

James L Hougland

List of Publications by Year in descending order

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42
papers

985
citations

516710

16
h-index

477307

29
g-index

46
all docs

46
docs citations

46
times ranked

1005
citing authors

#	ARTICLE	IF	CITATIONS
1	Using Selective Withdrawal to Coat Microparticles. <i>Science</i> , 2001, 292, 265-267.	12.6	142
2	Functional Identification of Catalytic Metal Ion Binding Sites within RNA. <i>PLoS Biology</i> , 2005, 3, e277.	5.6	67
3	Identification of Novel Peptide Substrates for Protein Farnesyltransferase Reveals Two Substrate Classes with Distinct Sequence Selectivities. <i>Journal of Molecular Biology</i> , 2010, 395, 176-190.	4.2	60
4	Identification of a Novel Class of Farnesylation Targets by Structure-Based Modeling of Binding Specificity. <i>PLoS Computational Biology</i> , 2011, 7, e1002170.	3.2	58
5	Ghrelin Signaling: GOAT and GHS-R1a Take a LEAP in Complexity. <i>Trends in Endocrinology and Metabolism</i> , 2020, 31, 107-117.	7.1	48
6	Structure-Activity Analysis of Human Ghrelin O-Acyltransferase Reveals Chemical Determinants of Ghrelin Selectivity and Acyl Group Recognition. <i>Biochemistry</i> , 2015, 54, 1100-1110.	2.5	44
7	Context-Dependent Substrate Recognition by Protein Farnesyltransferase. <i>Biochemistry</i> , 2009, 48, 1691-1701.	2.5	38
8	Efficient farnesylation of an extended C-terminal C(x)3X sequence motif expands the scope of the prenylated proteome. <i>Journal of Biological Chemistry</i> , 2018, 293, 2770-2785.	3.4	33
9	A fluorescent peptide substrate facilitates investigation of ghrelin recognition and acylation by ghrelin O-acyltransferase. <i>Analytical Biochemistry</i> , 2013, 437, 68-76.	2.4	31
10	Synthetic Triterpenoid Inhibition of Human Ghrelin O-Acyltransferase: The Involvement of a Functionally Required Cysteine Provides Mechanistic Insight into Ghrelin Acylation. <i>Biochemistry</i> , 2017, 56, 919-931.	2.5	28
11	Synthesis and screening of a CaaL peptide library versus FTase reveals a surprising number of substrates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 767-770.	2.2	26
12	The ghrelin O-acyltransferase structure reveals a catalytic channel for transmembrane hormone acylation. <i>Journal of Biological Chemistry</i> , 2019, 294, 14166-14174.	3.4	26
13	Simultaneous Site-Specific Dual Protein Labeling Using Protein Prenyltransferases. <i>Bioconjugate Chemistry</i> , 2015, 26, 2542-2553.	3.6	25
14	Ghrelin octanoylation by ghrelin O-acyltransferase: Unique protein biochemistry underlying metabolic signaling. <i>Biochemical Society Transactions</i> , 2019, 47, 169-178.	3.4	24
15	The 2'-Hydroxyl Group of the Guanosine Nucleophile Donates a Functionally Important Hydrogen Bond in the Tetrahymena Ribozyme Reaction. <i>Biochemistry</i> , 2008, 47, 7684-7694.	2.5	19
16	Formation and Anomalous Behavior of Aminonaphthalene-Cinnamionitrile Exciplexes. <i>Journal of Physical Chemistry A</i> , 2000, 104, 3261-3268.	2.5	18
17	Ghrelin Octanoylation Is Completely Stabilized in Biological Samples by Alkyl Fluorophosphonates. <i>Endocrinology</i> , 2016, 157, 4330-4338.	2.8	18
18	Engineering reversible cell-cell interactions using enzymatically lipidated chemically self-assembled nanorings. <i>Chemical Science</i> , 2021, 12, 331-340.	7.4	17

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19	A closer look at alcohol-induced changes in the ghrelin system: novel insights from preclinical and clinical data. <i>Addiction Biology</i> , 2022, 27, e13033.	2.6	17
20	Protein Isoprenylation in Yeast Targets COOH-Terminal Sequences Not Adhering to the CaaX Consensus. <i>Genetics</i> , 2018, 210, 1301-1316.	2.9	16
21	Protein Farnesyltransferase Catalyzes Unanticipated Farnesylation and Geranylgeranylation of Shortened Target Sequences. <i>Biochemistry</i> , 2020, 59, 1149-1162.	2.5	16
22	An Atomic Mutation Cycle for Exploring RNA's 2'-Hydroxyl Group. <i>Journal of the American Chemical Society</i> , 2004, 126, 13578-13579.	13.7	15
23	Expansion of Protein Farnesyltransferase Specificity Using Tunable Active Site Interactions. <i>Journal of Biological Chemistry</i> , 2012, 287, 38090-38100.	3.4	15
24	A new class of ghrelin O-acyltransferase inhibitors incorporating triazole-linked lipid mimetic groups. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 2800-2803.	2.2	15
25	An overview of ghrelin O-acyltransferase inhibitors: a literature and patent review for 2010-2019. <i>Expert Opinion on Therapeutic Patents</i> , 2020, 30, 581-593.	5.0	14
26	Improved synthesis of 2'-amino-2'-deoxyguanosine and its phosphoramidite. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 705-713.	3.0	13
27	Getting a handle on protein prenylation. <i>Nature Chemical Biology</i> , 2009, 5, 197-198.	8.0	13
28	The octanoylated energy regulating hormone ghrelin: An expanded view of ghrelin's biological interactions and avenues for controlling ghrelin signaling. <i>Molecular Membrane Biology</i> , 2016, 33, 111-124.	2.0	13
29	Ghrelin octanoylation by ghrelin O-acyltransferase: protein acylation impacting metabolic and neuroendocrine signalling. <i>Open Biology</i> , 2021, 11, 210080.	3.6	12
30	Targeted Reengineering of Protein Geranylgeranyltransferase Type I Selectivity Functionally Implicates Active-Site Residues in Protein-Substrate Recognition. <i>Biochemistry</i> , 2014, 53, 434-446.	2.5	11
31	Biochemical Assays for Ghrelin Acylation and Inhibition of Ghrelin O-Acyltransferase. <i>Methods in Molecular Biology</i> , 2019, 2009, 227-241.	0.9	11
32	Novel Regulator of Acylated Ghrelin, CF801, Reduces Weight Gain, Rebound Feeding after a Fast, and Adiposity in Mice. <i>Frontiers in Endocrinology</i> , 2015, 6, 144.	3.5	10
33	Functional group and stereochemical requirements for substrate binding by ghrelin O-acyltransferase revealed by unnatural amino acid incorporation. <i>Bioorganic Chemistry</i> , 2018, 79, 98-106.	4.1	10
34	Progress in Small Molecule and Biologic Therapeutics Targeting Ghrelin Signaling. <i>Mini-Reviews in Medicinal Chemistry</i> , 2016, 16, 465-480.	2.4	10
35	Synthesis of Frame-Shifted Farnesyl Diphosphate Analogs. <i>Organic Letters</i> , 2012, 14, 4038-4041.	4.6	7
36	Quantitative Determination of Cellular Farnesyltransferase Activity: Towards Defining the Minimum Substrate Reactivity for Biologically Relevant Protein Farnesylation. <i>ChemBioChem</i> , 2014, 15, 2205-2210.	2.6	7

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37	MALDI-MS Analysis of Peptide Libraries Expands the Scope of Substrates for Farnesyltransferase. International Journal of Molecular Sciences, 2021, 22, 12042.	4.1	7
38	Global Identification of Protein Prenyltransferase Substrates. The Enzymes, 2011, 29, 207-234.	1.7	6
39	Simultaneous Analysis of a Non-Lipidated Protein and Its Lipidated Counterpart: Enabling Quantitative Investigation of Protein Lipidation's Impact on Cellular Regulation. Analytical Chemistry, 2017, 89, 13502-13507.	6.5	6
40	Temperature-Responsive Nano-Biomaterials from Genetically Encoded Farnesylated Disordered Proteins. ACS Applied Bio Materials, 2022, 5, 1846-1856.	4.6	6
41	2'-Amino-Modified Ribonucleotides as Probes for Local Interactions Within RNA. Methods in Enzymology, 2009, 468, 107-125.	1.0	4
42	Mechanisms of CaaX Protein Processing: Protein Prenylation by FTase and GGTase-I. , 2020, , 497-527.		0