reports</i> , 2018, 10, 331-338.	2.3	16
43	Enhancing venetoclax activity in acute myeloid leukemia by co-targeting MCL1. <i>Leukemia</i> , 2018, 32, 303-312.	3.3	123
44	The Mitochondrial Apoptotic Effectors BAX/BAK Activate Caspase-3 and -7 to Trigger NLRP3 Inflammasome and Caspase-8 Driven IL-1 β Activation. <i>Cell Reports</i> , 2018, 25, 2339-2353.e4.	2.9	164
45	VDAC2 enables BAX to mediate apoptosis and limit tumor development. <i>Nature Communications</i> , 2018, 9, 4976.	5.8	110
46	AMG 176, a Selective MCL1 Inhibitor, Is Effective in Hematologic Cancer Models Alone and in Combination with Established Therapies. <i>Cancer Discovery</i> , 2018, 8, 1582-1597.	7.7	310
47	CARM1-mediated methylation of protein arginine methyltransferase 5 represses human β -globin gene expression in erythroleukemia cells. <i>Journal of Biological Chemistry</i> , 2018, 293, 17454-17463.	1.6	20
48	Infection with flaviviruses requires BCLXL for cell survival. <i>PLoS Pathogens</i> , 2018, 14, e1007299.	2.1	28
49	IMiDs prime myeloma cells for daratumumab-mediated cytotoxicity through loss of Ikaros and Aiolos. <i>Blood</i> , 2018, 132, 2166-2178.	0.6	65
50	Venetoclax in Patients with Previously Treated Chronic Lymphocytic Leukemia. <i>Clinical Cancer Research</i> , 2017, 23, 4527-4533.	3.2	56
51	DR5 and caspase-8 are dispensable in ER stress-induced apoptosis. <i>Cell Death and Differentiation</i> , 2017, 24, 944-950.	5.0	65
52	Clinicopathological features and outcomes of progression of CLL on the BCL2 inhibitor venetoclax. <i>Blood</i> , 2017, 129, 3362-3370.	0.6	150
53	Essential role for Bim in mediating the apoptotic and antitumor activities of immunotoxins. <i>Oncogene</i> , 2017, 36, 4953-4962.	2.6	10
54	Anti-apoptotic proteins BCL-2, MCL-1 and A1 summate collectively to maintain survival of immune cell populations both in vitro and in vivo. <i>Cell Death and Differentiation</i> , 2017, 24, 878-888.	5.0	103

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55	NatD promotes lung cancer progression by preventing histone H4 serine phosphorylation to activate Slug expression. <i>Nature Communications</i> , 2017, 8, 928.	5.8	69
56	Synergistic action of the MCL-1 inhibitor S63845 with current therapies in preclinical models of triple-negative and HER2-amplified breast cancer. <i>Science Translational Medicine</i> , 2017, 9, .	5.8	148
57	Design, Synthesis, and Biological Activity of 1,2,3-Triazolobenzodiazepine BET Bromodomain Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2017, 8, 1298-1303.	1.3	23
58	Targeting BCL2 With BH3 Mimetics: Basic Science and Clinical Application of Venetoclax in Chronic Lymphocytic Leukemia and Related B Cell Malignancies. <i>Clinical Pharmacology and Therapeutics</i> , 2017, 101, 89-98.	2.3	107
59	Defining a therapeutic window for kinase inhibitors in leukemia to avoid neutropenia. <i>Oncotarget</i> , 2017, 8, 57948-57963.	0.8	4
60	Identification of an activation site in Bak and mitochondrial Bax triggered by antibodies. <i>Nature Communications</i> , 2016, 7, 11734.	5.8	50
61	The BCL2 selective inhibitor venetoclax induces rapid onset apoptosis of CLL cells in patients via a TP53-independent mechanism. <i>Blood</i> , 2016, 127, 3215-3224.	0.6	242
62	Hepatocyte growth factor renders BRAF mutant human melanoma cell lines resistant to PLX4032 by downregulating the pro-apoptotic BH3-only proteins PUMA and BIM. <i>Cell Death and Differentiation</i> , 2016, 23, 2054-2062.	5.0	24
63	Hierarchy for targeting prosurvival BCL2 family proteins in multiple myeloma: pivotal role of MCL1. <i>Blood</i> , 2016, 128, 1834-1844.	0.6	127
64	MCL-1 is required throughout B-cell development and its loss sensitizes specific B-cell subsets to inhibition of BCL-2 or BCL-XL. <i>Cell Death and Disease</i> , 2016, 7, e2345-e2345.	2.7	53
65	Venetoclax responses of pediatric ALL xenografts reveal sensitivity of MLL-rearranged leukemia. <i>Blood</i> , 2016, 128, 1382-1395.	0.6	148
66	Eliminating Legionella by inhibiting BCL-XL to induce macrophage apoptosis. <i>Nature Microbiology</i> , 2016, 1, 15034.	5.9	75
67	The MCL1 inhibitor S63845 is tolerable and effective in diverse cancer models. <i>Nature</i> , 2016, 538, 477-482.	13.7	830
68	BET inhibition represses miR17-92 to drive BIM-initiated apoptosis of normal and transformed hematopoietic cells. <i>Leukemia</i> , 2016, 30, 1531-1541.	3.3	29
69	Small molecules targeting Mcl-1: the search for a silver bullet in cancer therapy. <i>MedChemComm</i> , 2016, 7, 778-787.	3.5	16
70	HSP90 activity is required for MLKL oligomerisation and membrane translocation and the induction of necroptotic cell death. <i>Cell Death and Disease</i> , 2016, 7, e2051-e2051.	2.7	123
71	Therapeutic Response to Non-genotoxic Activation of p53 by Nutlin3a Is Driven by PUMA-Mediated Apoptosis in Lymphoma Cells. <i>Cell Reports</i> , 2016, 14, 1858-1866.	2.9	35
72	Current challenges and novel treatment strategies in double hit lymphomas. <i>Therapeutic Advances in Hematology</i> , 2016, 7, 52-64.	1.1	20

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73	Clinicopathological Features and Outcomes of Progression for Chronic Lymphocytic Leukaemia (CLL) Treated with the BCL2 Inhibitor Venetoclax. <i>Blood</i> , 2016, 128, 3223-3223.	0.6	2
74	The <i>BIM</i> deletion polymorphism: A paradigm of a permissive interaction between germline and acquired TKI resistance factors in chronic myeloid leukemia. <i>Oncotarget</i> , 2016, 7, 2721-2733.	0.8	16
75	Targeting the Pro-Survival BCL2 Proteins with BH3 Mimetic Compounds for Treating Multiple Myeloma. <i>Blood</i> , 2016, 128, 3293-3293.	0.6	0
76	The Role of BAX/BAK-Mediated Apoptosis for the Cytotoxic Action of Anti-Myeloma Agents. <i>Blood</i> , 2016, 128, 5706-5706.	0.6	0
77	A Chemical Screening Approach to Identify Novel Key Mediators of Erythroid Enucleation. <i>PLoS ONE</i> , 2015, 10, e0142655.	1.1	8
78	Prosurvival Bcl-2 family members reveal a distinct apoptotic identity between conventional and plasmacytoid dendritic cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 4044-4049.	3.3	43
79	BCL2 inhibition in double hit lymphoma. <i>Leukemia and Lymphoma</i> , 2015, 56, 1928-1929.	0.6	1
80	Bcl-2 Antagonists Kill Plasmacytoid Dendritic Cells From Lupus-Prone Mice and Dampen Interferon- γ Production. <i>Arthritis and Rheumatology</i> , 2015, 67, 797-808.	2.9	43
81	Exploiting selective BCL-2 family inhibitors to dissect cell survival dependencies and define improved strategies for cancer therapy. <i>Science Translational Medicine</i> , 2015, 7, 279ra40.	5.8	430
82	A transgenic mouse model to inducibly target prosurvival Bcl2 proteins with selective BH3 peptides in vivo. <i>Cell Death and Disease</i> , 2015, 6, e1679-e1679.	2.7	1
83	A RIPK2 inhibitor delays NOD signalling events yet prevents inflammatory cytokine production. <i>Nature Communications</i> , 2015, 6, 6442.	5.8	112
84	Systematic Screening Identifies Dual PI3K and mTOR Inhibition as a Conserved Therapeutic Vulnerability in Osteosarcoma. <i>Clinical Cancer Research</i> , 2015, 21, 3216-3229.	3.2	58
85	Autoreactive T cells induce necrosis and not BCL-2-regulated or death receptor-mediated apoptosis or RIPK3-dependent necroptosis of transplanted islets in a mouse model of type 1 diabetes. <i>Diabetologia</i> , 2015, 58, 140-148.	2.9	32
86	Apoptotic Caspases Suppress mtDNA-Induced STING-Mediated Type I IFN Production. <i>Cell</i> , 2014, 159, 1549-1562.	13.5	698
87	Simplified Silvestrol Analogues with Potent Cytotoxic Activity. <i>ChemMedChem</i> , 2014, 9, 1556-1566.	1.6	16
88	Targeting BCL2 for the Treatment of Lymphoid Malignancies. <i>Seminars in Hematology</i> , 2014, 51, 219-227.	1.8	130
89	Targeting of MCL-1 kills MYC-driven mouse and human lymphomas even when they bear mutations in <i>p53</i> . <i>Genes and Development</i> , 2014, 28, 58-70.	2.7	156
90	Prosurvival Bcl-2 family members affect autophagy only indirectly, by inhibiting Bax and Bak. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 8512-8517.	3.3	166

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91	Enhanced stability of Mcl1, a prosurvival Bcl2 relative, blunts stress-induced apoptosis, causes male sterility, and promotes tumorigenesis. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 261-266.	3.3	43
92	Both leukaemic and normal peripheral B lymphoid cells are highly sensitive to the selective pharmacological inhibition of prosurvival Bcl-2 with ABT-199. Leukemia, 2014, 28, 1207-1215.	3.3	79
93	Evaluation of functional groups as acetyl-lysine mimetics for BET bromodomain inhibition. MedChemComm, 2014, 5, 1834-1842.	3.5	24
94	Further Insights into the Effects of Pre-organizing the BimBH3 Helix. ACS Chemical Biology, 2014, 9, 838-839.	1.6	26
95	De-Novo Designed Library of Benzoylureas as Inhibitors of BCL-X _L : Synthesis, Structural and Biochemical Characterization. Journal of Medicinal Chemistry, 2014, 57, 1323-1343.	2.9	33
96	Structure-Guided Rescaffolding of Selective Antagonists of BCL-X _L . ACS Medicinal Chemistry Letters, 2014, 5, 662-667.	1.3	37
97	197. Cytokine, 2014, 70, 75-76.	1.4	0
98	A Biosensor of Src Family Kinase Conformation by Exposable Tetracysteine Useful for Cell-Based Screening. ACS Chemical Biology, 2014, 9, 1426-1431.	1.6	9
99	Eradication of Acute Myeloid Leukemia Is Enhanced By Combined Bcl-2 and Mcl-1 Targeting. Blood, 2014, 124, 988-988.	0.6	2
100	ABT-199 (GDC-0199) in relapsed/refractory (R/R) chronic lymphocytic leukemia (CLL) and small lymphocytic lymphoma (SLL): High complete- response rate and durable disease control.. Journal of Clinical Oncology, 2014, 32, 7015-7015.	0.8	42
101	Targeting acute myeloid leukemia by dual inhibition of PI3K signaling and Cdk9-mediated Mcl-1 transcription. Blood, 2013, 122, 738-748.	0.6	53
102	ABT-199, a potent and selective BCL-2 inhibitor, achieves antitumor activity while sparing platelets. Nature Medicine, 2013, 19, 202-208.	15.2	2,426
103	Synthesis of Biotinylated Episilvestrol: Highly Selective Targeting of the Translation Factors eIF4AII. Organic Letters, 2013, 15, 1406-1409.	2.4	49
104	Bax Crystal Structures Reveal How BH3 Domains Activate Bax and Nucleate Its Oligomerization to Induce Apoptosis. Cell, 2013, 152, 519-531.	13.5	491
105	Structure-guided design of a selective BCL-XL inhibitor. Nature Chemical Biology, 2013, 9, 390-397.	3.9	324
106	Discovery of Potent and Selective Benzothiazole Hydrazone Inhibitors of Bcl-X _L . Journal of Medicinal Chemistry, 2013, 56, 5514-5540.	2.9	60
107	Stabilizing the Pro-Apoptotic BimBH3 Helix (BimSAHB) Does Not Necessarily Enhance Affinity or Biological Activity. ACS Chemical Biology, 2013, 8, 297-302.	1.6	123
108	BH3 mimetic therapy: an emerging and promising approach to treating chronic lymphocytic leukemia. Leukemia and Lymphoma, 2013, 54, 909-911.	0.6	2

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109	MCMV-mediated Inhibition of the Pro-apoptotic Bak Protein Is Required for Optimal In Vivo Replication. <i>PLoS Pathogens</i> , 2013, 9, e1003192.	2.1	21
110	Proapoptotic Bak and Bax guard against fatal systemic and organ-specific autoimmune disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 2599-2604.	3.3	43
111	Variability of Inducible Expression across the Hematopoietic System of Tetracycline Transactivator Transgenic Mice. <i>PLoS ONE</i> , 2013, 8, e54009.	1.1	26
112	Selective Bcl-2 Inhibition With ABT-199 Is Highly Active Against Chronic Lymphocytic Leukemia (CLL) Irrespective Of TP53 Mutation Or Dysfunction. <i>Blood</i> , 2013, 122, 1304-1304.	0.6	10
113	Abstract A19: The selective targeting of cell survival pathways in leukemia. , 2013, , .		0
114	Sensitization of BCL-2-expressing breast tumors to chemotherapy by the BH3 mimetic ABT-737. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 2766-2771.	3.3	173
115	Modulation of NOXA and MCL-1 as a Strategy for Sensitizing Melanoma Cells to the BH3-Mimetic ABT-737. <i>Clinical Cancer Research</i> , 2012, 18, 783-795.	3.2	98
116	Translation inhibitors induce cell death by multiple mechanisms and Mcl-1 reduction is only a minor contributor. <i>Cell Death and Disease</i> , 2012, 3, e409-e409.	2.7	42
117	Sheeppox Virus SPPV14 Encodes a Bcl-2-Like Cell Death Inhibitor That Counters a Distinct Set of Mammalian Proapoptotic Proteins. <i>Journal of Virology</i> , 2012, 86, 11501-11511.	1.5	41
118	Caspase-9 mediates the apoptotic death of megakaryocytes and platelets, but is dispensable for their generation and function. <i>Blood</i> , 2012, 119, 4283-4290.	0.6	70
119	Synthesis and biological evaluation of a potent salicylilalamide A lactam analogue. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 8147.	1.5	11
120	The restricted binding repertoire of Bcl-B leaves Bim as the universal BH3-only prosurvival Bcl-2 protein antagonist. <i>Cell Death and Disease</i> , 2012, 3, e443-e443.	2.7	61
121	A Cluster of Interferon- β -Inducible p65 GTPases Plays a Critical Role in Host Defense against <i>Toxoplasma gondii</i> . <i>Immunity</i> , 2012, 37, 302-313.	6.6	311
122	The Dendritic Cell Receptor Clec9A Binds Damaged Cells via Exposed Actin Filaments. <i>Immunity</i> , 2012, 36, 646-657.	6.6	272
123	Bcl-2, Bcl-xL, and Bcl-w are not equivalent targets of ABT-737 and navitoclax (ABT-263) in lymphoid and leukemic cells. <i>Blood</i> , 2012, 119, 5807-5816.	0.6	168
124	Total Synthesis of 2 β ,5 β -Diepisilvestrol and Its C1 β Epimer: Key Structure Activity Relationships at C1 β and C2 β . <i>Journal of Natural Products</i> , 2012, 75, 1500-1504.	1.5	19
125	Substantial Susceptibility of Chronic Lymphocytic Leukemia to BCL2 Inhibition: Results of a Phase I Study of Navitoclax in Patients With Relapsed or Refractory Disease. <i>Journal of Clinical Oncology</i> , 2012, 30, 488-496.	0.8	719
126	Megakaryocytes possess a functional intrinsic apoptosis pathway that must be restrained to survive and produce platelets. <i>Journal of Experimental Medicine</i> , 2011, 208, 2017-2031.	4.2	162

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127	Quinazoline Sulfonamides as Dual Binders of the Proteins B-Cell Lymphoma 2 and B-Cell Lymphoma Extra Long with Potent Proapoptotic Cell-Based Activity. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 1914-1926.	2.9	62
128	Bcl-xL-inhibitory BH3 mimetics can induce a transient thrombocytopeny that undermines the hemostatic function of platelets. <i>Blood</i> , 2011, 118, 1663-1674.	0.6	262
129	Overcoming blocks in apoptosis with BH3-mimetic therapy in haematological malignancies. <i>Pathology</i> , 2011, 43, 525-535.	0.3	36
130	CyclicAMP-dependent protein kinase A regulates apoptosis by stabilizing the BH3-only protein Bim. <i>EMBO Reports</i> , 2011, 12, 77-83.	2.0	52
131	Sensitivity to antitubulin chemotherapeutics is regulated by MCL1 and FBW7. <i>Nature</i> , 2011, 471, 110-114.	13.7	682
132	Deerpox Virus Encodes an Inhibitor of Apoptosis That Regulates Bak and Bax. <i>Journal of Virology</i> , 2011, 85, 1922-1934.	1.5	40
133	Evaluation of the Bcl-2 family antagonist ABT-737 in collagen-induced arthritis. <i>Journal of Leukocyte Biology</i> , 2011, 90, 819-829.	1.5	12
134	Induction of antigen-specific effector phase tolerance following vaccination against a previously ignored B-cell lymphoma. <i>Immunology and Cell Biology</i> , 2011, 89, 595-603.	1.0	13
135	Megakaryocytes possess a functional intrinsic apoptosis pathway that must be restrained to survive and produce platelets. <i>Journal of Cell Biology</i> , 2011, 194, i12-i12.	2.3	0
136	Transgenic, inducible RNAi in megakaryocytes and platelets in mice. <i>Journal of Thrombosis and Haemostasis</i> , 2010, 8, 2751-2756.	1.9	11
137	Deubiquitinase USP9X stabilizes MCL1 and promotes tumour cell survival. <i>Nature</i> , 2010, 463, 103-107.	13.7	529
138	Apoptosis and non-inflammatory phagocytosis can be induced by mitochondrial damage without caspases. <i>Cell Death and Differentiation</i> , 2010, 17, 821-832.	5.0	33
139	Glucose Induces Pancreatic Islet Cell Apoptosis That Requires the BH3-Only Proteins Bim and Puma and Multi-BH Domain Protein Bax. <i>Diabetes</i> , 2010, 59, 644-652.	0.3	103
140	BH3 mimetics antagonizing restricted prosurvival Bcl-2 proteins represent another class of selective immune modulatory drugs. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 10967-10971.	3.3	97
141	The BH3-Mimetic ABT-737 Induces Mast Cell Apoptosis In Vitro and In Vivo: Potential for Therapeutics. <i>Journal of Immunology</i> , 2010, 185, 2555-2562.	0.4	25
142	Structural Basis for Apoptosis Inhibition by Epstein-Barr Virus BHRF1. <i>PLoS Pathogens</i> , 2010, 6, e1001236.	2.1	99
143	Pro-apoptotic Bax is the major and Bak an auxiliary effector in cytokine deprivation-induced mast cell apoptosis. <i>Cell Death and Disease</i> , 2010, 1, e43-e43.	2.7	26
144	Megakaryocytes Possess a Functional Intrinsic Apoptosis Pathway That Must Be Restrained In Order to Survive and Produce Platelets. <i>Blood</i> , 2010, 116, 550-550.	0.6	0

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145	Novel Bcl-2 Homology-3 Domain-like Sequences Identified from Screening Randomized Peptide Libraries for Inhibitors of the Pro-survival Bcl-2 Proteins. <i>Journal of Biological Chemistry</i> , 2009, 284, 31315-31326.	1.6	29
146	MEK/ERK-Mediated Phosphorylation of Bim Is Required to Ensure Survival of T and B Lymphocytes during Mitogenic Stimulation. <i>Journal of Immunology</i> , 2009, 183, 261-269.	0.4	76
147	Correction for Fletcher et al., Inaugural Article: Apoptosis is triggered when prosurvival Bcl-2 proteins cannot restrain Bax. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 1678-1678.	3.3	0
148	The BH3 mimetic compound, ABT-737, synergizes with a range of cytotoxic chemotherapy agents in chronic lymphocytic leukemia. <i>Leukemia</i> , 2009, 23, 2034-2041.	3.3	91
149	XIAP discriminates between type I and type II FAS-induced apoptosis. <i>Nature</i> , 2009, 460, 1035-1039.	13.7	421
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