

Peter M Colman

List of Publications by Year in descending order

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93
papers

15,115
citations

41258

49
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43802

91
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129
all docs

129
docs citations

129
times ranked

13573
citing authors

#	ARTICLE	IF	CITATIONS
1	Rational design of potent sialidase-based inhibitors of influenza virus replication. <i>Nature</i> , 1993, 363, 418-423.	13.7	1,823
2	Differential Targeting of Prosurvival Bcl-2 Proteins by Their BH3-Only Ligands Allows Complementary Apoptotic Function. <i>Molecular Cell</i> , 2005, 17, 393-403.	4.5	1,639
3	Shape Complementarity at Protein/Protein Interfaces. <i>Journal of Molecular Biology</i> , 1993, 234, 946-950.	2.0	1,231
4	Conformations of immunoglobulin hypervariable regions. <i>Nature</i> , 1989, 342, 877-883.	13.7	1,199
5	BCL-2 family antagonists for cancer therapy. <i>Nature Reviews Drug Discovery</i> , 2008, 7, 989-1000.	21.5	549
6	Bax Crystal Structures Reveal How BH3 Domains Activate Bax and Nucleate Its Oligomerization to Induce Apoptosis. <i>Cell</i> , 2013, 152, 519-531.	13.5	491
7	Crystallographic structure studies of an IgG molecule and an Fc fragment. <i>Nature</i> , 1976, 264, 415-420.	13.7	423
8	The structural biology of type I viral membrane fusion. <i>Nature Reviews Molecular Cell Biology</i> , 2003, 4, 309-319.	16.1	408
9	The structure of the complex between influenza virus neuraminidase and sialic acid, the viral receptor. <i>Proteins: Structure, Function and Bioinformatics</i> , 1992, 14, 327-332.	1.5	399
10	Structural insights into the degradation of Mcl-1 induced by BH3 domains. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 6217-6222.	3.3	397
11	Structure-guided design of a selective BCL-XL inhibitor. <i>Nature Chemical Biology</i> , 2013, 9, 390-397.	3.9	324
12	To Trigger Apoptosis, Bak Exposes Its BH3 Domain and Homodimerizes via BH3:Groove Interactions. <i>Molecular Cell</i> , 2008, 30, 369-380.	4.5	296
13	The Dendritic Cell Receptor Clec9A Binds Damaged Cells via Exposed Actin Filaments. <i>Immunity</i> , 2012, 36, 646-657.	6.6	272
14	Electrostatic complementarity at protein/protein interfaces 1 Edited by B. Honig. <i>Journal of Molecular Biology</i> , 1997, 268, 570-584.	2.0	255
15	Discovery of a Potent and Selective BCL-X _L Inhibitor with <i>in Vivo</i> Activity. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 1088-1093.	1.3	242
16	Crystal and Molecular Structure of a Dimer Composed of the Variable Portions of the Bence-Jones Protein REI. <i>FEBS Journal</i> , 1974, 45, 513-524.	0.2	220
17	Drug design against a shifting target: a structural basis for resistance to inhibitors in a variant of influenza virus neuraminidase. <i>Structure</i> , 1998, 6, 735-746.	1.6	210
18	The Structure of the Fusion Glycoprotein of Newcastle Disease Virus Suggests a Novel Paradigm for the Molecular Mechanism of Membrane Fusion. <i>Structure</i> , 2001, 9, 255-266.	1.6	201

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19	Structure of the Haemagglutinin-neuraminidase from Human Parainfluenza Virus Type III. <i>Journal of Molecular Biology</i> , 2004, 335, 1343-1357.	2.0	200
20	Dihydropyranocarboxamides Related to Zanamivir: A New Series of Inhibitors of Influenza Virus Sialidases. 2. Crystallographic and Molecular Modeling Study of Complexes of 4-Amino-4H-pyran-6-carboxamides and Sialidase from Influenza Virus Types A and B. <i>Journal of Medicinal Chemistry</i> , 1998, 41, 798-807.	2.9	197
21	Three-dimensional structure of the complex of 4-guanidino-Neu5Ac2en and influenza virus neuraminidase. <i>Protein Science</i> , 1995, 4, 1081-1087.	3.1	176
22	Mutations in a Conserved Residue in the Influenza Virus Neuraminidase Active Site Decreases Sensitivity to Neu5Ac2en-Derived Inhibitors. <i>Journal of Virology</i> , 1998, 72, 2456-2462.	1.5	175
23	Structure of the human antibody molecule kol (immunoglobulin G1): An electron density map at 5 Å... resolution. <i>Journal of Molecular Biology</i> , 1976, 100, 257-278.	2.0	160
24	The structure of a complex between the NC10 antibody and influenza virus neuraminidase and comparison with the overlapping binding site of the NC41 antibody. <i>Structure</i> , 1994, 2, 733-746.	1.6	157
25	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, 341-355.	2.3	157
26	Generation and Characterization of an Influenza Virus Neuraminidase Variant with Decreased Sensitivity to the Neuraminidase-Specific Inhibitor 4-Guanidino-Neu5Ac2en. <i>Virology</i> , 1995, 214, 475-484.	1.1	155
27	Evaluation of Diverse \pm/\pm -Backbone Patterns for Functional \pm -Helix Mimicry: Analogues of the Bim BH3 Domain. <i>Journal of the American Chemical Society</i> , 2012, 134, 315-323.	6.6	144
28	High-Resolution Structural Characterization of a Helical \pm/\pm -Peptide Foldamer Bound to the Anti-Apoptotic Protein Bcl-2. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 4318-4322.	7.2	143
29	Bak Core and Latch Domains Separate during Activation, and Freed Core Domains Form Symmetric Homodimers. <i>Molecular Cell</i> , 2014, 55, 938-946.	4.5	140
30	Structures of BCL-2 in complex with venetoclax reveal the molecular basis of resistance mutations. <i>Nature Communications</i> , 2019, 10, 2385.	5.8	139
31	Structural Studies of the Resistance of Influenza Virus Neuraminidase to Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 2207-2212.	2.9	125
32	A Structural Viral Mimic of Prosurvival Bcl-2: A Pivotal Role for Sequestering Proapoptotic Bax and Bak. <i>Molecular Cell</i> , 2007, 25, 933-942.	4.5	125
33	The three-dimensional structure of N-acetylneuraminase lyase from Escherichia coli. <i>Structure</i> , 1994, 2, 361-369.	1.6	123
34	Crystallographic Structural Studies of a Human Fc Fragment. II. A Complete Model Based on a Fourier Map at 3.5 Å... Resolution. <i>Hoppe-Seyler's Zeitschrift für Physiologische Chemie</i> , 1976, 357, 1421-1434.	1.7	120
35	Recombinant anti-sialidase single-chain variable fragment antibody. Characterization, formation of dimer and higher-molecular-mass multimers and the solution of the crystal structure of the single-chain variable fragment/sialidase complex. <i>FEBS Journal</i> , 1994, 221, 151-157.	0.2	115
36	Targeting BCL-2-like Proteins to Kill Cancer Cells. <i>Trends in Cancer</i> , 2016, 2, 443-460.	3.8	114

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37	Structural Basis for Apoptosis Inhibition by Epstein-Barr Virus BHRF1. <i>PLoS Pathogens</i> , 2010, 6, e1001236.	2.1	99
38	Analysis of inhibitor binding in influenza virus neuraminidase. <i>Protein Science</i> , 2001, 10, 689-696.	3.1	97
39	Mutation to Bax beyond the BH3 Domain Disrupts Interactions with Pro-survival Proteins and Promotes Apoptosis. <i>Journal of Biological Chemistry</i> , 2011, 286, 7123-7131.	1.6	96
40	Conformational Changes in Bcl-2 Pro-survival Proteins Determine Their Capacity to Bind Ligands. <i>Journal of Biological Chemistry</i> , 2009, 284, 30508-30517.	1.6	79
41	New Antivirals and Drug Resistance. <i>Annual Review of Biochemistry</i> , 2009, 78, 95-118.	5.0	65
42	Crystal structure of the human Fab fragment Kol and its comparison with the intact Kol molecule. <i>Journal of Molecular Biology</i> , 1978, 121, 441-459.	2.0	62
43	Discovery of Potent and Selective Benzothiazole Hydrazone Inhibitors of Bcl-X _L . <i>Journal of Medicinal Chemistry</i> , 2013, 56, 5514-5540.	2.9	60
44	Recombinant antineuraminidase single chain antibody: Expression, characterization, and crystallization in complex with antigen. <i>Proteins: Structure, Function and Bioinformatics</i> , 1993, 16, 57-63.	1.5	58
45	Novel inhibitors of influenza sialidases related to GG167 structure-activity, crystallographic and Molecular dynamics studies with 4H-pyran-2-carboxylic acid 6-carboxamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1996, 6, 2931-2936.	1.0	58
46	Conversion of Bim-BH3 from Activator to Inhibitor of Bak through Structure-Based Design. <i>Molecular Cell</i> , 2017, 68, 659-672.e9.	4.5	57
47	Crystallographic Structural Studies of a Human Fc-Fragment. I. An Electron-Density Map at 4 Å... Resolution and a Partial Model. <i>Hoppe-Seyler's Zeitschrift für Physiologische Chemie</i> , 1976, 357, 435-446.	1.7	56
48	Structural Basis of Bcl-X _L Recognition by a BH3-Mimetic Peptide Generated by Sequence-Based Design. <i>ChemBioChem</i> , 2011, 12, 2025-2032.	1.3	56
49	Discovery and molecular characterization of a Bcl-2-regulated cell death pathway in schistosomes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 6999-7003.	3.3	53
50	BAK core dimers bind lipids and can be bridged by them. <i>Nature Structural and Molecular Biology</i> , 2020, 27, 1024-1031.	3.6	49
51	Sialidase inhibitors related to zanamivir. Further SAR studies of 4-amino-4H-pyran-2-carboxylic acid-6-propylamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2001, 11, 669-673.	1.0	46
52	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
53	Physiological restraint of Bak by Bcl-x _L is essential for cell survival. <i>Genes and Development</i> , 2016, 30, 1240-1250.	2.7	40
54	Structure of <i>Leishmania mexicana</i> Phosphomannomutase Highlights Similarities with Human Isoforms. <i>Journal of Molecular Biology</i> , 2006, 363, 215-227.	2.0	38

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55	Structural Insight into BH3 Domain Binding of Vaccinia Virus Antiapoptotic F1L. <i>Journal of Virology</i> , 2014, 88, 8667-8677.	1.5	37
56	Structure-Guided Rescaffolding of Selective Antagonists of BCL-X _L . <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 662-667.	1.3	37
57	Structural Insights into the Protease-like Antigen Plasmodium falciparum SERA5 and Its Noncanonical Active-Site Serine. <i>Journal of Molecular Biology</i> , 2009, 392, 154-165.	2.0	35
58	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
59	Ensemble Properties of Bax Determine Its Function. <i>Structure</i> , 2018, 26, 1346-1359.e5.	1.6	34
60	The Functional Differences between Pro-survival and Pro-apoptotic B Cell Lymphoma 2 (Bcl-2) Proteins Depend on Structural Differences in Their Bcl-2 Homology 3 (BH3) Domains. <i>Journal of Biological Chemistry</i> , 2014, 289, 36001-36017.	1.6	33
61	De-Novo Designed Library of Benzoylureas as Inhibitors of BCL-X _L : Synthesis, Structural and Biochemical Characterization. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 1323-1343.	2.9	33
62	Three-dimensional structures of single-chain Fv-neuraminidase complexes. <i>Journal of Molecular Biology</i> , 1998, 279, 901-910.	2.0	32
63	Cloning, Expression, and Crystallization of the Fusion Protein of Newcastle Disease Virus. <i>Virology</i> , 2001, 290, 290-299.	1.1	31
64	Preliminary crystallographic data for a basic copper-containing protein from cucumber seedlings. <i>Journal of Molecular Biology</i> , 1977, 112, 649-650.	2.0	30
65	The Structure of Cucurbitin: Subunit Symmetry and Organization in situ. <i>FEBS Journal</i> , 1980, 103, 585-588.	0.2	30
66	A small molecule interacts with VDAC2 to block mouse BAK-driven apoptosis. <i>Nature Chemical Biology</i> , 2019, 15, 1057-1066.	3.9	30
67	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
68	Modelling the structure of the fusion protein from human respiratory syncytial virus. <i>Protein Engineering, Design and Selection</i> , 2002, 15, 365-371.	1.0	28
69	Structure of glyceraldehyde-3-phosphate dehydrogenase from Plasmodium falciparum. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2005, 61, 1213-1221.	2.5	28
70	Dysglycemia and Index60 as Prediagnostic End Points for Type 1 Diabetes Prevention Trials. <i>Diabetes Care</i> , 2017, 40, 1494-1499.	4.3	28
71	Vaccinia Virus F1L Interacts with Bak Using Highly Divergent Bcl-2 Homology Domains and Replaces the Function of Mcl-1. <i>Journal of Biological Chemistry</i> , 2010, 285, 4695-4708.	1.6	26
72	Structure of detergent-activated BAK dimers derived from the inert monomer. <i>Molecular Cell</i> , 2021, 81, 2123-2134.e5.	4.5	26

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73	Crystal Structure of a BCL-W Domain-Swapped Dimer: Implications for the Function of BCL-2 Family Proteins. <i>Structure</i> , 2011, 19, 1467-1476.	1.6	25
74	Zanamivir: an influenza virus neuraminidase inhibitor. <i>Expert Review of Anti-Infective Therapy</i> , 2005, 3, 191-199.	2.0	24
75	Preliminary crystallographic data for a copper-containing protein, plastocyanin. <i>Journal of Molecular Biology</i> , 1977, 110, 187-189.	2.0	23
76	Tethered Neuraminidase Inhibitors That Bind an Influenza Virus: A First Step Towards a Diagnostic Method for Influenza. <i>Angewandte Chemie - International Edition</i> , 2003, 42, 3118-3121.	7.2	22
77	The Function of the Hypothalamicâ€Pituitaryâ€Adrenal Axis in Alzheimer's Disease. <i>British Journal of Psychiatry</i> , 1994, 165, 650-657.	1.7	18
78	The use of rotation and translation functions in the interpretation of low resolution electron density maps. <i>Journal of Molecular Biology</i> , 1976, 100, 278-282.	2.0	13
79	The disulphide bonds of an Asian influenza virus neuraminidase. <i>FEBS Letters</i> , 1983, 153, 29-33.	1.3	12
80	NMR studies of interactions between Bax and BH3 domain-containing peptides in the absence and presence of CHAPS. <i>Archives of Biochemistry and Biophysics</i> , 2014, 545, 33-43.	1.4	11
81	The Trypanosomal Trans-Sialidase. <i>Structure</i> , 2002, 10, 1466-1468.	1.6	10
82	Ion currents through Kir potassium channels are gated by anionic lipids. <i>Nature Communications</i> , 2022, 13, 490.	5.8	9
83	Insights into pituitary tumorigenesis: from Sanger sequencing to next-generation sequencing and beyond. <i>Expert Review of Endocrinology and Metabolism</i> , 2019, 14, 399-418.	1.2	8
84	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
85	Structure of the BAK-activating antibody 7D10 bound to BAK reveals an unexpected role for the $\pm 1\text{-}\hat{1}\pm 2$ loop in BAK activation. <i>Cell Death and Differentiation</i> , 2022, 29, 1757-1768.	5.0	4
86	Specificity and Promiscuity in Protein - Ligand and Protein - Protein Interactions. <i>Australian Journal of Chemistry</i> , 2003, 56, 763.	0.5	3
87	Early days in drug discovery by crystallography â€ personal recollections. <i>Acta Crystallographica Section A: Foundations and Advances</i> , 2013, 69, 60-62.	0.3	3
88	Optimization of Benzothiazole and Thiazole Hydrazones as Inhibitors of Schistosome BCL-2. <i>ACS Infectious Diseases</i> , 2021, 7, 1143-1163.	1.8	3
89	Crystallography and New Medicines: Examples from Influenza and Cell Death. <i>Australian Journal of Chemistry</i> , 2014, 67, 1720.	0.5	2
90	Intense focal pituitary FDG uptake due to intravascular large Bâ€cell lymphoma in pyrexia of unknown origin. <i>American Journal of Hematology</i> , 2016, 91, 1167-1168.	2.0	2

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91	The Bak core dimer focuses triacylglycerides in the membrane. <i>Biophysical Journal</i> , 2022, 121, 347-360.	0.2	1
92	Vale Colin Wardâ€™A Leader in Receptor Structural Biology. <i>Frontiers in Endocrinology</i> , 2017, 8, 95.	1.5	0
93	Colin Wesley Ward 1943â€™2017. <i>Historical Records of Australian Science</i> , 2018, 29, 191.	0.3	0