

James W Checco

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7738904/publications.pdf>

Version: 2024-02-01

19
papers

591
citations

840776
11
h-index

794594
19
g-index

19
all docs

19
docs citations

19
times ranked

726
citing authors

#	ARTICLE	IF	CITATIONS
1	Evaluation of endogenous peptide stereochemistry using liquid chromatography-mass spectrometry-based spiking experiments. <i>Methods in Enzymology</i> , 2022, 663, 205-234.	1.0	2
2	Peptidomics analysis reveals changes in small urinary peptides in patients with interstitial cystitis/bladder pain syndrome. <i>Scientific Reports</i> , 2022, 12, 8289.	3.3	4
3	Trimer-to-Monomer Disruption Mechanism for a Potent, Protease-Resistant Antagonist of Tumor Necrosis Factor- α Signaling. <i>Journal of the American Chemical Society</i> , 2022, 144, 9610-9617.	13.7	5
4	Mass Spectrometry Approaches Empowering Neuropeptide Discovery and Therapeutics. <i>Pharmacological Reviews</i> , 2022, 74, 662-679.	16.0	5
5	Advancing d-amino acid-containing peptide discovery in the metazoan. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2021, 1869, 140553.	2.3	17
6	Identifying Receptors for Neuropeptides and Peptide Hormones: Challenges and Recent Progress. <i>ACS Chemical Biology</i> , 2021, 16, 251-263.	3.4	16
7	Evaluating functional ligand-GPCR interactions in cell-based assays. <i>Methods in Cell Biology</i> , 2021, 166, 15-42.	1.1	3
8	Differential Post-Translational Amino Acid Isomerization Found among Neuropeptides in <i>Aplysia californica</i> . <i>ACS Chemical Biology</i> , 2020, 15, 272-281.	3.4	19
9	Tumor Necrosis Factor- α Trimer Disassembly and Inactivation via Peptide-Small Molecule Synergy. <i>ACS Chemical Biology</i> , 2020, 15, 2116-2124.	3.4	5
10	Molecular and Physiological Characterization of a Receptor for D-Amino Acid-Containing Neuropeptides. <i>ACS Chemical Biology</i> , 2018, 13, 1343-1352.	3.4	27
11	<i>Aplysia</i> allatotropin-related peptide and its newly identified d-amino acid-containing epimer both activate a receptor and a neuronal target. <i>Journal of Biological Chemistry</i> , 2018, 293, 16862-16873.	3.4	25
12	Conformational investigation of the structure-activity relationship of GdFFD and its analogues on an achatin-like neuropeptide receptor of <i>Aplysia californica</i> involved in the feeding circuit. <i>Physical Chemistry Chemical Physics</i> , 2018, 20, 22047-22057.	2.8	13
13	Non-targeted Identification of d-Amino Acid-Containing Peptides Through Enzymatic Screening, Chiral Amino Acid Analysis, and LC-MS. <i>Methods in Molecular Biology</i> , 2018, 1719, 107-118.	0.9	4
14	Iterative Nonproteinogenic Residue Incorporation Yields β -Peptides with a Helix-Loop-Helix Tertiary Structure and High Affinity for VEGF. <i>ChemBioChem</i> , 2017, 18, 291-299.	2.6	19
15	Targeting recognition surfaces on natural proteins with peptidic foldamers. <i>Current Opinion in Structural Biology</i> , 2016, 39, 96-105.	5.7	76
16	Targeting diverse protein-protein interaction interfaces with β -peptides derived from the Z-domain scaffold. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 4552-4557.	7.1	93
17	β -Peptide Foldamers Targeting Intracellular Protein-Protein Interactions with Activity in Living Cells. <i>Journal of the American Chemical Society</i> , 2015, 137, 11365-11375.	13.7	101
18	Structure-Guided Rational Design of β -Peptide Foldamers with High Affinity for BCL-2 Family Prosurvival Proteins. <i>ChemBioChem</i> , 2013, 14, 1564-1572.	2.6	65

#	ARTICLE	IF	CITATIONS
19	Extending Foldamer Design beyond α -Helix Mimicry: α / β -Peptide Inhibitors of Vascular Endothelial Growth Factor Signaling. Journal of the American Chemical Society, 2012, 134, 7652-7655.	13.7	92