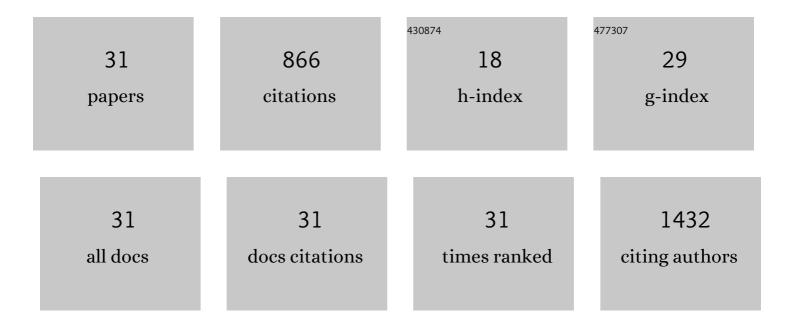
## **Robert Schnell**

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Siroheme- and [Fe4-S4]-dependent NirA from Mycobacterium tuberculosis Is a Sulfite Reductase with a Covalent Cys-Tyr Bond in the Active Site. Journal of Biological Chemistry, 2005, 280, 27319-27328.	3.4	97
2	Structural Insights into Catalysis and Inhibition of O-Acetylserine Sulfhydrylase from Mycobacterium tuberculosis. Journal of Biological Chemistry, 2007, 282, 23473-23481.	3.4	82
3	A secretagogin locus of the mammalian hypothalamus controls stress hormone release. EMBO Journal, 2015, 34, 36-54.	7.8	75
4	A <scp>TRPV</scp> 1â€toâ€secretagogin regulatory axis controls pancreatic βâ€cell survival by modulating protein turnover. EMBO Journal, 2017, 36, 2107-2125.	7.8	52
5	Peptidoglycan Remodeling in Mycobacterium tuberculosis: Comparison of Structures and Catalytic Activities of RipA and RipB. Journal of Molecular Biology, 2011, 413, 247-260.	4.2	50
6	Inhibitors of the Cysteine Synthase CysM with Antibacterial Potency against Dormant <i>Mycobacterium tuberculosis</i> . Journal of Medicinal Chemistry, 2016, 59, 6848-6859.	6.4	45
7	Discovery of an Allosteric Inhibitor Binding Site in 3-Oxo-acyl-ACP Reductase from <i>Pseudomonas aeruginosa</i> . ACS Chemical Biology, 2013, 8, 2518-2527.	3.4	38
8	Structure of Ldt <sub>Mt2</sub> , an <scp>L</scp> , <scp>D</scp> -transpeptidase from <i>Mycobacterium tuberculosis</i> . Acta Crystallographica Section D: Biological Crystallography, 2013, 69, 432-441.	2.5	36
9	Binding and processing of βâ€lactam antibiotics by the transpeptidase Ldt <sub>Mt2</sub> from <i>Mycobacterium tuberculosis</i> . FEBS Journal, 2017, 284, 725-741.	4.7	35
10	The AEROPATH project targetingPseudomonas aeruginosa: crystallographic studies for assessment of potential targets in early-stage drug discovery. Acta Crystallographica Section F: Structural Biology Communications, 2013, 69, 25-34.	0.7	30
11	Structural and Functional Characterization of the BcsG Subunit of the Cellulose Synthase in Salmonella typhimurium. Journal of Molecular Biology, 2018, 430, 3170-3189.	4.2	29
12	Pyridoxal-phosphate dependent mycobacterial cysteine synthases: Structure, mechanism and potential as drug targets. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2015, 1854, 1175-1183.	2.3	28
13	Engineering of Ancestors as a Tool to Elucidate Structure, Mechanism, and Specificity of Extant Terpene Cyclase. Journal of the American Chemical Society, 2021, 143, 3794-3807.	13.7	28
14	The structure of the Nâ€ŧerminal module of the cell wall hydrolase RipA and its role in regulating catalytic activity. Proteins: Structure, Function and Bioinformatics, 2018, 86, 912-923.	2.6	26
15	1.9â€Ã structure of the signal receiver domain of the putative response regulator NarL from <i>Mycobacterium tuberculosis</i> . Acta Crystallographica Section F: Structural Biology Communications, 2008, 64, 1096-1100.	0.7	24
16	Structural enzymology of sulphur metabolism in Mycobacterium tuberculosis. Biochemical and Biophysical Research Communications, 2010, 396, 33-38.	2.1	22
17	RipD (Rv1566c) from <i>Mycobacterium tuberculosis</i> : adaptation of an NlpC/p60 domain to a non-catalytic peptidoglycan-binding function. Biochemical Journal, 2014, 457, 33-41.	3.7	21
18	CysK2 from Mycobacterium tuberculosis Is an <i>O</i> -Phospho- <scp>l</scp> -Serine-Dependent <i>S</i> -Sulfocysteine Synthase. Journal of Bacteriology, 2014, 196, 3410-3420.	2.2	21

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19	Tetrahydrodipicolinate N-Succinyltransferase and Dihydrodipicolinate Synthase from Pseudomonas aeruginosa: Structure Analysis and Gene Deletion. PLoS ONE, 2012, 7, e31133.	2.5	20
20	Secretagogin protects Pdx1 from proteasomal degradation to control a transcriptional program required for β cell specification. Molecular Metabolism, 2018, 14, 108-120.	6.5	19
21	Structural characterization of substrate and inhibitor binding to farnesyl pyrophosphate synthase from <i>Pseudomonas aeruginosa</i> . Acta Crystallographica Section D: Biological Crystallography, 2015, 71, 721-731.	2.5	17
22	Profiling of in vitro activities of urea-based inhibitors against cysteine synthases from Mycobacterium tuberculosis. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 4582-4587.	2.2	13
23	A FabG inhibitor targeting an allosteric binding site inhibits several orthologs from Gram-negative ESKAPE pathogens. Bioorganic and Medicinal Chemistry, 2021, 30, 115898.	3.0	12
24	GABAergic Terminals Are a Source of Galanin to Modulate Cholinergic Neuron Development in the Neonatal Forebrain. Cerebral Cortex, 2014, 24, 3277-3288.	2.9	10
25	Crystal structure of NirD, the small subunit of the nitrite reductase NirbD from <i>Mycobacterium tuberculosis</i> at 2.0 Ã resolution. Proteins: Structure, Function and Bioinformatics, 2012, 80, 2799-2803.	2.6	9
26	Crystal Structures of the Kinase Domain of the Sulfate-Activating Complex in Mycobacterium tuberculosis. PLoS ONE, 2015, 10, e0121494.	2.5	9
27	N-Thio-β-lactams targeting L,D-transpeptidase-2, with activity against drug-resistant strains of Mycobacterium tuberculosis. Cell Chemical Biology, 2021, 28, 1321-1332.e5.	5.2	8
28	Substrate Channel Flexibility in Pseudomonas aeruginosa MurB Accommodates Two Distinct Substrates. PLoS ONE, 2013, 8, e66936.	2.5	5
29	Structures of <i>Pseudomonas aeruginosa</i> β-ketoacyl-(acyl-carrier-protein) synthase II (FabF) and a C164Q mutant provide templates for antibacterial drug discovery and identify a buried potassium ion and a ligand-binding site that is an artefact of the crystal form. Acta Crystallographica Section F, Structural Biology Communications, 2015, 71, 1020-1026.	0.8	4
30	Lead derivatization of ethyl 6-bromo-2-((dimethylamino)methyl)-5-hydroxy-1-phenyl-1H-indole-3-carboxylate and 5-bromo-2-(thiophene-2-carboxamido) benzoic acid as FabG inhibitors targeting ESKAPE pathogens. European Journal of Medicinal Chemistry, 2021, , 113976.	5.5	1
31	Crystal structure of the flavoenzyme PA4991 fromPseudomonas aeruginosa. Acta Crystallographica Section F, Structural Biology Communications, 2016, 72, 105-111.	0.8	0