List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Insights into the mechanism of action of the arbitrium communication system in SPbeta phages. Nature Communications, 2022, 13, .	12.8	6
2	The structural role of SARS-CoV-2 genetic background in the emergence and success of spike mutations: The case of the spike A222V mutation. PLoS Pathogens, 2022, 18, e1010631.	4.7	11
3	Structure-based analyses of Salmonella RcsB variants unravel new features of the Rcs regulon. Nucleic Acids Research, 2021, 49, 2357-2374.	14.5	10
4	Molecular Basis of Lysis–Lysogeny Decisions in Gram-Positive Phages. Annual Review of Microbiology, 2021, 75, 563-581.	7.3	31
5	The arbitrium system controls prophage induction. Current Biology, 2021, 31, 5037-5045.e3.	3.9	22
6	A regulatory cascade controls Staphylococcus aureus pathogenicity island activation. Nature Microbiology, 2021, 6, 1300-1308.	13.3	20
7	Evolutionary and Phenotypic Characterization of Two Spike Mutations in European Lineage 20E of SARS-CoV-2. MBio, 2021, 12, e0231521.	4.1	6
8	SARS-CoV-2 antibodies, serum inflammatory biomarkers and clinical severity of hospitalized COVID-19 patients. Journal of Clinical Virology, 2020, 131, 104611.	3.1	61
9	Beyond the CRISPR-Cas safeguard: PICI-encoded innate immune systems protect bacteria from bacteriophage predation. Current Opinion in Microbiology, 2020, 56, 52-58.	5.1	28
10	Revisiting the pH-gated conformational switch on the activities of HisKA-family histidine kinases. Nature Communications, 2020, 11, 769.	12.8	19
11	The SrrAB two-component system regulates <i>Staphylococcus aureus</i> pathogenicity through redox sensitive cysteines. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 10989-10999.	7.1	50
12	Structural and Functional Characterization of Autophosphorylation in Bacterial Histidine Kinases. Methods in Molecular Biology, 2020, 2077, 121-140.	0.9	1
13	The structure of a polygamous repressor reveals how phage-inducible chromosomal islands spread in nature. Nature Communications, 2019, 10, 3676.	12.8	11
14	Hijacking the Hijackers: Escherichia coli Pathogenicity Islands Redirect Helper Phage Packaging for Their Own Benefit. Molecular Cell, 2019, 75, 1020-1030.e4.	9.7	45
15	Unraveling the role of the secretor antigen in human rotavirus attachment to histo-blood group antigens. PLoS Pathogens, 2019, 15, e1007865.	4.7	41
16	Deciphering the Molecular Mechanism Underpinning Phage Arbitrium Communication Systems. Molecular Cell, 2019, 74, 59-72.e3.	9.7	42
17	Conformational dynamism for DNA interaction in the Salmonella RcsB response regulator. Nucleic Acids Research, 2018, 46, 456-472.	14.5	17
18	Structures of collagen IV globular domains: insight into associated pathologies, folding and network assembly. IUCrJ, 2018, 5, 765-779.	2.2	12

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19	Dissecting the link between the enzymatic activity and the SaPI inducing capacity of the phage 80α dUTPase. Scientific Reports, 2017, 7, 11234.	3.3	6
20	Broadening the antibacterial spectrum of histidine kinase autophosphorylation inhibitors via the use of ε-poly-L-lysine capped mesoporous silica-based nanoparticles. Nanomedicine: Nanotechnology, Biology, and Medicine, 2017, 13, 569-581.	3.3	19
21	Convergent evolution involving dimeric and trimeric dUTPases in pathogenicity island mobilization. PLoS Pathogens, 2017, 13, e1006581.	4.7	9
22	Sak and Sak4 recombinases are required for bacteriophage replication in Staphylococcus aureus. Nucleic Acids Research, 2017, 45, 6507-6519.	14.5	20
23	Pirating conserved phage mechanisms promotes promiscuous staphylococcal pathogenicity island transfer. ELife, 2017, 6, .	6.0	25
24	Convergent evolution of pathogenicity islands in helper <i>cos</i> phage interference. Philosophical Transactions of the Royal Society B: Biological Sciences, 2016, 371, 20150505.	4.0	29
25	Putative histidine kinase inhibitors with antibacterial effect against multi-drug resistant clinical isolates identified by in vitro and in silico screens. Scientific Reports, 2016, 6, 26085.	3.3	36
26	Another look at the mechanism involving trimeric dUTPases in <i>Staphylococcus aureus</i> pathogenicity island induction involves novel players in the party. Nucleic Acids Research, 2016, 44, 5457-5469.	14.5	20
27	The closed conformation of the LDL receptor is destabilized by the low Ca ⁺⁺ concentration but favored by the high Mg ⁺⁺ concentration in the endosome. FEBS Letters, 2015, 589, 3534-3540.	2.8	5
28	Bacterial Histidine Kinases as Novel Antibacterial Drug Targets. ACS Chemical Biology, 2015, 10, 213-224.	3.4	174
29	Virus Satellites Drive Viral Evolution and Ecology. PLoS Genetics, 2015, 11, e1005609.	3.5	49
30	Visualizing autophosphorylation in histidine kinases. Nature Communications, 2014, 5, 3258.	12.8	115
31	Staphylococcal pathogenicity island DNA packaging system involving <i>cos</i> -site packaging and phage-encoded HNH endonucleases. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 6016-6021.	7.1	73
32	Structural Basis of a Rationally Rewired Protein-Protein Interface Critical to Bacterial Signaling. Structure, 2013, 21, 1636-1647.	3.3	69
33	Phage dUTPases Control Transfer of Virulence Genes by a Proto-Oncogenic G Protein-like Mechanism. Molecular Cell, 2013, 49, 947-958.	9.7	51
34	dUTPases, the unexplored family of signalling molecules. Current Opinion in Microbiology, 2013, 16, 163-170.	5.1	32
35	WalK, the Path towards New Antibacterials with Low Potential for Resistance Development. ACS Medicinal Chemistry Letters, 2013, 4, 891-894.	2.8	15
36	Structural Basis of Rap Phosphatase Inhibition by Phr Peptides. PLoS Biology, 2013, 11, e1001511.	5.6	53

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37	Insights into the mechanism of activation of the phosphorylation-independent response regulator NbIR. Role of residues Cys69 and Cys96. Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms, 2012, 1819, 382-390.	1.9	4
38	Structural and Functional Insights into Endoglin Ligand Recognition and Binding. PLoS ONE, 2012, 7, e29948.	2.5	86
39	Laforin, a Dual Specificity Phosphatase Involved in Lafora Disease, Is Present Mainly as Monomeric Form with Full Phosphatase Activity. PLoS ONE, 2011, 6, e24040.	2.5	25
40	Laforin, a dual-specificity phosphatase involved in Lafora disease, is phosphorylated at Ser25 by AMP-activated protein kinase. Biochemical Journal, 2011, 439, 265-275.	3.7	29
41	Does the severity of the LGMD2A phenotype in compound heterozygotes depend on the combination of mutations?. Muscle and Nerve, 2011, 44, 710-714.	2.2	13
42	The mechanism of signal transduction by two-component systems. Current Opinion in Structural Biology, 2010, 20, 763-771.	5.7	206
43	The regulatory factor SipA is a highly stable βâ€I class protein with a SH3 fold. FEBS Letters, 2010, 584, 989-994.	2.8	8
44	Environmental control of phosphorylation pathways in a branched two omponent system. Molecular Microbiology, 2010, 78, 475-489.	2.5	46
45	Substrate Binding and Catalysis in Carbamate Kinase Ascertained by Crystallographic and Site-Directed Mutagenesis Studies: Movements and Significance of a Unique Globular Subdomain of This Key Enzyme for Fermentative ATP Production in Bacteria. Journal of Molecular Biology, 2010, 397, 1261-1275.	4.2	19
46	Structural Insight into Partner Specificity and Phosphoryl Transfer in Two-Component Signal Transduction. Cell, 2009, 139, 325-336.	28.9	351
47	Biochemical characterization of novel glucokinase mutations isolated from Spanish maturity-onset diabetes of the young (MODY2) patients. Journal of Human Genetics, 2008, 53, 460-466.	2.3	12
48	Phosphorylation-independent activation of the atypical response regulator NblR. Microbiology (United Kingdom), 2008, 154, 3002-3015.	1.8	44
49	Evolutionary Diversification in Polyamine Biosynthesis. Molecular Biology and Evolution, 2008, 25, 2119-2128.	8.9	150
50	Identification of a novel two component system in Thermotoga maritima. Complex stoichiometry and crystallization. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2007, 1774, 603-609.	2.3	11
51	The regulatory factor SipA provides a link between NblS and NblR signal transduction pathways in the cyanobacterium Synechococcus sp. PCC 7942. Molecular Microbiology, 2007, 66, 071120025032002-???.	2.5	32
52	Structure-function analysis of the α5 and the α13 helices of human glucokinase: Description of two novel activating mutations. Protein Science, 2005, 14, 2080-2086.	7.6	18
53	Structure of the entire cytoplasmic portion of a sensor histidine-kinase protein. EMBO Journal, 2005, 24, 4247-4259.	7.8	266
54	Gene Structure, Organization, Expression, and Potential Regulatory Mechanisms of Arginine Catabolism in Enterococcus faecalis. Journal of Bacteriology, 2002, 184, 6289-6300.	2.2	92

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55	Structure of Acetylglutamate Kinase, a Key Enzyme for Arginine Biosynthesis and a Prototype for the Amino Acid Kinase Enzyme Family, during Catalysis. Structure, 2002, 10, 329-342.	3.3	126
56	Mitochondrial DNA depletion and <i>dGK</i> gene mutations. Annals of Neurology, 2002, 52, 311-317.	5.3	152
57	A crystallographic glimpse of a nucleotide triphosphate (AMPPNP) bound to a protein surface: external and internal AMPPNP molecules in crystallineN-acetyl-L-glutamate kinase. Acta Crystallographica Section D: Biological Crystallography, 2002, 58, 1892-1895.	2.5	3
58	[21] Carbamoyl phosphate synthesis: Carbamate kinase from Pyrococcus furiosus. Methods in Enzymology, 2001, 331, 236-247.	1.0	8
59	Structural and Mutational Analysis of the PhoQ Histidine Kinase Catalytic Domain. Journal of Biological Chemistry, 2001, 276, 41182-41190.	3.4	111
60	The 1.5 Ã resolution crystal structure of the carbamate kinase-like carbamoyl phosphate synthetase from the hyperthermophilic archaeon Pyrococcus furiosus , bound to ADP, confirms that this thermostable enzyme is a carbamate kinase, and provides insight into substrate binding and stability in carbamate kinases 1 1Edited by R. Huber. Journal of Molecular Biology, 2000, 299, 463-476.	4.2	49
61	The Carbamoyl-phosphate Synthetase of Pyrococcus furiosus Is Enzymologically and Structurally a Carbamate Kinase. Journal of Biological Chemistry, 1999, 274, 16295-16303.	3.4	29
62	Crystallization and preliminary structural results of catalase from human erythrocytes. Acta Crystallographica Section D: Biological Crystallography, 1999, 55, 1066-1068.	2.5	2
63	N-Acetyl-L-glutamate kinase from Escherichia coli: cloning of the gene, purification and crystallization of the recombinant enzyme and preliminary X-ray analysis of the free and ligand-bound forms. Acta Crystallographica Section D: Biological Crystallography, 1999, 55, 1350-1352.	2.5	10
64	Carbamate kinase: New structural machinery for making carbamoyl phosphate, the common precursor of pyrimidines and arginine. Protein Science, 1999, 8, 934-940.	7.6	46
65	Carbamate kinase from Enterococcus faecalis and Enterococcus faecium . Cloning of the genes, studies on the enzyme expressed in Escherichia coli, and sequence similarity with N-acetyl- L-glutamate kinase. FEBS Journal, 1998, 253, 280-291.	0.2	29
66	Crystallization, characterization, and preliminary crystallographic studies of mitochondrial carbamoyl phosphate synthetase I ofRana catesbeiana. Proteins: Structure, Function and Bioinformatics, 1995, 22, 193-196.	2.6	3
67	Crystallization, Characterization and Preliminary Crystallographic Studies of Carbamate Kinase of Streptococcus faecium. Journal of Molecular Biology, 1994, 235, 1345-1347.	4.2	7