David C Huang

List of Publications by Year in Descending Order

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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.

86 32,438 179 230 h-index g-index citations papers 11.6 6.69 241 35,750 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
230	The Lck inhibitor, AMG-47a, blocks necroptosis and implicates RIPK1 in signalling downstream of MLKL <i>Cell Death and Disease</i> , 2022 , 13, 291	9.8	1
229	Comprehensive characterization of single-cell full-length isoforms in human and mouse with long-read sequencing. <i>Genome Biology</i> , 2021 , 22, 310	18.3	10
228	Mesenchymal stromal cell apoptosis is required for their therapeutic function. <i>Nature Communications</i> , 2021 , 12, 6495	17.4	11
227	The transcription factor IRF4 represses proapoptotic BMF and BIM to licence multiple myeloma survival. <i>Leukemia</i> , 2021 , 35, 2114-2118	10.7	8
226	Structure-Guided Development of Potent Benzoylurea Inhibitors of BCL-X and BCL-2. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 5447-5469	8.3	3
225	PRMT1-mediated H4R3me2a recruits SMARCA4 to promote colorectal cancer progression by enhancing EGFR signaling. <i>Genome Medicine</i> , 2021 , 13, 58	14.4	9
224	Intact TP-53 function is essential for sustaining durable responses to BH3-mimetic drugs in leukemias. <i>Blood</i> , 2021 , 137, 2721-2735	2.2	14
223	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	12.7	6
222	BCL2 and MCL1 inhibitors for hematologic malignancies. <i>Blood</i> , 2021 , 138, 1120-1136	2.2	7
221	Transcriptional silencing of fetal hemoglobin expression by NonO. <i>Nucleic Acids Research</i> , 2021 , 49, 97	11 <u>2</u> 972	3 1
220	Outcomes of patients with CLL sequentially resistant to both BCL2 and BTK inhibition. <i>Blood Advances</i> , 2021 , 5, 4054-4058	7.8	9
219	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	17.4	1
218	Cotargeting BCL-2 and MCL-1 in high-risk B-ALL. <i>Blood Advances</i> , 2020 , 4, 2762-2767	7.8	14
217	Loss of RIPK3 does not impact MYC-driven lymphomagenesis or chemotherapeutic drug-induced killing of malignant lymphoma cells. <i>Cell Death and Differentiation</i> , 2020 , 27, 2531-2533	12.7	3
216	MARCH5 requires MTCH2 to coordinate proteasomal turnover of the MCL1:NOXA complex. <i>Cell Death and Differentiation</i> , 2020 , 27, 2484-2499	12.7	16
215	Deep profiling of apoptotic pathways with mass cytometry identifies a synergistic drug combination for killing myeloma cells. <i>Cell Death and Differentiation</i> , 2020 , 27, 2217-2233	12.7	18
214	Acquired Mutations in BAX Confer Resistance to BH3 Mimetics in Acute Myeloid Leukemia. <i>Blood</i> , 2020 , 136, 7-8	2.2	5

(2019-2020)

213	BAX-Mutated Clonal Hematopoiesis in Patients on Long-Term Venetoclax for Relapsed/Refractory Chronic Lymphocytic Leukemia. <i>Blood</i> , 2020 , 136, 9-10	2.2	2
212	BH3 Mimetics for the Treatment of B-Cell Malignancies-Insights and Lessons from the Clinic. <i>Cancers</i> , 2020 , 12,	6.6	5
211	Defining the susceptibility of colorectal cancers to BH3-mimetic compounds. <i>Cell Death and Disease</i> , 2020 , 11, 735	9.8	6
210	Potent Inhibition of Necroptosis by Simultaneously Targeting Multiple Effectors of the Pathway. <i>ACS Chemical Biology</i> , 2020 , 15, 2702-2713	4.9	11
209	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	12.7	12
208	Molecular patterns of response and treatment failure after frontline venetoclax combinations in older patients with AML. <i>Blood</i> , 2020 , 135, 791-803	2.2	176
207	Multiple BCL2 mutations cooccurring with Gly101Val emerge in chronic lymphocytic leukemia progression on venetoclax. <i>Blood</i> , 2020 , 135, 773-777	2.2	55
206	Replication stress induces mitotic death through parallel pathways regulated by WAPL and telomere deprotection. <i>Nature Communications</i> , 2019 , 10, 4224	17.4	13
205	A small molecule interacts with VDAC2 to block mouse BAK-driven apoptosis. <i>Nature Chemical Biology</i> , 2019 , 15, 1057-1066	11.7	16
204	Characterization of a novel venetoclax resistance mutation (BCL2 Phe104Ile) observed in follicular lymphoma. <i>British Journal of Haematology</i> , 2019 , 186, e188-e191	4.5	24
203	Structures of BCL-2 in complex with venetoclax reveal the molecular basis of resistance mutations. <i>Nature Communications</i> , 2019 , 10, 2385	17.4	84
202	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	2.2	2
201	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	2.2	12
200	Venetoclax for the treatment of mantle cell lymphoma. <i>Annals of Lymphoma</i> , 2019 , 3, 4-4	1.8	O
199	Multiple myeloma with 1q21 amplification is highly sensitive to MCL-1 targeting. <i>Blood Advances</i> , 2019 , 3, 4202-4214	7.8	35
198	Combining BH3-mimetics to target both BCL-2 and MCL1 has potent activity in pre-clinical models of acute myeloid leukemia. <i>Leukemia</i> , 2019 , 33, 905-917	10.7	84
197	Dynamic molecular monitoring reveals that SWI-SNF mutations mediate resistance to ibrutinib plus venetoclax in mantle cell lymphoma. <i>Nature Medicine</i> , 2019 , 25, 119-129	50.5	94
196	Recipient BCL2 inhibition and NK cell ablation form part of a reduced intensity conditioning regime that improves allo-bone marrow transplantation outcomes. <i>Cell Death and Differentiation</i> , 2019 , 26, 1.	51 6-15 3	o ⁸

195	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	11.7	21
194	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	24.4	188
193	BAK/BAX-Mediated Apoptosis Is a Myc-Induced Roadblock to Reprogramming. <i>Stem Cell Reports</i> , 2018 , 10, 331-338	8	9
192	Enhancing venetoclax activity in acute myeloid leukemia by co-targeting MCL1. <i>Leukemia</i> , 2018 , 32, 303	-302/	96
191	The Mitochondrial Apoptotic Effectors BAX/BAK Activate Caspase-3 and -7 to Trigger NLRP3 Inflammasome and Caspase-8 Driven IL-1 (Activation. <i>Cell Reports</i> , 2018 , 25, 2339-2353.e4	10.6	102
190	VDAC2 enables BAX to mediate apoptosis and limit tumor development. <i>Nature Communications</i> , 2018 , 9, 4976	17.4	73
189	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	24.4	194
188	CARM1-mediated methylation of protein arginine methyltransferase 5 represses human Eglobin gene expression in erythroleukemia cells. <i>Journal of Biological Chemistry</i> , 2018 , 293, 17454-17463	5.4	11
187	Infection with flaviviruses requires BCLXL for cell survival. <i>PLoS Pathogens</i> , 2018 , 14, e1007299	7.6	18
186	IMiDs prime myeloma cells for daratumumab-mediated cytotoxicity through loss of Ikaros and Aiolos. <i>Blood</i> , 2018 , 132, 2166-2178	2.2	42
185	Venetoclax in Patients with Previously Treated Chronic Lymphocytic Leukemia. <i>Clinical Cancer Research</i> , 2017 , 23, 4527-4533	12.9	43
184	DR5 and caspase-8 are dispensable in ER stress-induced apoptosis. <i>Cell Death and Differentiation</i> , 2017 , 24, 944-950	12.7	51
183	Clinicopathological features and outcomes of progression of CLL on the BCL2 inhibitor venetoclax. <i>Blood</i> , 2017 , 129, 3362-3370	2.2	114
182	Essential role for Bim in mediating the apoptotic and antitumor activities of immunotoxins. <i>Oncogene</i> , 2017 , 36, 4953-4962	9.2	8
181	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	12.7	62
180	NatD promotes lung cancer progression by preventing histone H4 serine phosphorylation to activate Slug expression. <i>Nature Communications</i> , 2017 , 8, 928	17.4	46
179	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,	17.5	112
178	Design, Synthesis, and Biological Activity of 1,2,3-Triazolobenzodiazepine BET Bromodomain Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 1298-1303	4.3	17

(2016-2017)

177	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	6.1	74
176	Defining a therapeutic window for kinase inhibitors in leukemia to avoid neutropenia. <i>Oncotarget</i> , 2017 , 8, 57948-57963	3.3	3
175	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	9.8	42
174	Venetoclax responses of pediatric ALL xenografts reveal sensitivity of MLL-rearranged leukemia. <i>Blood</i> , 2016 , 128, 1382-95	2.2	100
173	Eliminating Legionella by inhibiting BCL-XL to induce macrophage apoptosis. <i>Nature Microbiology</i> , 2016 , 1, 15034	26.6	46
172	The MCL1 inhibitor S63845 is tolerable and effective in diverse cancer models. <i>Nature</i> , 2016 , 538, 477-4	1 83 0.4	617
171	BET inhibition represses miR17-92 to drive BIM-initiated apoptosis of normal and transformed hematopoietic cells. <i>Leukemia</i> , 2016 , 30, 1531-41	10.7	22
170	Small molecules targeting Mcl-1: the search for a silver bullet in cancer therapy. <i>MedChemComm</i> , 2016 , 7, 778-787	5	11
169	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	9.8	83
168	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, 1858-66	10.6	25
167	Current challenges and novel treatment strategies in double hit lymphomas. <i>Therapeutic Advances in Hematology</i> , 2016 , 7, 52-64	5.7	19
166	Clinicopathological Features and Outcomes of Progression for Chronic Lymphocytic Leukaemia (CLL) Treated with the BCL2 Inhibitor Venetoclax. <i>Blood</i> , 2016 , 128, 3223-3223	2.2	2
165	The BIM deletion polymorphism: A paradigm of a permissive interaction between germline and acquired TKI resistance factors in chronic myeloid leukemia. <i>Oncotarget</i> , 2016 , 7, 2721-33	3.3	13
164	Targeting the Pro-Survival BCL2 Proteins with BH3 Mimetic Compounds for Treating Multiple Myeloma. <i>Blood</i> , 2016 , 128, 3293-3293	2.2	
163	The Role of BAX/BAK-Mediated Apoptosis for the Cytotoxic Action of Anti-Myeloma Agents. <i>Blood</i> , 2016 , 128, 5706-5706	2.2	
162	Identification of an activation site in Bak and mitochondrial Bax triggered by antibodies. <i>Nature Communications</i> , 2016 , 7, 11734	17.4	37
161	The BCL2 selective inhibitor venetoclax induces rapid onset apoptosis of CLL cells in patients via a TP53-independent mechanism. <i>Blood</i> , 2016 , 127, 3215-24	2.2	181
160	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	12.7	18

159	Hierarchy for targeting prosurvival BCL2 family proteins in multiple myeloma: pivotal role of MCL1. <i>Blood</i> , 2016 , 128, 1834-1844	2.2	105
158	Bcl-2 antagonists kill plasmacytoid dendritic cells from lupus-prone mice and dampen interferon-⊟ production. <i>Arthritis and Rheumatology</i> , 2015 , 67, 797-808	9.5	35
157	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	17.5	344
156	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	9.8	1
155	A RIPK2 inhibitor delays NOD signalling events yet prevents inflammatory cytokine production. <i>Nature Communications</i> , 2015 , 6, 6442	17.4	74
154	Systematic Screening Identifies Dual PI3K and mTOR Inhibition as a Conserved Therapeutic Vulnerability in Osteosarcoma. <i>Clinical Cancer Research</i> , 2015 , 21, 3216-29	12.9	47
153	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, 140-8	10.3	24
152	A Chemical Screening Approach to Identify Novel Key Mediators of Erythroid Enucleation. <i>PLoS ONE</i> , 2015 , 10, e0142655	3.7	8
151	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, 4044-9	11.5	39
150	BCL2 inhibition in double hit lymphoma. <i>Leukemia and Lymphoma</i> , 2015 , 56, 1928-9	1.9	1
149	Targeting of MCL-1 kills MYC-driven mouse and human lymphomas even when they bear mutations in p53. <i>Genes and Development</i> , 2014 , 28, 58-70	12.6	121
148	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, 8512-7	11.5	141
147	Enhanced stability of Mcl1, a prosurvival Bcl2 relative, blunts stress-induced apoptosis, causes male sterility, and promotes tumorigenesis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 261-6	11.5	41
146	Both leukaemic and normal peripheral B lymphoid cells are highly sensitive to the selective pharmacological inhibition of prosurvival Bcl-2 with ABT-199. <i>Leukemia</i> , 2014 , 28, 1207-15	10.7	69
145	Evaluation of functional groups as acetyl-lysine mimetics for BET bromodomain inhibition. <i>MedChemComm</i> , 2014 , 5, 1834-1842	5	21
144	Further insights into the effects of pre-organizing the BimBH3 helix. ACS Chemical Biology, 2014, 9, 838	-9 .9	24
143	De-novo designed library of benzoylureas as inhibitors of BCL-XL: synthesis, structural and biochemical characterization. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 1323-43	8.3	31
142	Structure-Guided Rescaffolding of Selective Antagonists of BCL-XL. <i>ACS Medicinal Chemistry Letters</i> , 2014 , 5, 662-7	4.3	36

141	197. Cytokine, 2014 , 70, 75-76	4	
140	A biosensor of SRC family kinase conformation by exposable tetracysteine useful for cell-based screening. <i>ACS Chemical Biology</i> , 2014 , 9, 1426-31	4.9	9
139	Apoptotic caspases suppress mtDNA-induced STING-mediated type I IFN production. <i>Cell</i> , 2014 , 159, 1549-62	56.2	475
138	Simplified silvestrol analogues with potent cytotoxic activity. ChemMedChem, 2014, 9, 1556-66	3.7	14
137	Targeting BCL2 for the treatment of lymphoid malignancies. Seminars in Hematology, 2014, 51, 219-27	4	112
136	Eradication of Acute Myeloid Leukemia Is Enhanced By Combined Bcl-2 and Mcl-1 Targeting. <i>Blood</i> , 2014 , 124, 988-988	2.2	2
135	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 <i>Journal of Clinical Oncology</i> , 2014 , 32, 7015-7015	2.2	33
134	Targeting acute myeloid leukemia by dual inhibition of PI3K signaling and Cdk9-mediated Mcl-1 transcription. <i>Blood</i> , 2013 , 122, 738-48	2.2	47
133	ABT-199, a potent and selective BCL-2 inhibitor, achieves antitumor activity while sparing platelets. <i>Nature Medicine</i> , 2013 , 19, 202-8	50.5	1922
132	Synthesis of biotinylated episilvestrol: highly selective targeting of the translation factors eIF4AI/II. Organic Letters, 2013 , 15, 1406-9	6.2	44
131	Bax crystal structures reveal how BH3 domains activate Bax and nucleate its oligomerization to induce apoptosis. <i>Cell</i> , 2013 , 152, 519-31	56.2	402
130	Structure-guided design of a selective BCL-X(L) inhibitor. <i>Nature Chemical Biology</i> , 2013 , 9, 390-7	11.7	277
129	Discovery of potent and selective benzothiazole hydrazone inhibitors of Bcl-XL. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 5514-40	8.3	50
128	Stabilizing the pro-apoptotic BimBH3 helix (BimSAHB) does not necessarily enhance affinity or biological activity. <i>ACS Chemical Biology</i> , 2013 , 8, 297-302	4.9	109
127	BH3 mimetic therapy: an emerging and promising approach to treating chronic lymphocytic leukemia. <i>Leukemia and Lymphoma</i> , 2013 , 54, 909-11	1.9	2
126	MCMV-mediated inhibition of the pro-apoptotic Bak protein is required for optimal in vivo replication. <i>PLoS Pathogens</i> , 2013 , 9, e1003192	7.6	17
125	Proapoptotic Bak and Bax guard against fatal systemic and organ-specific autoimmune disease. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 2599-604	11.5	37
124	Variability of inducible expression across the hematopoietic system of tetracycline transactivator transgenic mice. <i>PLoS ONE</i> , 2013 , 8, e54009	3.7	19

123	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	2.2	8
122	Caspase-9 mediates the apoptotic death of megakaryocytes and platelets, but is dispensable for their generation and function. <i>Blood</i> , 2012 , 119, 4283-90	2.2	57
121	Synthesis and biological evaluation of a potent salicylihalamide A lactam analogue. <i>Organic and Biomolecular Chemistry</i> , 2012 , 10, 8147-53	3.9	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	9.8	41
119	A cluster of interferon-Inducible p65 GTPases plays a critical role in host defense against Toxoplasma gondii. <i>Immunity</i> , 2012 , 37, 302-13	32.3	230
118	The dendritic cell receptor Clec9A binds damaged cells via exposed actin filaments. <i>Immunity</i> , 2012 , 36, 646-57	32.3	224
117	Bcl-2, Bcl-x(L), and Bcl-w are not equivalent targets of ABT-737 and navitoclax (ABT-263) in lymphoid and leukemic cells. <i>Blood</i> , 2012 , 119, 5807-16	2.2	150
116	Total synthesis of 2∰@diepisilvestrol and its C1@epimer: key structure activity relationships at C1@end C2@Journal of Natural Products, 2012 , 75, 1500-4	4.9	17
115	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, 488-96	2.2	622
114	Sensitization of BCL-2-expressing breast tumors to chemotherapy by the BH3 mimetic ABT-737. Proceedings of the National Academy of Sciences of the United States of America, 2012 , 109, 2766-71	11.5	156
113	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, 783-95	12.9	92
112	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	9.8	36
111	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, 11501-11	6.6	32
110	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, 1914-2	2 ^{8.3}	55
109	Bcl-xL-inhibitory BH3 mimetics can induce a transient thrombocytopathy that undermines the hemostatic function of platelets. <i>Blood</i> , 2011 , 118, 1663-74	2.2	199
108	Overcoming blocks in apoptosis with BH3-mimetic therapy in haematological malignancies. <i>Pathology</i> , 2011 , 43, 525-35	1.6	33
107	Cyclic-AMP-dependent protein kinase A regulates apoptosis by stabilizing the BH3-only protein Bim. <i>EMBO Reports</i> , 2011 , 12, 77-83	6.5	49
106	Sensitivity to antitubulin chemotherapeutics is regulated by MCL1 and FBW7. <i>Nature</i> , 2011 , 471, 110-4	50.4	602

105	Megakaryocytes possess a functional intrinsic apoptosis pathway that must be restrained to survive and produce platelets. <i>Journal of Experimental Medicine</i> , 2011 , 208, 2017-31	16.6	139
104	Deerpox virus encodes an inhibitor of apoptosis that regulates Bak and Bax. <i>Journal of Virology</i> , 2011 , 85, 1922-34	6.6	33
103	Evaluation of the Bcl-2 family antagonist ABT-737 in collagen-induced arthritis. <i>Journal of Leukocyte Biology</i> , 2011 , 90, 819-29	6.5	11
102	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	5	9
101	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	7.3	
100	Deubiquitinase USP9X stabilizes MCL1 and promotes tumour cell survival. <i>Nature</i> , 2010 , 463, 103-7	50.4	485
99	Apoptosis and non-inflammatory phagocytosis can be induced by mitochondrial damage without caspases. <i>Cell Death and Differentiation</i> , 2010 , 17, 821-32	12.7	29
98	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, 644-52	0.9	90
97	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, 10967-71	11.5	88
96	The BH3-mimetic ABT-737 induces mast cell apoptosis in vitro and in vivo: potential for therapeutics. <i>Journal of Immunology</i> , 2010 , 185, 2555-62	5.3	24
95	Structural basis for apoptosis inhibition by Epstein-Barr virus BHRF1. <i>PLoS Pathogens</i> , 2010 , 6, e100123	6 7.6	74
94	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	9.8	22
93	Transgenic, inducible RNAi in megakaryocytes and platelets in mice. <i>Journal of Thrombosis and Haemostasis</i> , 2010 , 8, 2751-6	15.4	10
92	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	2.2	
91	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, 3131	5 -2 6	26
90	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, 261-9	5.3	66
89	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	11.5	78
88	The BH3 mimetic compound, ABT-737, synergizes with a range of cytotoxic chemotherapy agents in chronic lymphocytic leukemia. <i>Leukemia</i> , 2009 , 23, 2034-41	10.7	84

87	XIAP discriminates between type I and type II FAS-induced apoptosis. <i>Nature</i> , 2009 , 460, 1035-9	50.4	344
86	Virally mediated inhibition of Bax in leukocytes promotes dissemination of murine cytomegalovirus. <i>Cell Death and Differentiation</i> , 2009 , 16, 312-20	12.7	32
85	Bax activation by Bim?. Cell Death and Differentiation, 2009, 16, 1187-91	12.7	75
84	betaTrCP- and Rsk1/2-mediated degradation of BimEL inhibits apoptosis. <i>Molecular Cell</i> , 2009 , 33, 109-	16 7.6	138
83	BH3-mimeticsthe solution to chemoresistance?. <i>Leukemia and Lymphoma</i> , 2009 , 50, 1069-72	1.9	4
82	Two distinct pathways regulate platelet phosphatidylserine exposure and procoagulant function. <i>Blood</i> , 2009 , 114, 663-6	2.2	240
81	BH3-only proteins and their roles in programmed cell death. <i>Oncogene</i> , 2008 , 27 Suppl 1, S128-36	9.2	153
80	Controlling the cell death mediators Bax and Bak: puzzles and conundrums. <i>Cell Cycle</i> , 2008 , 7, 39-44	4.7	52
79	A novel BH3 ligand that selectively targets Mcl-1 reveals that apoptosis can proceed without Mcl-1 degradation. <i>Journal of Cell Biology</i> , 2008 , 180, 341-55	7.3	146
78	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> , 2008 , 105, 18081-7	11.5	141
77	Proapoptotic BH3-only protein Bid is essential for death receptor-induced apoptosis of pancreatic beta-cells. <i>Diabetes</i> , 2008 , 57, 1284-92	0.9	78
76	Vaccinia virus anti-apoptotic F1L is a novel Bcl-2-like domain-swapped dimer that binds a highly selective subset of BH3-containing death ligands. <i>Cell Death and Differentiation</i> , 2008 , 15, 1564-71	12.7	177
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