

Herwig SchÃ¼ler

List of Publications by Year in descending order

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80
papers

4,796
citations

101543

36
h-index

98798

67
g-index

88
all docs

88
docs citations

88
times ranked

5561
citing authors

#	ARTICLE	IF	CITATIONS
1	Toward a unified nomenclature for mammalian ADP-ribosyltransferases. <i>Trends in Biochemical Sciences</i> , 2010, 35, 208-219.	7.5	724
2	Family-wide chemical profiling and structural analysis of PARP and tankyrase inhibitors. <i>Nature Biotechnology</i> , 2012, 30, 283-288.	17.5	410
3	Structural Basis for Potency and Promiscuity in Poly(ADP-ribose) Polymerase (PARP) and Tankyrase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 1262-1271.	6.4	262
4	Structural basis for the allosteric inhibitory mechanism of human kidney-type glutaminase (KGA) and its regulation by Raf-Mek-Erk signaling in cancer cell metabolism. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 7705-7710.	7.1	178
5	PARP-3 Is a Mono-ADP-ribosylase That Activates PARP-1 in the Absence of DNA. <i>Journal of Biological Chemistry</i> , 2010, 285, 8054-8060.	3.4	135
6	Crystal Structures of the ATPase Domains of Four Human Hsp70 Isoforms: HSPA1L/Hsp70-hom, HSPA2/Hsp70-2, HSPA6/Hsp70B', and HSPA5/BiP/GRP78. <i>PLoS ONE</i> , 2010, 5, e8625.	2.5	123
7	The DEXD/H-box RNA Helicase DDX19 Is Regulated by an $\hat{\pm}$ -Helical Switch. <i>Journal of Biological Chemistry</i> , 2009, 284, 10296-10300.	3.4	119
8	Comparative Structural Analysis of Lipid Binding START Domains. <i>PLoS ONE</i> , 2011, 6, e19521.	2.5	117
9	Structural Basis for the Interaction between Tankyrase-2 and a Potent Wnt-Signaling Inhibitor. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 5352-5355.	6.4	110
10	Recognition of Mono-ADP-Ribosylated ARTD10 Substrates by ARTD8 Macrodomains. <i>Structure</i> , 2013, 21, 462-475.	3.3	107
11	Comparative Structural Analysis of Human DEAD-Box RNA Helicases. <i>PLoS ONE</i> , 2010, 5, e12791.	2.5	101
12	The role of MeH73 in actin polymerization and ATP hydrolysis 1 Edited by R. Huber. <i>Journal of Molecular Biology</i> , 2002, 317, 577-589.	4.2	95
13	Structural Basis for Inhibitor Specificity in Human Poly(ADP-ribose) Polymerase-3. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 3108-3111.	6.4	88
14	Identification of Structure-Activity Relationships from Screening a Structurally Compact DNA-Encoded Chemical Library. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 3927-3931.	13.8	86
15	A Focused DNA-Encoded Chemical Library for the Discovery of Inhibitors of NAD ⁺ -Dependent Enzymes. <i>Journal of the American Chemical Society</i> , 2019, 141, 5169-5181.	13.7	84
16	Unusual properties of <i>Plasmodium falciparum</i> actin: new insights into microfilament dynamics of apicomplexan parasites. <i>FEBS Letters</i> , 2005, 579, 655-660.	2.8	83
17	Critical role for a stage-specific actin in male exflagellation of the malaria parasite. <i>Cellular Microbiology</i> , 2011, 13, 1714-1730.	2.1	79
18	A <i>Plasmodium</i> Actin-depolymerizing Factor That Binds Exclusively to Actin Monomers. <i>Molecular Biology of the Cell</i> , 2005, 16, 4013-4023.	2.1	73

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19	Crystal Structure of the Catalytic Domain of Human PARP2 in Complex with PARP Inhibitor ABT-888. <i>Biochemistry</i> , 2010, 49, 1056-1058.	2.5	72
20	<sc>PARP</sc> inhibitors: polypharmacology versus selective inhibition. <i>FEBS Journal</i> , 2013, 280, 3563-3575.	4.7	70
21	Structural Basis for Lack of ADP-ribosyltransferase Activity in Poly(ADP-ribose) Polymerase-13/Zinc Finger Antiviral Protein. <i>Journal of Biological Chemistry</i> , 2015, 290, 7336-7344.	3.4	70
22	Regulation of Apicomplexan Microfilament Dynamics by a Minimal Set of Actin-Binding Proteins. <i>Traffic</i> , 2006, 7, 1433-1439.	2.7	69
23	Structural Basis for Parasite-Specific Functions of the Divergent Profilin of <i>Plasmodium falciparum</i> . <i>Structure</i> , 2008, 16, 1638-1648.	3.3	60
24	Tankyrase 1 Inhibitors with Drug-like Properties Identified by Screening a DNA-Encoded Chemical Library. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 5143-5149.	6.4	60
25	Thermal unfolding of G-actin monitored with the DNase- α -fl-inhibition assay. <i>FEBS Journal</i> , 2000, 267, 476-486.	0.2	59
26	Structural biology of the writers, readers, and erasers in mono- and poly(ADP-ribose) mediated signaling. <i>Molecular Aspects of Medicine</i> , 2013, 34, 1088-1108.	6.4	58
27	ATPase activity and conformational changes in the regulation of actin. <i>BBA - Proteins and Proteomics</i> , 2001, 1549, 137-147.	2.1	53
28	A Potent and Selective PARP11 Inhibitor Suggests Coupling between Cellular Localization and Catalytic Activity. <i>Cell Chemical Biology</i> , 2018, 25, 1547-1553.e12.	5.2	50
29	Engineering Af1521 improves ADP-ribose binding and identification of ADP-ribosylated proteins. <i>Nature Communications</i> , 2020, 11, 5199.	12.8	49
30	Crystal Structure of Human Inosine Triphosphatase. <i>Journal of Biological Chemistry</i> , 2007, 282, 3182-3187.	3.4	48
31	PARP Inhibitor with Selectivity Toward ADP-Ribosyltransferase ARTD3/PARP3. <i>ACS Chemical Biology</i> , 2013, 8, 1698-1703.	3.4	48
32	Towards small molecule inhibitors of mono-ADP-ribosyltransferases. <i>European Journal of Medicinal Chemistry</i> , 2015, 95, 546-551.	5.5	46
33	Actin regulation in the malaria parasite. <i>European Journal of Cell Biology</i> , 2011, 90, 966-971.	3.6	45
34	System-wide identification and prioritization of enzyme substrates by thermal analysis. <i>Nature Communications</i> , 2021, 12, 1296.	12.8	44
35	Vital role for the <i>Plasmodium</i> actin capping protein (CP) β -subunit in motility of malaria sporozoites. <i>Molecular Microbiology</i> , 2009, 74, 1356-1367.	2.5	42
36	Small Molecule Microarray Based Discovery of PARP14 Inhibitors. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 248-253.	13.8	38

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37	Discovery of Ligands for ADP-Ribosyltransferases via Docking-Based Virtual Screening. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 7706-7718.	6.4	37
38	14-3-3 proteins activate <i>Pseudomonas</i> exotoxins-S and -T by chaperoning a hydrophobic surface. <i>Nature Communications</i> , 2018, 9, 3785.	12.8	37
39	A DNA-Encoded Library of Chemical Compounds Based on Common Scaffolding Structures Reveals the Impact of Ligand Geometry on Protein Recognition. <i>ChemMedChem</i> , 2018, 13, 1303-1307.	3.2	37
40	Crystal Structure of Human RNA Helicase A (DHX9): Structural Basis for Unselective Nucleotide Base Binding in a DEAD-Box Variant Protein. <i>Journal of Molecular Biology</i> , 2010, 400, 768-782.	4.2	36
41	Cofactor mobility determines reaction outcome in the IMPDH and GMPT (2-2)8 barrel enzymes. <i>Nature Chemical Biology</i> , 2011, 7, 950-958.	8.0	35
42	Structure and Function of a G-actin Sequestering Protein with a Vital Role in Malaria Oocyst Development inside the Mosquito Vector. <i>Journal of Biological Chemistry</i> , 2010, 285, 11572-11583.	3.4	34
43	Crystal Structures Explain Functional Differences in the Two Actin Depolymerization Factors of the Malaria Parasite. <i>Journal of Biological Chemistry</i> , 2011, 286, 28256-28264.	3.4	34
44	Design, synthesis and evaluation of potent and selective inhibitors of mono-(ADP-ribosyl)transferases PARP10 and PARP14. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 2050-2054.	2.2	34
45	Structure of the synthetase domain of human CTP synthetase, a target for anticancer therapy. <i>Acta Crystallographica Section F: Structural Biology Communications</i> , 2006, 62, 613-617.	0.7	33
46	Mutational analysis of Ser14 and Asp157 in the nucleotide-binding site of beta-actin. <i>FEBS Journal</i> , 1999, 265, 210-220.	0.2	32
47	Design, Synthesis, Crystallographic Studies, and Preliminary Biological Appraisal of New Substituted Triazolo[4,3- <i>b</i>]pyridazin-8-amine Derivatives as Tankyrase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 2807-2812.	6.4	31
48	Crystal Structure of the ATPase Domain of the Human AAA+ Protein Paraplegin/SPG7. <i>PLoS ONE</i> , 2009, 4, e6975.	2.5	30
49	Biochemical Discrimination between Selenium and Sulfur 1: A Single Residue Provides Selenium Specificity to Human Selenocysteine Lyase. <i>PLoS ONE</i> , 2012, 7, e30581.	2.5	28
50	<i>Plasmodium</i> motility: actin not actin' like actin. <i>Trends in Parasitology</i> , 2006, 22, 146-147.	3.3	25
51	Crystal Structure of Human ADP-ribose Transferase ARTD15/PARP16 Reveals a Novel Putative Regulatory Domain. <i>Journal of Biological Chemistry</i> , 2012, 287, 24077-24081.	3.4	23
52	Design and synthesis of potent inhibitors of the mono(ADP-ribosyl)transferase, PARP14. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 2907-2911.	2.2	23
53	Identification of Poly(ADP-Ribose) Polymerase Macrodomein Inhibitors Using an AlphaScreen Protocol. <i>SLAS Discovery</i> , 2018, 23, 353-362.	2.7	23
54	Mutational analysis of arginine 177 in the nucleotide binding site of 2-actin. <i>FEBS Journal</i> , 2000, 267, 4054-4062.	0.2	21

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55	Sirtuins are Unaffected by PARP Inhibitors Containing Planar Nicotinamide Bioisosteres. <i>Chemical Biology and Drug Design</i> , 2016, 87, 478-482.	3.2	21
56	Sound attenuation of polymerizing actin reflects supramolecular structures: viscoelastic properties of actin gels modified by cytochalasin D, profilin and β -actinin. <i>Biochemical Journal</i> , 2001, 355, 771-778.	3.7	20
57	Crystal Structures Explain Functional Differences in the Two Actin Depolymerization Factors of the Malaria Parasite. <i>Journal of Biological Chemistry</i> , 2011, 286, 28256-28264.	3.4	19
58	Actin/Myosin-Based Gliding Motility in Apicomplexan Parasites. <i>Sub-Cellular Biochemistry</i> , 2008, 47, 110-120.	2.4	17
59	Pivotal and distinct role for Plasmodium actin capping protein alpha during blood infection of the malaria parasite. <i>Molecular Microbiology</i> , 2015, 96, 84-94.	2.5	16
60	Structural and functional characterization of human Iba proteins. <i>FEBS Journal</i> , 2008, 275, 4627-4640.	4.7	15
61	Sister Chromatid Cohesion Establishment Factor ESCO1 Operates by Substrate-Assisted Catalysis. <i>Structure</i> , 2016, 24, 789-796.	3.3	14
62	Structure-activity relationships for inhibitors of Pseudomonas aeruginosa exoenzyme S ADP-ribosyltransferase activity. <i>European Journal of Medicinal Chemistry</i> , 2018, 143, 568-576.	5.5	14
63	Covalent binding of ATP ^γ S to the nucleotide-binding site in S14C-actin. <i>FEBS Letters</i> , 2000, 476, 155-159.	2.8	13
64	MacroGreen, a simple tool for detection of ADP-ribosylated proteins. <i>Communications Biology</i> , 2021, 4, 919.	4.4	13
65	Comparative Structural Analysis of the Putative Mono-ADP-Ribosyltransferases of the ARTD/PARP Family. <i>Current Topics in Microbiology and Immunology</i> , 2014, 384, 153-166.	1.1	12
66	Identification of Inhibitors of Pseudomonas aeruginosa Exotoxin-S ADP-Ribosyltransferase Activity. <i>Journal of Biomolecular Screening</i> , 2016, 21, 590-595.	2.6	12
67	Structures of the hydrolase domain of human 10-formyltetrahydrofolate dehydrogenase and its complex with a substrate analogue. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2006, 62, 1294-1299.	2.5	10
68	The crystal structure of human cleavage and polyadenylation specific factor ⁵ reveals a dimeric Nudix protein with a conserved catalytic site. <i>Proteins: Structure, Function and Bioinformatics</i> , 2008, 73, 1047-1052.	2.6	10
69	Structure-function analysis of the filamentous actin binding domain of the neuronal scaffolding protein spinophilin. <i>FEBS Journal</i> , 2008, 275, 59-68.	4.7	10
70	Arp1, an actin-related protein, in Plasmodium berghei. <i>Molecular and Biochemical Parasitology</i> , 2010, 173, 88-96.	1.1	10
71	Chemical Probes to Study ADP-Ribosylation: Synthesis and Biochemical Evaluation of Inhibitors of the Human ADP-Ribosyltransferase ARTD3/PARP3. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 9556-9568.	6.4	9
72	The Connection Between Actin ATPase and Polymerization. <i>Advances in Molecular and Cell Biology</i> , 2006, 37, 49-66.	0.1	8

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73	PARP10 Multi-Site Auto- and Histone MARylation Visualized by Acid-Urea Gel Electrophoresis. <i>Cells</i> , 2021, 10, 654.	4.1	8
74	DNA binding to SMC ATPases is trapped for release. <i>EMBO Journal</i> , 2016, 35, 703-705.	7.8	5
75	The Corky Root Rot Pathogen <i>Pyrenochaeta lycopersici</i> Secretes a Proteinaceous Inducer of Cell Death Affecting Host Plants Differentially. <i>Phytopathology</i> , 2012, 102, 878-891.	2.2	4
76	Small Molecule Microarray Based Discovery of PARP14 Inhibitors. <i>Angewandte Chemie</i> , 2017, 129, 254-259.	2.0	4
77	Crystallization and preliminary structural characterization of the two actin-depolymerization factors of the malaria parasite. <i>Acta Crystallographica Section F: Structural Biology Communications</i> , 2010, 66, 583-587.	0.7	3
78	Pharmacology of ADP-ribosylation. <i>FEBS Journal</i> , 2013, 280, 3542-3542.	4.7	2
79	PARP-3 is a mono-ADP-ribosylase that activates PARP-1 in the absence of DNA. <i>Journal of Biological Chemistry</i> , 2012, 287, 34494.	3.4	1
80	Studies on the ATP-binding Site of Actin Using Site-directed Mutagenesis. , 1997, , 261-264.		0