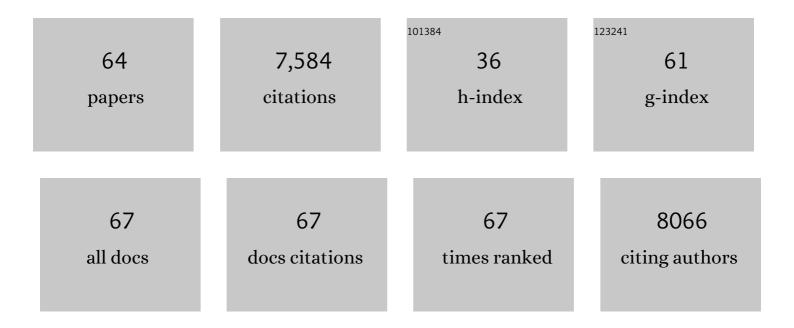
## Catherine L Day

List of Publications by Year in descending order

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CATHEDINE L DAV

#	Article	IF	CITATIONS
1	Differential Targeting of Prosurvival Bcl-2 Proteins by Their BH3-Only Ligands Allows Complementary Apoptotic Function. Molecular Cell, 2005, 17, 393-403.	4.5	1,639
2	The BH3 mimetic ABT-737 targets selective Bcl-2 proteins and efficiently induces apoptosis via Bak/Bax if Mcl-1 is neutralized. Cancer Cell, 2006, 10, 389-399.	7.7	1,149
3	HtrA2 Promotes Cell Death through Its Serine Protease Activity and Its Ability to Antagonize Inhibitor of Apoptosis Proteins. Journal of Biological Chemistry, 2002, 277, 445-454.	1.6	484
4	Structural insights into the degradation of Mcl-1 induced by BH3 domains. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 6217-6222.	3.3	397
5	The Bcl-2-regulated apoptotic pathway. Journal of Cell Science, 2003, 116, 4053-4056.	1.2	206
6	TRAF2 Must Bind to Cellular Inhibitors of Apoptosis for Tumor Necrosis Factor (TNF) to Efficiently Activate NF-κB and to Prevent TNF-induced Apoptosis. Journal of Biological Chemistry, 2009, 284, 35906-35915.	1.6	202
7	Bak Activation for Apoptosis Involves Oligomerization of Dimers via Their α6 Helices. Molecular Cell, 2009, 36, 696-703.	4.5	200
8	Solution Structure of Prosurvival Mcl-1 and Characterization of Its Binding by Proapoptotic BH3-only Ligands. Journal of Biological Chemistry, 2005, 280, 4738-4744.	1.6	187
9	Structure of the BH3 Domains from the p53-Inducible BH3-Only Proteins Noxa and Puma in Complex with Mcl-1. Journal of Molecular Biology, 2008, 380, 958-971.	2.0	178
10	RINGs hold the key to ubiquitin transfer. Trends in Biochemical Sciences, 2012, 37, 58-65.	3.7	168
11	Solution structure of a baculoviral inhibitor of apoptosis (IAP) repeat. Nature Structural Biology, 1999, 6, 648-651.	9.7	165
12	Structures of the cIAP2 RING Domain Reveal Conformational Changes Associated with Ubiquitin-conjugating Enzyme (E2) Recruitment. Journal of Biological Chemistry, 2008, 283, 31633-31640.	1.6	153
13	The structure of Bcl-w reveals a role for the C-terminal residues in modulating biological activity. EMBO Journal, 2003, 22, 1497-1507.	3.5	151
14	Birinapant, a Smac-Mimetic with Improved Tolerability for the Treatment of Solid Tumors and Hematological Malignancies. Journal of Medicinal Chemistry, 2014, 57, 3666-3677.	2.9	146
15	Smac Mimetics Activate the E3 Ligase Activity of cIAP1 Protein by Promoting RING Domain Dimerization. Journal of Biological Chemistry, 2011, 286, 17015-17028.	1.6	142
16	Determination of cell survival by RING-mediated regulation of inhibitor of apoptosis (IAP) protein abundance. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 16182-16187.	3.3	133
17	The anti-apoptotic activity of XIAP is retained upon mutation of both the caspase 3– and caspase 9–interacting sites. Journal of Cell Biology, 2002, 157, 115-124.	2.3	124
18	Proapoptotic BH3-only proteins trigger membrane integration of prosurvival Bcl-w and neutralize its activity. Journal of Cell Biology, 2003, 162, 877-888.	2.3	104

CATHERINE L DAY

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19	Structure of the bacterial type <scp>II NADH</scp> dehydrogenase: a monotopic membrane protein with an essential role in energy generation. Molecular Microbiology, 2014, 91, 950-964.	1.2	103
20	Structural Plasticity Underpins Promiscuous Binding of the Prosurvival Protein A1. Structure, 2008, 16, 818-829.	1.6	97
21	CARD-Mediated Autoinhibition of cIAP1's E3 Ligase Activity Suppresses Cell Proliferation and Migration. Molecular Cell, 2011, 42, 569-583.	4.5	89
22	Structure of the Recombinant N-Terminal Lobe of Human Lactoferrin at 2·0 à Resolution. Journal of Molecular Biology, 1993, 232, 1084-1100.	2.0	83
23	RING domain dimerization is essential for RNF4 function. Biochemical Journal, 2010, 431, 23-29.	1.7	80
24	Asymmetric Recruitment of cIAPs by TRAF2. Journal of Molecular Biology, 2010, 400, 8-15.	2.0	72
25	Intrinsically Disordered Proteins in Bcl-2 Regulated Apoptosis. International Journal of Molecular Sciences, 2010, 11, 1808-1824.	1.8	69
26	Localization of dynein light chains 1 and 2 and their pro-apoptotic ligands. Biochemical Journal, 2004, 377, 597-605.	1.7	65
27	Regulation of apoptosis: uncovering the binding determinants. Current Opinion in Structural Biology, 2005, 15, 690-699.	2.6	63
28	Altered Domain Closure and Iron Binding in Transferrins: The Crystal Structure of the Asp60Ser Mutant of the Amino-terminal Half-molecule of Human Lactoferrin. Journal of Molecular Biology, 1996, 256, 352-363.	2.0	62
29	Crystal structure of the amino-terminal coiled-coil domain of the APC tumor suppressor 1 1Edited by I. A. Wilson. Journal of Molecular Biology, 2000, 301, 147-156.	2.0	50
30	Regulation of ubiquitin transfer by XIAP, a dimeric RING E3 ligase. Biochemical Journal, 2013, 450, 629-638.	1.7	50
31	Reevaluation of Abscisic Acid-Binding Assays Shows That G-Protein-Coupled Receptor2 Does Not Bind Abscisic Acid. Plant Physiology, 2009, 150, 6-11.	2.3	48
32	Secondary ubiquitin-RING docking enhances Arkadia and Ark2C E3 ligase activity. Nature Structural and Molecular Biology, 2016, 23, 45-52.	3.6	46
33	The molecular basis of lysine 48 ubiquitin chain synthesis by Ube2K. Scientific Reports, 2015, 5, 16793.	1.6	43
34	IAPs: Modular regulators of cell signalling. Seminars in Cell and Developmental Biology, 2015, 39, 80-90.	2.3	43
35	The activity of TRAF RING homo- and heterodimers is regulated by zinc finger 1. Nature Communications, 2017, 8, 1788.	5.8	42
36	FCA does not bind abscisic acid. Nature, 2008, 456, E5-E6.	13.7	40

CATHERINE L DAY

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37	Solution Structure of Psb27 from Cyanobacterial Photosystem II,. Biochemistry, 2009, 48, 8771-8773.	1.2	40
38	The N-Terminal Extension of UBE2E Ubiquitin-Conjugating Enzymes Limits Chain Assembly. Journal of Molecular Biology, 2013, 425, 4099-4111.	2.0	39
39	Enhancing the peroxidase activity of cytochrome <i>c</i> by mutation of residue 41: implications for the peroxidase mechanism and cytochrome <i>c</i> release. Biochemical Journal, 2014, 458, 259-265.	1.7	38
40	Tumor Necrosis Factor (TNF) Signaling, but Not TWEAK (TNF-like Weak Inducer of Apoptosis)-triggered cIAP1 (Cellular Inhibitor of Apoptosis Protein 1) Degradation, Requires cIAP1 RING Dimerization and E2 Binding. Journal of Biological Chemistry, 2010, 285, 17525-17536.	1.6	37
41	A Direct Interaction with NEDD1 Regulates Î <sup>3</sup> -Tubulin Recruitment to the Centrosome. PLoS ONE, 2010, 5, e9618.	1.1	36
42	Mutation of Arginine 121 in Lactoferrin Destabilizes Iron Binding by Disruption of Anion Binding:Â Crystal Structures of R121S and R121E Mutantsâ€,â€j. Biochemistry, 1996, 35, 14473-14479.	1.2	34
43	FRIGIDA and related proteins have a conserved central domain and family specific N- and C- terminal regions that are functionally important. Plant Molecular Biology, 2010, 73, 493-505.	2.0	29
44	Noncovalent Ubiquitin Interactions Regulate the Catalytic Activity of Ubiquitin Writers. Trends in Biochemical Sciences, 2016, 41, 924-937.	3.7	27
45	A bidentate Polycomb Repressive-Deubiquitinase complex is required for efficient activity on nucleosomes. Nature Communications, 2018, 9, 3932.	5.8	25
46	The structure of Boo/Diva reveals a divergent Bcl-2 protein. Proteins: Structure, Function and Bioinformatics, 2010, 78, NA-NA.	1.5	24
47	Use of E2~Ubiquitin Conjugates for the Characterization of Ubiquitin Transfer by RING E3 Ligases Such as the Inhibitor of Apoptosis Proteins. Methods in Enzymology, 2014, 545, 243-263.	0.4	23
48	Structure and Function of the RING Domains of RNF20 and RNF40, Dimeric E3 Ligases that Monoubiquitylate Histone H2B. Journal of Molecular Biology, 2016, 428, 4073-4086.	2.0	23
49	The Structure and Ubiquitin Binding Properties of TRAF RING Heterodimers. Journal of Molecular Biology, 2021, 433, 166844.	2.0	20
50	Regulation of E2s: A Role for Additional Ubiquitin Binding Sites?. Journal of Molecular Biology, 2017, 429, 3430-3440.	2.0	17
51	Preliminary crystallographic studies of the amino terminal half of human lactoferrin in its iron-saturated and iron-free forms. Journal of Molecular Biology, 1992, 228, 973-974.	2.0	13
52	A cryptic tubulin-binding domain links MEKK1 to curved tubulin protomers. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 21308-21318.	3.3	12
53	Identification of Ubiquitin Variants That Inhibit the E2 Ubiquitin Conjugating Enzyme, Ube2k. ACS Chemical Biology, 2021, 16, 1745-1756.	1.6	12
54	Collaborative networks enable the rapid establishment of serological assays for SARS-CoV-2 during nationwide lockdown in New Zealand. PeerJ, 2020, 8, e9863.	0.9	12

CATHERINE L DAY

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55	HtrA—A Renaissance Protein. Structure, 2002, 10, 737-739.	1.6	10
56	Structure of a domain-opened mutant (R121D) of the human lactoferrin N-lobe refined from a merohedrally twinned crystal form. Acta Crystallographica Section D: Biological Crystallography, 2002, 58, 955-962.	2.5	10
57	The Ubiquitin-associated Domain of Cellular Inhibitor of Apoptosis Proteins Facilitates Ubiquitylation. Journal of Biological Chemistry, 2014, 289, 25721-25736.	1.6	10
58	The RING domain of RING Finger 11 ( RNF 11) protein binds Ubc13 and inhibits formation of polyubiquitin chains. FEBS Letters, 2018, 592, 1434-1444.	1.3	9
59	Ubiquitin and a charged loop regulate the ubiquitin E3 ligase activity of Ark2C. Nature Communications, 2022, 13, 1181.	5.8	8
60	E2 enzymes: lessons in ubiquitin transfer from XLID patients. Nature Chemical Biology, 2019, 15, 6-7.	3.9	2
61	Ubiquitin Variant Inhibitors Meet the Deubiquitinase USP15. Structure, 2019, 27, 564-565.	1.6	1
62	Solution structure of Mclâ€1 and its complexes. FASEB Journal, 2007, 21, A638.	0.2	0
63	Solution Structure and Physiological Requirements for Psb27 in Synechocystis sp. PCC 6803. Advanced Topics in Science and Technology in China, 2013, , 432-435.	0.0	Ο
64	Altered Domain Closure and Iron Binding in Lactoferrin Mutants. , 1997, , 25-38.		0