

# Jonathan S Rosenblum

## List of Publications by Year in descending order

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16  
papers

934  
citations

623734

14  
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996975

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g-index

16  
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16  
docs citations

16  
times ranked

1336  
citing authors

#	ARTICLE	IF	CITATIONS
1	Prolyl peptidases: a serine protease subfamily with high potential for drug discovery. <i>Current Opinion in Chemical Biology</i> , 2003, 7, 496-504.	6.1	280
2	Wortmannin, a Widely Used Phosphoinositide 3-Kinase Inhibitor, also Potently Inhibits Mammalian Polo-like Kinase. <i>Chemistry and Biology</i> , 2005, 12, 99-107.	6.0	184
3	A Nuclear Import Pathway for a Protein Involved in tRNA Maturation. <i>Journal of Cell Biology</i> , 1997, 139, 1655-1661.	5.2	82
4	ERK5 kinase activity is dispensable for cellular immune response and proliferation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, 11865-11870.	7.1	74
5	Polo-like Kinases Inhibited by Wortmannin. <i>Journal of Biological Chemistry</i> , 2007, 282, 2505-2511.	3.4	69
6	Nuclear Import and the Evolution of a Multifunctional RNA-binding Protein. <i>Journal of Cell Biology</i> , 1998, 143, 887-899.	5.2	59
7	Synergistic Computational and Experimental Proteomics Approaches for More Accurate Detection of Active Serine Hydrolases in Yeast. <i>Molecular and Cellular Proteomics</i> , 2004, 3, 209-225.	3.8	46
8	Synthesis and structure-activity relationship of N-alkyl Gly-boro-Pro inhibitors of DPP4, FAP, and DPP7. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 4239-4242.	2.2	32
9	Synthesis and structure-activity relationship of 4-quinolone-3-carboxylic acid based inhibitors of glycogen synthase kinase-3 <sup>β</sup> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 5948-5951.	2.2	18
10	Boro-norleucine as a P1 residue for the design of selective and potent DPP7 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 4256-4260.	2.2	17
11	DPP8 and DPP9 expression in cynomolgus monkey and Sprague Dawley rat tissues. <i>Regulatory Peptides</i> , 2013, 186, 26-35.	1.9	17
12	Synthesis and activity of a potent, specific azabicyclo[3.3.0]-octane-based DPP II inhibitor. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 507-510.	2.2	16
13	Functional interrogation of kinases and other nucleotide-binding proteins. <i>FEBS Letters</i> , 2013, 587, 1870-1877.	2.8	16
14	Determination of enzymatic source of alanine aminotransferase activity in serum from dogs with liver injury. <i>Journal of Pharmacological and Toxicological Methods</i> , 2009, 60, 307-315.	0.7	15
15	Synthesis and structure-activity relationship of (1-halo-2-naphthyl) carbamate-based inhibitors of KIAA1363 (NCEH1/AADACL1). <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 5748-5751.	2.2	9
16	Efficient Nuclear Transport of Structurally Disturbed Cargo: Mutations in a Cargo Protein Switch Its Cognate Karyopherin. <i>PLoS ONE</i> , 2011, 6, e16846.	2.5	0