## 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

#	Article	IF	CITATIONS
1	Identification and Structural Characterization of the ATP/ADP-Binding Site in the Hsp90 Molecular Chaperone. Cell, 1997, 90, 65-75.	28.9	1,203
2	Structure and Mechanism of the Hsp90 Molecular Chaperone Machinery. Annual Review of Biochemistry, 2006, 75, 271-294.	11.1	988
3	Structural Basis for Inhibition of the Hsp90 Molecular Chaperone by the Antitumor Antibiotics Radicicol and Geldanamycin. Journal of Medicinal Chemistry, 1999, 42, 260-266.	6.4	948
4	Crystal structure of an Hsp90–nucleotide–p23/Sba1 closed chaperone complex. Nature, 2006, 440, 1013-1017.	27.8	857
5	CSK-3-Selective Inhibitors Derived from Tyrian Purple Indirubins. Chemistry and Biology, 2003, 10, 1255-1266.	6.0	720
6	ATP binding and hydrolysis are essential to the function of the Hsp90 molecular chaperone invivo. EMBO Journal, 1998, 17, 4829-4836.	7.8	662
7	A structural model for the retroviral proteases. Nature, 1987, 329, 351-354.	27.8	638
8	Crystal Structure of Glycogen Synthase Kinase 3β. Cell, 2001, 105, 721-732.	28.9	610
9	DNA repair, genome stability and cancer: a historical perspective. Nature Reviews Cancer, 2016, 16, 35-42.	28.4	575
10	Activation of the ATPase Activity of Hsp90 by the Stress-Regulated Cochaperone Aha1. Molecular Cell, 2002, 10, 1307-1318.	9.7	487
11	Therapeutic opportunities within the DNA damage response. Nature Reviews Cancer, 2015, 15, 166-180.	28.4	442
12	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
13	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
14	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
15	The structural basis of specific base-excision repair by uracil–DNA glycosylase. Nature, 1995, 373, 487-493.	27.8	413
16	The Hsp90 molecular chaperone: an open and shut case for treatment. Biochemical Journal, 2008, 410, 439-453.	3.7	410
17	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
18	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

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19	Regulation of Hsp90 ATPase activity by tetratricopeptide repeat (TPR)-domain co-chaperones. EMBO Journal, 1999, 18, 754-762.	7.8	376
20	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
21	The Mechanism of Hsp90 Regulation by the Protein Kinase-Specific Cochaperone p50cdc37. Cell, 2004, 116, 87-98.	28.9	319
22	Structure and in vivo function of Hsp90. Current Opinion in Structural Biology, 2000, 10, 46-51.	5.7	294
23	Structure of an Hsp90-Cdc37-Cdk4 Complex. Molecular Cell, 2006, 23, 697-707.	9.7	288
24	Structural basis for recruitment of glycogen synthase kinase 3beta to the axin-APC scaffold complex. EMBO Journal, 2003, 22, 494-501.	7.8	269
25	Structure and function in the uracil-DNA glycosylase superfamily. Mutation Research DNA Repair, 2000, 460, 165-181.	3.7	268
26	Regulation of Hsp90 ATPase Activity by the Co-chaperone Cdc37p/p50. Journal of Biological Chemistry, 2002, 277, 20151-20159.	3.4	246
27	The active site of aspartic proteinases. FEBS Letters, 1984, 174, 96-101.	2.8	242
28	Structure and Functional Relationships of Hsp90. Current Cancer Drug Targets, 2003, 3, 301-323.	1.6	242
29	Crystal Structure of a G:T/U Mismatch-Specific DNA Glycosylase. Cell, 1998, 92, 117-129.	28.9	240
30	Hsp90 and Cdc37 – a chaperone cancer conspiracy. Current Opinion in Genetics and Development, 2005, 15, 55-61.	3.3	238
31	Crystal structure of the β-glycosidase from the hyperthermophilic archeon Sulfolobus solfataricus: resilience as a key factor in thermostability. Journal of Molecular Biology, 1997, 271, 789-802.	4.2	235
32	Three-dimensional structure, specificity and catalytic mechanism of renin. Nature, 1983, 304, 273-275.	27.8	229
33	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
34	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
35	Human BRCA1–BARD1 ubiquitin ligase activity counteracts chromatin barriers to DNA resection. Nature Structural and Molecular Biology, 2016, 23, 647-655.	8.2	222
36	Recursive PCR: a novel technique for total gene synthesis. Protein Engineering, Design and Selection, 1992, 5, 827-829.	2.1	217

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37	A molecular clamp in the crystal structure of the N-terminal domain of the yeast Hsp90 chaperone. Nature Structural Biology, 1997, 4, 477-482.	9.7	214
38	The zinc-finger domains of PARP1 cooperate to recognize DNA strand breaks. Nature Structural and Molecular Biology, 2012, 19, 685-692.	8.2	214
39	Structural basis for recruitment of translesion DNA polymerase Pol IV/DinB to the Â-clamp. EMBO Journal, 2003, 22, 5883-5892.	7.8	212
40	Structure of the Ire1 autophosphorylation complex and implications for the unfolded protein response. EMBO Journal, 2011, 30, 894-905.	7.8	201
41	High-throughput screening assay for inhibitors of heat-shock protein 90 ATPase activity. Analytical Biochemistry, 2004, 327, 176-183.	2.4	192
42	Hsp90-Dependent Activation of Protein Kinases Is Regulated by Chaperone-Targeted Dephosphorylation of Cdc37. Molecular Cell, 2008, 31, 886-895.	9.7	184
43	Co-chaperone Regulation of Conformational Switching in the Hsp90 ATPase Cycle. Journal of Biological Chemistry, 2004, 279, 51989-51998.	3.4	183
44	Structural basis for recruitment of the ATPase activator Aha1 to the Hsp90 chaperone machinery. EMBO Journal, 2004, 23, 1402-1410.	7.8	179
45	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
46	Structure, function, and mechanism of the Hsp90 molecular chaperone. Advances in Protein Chemistry, 2001, 59, 157-186.	4.4	172
47	Swe1Wee1-Dependent Tyrosine Phosphorylation of Hsp90 Regulates Distinct Facets of Chaperone Function. Molecular Cell, 2010, 37, 333-343.	9.7	165
48	Structural basis for recruitment of the ATPase activator Aha1 to the Hsp90 chaperone machinery. EMBO Journal, 2004, 23, 511-519.	7.8	164
49	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
50	Structural basis for recruitment of BRCA2 by PALB2. EMBO Reports, 2009, 10, 990-996.	4.5	154
51	Activation segment dimerization: a mechanism for kinase autophosphorylation of non-consensus sites. EMBO Journal, 2008, 27, 704-714.	7.8	147
52	Structural basis for uracil recognition by archaeal family B DNA polymerases. Nature Structural Biology, 2002, 9, 922-927.	9.7	146
53	Threonine 22 Phosphorylation Attenuates Hsp90 Interaction with Cochaperones and Affects Its Chaperone Activity. Molecular Cell, 2011, 41, 672-681.	9.7	146
54	Review: The HSP90 molecular chaperone—an enigmatic ATPase. Biopolymers, 2016, 105, 594-607.	2.4	144

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55	High resolution X-ray analyses of renin inhibitor-aspartic proteinase complexes. Nature, 1987, 327, 349-352.	27.8	143
56	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
57	Regulation of DNA Replication through Sld3-Dpb11 Interaction Is Conserved from Yeast to Humans. Current Biology, 2011, 21, 1152-1157.	3.9	135
58	Molecular Recognition of Transcriptional Repressor Motifs by the WD Domain of the Groucho/TLE Corepressor. Molecular Cell, 2006, 22, 645-655.	9.7	134
59	ATP-competitive inhibitors block protein kinase recruitment to the Hsp90-Cdc37 system. Nature Chemical Biology, 2013, 9, 307-312.	8.0	132
60	Trans-activation of the DNA-damage signalling protein kinase Chk2 by T-loop exchange. EMBO Journal, 2006, 25, 3179-3190.	7.8	131
61	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
62	Structural and Mechanistic Insights into Ras Association Domains of Phospholipase C Epsilon. Molecular Cell, 2006, 21, 495-507.	9.7	129
63	Structure and Specificity of the Vertebrate Anti-Mutator Uracil-DNA Glycosylase SMUG1. Molecular Cell, 2003, 11, 1647-1659.	9.7	127
64	Crystal Structure of an Octameric RuvA–Holliday Junction Complex. Molecular Cell, 1998, 2, 361-372.	9.7	126
65	Crystal structure of a thwarted mismatch glycosylase DNA repair complex. EMBO Journal, 1999, 18, 6599-6609.	7.8	122
66	X-ray analyses of aspartic proteinases. Journal of Molecular Biology, 1990, 211, 919-941.	4.2	120
67	Insights into histone code syntax from structural and biochemical studies of CARM1 methyltransferase. EMBO Journal, 2007, 26, 4402-4412.	7.8	117
68	Identification of the Axin and Frat Binding Region of Glycogen Synthase Kinase-3. Journal of Biological Chemistry, 2002, 277, 2176-2185.	3.4	112
69	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
70	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
71	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
72	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

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73	Structural and functional coupling of Hsp90- and Sgt1-centred multi-protein complexes. EMBO Journal, 2008, 27, 2789-2798.	7.8	104
74	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
75	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
76	Crystal structure of the catalytic fragment of murine poly(ADP-ribose) polymerase-2. Nucleic Acids Research, 2004, 32, 456-464.	14.5	101
77	Three-Dimensional Structure and Regulation of the DNA-Dependent Protein Kinase Catalytic Subunit (DNA-PKcs). Structure, 2005, 13, 243-255.	3.3	98
78	MDC1 Interacts with TOPBP1 to Maintain Chromosomal Stability during Mitosis. Molecular Cell, 2019, 74, 571-583.e8.	9.7	97
79	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
80	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
81	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
82	The crystal structure of yeast CCT reveals intrinsic asymmetry of eukaryotic cytosolic chaperonins. EMBO Journal, 2011, 30, 3078-3090.	7.8	94
83	A Mechanism for the Inhibition of DNA-PK-Mediated DNA Sensing by a Virus. PLoS Pathogens, 2013, 9, e1003649.	4.7	94
84	Visualization of a DNA-PK/PARP1 complex. Nucleic Acids Research, 2012, 40, 4168-4177.	14.5	89
85	Nucleotide mimicry in the crystal structure of the uracil-DNA glycosylase–uracil glycosylase inhibitor protein complex. Nature Structural and Molecular Biology, 1995, 2, 752-757.	8.2	87
86	Activation segment exchange: a common mechanism of kinase autophosphorylation?. Trends in Biochemical Sciences, 2007, 32, 351-356.	7.5	86
87	Structural Basis for Phosphorylation-Dependent Recruitment of Tel2 to Hsp90 by Pih1. Structure, 2014, 22, 805-818.	3.3	86
88	Crystal structure and induction mechanism of AmiC-AmiR: a ligand-regulated transcription antitermination complex. EMBO Journal, 1999, 18, 5175-5186.	7.8	83
89	Optimizing Natural Products by Biosynthetic Engineering: Discovery of Nonquinone Hsp90 Inhibitors. Journal of Medicinal Chemistry, 2008, 51, 5494-5497.	6.4	79
90	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

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91	Crystal Structure of the CCTÎ <sup>3</sup> Apical Domain: Implications for Substrate Binding to the Eukaryotic Cytosolic Chaperonin. Journal of Molecular Biology, 2002, 318, 1367-1379.	4.2	72
92	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
93	Enzyme thermostability and thermoactivity. Protein Engineering, Design and Selection, 1996, 9, 629-630.	2.1	68
94	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
95	Cooperation of local motions in the Hsp90 molecular chaperone ATPase mechanism. Nature Chemical Biology, 2016, 12, 628-635.	8.0	68
96	Visualization of DNA-induced conformational changes in the DNA repair kinase DNA-PKcs. EMBO Journal, 2003, 22, 5875-5882.	7.8	67
97	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
98	An Iron-Sulfur Cluster in the Family 4 Uracil-DNA Glycosylases. Journal of Biological Chemistry, 2002, 277, 16936-16940.	3.4	66
99	Regulation of protein kinases in insulin, growth factor and Wnt signalling. Current Opinion in Structural Biology, 2002, 12, 761-767.	5.7	66
100	Structural and functional analysis of the Crb2–BRCT <sub>2</sub> domain reveals distinct roles in checkpoint signaling and DNA damage repair. Genes and Development, 2008, 22, 2034-2047.	5.9	65
101	Destabilized SMC5/6 complex leads to chromosome breakage syndrome with severe lung disease. Journal of Clinical Investigation, 2016, 126, 2881-2892.	8.2	65
102	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
103	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
104	Crystal Structure of the C-Terminal WD40 Repeat Domain of the Human Groucho/TLE1 Transcriptional Corepressor. Structure, 2002, 10, 751-761.	3.3	63
105	Crystal structure of the fission yeast mitochondrial Holliday junction resolvase Ydc2. EMBO Journal, 2001, 20, 6601-6611.	7.8	62
106	ATM Localization and Heterochromatin Repair Depend on Direct Interaction of the 53BP1-BRCT 2 Domain with Î <sup>3</sup> H2AX. Cell Reports, 2015, 13, 2081-2089.	6.4	61
107	Specialized interfaces of Smc5/6 control hinge stability and DNA association. Nature Communications, 2017, 8, 14011.	12.8	61
108	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

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109	PARP3 is a sensor of nicked nucleosomes and monoribosylates histone H2BGlu2. Nature Communications, 2016, 7, 12404.	12.8	60
110	Molecular modelling of the interactions of tetra-(4-N-methylpyridyl) porphin with TA and CG sites on DNA. Nucleic Acids Research, 1987, 15, 6553-6562.	14.5	59
111	The catalytic mechanism of aspartic proteinases. FEBS Letters, 1987, 214, 8-12.	2.8	59
112	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
113	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
114	Molecular Characterization of Macbecin as an Hsp90 Inhibitor. Journal of Medicinal Chemistry, 2008, 51, 2853-2857.	6.4	56
115	The 3D Solution Structure of the C-terminal Region of Ku86 (Ku86CTR). Journal of Molecular Biology, 2004, 335, 573-582.	4.2	55
116	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
117	RPAP3 provides a flexible scaffold for coupling HSP90 to the human R2TP co-chaperone complex. Nature Communications, 2018, 9, 1501.	12.8	54
118	A second front against AIDS. Nature, 1989, 337, 596-597.	27.8	53
119	Structural Basis of the Radicicol Resistance Displayed by a Fungal Hsp90. ACS Chemical Biology, 2009, 4, 289-297.	3.4	53
120	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
121	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
122	MoKCa database—mutations of kinases in cancer. Nucleic Acids Research, 2009, 37, D824-D831.	14.5	51
123	A common conformationally coupled ATPase mechanism for yeast and human cytoplasmic HSP90s. FEBS Journal, 2009, 276, 199-209.	4.7	51
124	Generation of molecular surfaces for graphic display. Journal of Molecular Graphics, 1983, 1, 9-12.	1.1	50
125	Identification and characterisation of 2-aminopyridine inhibitors of checkpoint kinase 2. Bioorganic and Medicinal Chemistry, 2010, 18, 707-718.	3.0	50
126	Differential Regulation of G1 CDK Complexes by the Hsp90-Cdc37 Chaperone System. Cell Reports, 2017, 21, 1386-1398.	6.4	49

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127	Direct Measurement of the Substrate Preference of Uracil-DNA Glycosylase. Journal of Biological Chemistry, 1998, 273, 45-50.	3.4	48
128	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
129	The crystal structure of the DNA-binding drug berenil: molecular modelling studies of berenil-DNA complexes. Nucleic Acids Research, 1987, 15, 3469-3478.	14.5	46
130	Chaperone ligand-discrimination by the TPR-domain protein Tah1. Biochemical Journal, 2008, 413, 261-268.	3.7	46
131	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
132	Uracil-DNA glycosylase activities in hyperthermophilic micro-organisms. FEMS Microbiology Letters, 1996, 143, 267-271.	1.8	44
133	Qri2/Nse4, a component of the essential Smc5/6 DNA repair complex. Molecular Microbiology, 2005, 55, 1735-1750.	2.5	43
134	Selective cleavage of glycyl bonds by papaya proteinase IV. FEBS Letters, 1990, 260, 195-197.	2.8	42
135	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
136	Phosphorylation-mediated interactions with TOPBP1 couple 53BP1 and 9-1-1 to control the G1 DNA damage checkpoint. ELife, 2019, 8, .	6.0	40
137	Power frequency magnetic field and illness in multistorey blocks. Public Health, 1988, 102, 11-18.	2.9	39
138	The problem with pyrimidines. Nature Structural and Molecular Biology, 1996, 3, 485-487.	8.2	39
139	Restricting direct interaction of CDC37 with HSP90 does not compromise chaperoning of client proteins. Oncogene, 2015, 34, 15-26.	5.9	39
140	Towards meeting the paracelsus challenge: The design, synthesis, and characterization of paracelsin-43, an α-helical protein with over 50% sequence identity to an all-β protein. , 1996, 24, 502-513.		36
141	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
142	Power frequency magnetic field; depressive illness and myocardial infarction. Public Health, 1989, 103, 177-180.	2.9	35
143	Thermostable β-Clycosidase fromSulfolobus Solfataricus. Biocatalysis, 1994, 11, 89-103.	0.9	34
144	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

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145	BRCT domains of the DNA damage checkpoint proteins TOPBP1/Rad4 display distinct specificities for phosphopeptide ligands. ELife, 2018, 7, .	6.0	34
146	A novel expression system of domain I of human beta2 glycoprotein I in Escherichia coli. BMC Biotechnology, 2006, 6, 8.	3.3	33
147	An Artemis polymorphic variant reduces Artemis activity and confers cellular radiosensitivity. DNA Repair, 2010, 9, 1003-1010.	2.8	33
148	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
149	The Structural Basis for Substrate Recognition by Mammalian Polynucleotide Kinase 3′ Phosphatase. Molecular Cell, 2011, 44, 385-396.	9.7	32
150	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
151	The BAH domain of Rsc2 is a histone H3 binding domain. Nucleic Acids Research, 2013, 41, 9168-9182.	14.5	31
152	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
153	Mode of action of DNA-competitive small molecule inhibitors of tyrosyl DNA phosphodiesterase 2. Biochemical Journal, 2016, 473, 1869-1879.	3.7	30
154	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
155	Targeting the Hsp90 Molecular Chaperone with Novel Macrolactams. Synthesis, Structural, Binding, and Cellular Studies. ACS Chemical Biology, 2011, 6, 1339-1347.	3.4	27
156	Phosphorylation-Dependent Assembly and Coordination of the DNA Damage Checkpoint Apparatus by Rad4TopBP1. Molecular Cell, 2013, 51, 723-736.	9.7	27
157	In vitro inhibition of HIV-1 proteinase by cerulenin. FEBS Letters, 1990, 261, 373-377.	2.8	26
158	Recurrence of a binding motif?. Nature, 1993, 362, 299-299.	27.8	26
159	Identification of Two New Genes in the Pseudomonasaeruginosa Amidase Operon, Encoding an ATPase (AmiB) and a Putative Integral Membrane Protein (AmiS). Journal of Biological Chemistry, 1995, 270, 18818-18824.	3.4	26
160	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
161	The structure of FKBP38 in complex with the MEEVD tetratricopeptide binding-motif of Hsp90. PLoS ONE, 2017, 12, e0173543.	2.5	25
162	Electron microscopy of Xrcc4 and the DNA ligase IV–Xrcc4 DNA repair complex. DNA Repair, 2009, 8, 1380-1389.	2.8	24

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163	The Hsp90 of Candida albicans can confer Hsp90 functions in Saccharomyces cerevisiae: a potential model for the processes that generate immunogenic fragments of this molecular chaperone in C. albicans infections. Microbiology (United Kingdom), 1999, 145, 3455-3463.	1.8	24
164	Crystal structure of the proximal BAH domain of the polybromo protein. Biochemical Journal, 2005, 389, 657-664.	3.7	23
165	HECTD3 Mediates an HSP90-Dependent Degradation Pathway for Protein Kinase Clients. Cell Reports, 2017, 19, 2515-2528.	6.4	23
166	Inhibition of Hsp90 with Resorcylic Acid Macrolactones: Synthesis and Binding Studies. Chemistry - A European Journal, 2010, 16, 10366-10372.	3.3	22
167	Phosphorylation-dependent assembly of DNA damage response systems and the central roles of TOPBP1. DNA Repair, 2021, 108, 103232.	2.8	21
168	Crystallization and Preliminary X-ray Analysis of the β-Galactosidase from the Extreme Thermophilic Archaebacterium Sulfolobus solfataricus. Journal of Molecular Biology, 1993, 229, 561-563.	4.2	20
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