Laurence H Pearl

List of Publications by Year in descending order

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		4960	6300
223	26,920	84	158
papers	citations	h-index	g-index
251	251	251	23432
all docs	docs citations	times ranked	citing authors

LAUDENCE H DEADL

#	Article	IF	CITATIONS
1	Structure of the human RAD17–RFC clamp loader and 9–1–1 checkpoint clamp bound to a dsDNA–ssDNA junction. Nucleic Acids Research, 2022, 50, 8279-8289.	14.5	13
2	Uncovering an allosteric mode of action for a selective inhibitor of human Bloom syndrome protein. ELife, 2021, 10, .	6.0	18
3	Structural basis for recruitment of the CHK1 DNA damage kinase by the CLASPIN scaffold protein. Structure, 2021, 29, 531-539.e3.	3.3	8
4	Structure of the TELO2-TTI1-TTI2 complex and its function in TOR recruitment to the R2TP chaperone. Cell Reports, 2021, 36, 109317.	6.4	20
5	Phosphorylation-dependent assembly of DNA damage response systems and the central roles of TOPBP1. DNA Repair, 2021, 108, 103232.	2.8	21
6	CK2 Phosphorylation of Human Papillomavirus 16 E2 on Serine 23 Promotes Interaction with TopBP1 and Is Critical for E2 Interaction with Mitotic Chromatin and the Viral Life Cycle. MBio, 2021, 12, e0116321.	4.1	16
7	Solution structure of the Hop TPR2A domain and investigation of target druggability by NMR, biochemical and in silico approaches. Scientific Reports, 2020, 10, 16000.	3.3	8
8	The structure-function relationship of oncogenic LMTK3. Science Advances, 2020, 6, .	10.3	18
9	Modeling of a 14 kDa RUVBL2-Binding Domain with Medium Resolution Cryo-EM Density. Journal of Chemical Information and Modeling, 2020, 60, 2541-2551.	5.4	3
10	Efficient Single-Strand Break Repair Requires Binding to Both Poly(ADP-Ribose) and DNA by the Central BRCT Domain of XRCC1. Cell Reports, 2019, 26, 573-581.e5.	6.4	58
11	Structural mechanism for regulation of the AAA-ATPases RUVBL1-RUVBL2 in the R2TP co-chaperone revealed by cryo-EM. Science Advances, 2019, 5, eaaw1616.	10.3	33
12	MDC1 Interacts with TOPBP1 to Maintain Chromosomal Stability during Mitosis. Molecular Cell, 2019, 74, 571-583.e8.	9.7	97
13	Phosphorylation-mediated interactions with TOPBP1 couple 53BP1 and 9-1-1 to control the G1 DNA damage checkpoint. ELife, 2019, 8, .	6.0	40
14	RPAP3 provides a flexible scaffold for coupling HSP90 to the human R2TP co-chaperone complex. Nature Communications, 2018, 9, 1501.	12.8	54
15	Advances on the Structure of the R2TP/Prefoldin-like Complex. Advances in Experimental Medicine and Biology, 2018, 1106, 73-83.	1.6	15
16	BRCT domains of the DNA damage checkpoint proteins TOPBP1/Rad4 display distinct specificities for phosphopeptide ligands. ELife, 2018, 7, .	6.0	34
17	Specialized interfaces of Smc5/6 control hinge stability and DNA association. Nature Communications, 2017, 8, 14011.	12.8	61
18	HECTD3 Mediates an HSP90-Dependent Degradation Pathway for Protein Kinase Clients. Cell Reports, 2017, 19, 2515-2528.	6.4	23

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19	Differential Regulation of G1 CDK Complexes by the Hsp90-Cdc37 Chaperone System. Cell Reports, 2017, 21, 1386-1398.	6.4	49
20	The Structure of the R2TP Complex Defines a Platform for Recruiting Diverse Client Proteins to the HSP90 Molecular Chaperone System. Structure, 2017, 25, 1145-1152.e4.	3.3	48
21	The structure of FKBP38 in complex with the MEEVD tetratricopeptide binding-motif of Hsp90. PLoS ONE, 2017, 12, e0173543.	2.5	25
22	A first generation inhibitor of human Greatwall kinase, enabled by structural and functional characterisation of a minimal kinase domain construct. Oncotarget, 2016, 7, 71182-71197.	1.8	30
23	Mode of action of DNA-competitive small molecule inhibitors of tyrosyl DNA phosphodiesterase 2. Biochemical Journal, 2016, 473, 1869-1879.	3.7	30
24	Review: The HSP90 molecular chaperone—an enigmatic ATPase. Biopolymers, 2016, 105, 594-607.	2.4	144
25	The Ku-binding motif is a conserved module for recruitment and stimulation of non-homologous end-joining proteins. Nature Communications, 2016, 7, 11242.	12.8	57
26	PARP3 is a sensor of nicked nucleosomes and monoribosylates histone H2BGlu2. Nature Communications, 2016, 7, 12404.	12.8	60
27	Cooperation of local motions in the Hsp90 molecular chaperone ATPase mechanism. Nature Chemical Biology, 2016, 12, 628-635.	8.0	68
28	Human BRCA1–BARD1 ubiquitin ligase activity counteracts chromatin barriers to DNA resection. Nature Structural and Molecular Biology, 2016, 23, 647-655.	8.2	222
29	DNA repair, genome stability and cancer: a historical perspective. Nature Reviews Cancer, 2016, 16, 35-42.	28.4	575
30	Destabilized SMC5/6 complex leads to chromosome breakage syndrome with severe lung disease. Journal of Clinical Investigation, 2016, 126, 2881-2892.	8.2	65
31	ATM Localization and Heterochromatin Repair Depend on Direct Interaction of the 53BP1-BRCT 2 Domain with Î ³ H2AX. Cell Reports, 2015, 13, 2081-2089.	6.4	61
32	Restricting direct interaction of CDC37 with HSP90 does not compromise chaperoning of client proteins. Oncogene, 2015, 34, 15-26.	5.9	39
33	Therapeutic opportunities within the DNA damage response. Nature Reviews Cancer, 2015, 15, 166-180.	28.4	442
34	Tah1 helix-swap dimerization prevents mixed Hsp90 co-chaperone complexes. Acta Crystallographica Section D: Biological Crystallography, 2015, 71, 1197-1206.	2.5	13
35	Design and discovery of 3-aryl-5-substituted-isoquinolin-1-ones as potent tankyrase inhibitors. MedChemComm, 2015, 6, 1687-1692.	3.4	11
36	Development of an oligonucleotide-based fluorescence assay for the identification of tyrosyl-DNA phosphodiesterase 1 (TDP1) inhibitors. Analytical Biochemistry, 2014, 454, 17-22.	2.4	14

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37	Structural Basis for Phosphorylation-Dependent Recruitment of Tel2 to Hsp90 by Pih1. Structure, 2014, 22, 805-818.	3.3	86
38	Phosphorylation-Dependent Assembly and Coordination of the DNA Damage Checkpoint Apparatus by Rad4TopBP1. Molecular Cell, 2013, 51, 723-736.	9.7	27
39	3,6-Diamino-4-(2-halophenyl)-2-benzoylthieno[2,3- <i>b</i>]pyridine-5-carbonitriles Are Selective Inhibitors of Plasmodium falciparum Glycogen Synthase Kinase-3. Journal of Medicinal Chemistry, 2013, 56, 264-275.	6.4	54
40	ATP-competitive inhibitors block protein kinase recruitment to the Hsp90-Cdc37 system. Nature Chemical Biology, 2013, 9, 307-312.	8.0	132
41	A Mechanism for the Inhibition of DNA-PK-Mediated DNA Sensing by a Virus. PLoS Pathogens, 2013, 9, e1003649.	4.7	94
42	The BAH domain of Rsc2 is a histone H3 binding domain. Nucleic Acids Research, 2013, 41, 9168-9182.	14.5	31
43	Visualization of a DNA-PK/PARP1 complex. Nucleic Acids Research, 2012, 40, 4168-4177.	14.5	89
44	The zinc-finger domains of PARP1 cooperate to recognize DNA strand breaks. Nature Structural and Molecular Biology, 2012, 19, 685-692.	8.2	214
45	Engineering human MEK-1 for structural studies: A case study of combinatorial domain hunting. Journal of Structural Biology, 2012, 177, 329-334.	2.8	19
46	Selectivity, Cocrystal Structures, and Neuroprotective Properties of Leucettines, a Family of Protein Kinase Inhibitors Derived from the Marine Sponge Alkaloid Leucettamine B. Journal of Medicinal Chemistry, 2012, 55, 9312-9330.	6.4	174
47	CCT241533 Is a Potent and Selective Inhibitor of CHK2 that Potentiates the Cytotoxicity of PARP Inhibitors. Cancer Research, 2011, 71, 463-472.	0.9	96
48	Targeting the Hsp90 Molecular Chaperone with Novel Macrolactams. Synthesis, Structural, Binding, and Cellular Studies. ACS Chemical Biology, 2011, 6, 1339-1347.	3.4	27
49	Threonine 22 Phosphorylation Attenuates Hsp90 Interaction with Cochaperones and Affects Its Chaperone Activity. Molecular Cell, 2011, 41, 672-681.	9.7	146
50	The Structural Basis for Substrate Recognition by Mammalian Polynucleotide Kinase 3′ Phosphatase. Molecular Cell, 2011, 44, 385-396.	9.7	32
51	Structure of the Ire1 autophosphorylation complex and implications for the unfolded protein response. EMBO Journal, 2011, 30, 894-905.	7.8	201
52	Regulation of DNA Replication through Sld3-Dpb11 Interaction Is Conserved from Yeast to Humans. Current Biology, 2011, 21, 1152-1157.	3.9	135
53	Structure-Based Design of Potent and Selective 2-(Quinazolin-2-yl)phenol Inhibitors of Checkpoint Kinase 2. Journal of Medicinal Chemistry, 2011, 54, 580-590.	6.4	46
54	Features of the <i>Streptomyces hygroscopicus</i> HtpG reveal how partial geldanamycin resistance can arise with mutation to the ATP binding pocket of a eukaryotic Hsp90. FASEB Journal, 2011, 25, 3828-3837.	0.5	32

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55	A combinatorial method to enable detailed investigation of protein–protein interactions. Future Medicinal Chemistry, 2011, 3, 271-282.	2.3	5
56	Structure and function of the Rad9-binding region of the DNA-damage checkpoint adaptor TopBP1. Nucleic Acids Research, 2011, 39, 313-324.	14.5	72
57	Cyclin-Cyclin-dependent Kinase Regulatory Response Is Linked to Substrate Recognition. Journal of Biological Chemistry, 2011, 286, 9713-9725.	3.4	3
58	p185, an Immunodominant Epitope, Is an Autoantigen Mimotope. Journal of Biological Chemistry, 2011, 286, 26220-26227.	3.4	5
59	Evidence for a remodelling of DNA-PK upon autophosphorylation from electron microscopy studies. Nucleic Acids Research, 2011, 39, 5757-5767.	14.5	20
60	The crystal structure of yeast CCT reveals intrinsic asymmetry of eukaryotic cytosolic chaperonins. EMBO Journal, 2011, 30, 3078-3090.	7.8	94
61	An Artemis polymorphic variant reduces Artemis activity and confers cellular radiosensitivity. DNA Repair, 2010, 9, 1003-1010.	2.8	33
62	Inhibition of Hsp90 with Resorcylic Acid Macrolactones: Synthesis and Binding Studies. Chemistry - A European Journal, 2010, 16, 10366-10372.	3.3	22
63	Identification and characterisation of 2-aminopyridine inhibitors of checkpoint kinase 2. Bioorganic and Medicinal Chemistry, 2010, 18, 707-718.	3.0	50
64	Detection of the ATPase Activity of the Molecular Chaperones Hsp90 and Hsp72 Using the Transcreenerâ"¢ ADP Assay Kit. Journal of Biomolecular Screening, 2010, 15, 279-286.	2.6	29
65	Swe1Wee1-Dependent Tyrosine Phosphorylation of Hsp90 Regulates Distinct Facets of Chaperone Function. Molecular Cell, 2010, 37, 333-343.	9.7	165
66	Structural Basis for Assembly of Hsp90-Sgt1-CHORD Protein Complexes: Implications for Chaperoning of NLR Innate Immunity Receptors. Molecular Cell, 2010, 39, 269-281.	9.7	108
67	Specific recognition of a multiply phosphorylated motif in the DNA repair scaffold XRCC1 by the FHA domain of human PNK. Nucleic Acids Research, 2009, 37, 1701-1712.	14.5	75
68	MoKCa database—mutations of kinases in cancer. Nucleic Acids Research, 2009, 37, D824-D831.	14.5	51
69	Electron microscopy of Xrcc4 and the DNA ligase IV–Xrcc4 DNA repair complex. DNA Repair, 2009, 8, 1380-1389.	2.8	24
70	Structural basis for recruitment of BRCA2 by PALB2. EMBO Reports, 2009, 10, 990-996.	4.5	154
71	A common conformationally coupled ATPase mechanism for yeast and human cytoplasmic HSP90s. FEBS Journal, 2009, 276, 199-209.	4.7	51
72	Structural–Thermodynamic Relationships of Interactions in the N-Terminal ATP-Binding Domain of Hsp90. Journal of Molecular Biology, 2009, 392, 923-936.	4.2	19

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73	Structural Insights into Formation of an Active Signaling Complex between Rac and Phospholipase C Gamma 2. Molecular Cell, 2009, 34, 223-233.	9.7	67
74	Crystal Structure of the Rad9-Rad1-Hus1 DNA Damage Checkpoint Complex—Implications for Clamp Loading and Regulation. Molecular Cell, 2009, 34, 735-745.	9.7	112
75	Structural Basis of the Radicicol Resistance Displayed by a Fungal Hsp90. ACS Chemical Biology, 2009, 4, 289-297.	3.4	53
76	Structural and functional coupling of Hsp90- and Sgt1-centred multi-protein complexes. EMBO Journal, 2008, 27, 2789-2798.	7.8	104
77	Activation segment dimerization: a mechanism for kinase autophosphorylation of non-consensus sites. EMBO Journal, 2008, 27, 704-714.	7.8	147
78	4,5-Diarylisoxazole Hsp90 Chaperone Inhibitors: Potential Therapeutic Agents for the Treatment of Cancer. Journal of Medicinal Chemistry, 2008, 51, 196-218.	6.4	386
79	Hsp90-Dependent Activation of Protein Kinases Is Regulated by Chaperone-Targeted Dephosphorylation of Cdc37. Molecular Cell, 2008, 31, 886-895.	9.7	184
80	Optimizing Natural Products by Biosynthetic Engineering: Discovery of Nonquinone Hsp90 Inhibitors. Journal of Medicinal Chemistry, 2008, 51, 5494-5497.	6.4	79
81	Molecular Characterization of Macbecin as an Hsp90 Inhibitor. Journal of Medicinal Chemistry, 2008, 51, 2853-2857.	6.4	56
82	The Hsp90 molecular chaperone: an open and shut case for treatment. Biochemical Journal, 2008, 410, 439-453.	3.7	410
83	The ATPase-dependent chaperoning activity of Hsp90a regulates thick filament formation and integration during skeletal muscle myofibrillogenesis. Development (Cambridge), 2008, 135, 1147-1156.	2.5	94
84	Structural and functional analysis of the Crb2–BRCT ₂ domain reveals distinct roles in checkpoint signaling and DNA damage repair. Genes and Development, 2008, 22, 2034-2047.	5.9	65
85	NVP-AUY922: A Novel Heat Shock Protein 90 Inhibitor Active against Xenograft Tumor Growth, Angiogenesis, and Metastasis. Cancer Research, 2008, 68, 2850-2860.	0.9	433
86	Chaperone ligand-discrimination by the TPR-domain protein Tah1. Biochemical Journal, 2008, 413, 261-268.	3.7	46
87	Inhibition of the heat shock protein 90 molecular chaperone in vitro and in vivo by novel, synthetic, potent resorcinylic pyrazole/isoxazole amide analogues. Molecular Cancer Therapeutics, 2007, 6, 1198-1211.	4.1	141
88	Crystal Structure of the Retinoblastoma Protein N Domain Provides Insight into Tumor Suppression, Ligand Interaction, and Holoprotein Architecture. Molecular Cell, 2007, 28, 371-385.	9.7	68
89	In vitro Biological Characterization of a Novel, Synthetic Diaryl Pyrazole Resorcinol Class of Heat Shock Protein 90 Inhibitors. Cancer Research, 2007, 67, 2206-2216.	0.9	111
90	Insights into histone code syntax from structural and biochemical studies of CARM1 methyltransferase. EMBO Journal, 2007, 26, 4402-4412.	7.8	117

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91	Structural model of fullâ€length human Ku70–Ku80 heterodimer and its recognition of DNA and DNAâ€PKcs. EMBO Reports, 2007, 8, 56-62.	4.5	111
92	Activation segment exchange: a common mechanism of kinase autophosphorylation?. Trends in Biochemical Sciences, 2007, 32, 351-356.	7.5	86
93	Structure and Mechanism of the Hsp90 Molecular Chaperone Machinery. Annual Review of Biochemistry, 2006, 75, 271-294.	11.1	988
94	Structural and Mechanistic Insights into Ras Association Domains of Phospholipase C Epsilon. Molecular Cell, 2006, 21, 495-507.	9.7	129
95	Three-Dimensional Structure of the Human DNA-PKcs/Ku70/Ku80 Complex Assembled on DNA and Its Implications for DNA DSB Repair. Molecular Cell, 2006, 22, 511-519.	9.7	223
96	Molecular Recognition of Transcriptional Repressor Motifs by the WD Domain of the Groucho/TLE Corepressor. Molecular Cell, 2006, 22, 645-655.	9.7	134
97	Structure of an Hsp90-Cdc37-Cdk4 Complex. Molecular Cell, 2006, 23, 697-707.	9.7	288
98	Structure of an archaeal PCNA1–PCNA2–FEN1 complex: elucidating PCNA subunit and client enzyme specificity. Nucleic Acids Research, 2006, 34, 4515-4526.	14.5	64
99	A novel expression system of domain I of human beta2 glycoprotein I in Escherichia coli. BMC Biotechnology, 2006, 6, 8.	3.3	33
100	Crystal structure of an Hsp90–nucleotide–p23/Sba1 closed chaperone complex. Nature, 2006, 440, 1013-1017.	27.8	857
101	Trans-activation of the DNA-damage signalling protein kinase Chk2 by T-loop exchange. EMBO Journal, 2006, 25, 3179-3190.	7.8	131
102	Inhibition of Hsp90 with Synthetic Macrolactones: Synthesis and Structural and Biological Evaluation of Ring and Conformational Analogs of Radicicol. Chemistry and Biology, 2006, 13, 1203-1215.	6.0	64
103	Combinatorial Domain Hunting: An effective approach for the identification of soluble protein domains adaptable to high-throughput applications. Protein Science, 2006, 15, 2356-2365.	7.6	34
104	Expressed in the Yeast Saccharomyces cerevisiae , Human ERK5 Is a Client of the Hsp90 Chaperone That Complements Loss of the Slt2p (Mpk1p) Cell Integrity Stress-Activated Protein Kinase. Eukaryotic Cell, 2006, 5, 1914-1924.	3.4	60
105	Crystal structure of the proximal BAH domain of the polybromo protein. Biochemical Journal, 2005, 389, 657-664.	3.7	23
106	The identification, synthesis, protein crystal structure and in vitro biochemical evaluation of a new 3,4-diarylpyrazole class of Hsp90 inhibitors. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 3338-3343.	2.2	228
107	Qri2/Nse4, a component of the essential Smc5/6 DNA repair complex. Molecular Microbiology, 2005, 55, 1735-1750.	2.5	43
108	Lupus autoantibodies to native DNA preferentially bind DNA presented on PolIV. Immunology, 2005, 114, 418-427.	4.4	7

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109	Three-Dimensional Structure and Regulation of the DNA-Dependent Protein Kinase Catalytic Subunit (DNA-PKcs). Structure, 2005, 13, 243-255.	3.3	98
110	A Two-Hybrid Screen of the Yeast Proteome for Hsp90 Interactors Uncovers a Novel Hsp90 Chaperone Requirement in the Activity of a Stress-Activated Mitogen-Activated Protein Kinase, Slt2p (Mpk1p). Eukaryotic Cell, 2005, 4, 849-860.	3.4	159
111	Chaperoned Ubiquitylation—Crystal Structures of the CHIP U Box E3 Ubiquitin Ligase and a CHIP-Ubc13-Uev1a Complex. Molecular Cell, 2005, 20, 525-538.	9.7	382
112	Beta-2-glycoprotein specificity of human anti-phospholipid antibody resides on the light chain: a novel mechanism for acquisition of cross-reactivity by an autoantibody. Molecular Immunology, 2005, 42, 39-48.	2.2	3
113	Hsp90 and Cdc37 – a chaperone cancer conspiracy. Current Opinion in Genetics and Development, 2005, 15, 55-61.	3.3	238
114	Investigating the protein-protein interactions of the yeast Hsp90 chaperone system by two-hybrid analysis: potential uses and limitations of this approach. Cell Stress and Chaperones, 2004, 9, 359.	2.9	41
115	Crystal structure of the catalytic fragment of murine poly(ADP-ribose) polymerase-2. Nucleic Acids Research, 2004, 32, 456-464.	14.5	101
116	Co-chaperone Regulation of Conformational Switching in the Hsp90 ATPase Cycle. Journal of Biological Chemistry, 2004, 279, 51989-51998.	3.4	183
117	Structural basis for recruitment of the ATPase activator Aha1 to the Hsp90 chaperone machinery. EMBO Journal, 2004, 23, 511-519.	7.8	164
118	Structural basis for recruitment of the ATPase activator Aha1 to the Hsp90 chaperone machinery. EMBO Journal, 2004, 23, 1402-1410.	7.8	179
119	High-throughput screening assay for inhibitors of heat-shock protein 90 ATPase activity. Analytical Biochemistry, 2004, 327, 176-183.	2.4	192
120	Electron microscopy studies on DNA recognition by DNA-PK. Micron, 2004, 35, 625-633.	2.2	14
121	Plasmodium falciparum glycogen synthase kinase-3: molecular model, expression, intracellular localisation and selective inhibitors. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2004, 1697, 181-196.	2.3	95
122	Structural Basis for the Synthesis of Indirubins as Potent and Selective Inhibitors of Glycogen Synthase Kinase-3 and Cyclin-Dependent Kinases. Journal of Medicinal Chemistry, 2004, 47, 935-946.	6.4	343
123	Fine binding characteristics of human autoantibodies?partial molecular characterization. Molecular Immunology, 2004, 41, 495-495.	2.2	0
124	Fine binding characteristics of human autoantibodies—partial molecular characterization. Molecular Immunology, 2004, 41, 495-510.	2.2	4
125	The 3D Solution Structure of the C-terminal Region of Ku86 (Ku86CTR). Journal of Molecular Biology, 2004, 335, 573-582.	4.2	55
126	The Mechanism of Hsp90 Regulation by the Protein Kinase-Specific Cochaperone p50cdc37. Cell, 2004, 116, 87-98.	28.9	319

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127	Structural basis for recruitment of glycogen synthase kinase 3beta to the axin-APC scaffold complex. EMBO Journal, 2003, 22, 494-501.	7.8	269
128	Structural basis for recruitment of translesion DNA polymerase Pol IV/DinB to the Â-clamp. EMBO Journal, 2003, 22, 5883-5892.	7.8	212
129	GSK-3-Selective Inhibitors Derived from Tyrian Purple Indirubins. Chemistry and Biology, 2003, 10, 1255-1266.	6.0	720
130	Sensitivity to Hsp90-targeting drugs can arise with mutation to the Hsp90 chaperone, cochaperones and plasma membrane ATP binding cassette transporters of yeast. FEBS Journal, 2003, 270, 4689-4695.	0.2	52
131	Assays for HSP90 and Inhibitors. , 2003, 85, 149-162.		14
132	Visualization of DNA-induced conformational changes in the DNA repair kinase DNA-PKcs. EMBO Journal, 2003, 22, 5875-5882.	7.8	67
133	Crystal Structure of Alanine:Glyoxylate Aminotransferase and the Relationship Between Genotype and Enzymatic Phenotype in Primary Hyperoxaluria Type 1. Journal of Molecular Biology, 2003, 331, 643-652.	4.2	129
134	Yeast is selectively hypersensitised to heat shock protein 90 (Hsp90)-targetting drugs with heterologous expression of the human Hsp90β, a property that can be exploited in screens for new Hsp90 chaperone inhibitors. Gene, 2003, 302, 165-170.	2.2	51
135	Anti-cardiolipin/β-2 glycoprotein activities co-exist on human anti-DNA antibody light chains. Molecular Immunology, 2003, 40, 517-530.	2.2	7
136	Structural and Functional Analysis of the Middle Segment of Hsp90: Implications for ATP Hydrolysis and Client Protein and Cochaperone Interactions. Molecular Cell, 2003, 11, 647-658.	9.7	434
137	Structure and Specificity of the Vertebrate Anti-Mutator Uracil-DNA Glycosylase SMUG1. Molecular Cell, 2003, 11, 1647-1659.	9.7	127
138	Crystal structure of the Escherichia coli dcm very-short-patch DNA repair endonuclease bound to its reaction product-site in a DNA superhelix. Nucleic Acids Research, 2003, 31, 1633-1639.	14.5	26
139	Structure and Functional Relationships of Hsp90. Current Cancer Drug Targets, 2003, 3, 301-323.	1.6	242
140	An Iron-Sulfur Cluster in the Family 4 Uracil-DNA Glycosylases. Journal of Biological Chemistry, 2002, 277, 16936-16940.	3.4	66
141	Regulation of Hsp90 ATPase Activity by the Co-chaperone Cdc37p/p50. Journal of Biological Chemistry, 2002, 277, 20151-20159.	3.4	246
142	Identification of the Axin and Frat Binding Region of Glycogen Synthase Kinase-3. Journal of Biological Chemistry, 2002, 277, 2176-2185.	3.4	112
143	Reciprocal "flipping―underlies substrate recognition and catalytic activation by the human 8-oxo-guanine DNA glycosylase. Journal of Molecular Biology, 2002, 317, 171-177.	4.2	101
144	Glycogen Synthase Kinase-3 Inhibition by Lithium and Beryllium Suggests the Presence of Two Magnesium Binding Sites. Biochemical and Biophysical Research Communications, 2002, 290, 967-972.	2.1	102

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145	Crystal Structure of the CCTÎ ³ Apical Domain: Implications for Substrate Binding to the Eukaryotic Cytosolic Chaperonin. Journal of Molecular Biology, 2002, 318, 1367-1379.	4.2	72
146	Activation of the ATPase Activity of Hsp90 by the Stress-Regulated Cochaperone Aha1. Molecular Cell, 2002, 10, 1307-1318.	9.7	487
147	Crystal Structure of the C-Terminal WD40 Repeat Domain of the Human Groucho/TLE1 Transcriptional Corepressor. Structure, 2002, 10, 751-761.	3.3	63
148	Regulation of protein kinases in insulin, growth factor and Wnt signalling. Current Opinion in Structural Biology, 2002, 12, 761-767.	5.7	66
149	Structural basis for uracil recognition by archaeal family B DNA polymerases. Nature Structural Biology, 2002, 9, 922-927.	9.7	146
150	Backbone 1H, 13C, and 15N resonance assignments for the C-terminal region of Ku86 (Ku86CTR). Journal of Biomolecular NMR, 2002, 22, 373-374.	2.8	1
151	Backbone resonance assignments of the 25kD N-terminal ATPase domain from the Hsp90 chaperone. Journal of Biomolecular NMR, 2002, 23, 327-328.	2.8	19
152	Expression of the Fabs of human auto-antibodies in Escherichia coli: optimization and determination of their fine binding characteristics and cross-reactivity. Journal of Molecular Biology, 2001, 308, 527-539.	4.2	18
153	Crystal Structure of Glycogen Synthase Kinase 3β. Cell, 2001, 105, 721-732.	28.9	610
154	Crystallization and preliminary crystallographic analysis of human alanine:glyoxylate aminotransferase and its polymorphic variants. Acta Crystallographica Section D: Biological Crystallography, 2001, 57, 1936-1937.	2.5	14
155	Crystal structure of the fission yeast mitochondrial Holliday junction resolvase Ydc2. EMBO Journal, 2001, 20, 6601-6611.	7.8	62
156	Structure, function, and mechanism of the Hsp90 molecular chaperone. Advances in Protein Chemistry, 2001, 59, 157-186.	4.4	172
157	Structure and in vivo function of Hsp90. Current Opinion in Structural Biology, 2000, 10, 46-51.	5.7	294
158	The ATPase cycle of Hsp90 drives a molecular clamp' via transient dimerization of the N-terminal domains. EMBO Journal, 2000, 19, 4383-4392.	7.8	418
159	Steric Hindrance Regulation of the Pseudomonas aeruginosa Amidase Operon. Journal of Biological Chemistry, 2000, 275, 30660-30667.	3.4	18
160	Improving dideoxynucleotide-triphosphate utilisation by the hyper-thermophilic DNA polymerase from the archaeon Pyrococcus furiosus. Nucleic Acids Research, 2000, 28, 1059-1066.	14.5	36
161	The Mechanism of Dna Repair by Uracil-Dna Glycosylase: Studies Using Nucleotide Analogues. Nucleosides, Nucleotides and Nucleic Acids, 2000, 19, 1505-1516.	1.1	2
162	Molecular Cloning and Expression of the Fabs of Human Autoantibodies in Escherichia coli. Journal of Biological Chemistry, 2000, 275, 35129-35136.	3.4	19

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163	Structure and function in the uracil-DNA glycosylase superfamily. Mutation Research DNA Repair, 2000, 460, 165-181.	3.7	268
164	Crystal structure and induction mechanism of AmiC-AmiR: a ligand-regulated transcription antitermination complex. EMBO Journal, 1999, 18, 5175-5186.	7.8	83
165	Crystal structure of a thwarted mismatch glycosylase DNA repair complex. EMBO Journal, 1999, 18, 6599-6609.	7.8	122
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