

John A Karas

List of Publications by Year in descending order

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45
papers

1,322
citations

331670

21
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361022

35
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46
all docs

46
docs citations

46
times ranked

1888
citing authors

#	ARTICLE	IF	CITATIONS
1	The Chemical Synthesis of Insulin: An Enduring Challenge. <i>Chemical Reviews</i> , 2021, 121, 4531-4560.	47.7	36
2	Modular Synthesis of Trifunctional Peptide-oligonucleotide Conjugates via Native Chemical Ligation. <i>Frontiers in Chemistry</i> , 2021, 9, 627329.	3.6	9
3	Endosomal escape cell-penetrating peptides significantly enhance pharmacological effectiveness and CNS activity of systemically administered antisense oligonucleotides. <i>International Journal of Pharmaceutics</i> , 2021, 599, 120398.	5.2	10
4	Total Chemical Synthesis of a Nonfibrillating Human Glycoinsulin. <i>Journal of the American Chemical Society</i> , 2020, 142, 1164-1169.	13.7	41
5	Synthesis and structure-activity relationships of teixobactin. <i>Annals of the New York Academy of Sciences</i> , 2020, 1459, 86-105.	3.8	26
6	Structure-Activity Relationships of Daptomycin Lipopeptides. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 13266-13290.	6.4	30
7	The Antimicrobial Activity of Cannabinoids. <i>Antibiotics</i> , 2020, 9, 406.	3.7	64
8	The Killing Mechanism of Teixobactin against Methicillin-Resistant <i>Staphylococcus aureus</i> : an Untargeted Metabolomics Study. <i>MSystems</i> , 2020, 5, .	3.8	33
9	1,3-Dichloroacetone: A Robust Reagent for Preparing Bicyclic Peptides. <i>ACS Omega</i> , 2020, 5, 1840-1850.	3.5	7
10	The impact of backbone N-methylation on the structure-activity relationship of Leu 10-teixobactin. <i>Journal of Peptide Science</i> , 2019, 25, e3206.	1.4	6
11	Rapid Photolysis-Mediated Folding of Disulfide-Rich Peptides. <i>Chemistry - A European Journal</i> , 2019, 25, 8599-8603.	3.3	8
12	Thiol-Cyanobenzothiazole Ligation for the Efficient Preparation of Peptide-PNA Conjugates. <i>Bioconjugate Chemistry</i> , 2019, 30, 793-799.	3.6	20
13	Sputum Active Polymyxin Lipopeptides: Activity against Cystic Fibrosis <i>Pseudomonas aeruginosa</i> Isolates and Their Interactions with Sputum Biomolecules. <i>ACS Infectious Diseases</i> , 2018, 4, 646-655.	3.8	19
14	The Assembly of Fluorescently Labeled Peptide-Oligonucleotide Conjugates via Orthogonal Ligation Strategies. <i>Methods in Molecular Biology</i> , 2018, 1828, 355-363.	0.9	3
15	Rhenium and Technetium-oxo Complexes with Thioamide Derivatives of Pyridylhydrazine Bifunctional Chelators Conjugated to the Tumour Targeting Peptides Octreotate and Cyclic-RGDfK. <i>Inorganic Chemistry</i> , 2017, 56, 9725-9741.	4.0	19
16	Chemical Synthesis and Characterization of an Equinatoxin II (1-85) Analogue. <i>Molecules</i> , 2017, 22, 559.	3.8	2
17	The efficient synthesis and purification of amyloid- β (1-42) using an oligoethylene glycol-containing photocleavable lysine tag. <i>Chemical Communications</i> , 2017, 53, 6903-6905.	4.1	14
18	Innenteilbild: A One-Pot Chemically Cleavable Bis-Linker Tether Strategy for the Synthesis of Heterodimeric Peptides (<i>Angew. Chem.</i> 47/2016). <i>Angewandte Chemie</i> , 2016, 128, 14688-14688.	2.0	0

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19	A One-Pot Chemically Cleavable Bis-Linker Tether Strategy for the Synthesis of Heterodimeric Peptides. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 14552-14556.	13.8	21
20	A One-Pot Chemically Cleavable Bis-Linker Tether Strategy for the Synthesis of Heterodimeric Peptides. <i>Angewandte Chemie</i> , 2016, 128, 14772-14776.	2.0	5
21	Total Chemical Synthesis of an Intra-Chain Cystathionine Human Insulin Analogue with Enhanced Thermal Stability. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 14743-14747.	13.8	45
22	Total Chemical Synthesis of an Intra-Chain Cystathionine Human Insulin Analogue with Enhanced Thermal Stability. <i>Angewandte Chemie</i> , 2016, 128, 14963-14967.	2.0	18
23	2-Nitroveratryl as a Photocleavable Thiol-Protecting Group for Directed Disulfide Bond Formation in the Chemical Synthesis of Insulin. <i>Chemistry - A European Journal</i> , 2014, 20, 9549-9552.	3.3	48
24	Chemical Synthesis of a Fluorescent IGF-II Analogue. <i>International Journal of Peptide Research and Therapeutics</i> , 2013, 19, 61-69.	1.9	5
25	Synthetic dityrosine-linked I ² -amyloid dimers form stable, soluble, neurotoxic oligomers. <i>Chemical Science</i> , 2013, 4, 4449.	7.4	44
26	Total Chemical Synthesis of a Heterodimeric Interchain Bis-Lactam-Linked Peptide: Application to an Analogue of Human Insulin-Like Peptide 3. <i>International Journal of Peptides</i> , 2013, 2013, 1-8.	0.7	13
27	One-step radiosynthesis of 4-nitrophenyl 2-[¹⁸ F]fluoropropionate ([¹⁸ F]NFP); improved preparation of radiolabeled peptides for PET imaging. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2013, 56, 726-730.	1.0	21
28	Modulation of Conotoxin Structure and Function Is Achieved through a Multienzyme Complex in the Venom Glands of Cone Snails. <i>Journal of Biological Chemistry</i> , 2012, 287, 34288-34303.	3.4	41
29	A Cyclic Peptide Inhibitor of ApoC-II Peptide Fibril Formation: Mechanistic Insight from NMR and Molecular Dynamics Analysis. <i>Journal of Molecular Biology</i> , 2012, 416, 642-655.	4.2	16
30	Microwave Synthesis of Prion Protein Fragments up to 111 Amino Acids in Length Generates Biologically Active Peptides. <i>International Journal of Peptide Research and Therapeutics</i> , 2012, 18, 21-29.	1.9	11
31	Total Synthesis of the Antifungal Depsipeptide Petriellin A. <i>Journal of Organic Chemistry</i> , 2011, 76, 6686-6693.	3.2	27
32	Gallium-68 Complex of a Macrobicyclic Cage Amine Chelator Tethered to Two Integrin-Targeting Peptides for Diagnostic Tumor Imaging. <i>Bioconjugate Chemistry</i> , 2011, 22, 2093-2103.	3.6	49
33	Macrobicyclic Cage Amine Ligands for Copper Radiopharmaceuticals: A Single Bivalent Cage Amine Containing Two Lys3-bombesin Targeting Peptides. <i>Inorganic Chemistry</i> , 2011, 50, 6701-6710.	4.0	49
34	Synthesis of Peptide Sequences Derived from Fibril-Forming Proteins. <i>Methods in Molecular Biology</i> , 2011, 752, 29-43.	0.9	0
35	Embryonic Toxin Expression in the Cone Snail <i>Conus victoriae</i> . <i>Journal of Biological Chemistry</i> , 2011, 286, 22546-22557.	3.4	31
36	Novel Venom Proteins Produced by Differential Domain-Expression Strategies in Beaded Lizards and Gila Monsters (genus <i>Heloderma</i>). <i>Molecular Biology and Evolution</i> , 2010, 27, 395-407.	8.9	85

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37	Effects of mutation on the amyloidogenic propensity of apolipoprotein C-II60â€“70 peptide. <i>Physical Chemistry Chemical Physics</i> , 2010, 12, 14762.	2.8	15
38	Functional and Structural Diversification of the Anguimorpha Lizard Venom System. <i>Molecular and Cellular Proteomics</i> , 2010, 9, 2369-2390.	3.8	70
39	Versatile New Bis(thiosemicarbazone) Bifunctional Chelators: Synthesis, Conjugation to Bombesin(7âˆ“14)-NH ₂ , and Copper-64 Radiolabeling. <i>Inorganic Chemistry</i> , 2010, 49, 1884-1893.	4.0	76
40	Aromatic residues in the C-terminal helix of human apoC-I mediate phospholipid interactions and particle morphology. <i>Journal of Lipid Research</i> , 2009, 50, 1384-1394.	4.2	13
41	Rapid Optimization of a Peptide Inhibitor of Malaria Parasite Invasion by Comprehensive N-Methyl Scanning. <i>Journal of Biological Chemistry</i> , 2009, 284, 9361-9371.	3.4	54
42	A new bifunctional chelator for copper radiopharmaceuticals: a cage amine ligand with a carboxylate functional group for conjugation to peptides. <i>Chemical Communications</i> , 2009, , 3237.	4.1	55
43	A central role for venom in predation by <i>Varanus komodoensis</i> (Komodo Dragon) and the extinct giant <i>Varanus</i> (<i>Megalania</i>) <i>priscus</i> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 8969-8974.	7.1	120
44	Solid-phase synthesis of homodimeric peptides: preparation of covalently-linked dimers of amyloid Î² peptide. <i>Chemical Communications</i> , 2009, , 6228.	4.1	39
45	Comprehensive N-Methyl Scanning of a Potent Peptide Inhibitor of Malaria Invasion into Erythrocytes Leads to Pharmacokinetic Optimization of the Molecule. <i>International Journal of Peptide Research and Therapeutics</i> , 2008, 14, 381-386.	1.9	4