

Enrico Stura

List of Publications by Year in descending order

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140
papers

11,166
citations

36203

51
h-index

30010

103
g-index

140
all docs

140
docs citations

140
times ranked

10183
citing authors

#	ARTICLE	IF	CITATIONS
1	Crystal structures of two viral peptides in complex with murine MHC class I H-2Kb. <i>Science</i> , 1992, 257, 919-927.	6.0	877
2	Functional Mimicry of a Protein Hormone by a Peptide Agonist: The EPO Receptor Complex at 2.8 Å. <i>Science</i> , 1996, 273, 464-471.	6.0	621
3	Crystal Structure of Mouse CD1: An MHC-Like Fold with a Large Hydrophobic Binding Groove. <i>Science</i> , 1997, 277, 339-345.	6.0	596
4	Crystallographic Evidence for Preformed Dimers of Erythropoietin Receptor Before Ligand Activation. <i>Science</i> , 1999, 283, 987-990.	6.0	579
5	Structure of a flavivirus envelope glycoprotein in its low-pH-induced membrane fusion conformation. <i>EMBO Journal</i> , 2004, 23, 728-738.	3.5	526
6	Crystal structure of a <i>Staphylococcus aureus</i> protein A domain complexed with the Fab fragment of a human IgM antibody: Structural basis for recognition of B-cell receptors and superantigen activity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2000, 97, 5399-5404.	3.3	443
7	Immune Versus Natural Selection: Antibody Aldolases with Enzymic Rates But Broader Scope. <i>Science</i> , 1997, 278, 2085-2092.	6.0	402
8	Crystal structure of the principal neutralization site of HIV-1. <i>Science</i> , 1994, 264, 82-85.	6.0	265
9	Crystal structure of an H-2Kb-ovalbumin peptide complex reveals the interplay of primary and secondary anchor positions in the major histocompatibility complex binding groove. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1995, 92, 2479-2483.	3.3	240
10	Crystal Structure of Alkaline Phosphatase from Human Placenta at 1.8 Å... Resolution. <i>Journal of Biological Chemistry</i> , 2001, 276, 9158-9165.	1.6	234
11	Crystal structure of the human urokinase plasminogen activator receptor bound to an antagonist peptide. <i>EMBO Journal</i> , 2005, 24, 1655-1663.	3.5	213
12	Crystal structure of a human immunodeficiency virus type 1 neutralizing antibody, 50.1, in complex with its V3 loop peptide antigen. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1993, 90, 6325-6329.	3.3	211
13	An antagonist peptide-EPO receptor complex suggests that receptor dimerization is not sufficient for activation. <i>Nature Structural Biology</i> , 1998, 5, 993-1004.	9.7	204
14	Denmotoxin, a Three-finger Toxin from the Colubrid Snake <i>Boiga dendrophila</i> (Mangrove Catsnake) with Bird-specific Activity. <i>Journal of Biological Chemistry</i> , 2006, 281, 29030-29041.	1.6	183
15	Routes to catalysis: structure of a catalytic antibody and comparison with its natural counterpart. <i>Science</i> , 1994, 263, 646-652.	6.0	182
16	Structural Evidence for a Functional Role of Human Tissue Nonspecific Alkaline Phosphatase in Bone Mineralization. <i>Journal of Biological Chemistry</i> , 2001, 276, 31171-31178.	1.6	182
17	Structure and mutational analysis of Rab GDP-dissociation inhibitor. <i>Nature</i> , 1996, 381, 42-48.	13.7	169
18	Irditoxin, a novel covalently linked heterodimeric three-finger toxin with high taxon-specific neurotoxicity. <i>FASEB Journal</i> , 2009, 23, 534-545.	0.2	165

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19	Three-dimensional Structure of an Anti-steroid Fab ϵ^2 and Progesterone-Fab ϵ^2 Complex. <i>Journal of Molecular Biology</i> , 1993, 231, 103-118.	2.0	154
20	Applications of the streak seeding technique in protein crystallization. <i>Journal of Crystal Growth</i> , 1991, 110, 270-282.	0.7	147
21	Dual conformations for the HIV-1 gp120 V3 loop in complexes with different neutralizing Fabs. <i>Structure</i> , 1999, 7, 131-142.	1.6	145
22	Crystal structure of <i>Aplysia</i> ADP ribosyl cyclase, a homologue of the bifunctional ectozyme CD38. <i>Nature Structural Biology</i> , 1996, 3, 957-964.	9.7	142
23	An Antibody exo Diels-Alderase Inhibitor Complex at 1.95Å Resolution. <i>Science</i> , 1998, 279, 1934-1940.	6.0	141
24	T cell receptor interactions with syngeneic and allogeneic ligands: Affinity measurements and crystallization. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1997, 94, 13838-13843.	3.3	135
25	The CuA domain of <i>Thermus thermophilus</i> ba3-type cytochrome c oxidase at 1.6 Å resolution. <i>Nature Structural Biology</i> , 1999, 6, 509-516.	9.7	131
26	Complex between <i>Peptostreptococcus magnus</i> Protein L and a Human Antibody Reveals Structural Convergence in the Interaction Modes of Fab Binding Proteins. <i>Structure</i> , 2001, 9, 679-687.	1.6	121
27	Strategies in the crystallization of glycoproteins and protein complexes. <i>Journal of Crystal Growth</i> , 1992, 122, 273-285.	0.7	117
28	Structural Analysis of Antibody Specificity. <i>Journal of Molecular Biology</i> , 1994, 241, 663-690.	2.0	113
29	The mechanism of an inhibitory antibody on TF-initiated blood coagulation revealed by the crystal structures of human tissue factor, Fab 5G9 and TF \cdot 5G9 complex. Edited by D. C. Rees. <i>Journal of Molecular Biology</i> , 1998, 275, 873-894.	2.0	113
30	X-ray structure of the arenavirus glycoprotein GP2 in its postfusion hairpin conformation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 19967-19972.	3.3	101
31	Identification of a 13 Amino Acid Peptide Mimetic of Erythropoietin and Description of Amino Acids Critical for the Mimetic Activity of EMP1. <i>Biochemistry</i> , 1998, 37, 3699-3710.	1.2	99
32	Analytical and production seeding techniques. <i>Methods</i> , 1990, 1, 38-49.	1.9	98
33	Third generation of matrix metalloprotease inhibitors: Gain in selectivity by targeting the depth of the S1 ϵ^2 cavity. <i>Biochimie</i> , 2010, 92, 1501-1508.	1.3	88
34	Structural Studies of Human Placental Alkaline Phosphatase in Complex with Functional Ligands. <i>Journal of Molecular Biology</i> , 2005, 350, 441-451.	2.0	85
35	Structure of the complex between the Fab fragment of a neutralizing antibody for type 1 poliovirus and its viral epitope. <i>Nature Structural and Molecular Biology</i> , 1995, 2, 232-243.	3.6	83
36	Structural insights into the neutralization mechanism of a higher primate antibody against dengue virus. <i>EMBO Journal</i> , 2012, 31, 767-779.	3.5	81

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37	Crystal Structures of Mite Allergens Der f 1 and Der p 1 Reveal Differences in Surface-Exposed Residues that May Influence Antibody Binding. <i>Journal of Molecular Biology</i> , 2009, 386, 520-530.	2.0	79
38	Crystal structure of dUTP pyrophosphatase from feline immunodeficiency virus. <i>Protein Science</i> , 1996, 5, 2429-2437.	3.1	78
39	Crystal Structure of a Ternary Complex between Human Prostate-specific Antigen, Its Substrate Acyl Intermediate and an Activating Antibody. <i>Journal of Molecular Biology</i> , 2008, 376, 1021-1033.	2.0	78
40	Towards Structure-based Drug Design: Crystal Structure of a Multisubstrate Adduct Complex of Glycinamide Ribonucleotide Transformylase at 1.96 Å... Resolution. <i>Journal of Molecular Biology</i> , 1995, 249, 153-175.	2.0	77
41	Three-dimensional structure and IgE-binding properties of mature fully active Der p 1, a clinically relevant major allergen. <i>Journal of Allergy and Clinical Immunology</i> , 2006, 117, 571-576.	1.5	76
42	Crystal structure of glycinamide ribonucleotide transformylase from <i>Escherichia coli</i> at 3.0 Å... resolution. <i>Journal of Molecular Biology</i> , 1992, 227, 283-292.	2.0	65
43	Crystal Structure of a Peptide Complex of Anti-influenza Peptide Antibody Fab 26/9. <i>Journal of Molecular Biology</i> , 1994, 241, 534-556.	2.0	65
44	Crystal Structure of a Human Autoimmune Complex between IgM Rheumatoid Factor RF61 and IgG1 Fc Reveals a Novel Epitope and Evidence for Affinity Maturation. <i>Journal of Molecular Biology</i> , 2007, 368, 1321-1331.	2.0	61
45	NAD Binding Induces Conformational Changes in Rho ADP-ribosylating <i>Clostridium botulinum</i> C3 Exoenzyme. <i>Journal of Biological Chemistry</i> , 2002, 277, 30950-30957.	1.6	60
46	Proposals for the catalytic mechanism of glycogen phosphorylase b prompted by crystallographic studies on glucose 1-phosphate binding. <i>Journal of Molecular Biology</i> , 1980, 140, 565-580.	2.0	59
47	On the role of the cis-proline residue in the active site of DsbA. <i>Protein Science</i> , 1999, 8, 96-105.	3.1	58
48	Nucleotide binding to glycogen phosphorylase b in the crystal. <i>Journal of Molecular Biology</i> , 1979, 134, 639-653.	2.0	57
49	Crystal Structure of Human Prostate-Specific Antigen in a Sandwich Antibody Complex. <i>Journal of Molecular Biology</i> , 2011, 414, 530-544.	2.0	54
50	<i>N</i> -Isopropyl Sulfonamido-Based Hydroxamates as Matrix Metalloproteinase Inhibitors: Hit Selection and in Vivo Antiangiogenic Activity. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 7224-7240.	2.9	54
51	Comparison of AMP and NADH binding to glycogen phosphorylase b. <i>Journal of Molecular Biology</i> , 1983, 170, 529-565.	2.0	53
52	Crystal Structure of the Complex between the Monomeric Form of <i>Toxoplasma gondii</i> Surface Antigen 1 (SAG1) and a Monoclonal Antibody that Mimics the Human Immune Response. <i>Journal of Molecular Biology</i> , 2005, 354, 447-458.	2.0	50
53	Structure of <i>Azotobacter vinelandii</i> 7Fe ferredoxin at 1.35 Å... resolution and determination of the [Fe-S] bonds with 0.01 Å... accuracy. <i>Journal of Molecular Biology</i> , 1998, 278, 629-639.	2.0	48
54	Insights from Selective Non-phosphinic Inhibitors of MMP-12 Tailored to Fit with an S1 ^h Loop Canonical Conformation. <i>Journal of Biological Chemistry</i> , 2010, 285, 35900-35909.	1.6	48

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55	Molecular aspects of human Fcγ3R interactions with IgG: Functional and therapeutic consequences. <i>Immunology Letters</i> , 2006, 106, 111-118.	1.1	47
56	Crystallization of bi-functional ligand protein complexes. <i>Journal of Structural Biology</i> , 2013, 182, 246-254.	1.3	45
57	[7] X-ray crystallographic analysis of free and antigen-complexed Fab fragments to investigate structural basis of immune recognition. <i>Methods in Enzymology</i> , 1991, 203, 153-176.	0.4	41
58	Reverse screening. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 1994, 50, 448-455.	2.5	41
59	Mambalgin-1 Pain-relieving Peptide, Stepwise Solid-phase Synthesis, Crystal Structure, and Functional Domain for Acid-sensing Ion Channel 1a Inhibition. <i>Journal of Biological Chemistry</i> , 2016, 291, 2616-2629.	1.6	41
60	Discovery of a new selective inhibitor of A Disintegrin And Metalloprotease 10 (ADAM-10) able to reduce the shedding of NKG2D ligands in Hodgkin's lymphoma cell models. <i>European Journal of Medicinal Chemistry</i> , 2016, 111, 193-201.	2.6	40
61	Evidence for Plasticity and Structural Mimicry at the Immunoglobulin Light Chain-Protein L Interface. <i>Journal of Biological Chemistry</i> , 2002, 277, 47500-47506.	1.6	39
62	Fine Tuning of the Specificity of an Anti-progesterone Antibody by First and Second Sphere Residue Engineering. <i>Journal of Biological Chemistry</i> , 2005, 280, 24880-24887.	1.6	37
63	Molecular Determinants of a Selective Matrix Metalloprotease-12 Inhibitor: Insights from Crystallography and Thermodynamic Studies. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 1149-1159.	2.9	37
64	Immunoglobulin-binding domains: Protein L from <i>Peptostreptococcus magnus</i> . <i>Biochemical Society Transactions</i> , 2003, 31, 716-718.	1.6	36
65	Sugar-Based Arylsulfonamide Carboxylates as Selective and Water-Soluble Matrix Metalloproteinase-12 Inhibitors. <i>ChemMedChem</i> , 2016, 11, 1626-1637.	1.6	36
66	Simple Pseudo-dipeptides with a P2-Glutamate. <i>Journal of Biological Chemistry</i> , 2012, 287, 26647-26656.	1.6	35
67	Highly selective inhibition of myosin motors provides the basis of potential therapeutic application. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E7448-E7455.	3.3	34
68	Development of Thioaryl-Based Matrix Metalloproteinase-12 Inhibitors with Alternative Zinc-Binding Groups: Synthesis, Potentiometric, NMR, and Crystallographic Studies. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 4421-4435.	2.9	34
69	Crystallization of murine major histocompatibility complex class I H-2Kb with single peptides. <i>Journal of Molecular Biology</i> , 1992, 228, 975-982.	2.0	33
70	Transthyretin complexes with curcumin and bromo-estradiol: evaluation of solubilizing multicomponent mixtures. <i>New Biotechnology</i> , 2015, 32, 54-64.	2.4	33
71	Green mamba peptide targets type-2 vasopressin receptor against polycystic kidney disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, 7154-7159.	3.3	33
72	Structure of the 16S rRNA pseudouridine synthase RsuA bound to uracil and UMP. <i>Nature Structural Biology</i> , 2002, 9, 353-8.	9.7	32

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73	Structures of feline immunodeficiency virus dUTP pyrophosphatase and its nucleotide complexes in three crystal forms. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2000, 56, 1100-1109.	2.5	31
74	Crystal structure of full-length human collagenase 3 (MMP-13) with peptides in the active site defines exosites in the catalytic domain. <i>FASEB Journal</i> , 2013, 27, 4395-4405.	0.2	31
75	Macromolecular crystallography with synchrotron radiation. II. Results. <i>Journal of Applied Crystallography</i> , 1983, 16, 28-41.	1.9	30
76	Highly specific anti-estradiol antibodies: structural characterisation and binding diversity 1 Edited by I. A. Wilson. <i>Journal of Molecular Biology</i> , 2002, 315, 699-712.	2.0	30
77	Strategies for Protein Cryocrystallography. <i>Crystal Growth and Design</i> , 2014, 14, 427-435.	1.4	29
78	Structural studies of human alkaline phosphatase in complex with strontium: Implication for its secondary effect in bones. <i>Protein Science</i> , 2006, 15, 1691-1700.	3.1	28
79	Different Interactions between MT7 Toxin and the Human Muscarinic M ₁ Receptor in Its Free and N-Methylscopolamine-Occupied States. <i>Molecular Pharmacology</i> , 2008, 74, 1554-1563.	1.0	28
80	Crystal Structure of a Hydrophobic Immunodominant Antigenic Site on Hepatitis C Virus Core Protein Complexed to Monoclonal Antibody 19D9D6. <i>Journal of Immunology</i> , 2003, 170, 1917-1924.	0.4	27
81	Engineering of Three-Finger Fold Toxins Creates Ligands with Original Pharmacological Profiles for Muscarinic and Adrenergic Receptors. <i>PLoS ONE</i> , 2012, 7, e39166.	1.1	27
82	Comparison of helical scan and standard rotation methods in single-crystal X-ray data collection strategies. <i>Journal of Synchrotron Radiation</i> , 2017, 24, 42-52.	1.0	27
83	Zinc-Metalloproteinase Inhibitors: Evaluation of the Complex Role Played by the Zinc-Binding Group on Potency and Selectivity. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 403-414.	2.9	27
84	Synthesis and in Vitro and in Vivo Evaluation of MMP-12 Selective Optical Probes. <i>Bioconjugate Chemistry</i> , 2016, 27, 2407-2417.	1.8	26
85	Copper mediated amyloid- β^2 binding to Transthyretin. <i>Scientific Reports</i> , 2018, 8, 13744.	1.6	26
86	Practical Use of Glycerol in Protein Crystallization. <i>Crystal Growth and Design</i> , 2011, 11, 2755-2762.	1.4	25
87	Structural basis for the NAD ⁺ hydrolysis mechanism and the ARTT-loop plasticity of C3 exoenzymes. <i>Protein Science</i> , 2008, 17, 878-886.	3.1	24
88	Halogen Bonding Controls Selectivity of FRET Substrate Probes for MMP-9. <i>Chemistry and Biology</i> , 2014, 21, 408-413.	6.2	24
89	Crystallization of Antibodies and Antibody-Antigen Complexes. <i>ImmunoMethods</i> , 1993, 3, 164-179.	0.8	22
90	Multicomponent mixtures for cryoprotection and ligand solubilization. <i>Biotechnology Reports (Amsterdam, Netherlands)</i> , 2015, 7, 120-127.	2.1	22

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91	Structural Aspects of Antibodies and Antibody-Antigen Complexes. Novartis Foundation Symposium, 1991, 159, 13-39.	1.2	21
92	Observation and Characterization of the Interaction between a Single Immunoglobulin Binding Domain of Protein L and Two Equivalents of Human λ Light Chains. Journal of Biological Chemistry, 2004, 279, 9370-9378.	1.6	20
93	A Tandem of SH3-like Domains Participates in RNA Binding in KIN17, a Human Protein Activated in Response to Genotoxics. Journal of Molecular Biology, 2006, 364, 764-776.	2.0	20
94	Purification, sequence and crystallization of an anti-tissue factor Fab and its use for the crystallization of tissue factor. Journal of Crystal Growth, 1992, 122, 253-264.	0.7	19
95	Crystallization of an intact monoclonal antibody (4B7) against Plasmodium falciparum malaria with peptides from the Pfs25 protein antigen. Acta Crystallographica Section D: Biological Crystallography, 1994, 50, 556-562.	2.5	19
96	Engineering protein for X-ray crystallography: The murine major histocompatibility complex class II molecule λ A ^d . Protein Science, 1998, 7, 413-418.	3.1	18
97	A new crystal form of human transthyretin obtained with a curcumin derived ligand. Journal of Structural Biology, 2016, 194, 8-17.	1.3	18
98	Crystallization and preliminary crystallographic data for an antiprogesterone monoclonal antibody Fab ² and steroid-Fab ² complexes. Journal of Molecular Biology, 1987, 193, 229-231.	2.0	17
99	Protein L mutants for the crystallization of antibody fragments. Acta Crystallographica Section D: Biological Crystallography, 2002, 58, 1744-1748.	2.5	17
100	Structure-Based Secondary Structure-Independent Approach To Design Protein Ligands: Application to the Design of Kv1.2 Potassium Channel Blockers. Journal of the American Chemical Society, 2006, 128, 16190-16205.	6.6	17
101	Human TTR conformation altered by rhenium tris-carbonyl derivatives. Journal of Structural Biology, 2016, 195, 353-364.	1.3	17
102	Structural Framework for Covalent Inhibition of Clostridium botulinum Neurotoxin A by Targeting Cys165. Journal of Biological Chemistry, 2012, 287, 33607-33614.	1.6	16
103	Epitaxial jumps. Journal of Crystal Growth, 1999, 196, 250-260.	0.7	15
104	Synthesis and structural analysis of halogen substituted fibril formation inhibitors of Human Transthyretin (TTR). Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 40-51.	2.5	15
105	Ancestral protein resurrection and engineering opportunities of the mamba aminergic toxins. Scientific Reports, 2017, 7, 2701.	1.6	15
106	Screening Using Polymorphs for the Crystallization of Protein-Ligand Complexes. Crystal Growth and Design, 2013, 13, 1878-1888.	1.4	14
107	Bifunctional Inhibitors as a New Tool To Reduce Cancer Cell Invasion by Impairing MMP-9 Homodimerization. ACS Medicinal Chemistry Letters, 2017, 8, 293-298.	1.3	13
108	Crystallization of macromolecular complexes: combinatorial complex crystallization. Journal of Crystal Growth, 2001, 232, 573-579.	0.7	12

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109	In Vitro Affinity Maturation of an Anti-PSA Antibody for Prostate Cancer Diagnostic Assay. <i>Journal of Molecular Biology</i> , 2011, 414, 545-562.	2.0	12
110	Scaffolds for protein crystallisation. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2002, 58, 1715-1721.	2.5	11
111	Grafting of functional motifs onto protein scaffolds identified by <scp>PDB</scp> screening â€œ an efficient route to design optimizable protein binders. <i>FEBS Journal</i> , 2013, 280, 139-159.	2.2	11
112	Diversified targets of FKBP25 and its complex with rapamycin. <i>International Journal of Biological Macromolecules</i> , 2014, 69, 344-352.	3.6	11
113	A B-Cell Superantigen that Targets B-1 Lymphocytes. <i>Current Topics in Microbiology and Immunology</i> , 2000, 252, 251-263.	0.7	11
114	Crystallization, sequence and preliminary crystallographic data for transmission-blocking anti-malaria Fab 4B7 with cyclic peptides from the Pfs25 protein of <i>P. falciparum</i> . <i>Acta Crystallographica Section D: Biological Crystallography</i> , 1994, 50, 535-542.	2.5	10
115	Effect of zinc on human IgG1 and its FcÎ³R interactions. <i>Immunology Letters</i> , 2012, 143, 60-69.	1.1	10
116	X-ray crystal structure and activity of fluorenyl-based compounds as transthyretin fibrillogenesis inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 824-833.	2.5	10
117	Crystallization studies of glycosylated and unglycosylated human recombinant interleukin-2. <i>Proteins: Structure, Function and Bioinformatics</i> , 1992, 12, 24-30.	1.5	9
118	Crystallization and preliminary crystallographic data for class I deoxyribose-5-phosphate aldolase from <i>Escherichia coli</i> : An Application of Reverse Screening. <i>Proteins: Structure, Function and Bioinformatics</i> , 1995, 22, 67-72.	1.5	8
119	Crystallization of macromolecular complexes. <i>Journal of Crystal Growth</i> , 2001, 232, 580-590.	0.7	8
120	A simple modification of the Q-plate for parallel screening and combinatorial crystallization. <i>Journal of Crystal Growth</i> , 2001, 232, 545-552.	0.7	8
121	Crystallization and preliminary structural studies of a chorismate mutase catalytic antibody complexed with a transition state analog. <i>Proteins: Structure, Function and Bioinformatics</i> , 1994, 18, 198-200.	1.5	7
122	Crystallization and halide phasing of the C-terminal domain of human KIN17. <i>Acta Crystallographica Section F: Structural Biology Communications</i> , 2006, 62, 245-248.	0.7	7
123	Acid pH crystallization of the basic protein lysin from the spermatozoa of red abalone (<i>Haliotis</i>) Tj ETQq1 1 0.784314.rgBT /Overlock 10	2.5	6
124	Heparin-aggregated RANTES can be crystallised. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2002, 58, 1670-1673.	2.5	6
125	Crystallization and Preliminary Crystallographic Data for Rab Guanine Nucleotide Dissociation Inhibitor (RabGDI) from Bovine Brain. <i>Journal of Molecular Biology</i> , 1994, 244, 469-473.	2.0	5
126	Comparison of the crystallization and crystal packing of two Fab single-site mutant protein L complexes. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2005, 61, 750-754.	2.5	5

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127	Crystallization and preliminary X-ray diffraction data of the complex between human centrin 2 and a peptide from the protein XPC. <i>Acta Crystallographica Section F: Structural Biology Communications</i> , 2006, 62, 649-651.	0.7	5
128	Improving Diffraction from 3 to 2 Å... for a Complex between a Small GTPase and Its Effector by Analysis of Crystal Contacts and Use of Reverse Screening. <i>Crystal Growth and Design</i> , 2007, 7, 2140-2146.	1.4	5
129	Conformational Exchange Is Critical for the Productivity of an Oxidative Folding Intermediate with Buried Free Cysteines. <i>Journal of Molecular Biology</i> , 2010, 403, 299-312.	2.0	5
130	Crystallization of recombinant green mamba α -D α 1a toxin during a lyophilization procedure and its structure determination. <i>Acta Crystallographica Section F: Structural Biology Communications</i> , 2013, 69, 704-709.	0.7	5
131	Crystallization and preliminary crystallographic data for a ternary complex between tissue factor, factor VIIa and a BPTI-derived inhibitor. <i>Journal of Crystal Growth</i> , 1996, 168, 260-269.	0.7	4
132	Sequence, specificity and crystallization of an oestrone- β -glucuronide antibody (3910). <i>Immunology</i> , 1997, 90, 632-639.	2.0	4
133	Crystallization of human complement component C5. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 1998, 54, 643-646.	2.5	4
134	Crystallization and preliminary crystallographic investigations of avian 5-aminoimidazole-4-carboxamide ribonucleotide transformylase- α -inosine monophosphate cyclohydrolase expressed in <i>Escherichia coli</i> . <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2000, 56, 1051-1054.	2.5	4
135	Different crystal packing in Fab- α -protein L semi-disordered peptide complex. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2005, 61, 744-749.	2.5	3
136	MT9, a natural peptide from black mamba venom antagonizes the muscarinic type 2 receptor and reverses the M2R-agonist-induced relaxation in rat and human arteries. <i>Biomedicine and Pharmacotherapy</i> , 2022, 150, 113094.	2.5	3
137	Disulphide dimerised peptide creates a crystal contact in an anti-peptide antibody. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2002, 58, 1740-1743.	2.5	1
138	Protein crystallization for drug design in the last 50 years. <i>Arbor</i> , 2015, 191, a222.	0.1	1
139	A peptide mimetic of erythropoietin: Critical residues and description of a minimal functional epitope. , 2002, , 501-503.		0
140	Peptide mimetics of erythropoietin are powerful probes of receptor activation mechanisms. , 2002, , 536-538.		0