## Pan Pan

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11371268/publications.pdf

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11	352	8	10
papers	citations	h-index	g-index
13	13	13	574
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Selectivity of Pyridone- and Diphenyl Ether-Based Inhibitors for the <i>Yersinia pestis</i> FabV Enoyl-ACP Reductase. Biochemistry, 2016, 55, 2992-3006.	2.5	6
2	Formulation studies of InhA inhibitors and combination therapy to improve efficacy against Mycobacterium tuberculosis. Tuberculosis, 2016, 101, 8-14.	1.9	4
3	Radiolabelling and positron emission tomography of PT70, a time-dependent inhibitor of InhA, the Mycobacterium tuberculosis enoyl-ACP reductase. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 4782-4786.	2.2	9
4	Timeâ€Dependent Diaryl Ether Inhibitors of InhA: Structure–Activity Relationship Studies of Enzyme Inhibition, Antibacterial Activity, and in vivo Efficacy. ChemMedChem, 2014, 9, 776-791.	3.2	48
5	A Structural and Energetic Model for the Slow-Onset Inhibition of the <i>Mycobacterium tuberculosis</i> Enoyl-ACP Reductase InhA. ACS Chemical Biology, 2014, 9, 986-993.	3.4	63
6	Rational Design of Broad Spectrum Antibacterial Activity Based on a Clinically Relevant Enoyl-Acyl Carrier Protein (ACP) Reductase Inhibitor. Journal of Biological Chemistry, 2014, 289, 15987-16005.	3.4	63
7	Rational Optimization of Drug-Target Residence Time: Insights from Inhibitor Binding to the <i>Staphylococcus aureus</i> Fabl Enzyme–Product Complex. Biochemistry, 2013, 52, 4217-4228.	2.5	58
8	Targeting InhA, the FASII Enoyl-ACP Reductase: SAR Studies on Novel Inhibitor Scaffolds. Current Topics in Medicinal Chemistry, 2012, 12, 672-693.	2.1	76
9	Residence Time and in vivo Antibacterial Activity ―A Critical Aspect of Lead Compound Optimization. FASEB Journal, 2010, 24, 680.3.	0.5	O
10	Selective deprotection of the Cbz amine protecting group for the facile synthesis of kanamycin A dimers linked at N-3″ position. Tetrahedron, 2009, 65, 5922-5927.	1.9	10
11	Regioselective modification of amino groups in aminoglycosides based on cyclic carbamate formation. Tetrahedron, 2008, 64, 9078-9087.	1.9	15