

# Harichandra D Tagad

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7944988/publications.pdf>

Version: 2024-02-01

12  
papers

246  
citations

1040056

9  
h-index

1199594

12  
g-index

12  
all docs

12  
docs citations

12  
times ranked

420  
citing authors

#	ARTICLE	IF	CITATIONS
1	Characterization of the p300 Taz2â€™p53 TAD2 Complex and Comparison with the p300 Taz2â€™p53 TAD1 Complex. <i>Biochemistry</i> , 2015, 54, 2001-2010.	2.5	47
2	Binding of a Third Metal Ion by the Human Phosphatases PP2CÎ± and Wip1 Is Required for Phosphatase Activity. <i>Biochemistry</i> , 2013, 52, 5830-5843.	2.5	28
3	Design of pentapeptidic BACE1 inhibitors with carboxylic acid bioisosteres at P1 and P4 positions. <i>Bioorganic and Medicinal Chemistry</i> , 2010, 18, 3175-3186.	3.0	21
4	Intensity and duration of TCR signaling is limited by p38 phosphorylation of ZAP-70 and destabilization of the signalosome. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, 2174-2179.	7.1	27
5	Tripeptidic BACE1 inhibitors devised by in-silico conformational structure-based design. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 1130-1135.	2.2	24
6	Cooperative assembly of a four-molecule signaling complex formed upon T cell antigen receptor activation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E11914-E11923. <a href="#">design and synthesis of</a>	7.1	24
7	<a href="#">position 1 phenylcycloalkylamine-derived pentapeptidic BACE1 inhibitors.</a> <i>Bioorganic and Medicinal Chemistry</i> , 2010, 18, 3175-3186.	3.0	21
8	A trapped human PPM1Aâ€™phosphopeptide complex reveals structural features critical for regulation of PPM protein phosphatase activity. <i>Journal of Biological Chemistry</i> , 2018, 293, 7993-8008.	3.4	19
9	Unique properties of TCR-activated p38 are necessary for NFAT-dependent T-cell activation. <i>PLoS Biology</i> , 2018, 16, e2004111.	5.6	10
10	Chemical Features Important for Activity in a Class of Inhibitors Targeting the Wip1 Flap Subdomain. <i>ChemMedChem</i> , 2018, 13, 894-901.	3.2	8
11	Physiologically relevant orthogonal assays for the discovery of small-molecule modulators of WIP1 phosphatase in high-throughput screens. <i>Journal of Biological Chemistry</i> , 2019, 294, 17654-17668.	3.4	6
12	Tetrapeptides, as smallâ€™sized peptidic inhibitors; synthesis and their inhibitory activity against BACE1. <i>Journal of Peptide Science</i> , 2010, 16, 257-262.	1.4	5