

Vladimir S Borodkin

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

1,152
citations

516710

16
h-index

677142

22
g-index

22
all docs

22
docs citations

22
times ranked

1004
citing authors

#	ARTICLE	IF	CITATIONS
1	Proteome Wide Purification and Identification of <i>O</i> -GlcNAc-Modified Proteins Using Click Chemistry and Mass Spectrometry. <i>Journal of Proteome Research</i> , 2013, 12, 927-936.	3.7	151
2	GlcNAcstatin: A Picomolar, Selective <i>O</i> -GlcNAcase Inhibitor That Modulates Intracellular <i>O</i> -GlcNAcylation Levels. <i>Journal of the American Chemical Society</i> , 2006, 128, 16484-16485.	13.7	136
3	<i>O</i> -GlcNAc transferase invokes nucleotide sugar pyrophosphate participation in catalysis. <i>Nature Chemical Biology</i> , 2012, 8, 969-974.	8.0	123
4	The active site of <i>O</i> -GlcNAc transferase imposes constraints on substrate sequence. <i>Nature Structural and Molecular Biology</i> , 2015, 22, 744-750.	8.2	114
5	GlcNAcstatins are nanomolar inhibitors of human <i>O</i> -GlcNAcase inducing cellular hyper- <i>O</i> -GlcNAcylation. <i>Biochemical Journal</i> , 2009, 420, 221-227.	3.7	83
6	Human OGA binds substrates in a conserved peptide recognition groove. <i>Biochemical Journal</i> , 2010, 432, 1-12.	3.7	58
7	Catalytic deficiency of <i>O</i> -GlcNAc transferase leads to X-linked intellectual disability. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 14961-14970.	7.1	58
8	Bisubstrate UDP-peptide conjugates as human <i>O</i> -GlcNAc transferase inhibitors. <i>Biochemical Journal</i> , 2014, 457, 497-502.	3.7	57
9	Cell-Penetrant, Nanomolar <i>O</i> -GlcNAcase Inhibitors Selective against Lysosomal Hexosaminidases. <i>Chemistry and Biology</i> , 2010, 17, 1250-1255.	6.0	52
10	Genetic recoding to dissect the roles of site-specific protein <i>O</i> -GlcNAcylation. <i>Nature Structural and Molecular Biology</i> , 2019, 26, 1071-1077.	8.2	50
11	Synergy of Peptide and Sugar in <i>O</i> -GlcNAcase Substrate Recognition. <i>Chemistry and Biology</i> , 2012, 19, 173-178.	6.0	48
12	Recognition of a glycosylation substrate by the <i>O</i> -GlcNAc transferase TPR repeats. <i>Open Biology</i> , 2017, 7, 170078.	3.6	48
13	A Structural and Biochemical Model of Processive Chitin Synthesis. <i>Journal of Biological Chemistry</i> , 2014, 289, 23020-23028.	3.4	46
14	Thio-Linked UDP-peptide Conjugates as <i>O</i> -GlcNAc Transferase Inhibitors. <i>Bioconjugate Chemistry</i> , 2018, 29, 1834-1840.	3.6	34
15	Proteolysis of HCF-1 by Ser/Thr glycosylation-incompetent <i>O</i> -GlcNAc transferase:UDP-GlcNAc complexes. <i>Genes and Development</i> , 2016, 30, 960-972.	5.9	21
16	A mechanism-inspired UDP-N-acetylglucosamine pyrophosphorylase inhibitor. <i>RSC Chemical Biology</i> , 2020, 1, 13-25.	4.1	20
17	Synthesis of potential bisubstrate inhibitors of <i>Leishmania</i> elongating \pm -d-mannosyl phosphate transferase. <i>Tetrahedron Letters</i> , 2004, 45, 857-862.	1.4	16
18	An efficient and versatile synthesis of GlcNAcstatins-potent and selective <i>O</i> -GlcNAcase inhibitors built on the tetrahydroimidazo[1,2-a]pyridine scaffold. <i>Tetrahedron</i> , 2010, 66, 7838-7849.	1.9	9

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19	O-GlcNAcase Fragment Discovery with Fluorescence Polarimetry. ACS Chemical Biology, 2018, 13, 1353-1360.	3.4	8
20	The conserved threonine-rich region of the HCF-1PRO repeat activates promiscuous OGT:UDP-GlcNAc glycosylation and proteolysis activities. Journal of Biological Chemistry, 2018, 293, 17754-17768.	3.4	7
21	The Chemical Synthesis of Glycosylphosphatidylinositol Anchors from Trypanosoma cruzi Trypomastigote Mucins. ACS Symposium Series, 2007, , 285-306.	0.5	4