

Michael J Ferracane

List of Publications by Year in descending order

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Version: 2024-02-01

10
papers

370
citations

1163117

8
h-index

1372567

10
g-index

11
all docs

11
docs citations

11
times ranked

590
citing authors

#	ARTICLE	IF	CITATIONS
1	The mucin-selective protease StcE enables molecular and functional analysis of human cancer-associated mucins. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 7278-7287.	7.1	186
2	Classification, structural biology, and applications of mucin domain-targeting proteases. <i>Biochemical Journal</i> , 2021, 478, 1585-1603.	3.7	37
3	Identification and Characterization of Complex Glycosylated Peptides Presented by the MHC Class II Processing Pathway in Melanoma. <i>Journal of Proteome Research</i> , 2017, 16, 228-237.	3.7	34
4	The Glycoprotease CpaA Secreted by Medically Relevant <i>Acinetobacter</i> Species Targets Multiple <i>O</i> -Linked Host Glycoproteins. <i>MBio</i> , 2020, 11, .	4.1	31
5	Glycosylation of α -amino acids by sugar acetate donors with InBr ₃ . Minimally competent Lewis acids. <i>Carbohydrate Research</i> , 2012, 351, 121-125.	2.3	27
6	Multifunctional opioid receptor agonism and antagonism by a novel macrocyclic tetrapeptide prevents reinstatement of morphine-seeking behaviour. <i>British Journal of Pharmacology</i> , 2020, 177, 4209-4222.	5.4	21
7	Design, Synthesis, and Characterization of the Macrocyclic Tetrapeptide <i>cyclo</i> [Pro-Sar-Phe- <i>d</i> -Phe]: A Mixed Opioid Receptor Agonist-Antagonist Following Oral Administration. <i>ACS Chemical Neuroscience</i> , 2020, 11, 1324-1336.	3.5	12
8	Structure-guided mutagenesis of a mucin-selective metalloprotease from <i>Akkermansia muciniphila</i> alters substrate preferences. <i>Journal of Biological Chemistry</i> , 2022, 298, 101917.	3.4	11
9	Mass Spectrometric Identification and Molecular Modeling of Glycopeptides Presented by MHC Class I and All Processing Pathways. <i>Methods in Molecular Biology</i> , 2019, 2024, 269-285.	0.9	5
10	Conformational Constraint between Aromatic Residue Side Chains in the α -Message-Sequence of the Peptide Arodyn Using Ring Closing Metathesis Results in a Potent and Selective Kappa Opioid Receptor Antagonist. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 3153-3164.	6.4	5