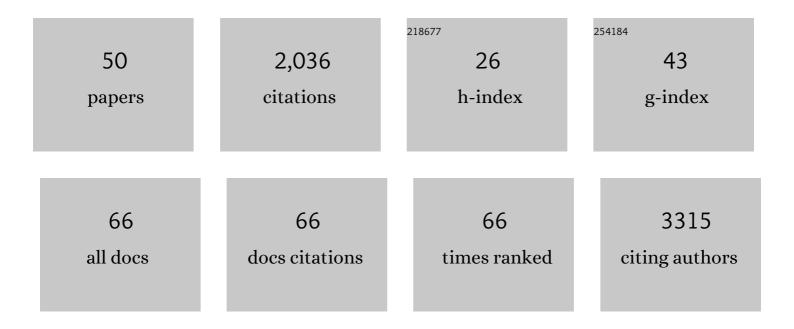
Haiming Dai

List of Publications by Year in descending order

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HAIMING DAL

#	Article	IF	CITATIONS
1	COMMD1 is linked to the WASH complex and regulates endosomal trafficking of the copper transporter ATP7A. Molecular Biology of the Cell, 2015, 26, 91-103.	2.1	200
2	Chemotherapy-induced pyroptosis is mediated by BAK/BAX-caspase-3-GSDME pathway and inhibited by 2-bromopalmitate. Cell Death and Disease, 2020, 11, 281.	6.3	149
3	Transient binding of an activator BH3 domain to the Bak BH3-binding groove initiates Bak oligomerization. Journal of Cell Biology, 2011, 194, 39-48.	5.2	139
4	Emerging understanding of Bcl-2 biology: Implications for neoplastic progression and treatment. Biochimica Et Biophysica Acta - Molecular Cell Research, 2015, 1853, 1658-1671.	4.1	122
5	BCL2 mutations are associated with increased risk of transformation and shortened survival in follicular lymphoma. Blood, 2015, 125, 658-667.	1.4	108
6	MCL-1 as a Buffer for Proapoptotic BCL-2 Family Members during TRAIL-induced Apoptosis. Journal of Biological Chemistry, 2007, 282, 29831-29846.	3.4	104
7	Prime, Shock, and Kill: Priming CD4 T Cells from HIV Patients with a BCL-2 Antagonist before HIV Reactivation Reduces HIV Reservoir Size. Journal of Virology, 2016, 90, 4032-4048.	3.4	85
8	Contribution of Bcl-2 Phosphorylation to Bak Binding and Drug Resistance. Cancer Research, 2013, 73, 6998-7008.	0.9	81
9	Noxa/Bcl-2 Protein Interactions Contribute to Bortezomib Resistance in Human Lymphoid Cells. Journal of Biological Chemistry, 2011, 286, 17682-17692.	3.4	80
10	Evaluation of the BH3-only Protein Puma as a Direct Bak Activator. Journal of Biological Chemistry, 2014, 289, 89-99.	3.4	65
11	CXCR4 Chemokine Receptor Signaling Induces Apoptosis in Acute Myeloid Leukemia Cells via Regulation of the Bcl-2 Family Members Bcl-XL, Noxa, and Bak. Journal of Biological Chemistry, 2013, 288, 22899-22914.	3.4	59
12	Solution structure of BRD7 bromodomain and its interaction with acetylated peptides from histone H3 and H4. Biochemical and Biophysical Research Communications, 2007, 358, 435-441.	2.1	54
13	4EBP1/c-MYC/PUMA and NF-κB/EGR1/BIM pathways underlie cytotoxicity of mTOR dual inhibitors in malignant lymphoid cells. Blood, 2016, 127, 2711-2722.	1.4	49
14	Context-dependent Bcl-2/Bak Interactions Regulate Lymphoid Cell Apoptosis. Journal of Biological Chemistry, 2009, 284, 18311-18322.	3.4	47
15	Platelet-derived Growth Factor Primes Cancer-associated Fibroblasts for Apoptosis. Journal of Biological Chemistry, 2014, 289, 22835-22849.	3.4	47
16	DHHC protein family targets different subsets of glioma stem cells in specific niches. Journal of Experimental and Clinical Cancer Research, 2019, 38, 25.	8.6	44
17	BCL2 Family, Mitochondrial Apoptosis, and Beyond. Cancer Translational Medicine, 2016, 2, 7.	0.2	44
18	Bak Conformational Changes Induced by Ligand Binding: Insight into BH3 Domain Binding and Bak Homo-Oligomerization. Scientific Reports, 2012, 2, 257.	3.3	41

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19	Constitutive BAK activation as a determinant of drug sensitivity in malignant lymphohematopoietic cells. Genes and Development, 2015, 29, 2140-2152.	5.9	38
20	Matrix Metalloproteinase 11 Is a Potential Therapeutic Target in Lung Adenocarcinoma. Molecular Therapy - Oncolytics, 2019, 14, 82-93.	4.4	35
21	Targeted Inhibition of LPL/FABP4/CPT1 fatty acid metabolic axis can effectively prevent the progression of nonalcoholic steatohepatitis to liver cancer. International Journal of Biological Sciences, 2021, 17, 4207-4222.	6.4	35
22	Mitochondrial apoptosis and BH3 mimetics. F1000Research, 2016, 5, 2804.	1.6	33
23	Characterization of an alternative BAK-binding site for BH3 peptides. Nature Communications, 2020, 11, 3301.	12.8	31
24	Casp8p41 generated by HIV protease kills CD4 T cells through direct Bak activation. Journal of Cell Biology, 2014, 206, 867-876.	5.2	28
25	High Cell Surface Death Receptor Expression Determines Type I Versus Type II Signaling*. Journal of Biological Chemistry, 2011, 286, 35823-35833.	3.4	27
26	Cytotoxicity of farnesyltransferase inhibitors in lymphoid cells mediated by MAPK pathway inhibition and Bim up-regulation. Blood, 2011, 118, 4872-4881.	1.4	27
27	Measurement of BH3-only protein tolerance. Cell Death and Differentiation, 2018, 25, 282-293.	11.2	27
28	Systematic expression analysis of WEE family kinases reveals the importance of PKMYT1 in breast carcinogenesis. Cell Proliferation, 2020, 53, e12741.	5.3	27
29	Matrix Metalloproteinase Expressions Play Important role in Prediction of Ovarian Cancer Outcome. Scientific Reports, 2019, 9, 11677.	3.3	25
30	Fluorine Pseudocontact Shifts Used for Characterizing the Protein–Ligand Interaction Mode in the Limit of NMR Intermediate Exchange. Angewandte Chemie - International Edition, 2017, 56, 12982-12986.	13.8	23
31	BCL2L13: physiological and pathological meanings. Cellular and Molecular Life Sciences, 2021, 78, 2419-2428.	5.4	22
32	Binding model of human coactosin-like protein with filament actin revealed by mutagenesis. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2006, 1764, 1688-1700.	2.3	18
33	Solution structure of calponin homology domain of Human MICAL-1. Journal of Biomolecular NMR, 2006, 36, 295-300.	2.8	18
34	Selective binding of mitophagy receptor protein Bcl-rambo to LC3/GABARAP family proteins. Biochemical and Biophysical Research Communications, 2020, 530, 292-300.	2.1	16
35	Protein Kinase Cβ Modulates Ligand-induced Cell Surface Death Receptor Accumulation. Journal of Biological Chemistry, 2010, 285, 888-902.	3.4	15
36	Combination treatment of RAD001 and BEZ235 exhibits synergistic antitumor activity via down-regulation of p-4E-BP1/Mcl-1 in small cell lung cancer. Oncotarget, 2017, 8, 106486-106498.	1.8	12

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37	Identification of FPR3 as a Unique Biomarker for Targeted Therapy in the Immune Microenvironment of Breast Cancer. Frontiers in Pharmacology, 2020, 11, 593247.	3.5	9
38	Structure of human upstream binding factor HMG box 5 and site for binding of the cell-cycle regulatory factor TAF1. Acta Crystallographica Section D: Biological Crystallography, 2007, 63, 730-737.	2.5	8
39	Investigation of the four cooperative unfolding units existing in the MICAL-1 CH domain. Biophysical Chemistry, 2007, 129, 269-278.	2.8	7
40	Enrichment and detection of circulating tumor cells by immunomagnetic beads and flow cytometry. Biotechnology Letters, 2021, 43, 25-34.	2.2	7
41	Cell Surface Protein Disulfide Isomerase Regulates Natriuretic Peptide Generation of Cyclic Guanosine Monophosphate. PLoS ONE, 2014, 9, e112986.	2.5	6
42	Letter to the Editor:1H,13C and15N Resonance Assignments and the Secondary Structures of Human Coactosin Like Protein (hCLP) D123N. Journal of Biomolecular NMR, 2004, 29, 455-456.	2.8	5
43	An Immune-Associated Genomic Signature Effectively Predicts Pathologic Complete Response to Neoadjuvant Paclitaxel and Anthracycline-Based Chemotherapy in Breast Cancer. Frontiers in Immunology, 2021, 12, 704655.	4.8	5
44	Constitutive BAK/MCL1 complexes predict paclitaxel and S63845 sensitivity of ovarian cancer. Cell Death and Disease, 2021, 12, 789.	6.3	4
45	Down-regulation of BCL2L13 renders poor prognosis in clear cell and papillary renal cell carcinoma. Cancer Cell International, 2021, 21, 332.	4.1	3
46	Therapeutics targeting BCL2 family proteins. , 2022, , 197-260.		3
47	Selective Inhibition of BFL1: It's All about Finding the Right Partner. Cell Chemical Biology, 2020, 27, 639-642.	5.2	2
48	Crystal structure of the WD40 domain of human PLRG1. Biochemical and Biophysical Research Communications, 2021, 534, 474-477.	2.1	1
49	2P060 Solution Structure of the Bromodomain of Brd2 and Its Interaction with Acetylated Peptide from Histone H4 and CtBP(29. Protein structure and dynamics (II),Poster Session,Abstract,Meeting) Tj ETQq1 1	0.78.4314	∙rgƁT /Overlo
50	Fluorine Pseudocontact Shifts Used for Characterizing the Protein–Ligand Interaction Mode in the Limit of NMR Intermediate Exchange. Angewandte Chemie, 2017, 129, 13162-13166.	2.0	0