

# Liskin Swint-Kruse

## List of Publications by Year in descending order

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59  
papers

1,898  
citations

279798

23  
h-index

276875

41  
g-index

66  
all docs

66  
docs citations

66  
times ranked

1559  
citing authors

#	ARTICLE	IF	CITATIONS
1	Allosteric regulation within the highly interconnected structural scaffold of <scp>AraC</scp>/<scp>XylS</scp> homologs tolerates a wide range of amino acid changes. <i>Proteins: Structure, Function and Bioinformatics</i> , 2022, 90, 186-199.	2.6	0
2	Substitutions at a rheostat position in human aldolase A cause a shift in the conformational population. <i>Protein Science</i> , 2022, 31, 357-370.	7.6	7
3	Structural Plasticity Is a Feature of Rheostat Positions in the Human Na <sup>+</sup> /Taurocholate Cotransporting Polypeptide (NTCP). <i>International Journal of Molecular Sciences</i> , 2022, 23, 3211.	4.1	4
4	Substitutions at Nonconserved Rheostat Positions Modulate Function by Rewiring Long-Range, Dynamic Interactions. <i>Molecular Biology and Evolution</i> , 2021, 38, 201-214.	8.9	30
5	Transcription   lac Operon Regulation. , 2021, , 455-465.		0
6	A clinically relevant polymorphism in the Na <sup>+</sup> /taurocholate cotransporting polypeptide (NTCP) occurs at a rheostat position. <i>Journal of Biological Chemistry</i> , 2021, 296, 100047.	3.4	19
7	Rheostat functional outcomes occur when substitutions are introduced at nonconserved positions that diverge with speciation. <i>Protein Science</i> , 2021, 30, 1833-1853.	7.6	12
8	Spectroscopic evidence of tetanus toxin translocation domain bilayer-induced refolding and insertion. <i>Biophysical Journal</i> , 2021, 120, 4763-4776.	0.5	3
9	Identification of biochemically neutral positions in liver pyruvate kinase. <i>Proteins: Structure, Function and Bioinformatics</i> , 2020, 88, 1340-1350.	2.6	14
10	Rheostat positions: A new classification of protein positions relevant to pharmacogenomics. <i>Medicinal Chemistry Research</i> , 2020, 29, 1133-1146.	2.4	16
11	The strengths and limitations of using biolayer interferometry to monitor equilibrium titrations of biomolecules. <i>Protein Science</i> , 2020, 29, 1004-1020.	7.6	19
12	Functional Characterization of Position 271 in NTCP, a Predicted Rheostat Location. <i>FASEB Journal</i> , 2020, 34, 1-1.	0.5	0
13	Homolog comparisons further reconcile in vitro and in vivo correlations of protein activities by revealing overlooked physiological factors. <i>Protein Science</i> , 2019, 28, 1806-1818.	7.6	15
14	Functional tunability from a distance: Rheostat positions influence allosteric coupling between two distant binding sites. <i>Scientific Reports</i> , 2019, 9, 16957.	3.3	15
15	Characterization of the Expression and Function of Rheostat Locations within the Na <sup>+</sup> /Taurocholate Cotransporting Polypeptide. <i>FASEB Journal</i> , 2019, 33, 507.10.	0.5	0
16	RheoScale: A tool to aggregate and quantify experimentally determined substitution outcomes for multiple variants at individual protein positions. <i>Human Mutation</i> , 2018, 39, 1814-1826.	2.5	23
17	Using Evolution to Guide Protein Engineering: The Devil IS in the Details. <i>Biophysical Journal</i> , 2016, 111, 10-18.	0.5	36
18	Data on publications, structural analyses, and queries used to build and utilize the AlloRep database. <i>Data in Brief</i> , 2016, 8, 948-957.	1.0	2

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19	A New Pattern in Protein Evolutionary Sequence Information Robustly Identifies Functionally-Important Amino Acid Positions. <i>Biophysical Journal</i> , 2016, 110, 188a.	0.5	0
20	AlloRep: A Repository of Sequence, Structural and Mutagenesis Data for the LacI/GalR Transcription Regulators. <i>Journal of Molecular Biology</i> , 2016, 428, 671-678.	4.2	18
21	Amino acid positions subject to multiple coevolutionary constraints can be robustly identified by their eigenvector network centrality scores. <i>Proteins: Structure, Function and Bioinformatics</i> , 2015, 83, 2293-2306.	2.6	23
22	Flexibility and Disorder in Gene Regulation: LacI/GalR and Hox Proteins. <i>Journal of Biological Chemistry</i> , 2015, 290, 24669-24677.	3.4	19
23	A Tale of Two Proteins. <i>FASEB Journal</i> , 2015, 29, 232.1.	0.5	1
24	Modular, Multi-Input Transcriptional Logic Gating with Orthogonal LacI/GalR Family Chimeras. <i>ACS Synthetic Biology</i> , 2014, 3, 645-651.	3.8	79
25	Rheostats and Toggle Switches for Modifying Protein Function. <i>Biophysical Journal</i> , 2014, 106, 207a.	0.5	1
26	In Vitro Thermodynamics of DNA Binding Correlate with In Vivo Transcription Repression by a Synthetic LacI/GalR Paralog. <i>Biophysical Journal</i> , 2013, 104, 576a.	0.5	0
27	Rheostats and Toggle Switches for Modulating Protein Function. <i>PLoS ONE</i> , 2013, 8, e83502.	2.5	51
28	Multiple Co-Evolutionary Networks Are Supported by the Common Tertiary Scaffold of the LacI/GalR Proteins. <i>PLoS ONE</i> , 2013, 8, e84398.	2.5	26
29	Novel insights from hybrid LacI/GalR proteins: family-wide functional attributes and biologically significant variation in transcription repression. <i>Nucleic Acids Research</i> , 2012, 40, 11139-11154.	14.5	74
30	Multiple Co-Evolutionary Networks have Evolved on the Common Tertiary Scaffold of the LacI/GalR Proteins. <i>Biophysical Journal</i> , 2012, 102, 184a.	0.5	0
31	Correlating in Vitro Measurements of Protein-DNA Binding Affinities with in Vivo Repression and Impact on the Growth Rate of the Host Organism. <i>Biophysical Journal</i> , 2011, 100, 321a.	0.5	0
32	In vivo tests of thermodynamic models of transcription repressor function. <i>Biophysical Chemistry</i> , 2011, 159, 142-151.	2.8	23
33	Functionally important positions can comprise the majority of a protein's architecture. <i>Proteins: Structure, Function and Bioinformatics</i> , 2011, 79, 1589-1608.	2.6	32
34	Comparing the Functional Roles of Nonconserved Sequence Positions in Homologous Transcription Repressors: Implications for Sequence/Function Analyses. <i>Journal of Molecular Biology</i> , 2010, 395, 785-802.	4.2	31
35	Allosteric transition pathways in the lactose repressor protein core domains: Asymmetric motions in a homodimer. <i>Protein Science</i> , 2009, 12, 2523-2541.	7.6	54
36	Allostery in the LacI/GalR family: variations on a theme. <i>Current Opinion in Microbiology</i> , 2009, 12, 129-137.	5.1	128

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37	Experimental identification of specificity determinants in the domain linker of a LacI/GalR protein: Bioinformatics-based predictions generate true positives and false negatives. <i>Proteins: Structure, Function and Bioinformatics</i> , 2008, 73, 941-957.	2.6	30
38	Enzymatic reaction sequences as coupled multiple traces on a multidimensional landscape. <i>Trends in Biochemical Sciences</i> , 2008, 33, 104-112.	7.5	23
39	Ligand-induced Conformational Changes and Conformational Dynamics in the Solution Structure of the Lactose Repressor Protein. <i>Journal of Molecular Biology</i> , 2008, 376, 466-481.	4.2	61
40	Subdividing Repressor Function: DNA Binding Affinity, Selectivity, and Allostery Can Be Altered by Amino Acid Substitution of Nonconserved Residues in a LacI/GalR Homologue. <i>Biochemistry</i> , 2008, 47, 8058-8069.	2.5	27
41	Linker Regions of the RhaS and RhaR Proteins. <i>Journal of Bacteriology</i> , 2007, 189, 269-271.	2.2	12
42	Ligand interactions with lactose repressor protein and the repressor-operator complex: The effects of ionization and oligomerization on binding. <i>Biophysical Chemistry</i> , 2007, 126, 94-105.	2.8	12
43	Functional consequences of exchanging domains between LacI and PurR are mediated by the intervening linker sequence. <i>Proteins: Structure, Function and Bioinformatics</i> , 2007, 68, 375-388.	2.6	35
44	The lactose repressor system: paradigms for regulation, allosteric behavior and protein folding. <i>Cellular and Molecular Life Sciences</i> , 2007, 64, 3-16.	5.4	151
45	Extrinsic Interactions Dominate Helical Propensity in Coupled Binding and Folding of the Lactose Repressor Protein Hinge Helix. <i>Biochemistry</i> , 2006, 45, 5896-5906.	2.5	39
46	Resmap: automated representation of macromolecular interfaces as two-dimensional networks. <i>Bioinformatics</i> , 2005, 21, 3327-3328.	4.1	99
47	Integrated Insights from Simulation, Experiment, and Mutational Analysis Yield New Details of LacI Function. <i>Biochemistry</i> , 2005, 44, 11201-11213.	2.5	32
48	Thermodynamics, Protein Modification, and Molecular Dynamics in Characterizing Lactose Repressor Protein: Strategies for Complex Analyses of Protein Structure-Function. <i>Methods in Enzymology</i> , 2004, 379, 188-209.	1.0	14
49	Using Networks To Identify Fine Structural Differences between Functionally Distinct Protein States. <i>Biochemistry</i> , 2004, 43, 10886-10895.	2.5	36
50	lac Operon. , 2004, , 529-534.		0
51	Perturbation from a Distance: Mutations that Alter LacI Function through Long-Range Effects. <i>Biochemistry</i> , 2003, 42, 14004-14016.	2.5	39
52	Fine-tuning function: Correlation of hinge domain interactions with functional distinctions between LacI and PurR. <i>Protein Science</i> , 2002, 11, 778-794.	7.6	40
53	Plasticity of quaternary structure: Twenty-two ways to form a LacI dimer. <i>Protein Science</i> , 2001, 10, 262-276.	7.6	43
54	Relieving repression. <i>Nature Structural Biology</i> , 2000, 7, 184-187.	9.7	29

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55	Comparison of Simulated and Experimentally Determined Dynamics for a Variant of the LacI DNA-Binding Domain, Nlac-P. Biophysical Journal, 1998, 74, 413-421.	0.5	23
56	Designed Disulfide between N-terminal Domains of Lactose Repressor Disrupts Allosteric Linkage. Journal of Biological Chemistry, 1997, 272, 26818-26821.	3.4	26
57	Temperature and pH Dependences of Hydrogen Exchange and Global Stability for Ovomuroid Third Domain. Biochemistry, 1996, 35, 171-180.	2.5	103
58	Hydrogen Bonds and the pH dependence of Ovomuroid Third Domain Stability. Biochemistry, 1995, 34, 4724-4732.	2.5	99
59	Thermodynamics of unfolding for turkey ovomuroid third domain: Thermal and chemical denaturation. Protein Science, 1993, 2, 2037-2049.	7.6	137