

Jing Shi

List of Publications by Year in descending order

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43
papers

848
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516710

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1071
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#	ARTICLE	IF	CITATIONS
1	Selective conversion of furfural to cyclopentanone or cyclopentanol using different preparation methods of Cu–Co catalysts. <i>Green Chemistry</i> , 2015, 17, 1038-1046.	9.0	168
2	Cysteine-Aminoethylation-Assisted Chemical Ubiquitination of Recombinant Histones. <i>Journal of the American Chemical Society</i> , 2019, 141, 3654-3663.	13.7	62
3	Aerobic oxidative esterification of 5-hydroxymethylfurfural to dimethyl furan-2,5-dicarboxylate by using homogeneous and heterogeneous PdCoBi/C catalysts under atmospheric oxygen. <i>Green Chemistry</i> , 2018, 20, 3050-3058.	9.0	58
4	Mechanistic Origin of Regioselectivity in Nickel-Catalyzed Olefin Hydroheteroarylation through C–H Activation. <i>Organometallics</i> , 2012, 31, 4356-4366.	2.3	56
5	A Theoretical Study on C–COOH Homolytic Bond Dissociation Enthalpies. <i>Journal of Physical Chemistry A</i> , 2010, 114, 6263-6272.	2.5	42
6	Mechanism and Origin of the Stereoselectivity in the Palladium-Catalyzed <i>trans</i> Hydroboration of Internal 1,3-Enynes with an Azaborine-Based Phosphine Ligand. <i>Chemistry - A European Journal</i> , 2018, 24, 178-186.	3.3	35
7	Chemical synthesis of a cyclotide via intramolecular cyclization of peptide O-esters. <i>Science China Chemistry</i> , 2012, 55, 64-69.	8.2	32
8	Selective modification of natural nucleophilic residues in peptides and proteins using arylpalladium complexes. <i>Organic Chemistry Frontiers</i> , 2018, 5, 3186-3193.	4.5	30
9	An activity-based probe developed by a sequential dehydroalanine formation strategy targets HECT E3 ubiquitin ligases. <i>Chemical Communications</i> , 2019, 55, 7109-7112.	4.1	25
10	Diaminodiacid-based solid-phase synthesis of all-hydrocarbon stapled α -helical peptides. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 6286-6290.	2.8	24
11	An E1-Catalyzed Chemoenzymatic Strategy to Isopeptide-N-Ethylated Deubiquitylase-Resistant Ubiquitin Probes. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 13496-13501.	13.8	23
12	Synthesis of Peptide Disulfide-Bond Mimics by Using Fully Orthogonally Protected Diaminodiacids. <i>Organic Letters</i> , 2018, 20, 6074-6078.	4.6	20
13	Chemical synthesis and biological activity of peptides incorporating an ether bridge as a surrogate for a disulfide bond. <i>Chemical Science</i> , 2020, 11, 7927-7932.	7.4	20
14	Chemical Synthesis of Natural Polyubiquitin Chains through Auxiliary-Mediated Ligation of an Expressed Ubiquitin Isomer. <i>Organic Letters</i> , 2018, 20, 329-332.	4.6	19
15	A computational study of C–X (X = H, C, F, Cl) bond dissociation enthalpies (BDEs) in polyhalogenated methanes and ethanes. <i>Journal of Physical Organic Chemistry</i> , 2011, 24, 65-73.	1.9	18
16	Engineered fluorescence tags for in vivo protein labelling. <i>RSC Advances</i> , 2014, 4, 7235-7245.	3.6	18
17	Dmb/ivDde protected diaminodiacids for solid-phase synthesis of peptide disulfide-bond mimics. <i>Tetrahedron Letters</i> , 2017, 58, 1677-1680.	1.4	17
18	Hydride Dissociation Energies of Six-Membered Heterocyclic Organic Hydrides Predicted by ONIOM-G4Method. <i>Journal of Chemical Information and Modeling</i> , 2012, 52, 63-75.	5.4	16

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19	Mechanism for the enhanced reactivity of 4-mercaptopropyl thioesters in native chemical ligation. <i>RSC Advances</i> , 2016, 6, 68312-68321.	3.6	15
20	Design of new neutral organic super π -electron donors: a theoretical study. <i>Journal of Physical Organic Chemistry</i> , 2010, 23, 75-83.	1.9	14
21	Photocaging of Activity β -Based Ubiquitin Probes via a C α -Terminal Backbone Modification Strategy. <i>Angewandte Chemie - International Edition</i> , 2022, 61, .	13.8	14
22	Efficient synthesis of hydrocarbon-bridged diaminiodiacids through nickel-catalyzed reductive cross-coupling. <i>Tetrahedron Letters</i> , 2017, 58, 3970-3973.	1.4	12
23	Mechanistic Study of Copper-Catalyzed Decarboxylative C α -N Cross-Coupling with Hypervalent Iodine Oxidant. <i>Organometallics</i> , 2017, 36, 2081-2087.	2.3	11
24	Efficient semi-synthesis of ubiquitin-7-amino-4-methylcoumarin. <i>Tetrahedron</i> , 2018, 74, 3931-3935.	1.9	10
25	Robust synthesis of C-terminal cysteine-containing peptide acids through a peptide hydrazide-based strategy. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 5698-5702.	2.8	10
26	A mechanistic study on the regioselective Ni-catalyzed methylation α -alkenylation of alkyne with AlMe ₃ and allylic alcohol. <i>Organic Chemistry Frontiers</i> , 2021, 9, 163-172.	4.5	9
27	Heterocyclic analogs of phenol as novel potential antioxidants. <i>Journal of Physical Organic Chemistry</i> , 2009, 22, 1038-1047.	1.9	7
28	Efficient chemical synthesis for the analogue of ubiquitin-based probe Ub α -AMC with native bioactivity. <i>RSC Advances</i> , 2016, 6, 47926-47930.	3.6	7
29	A mechanistic study on Cu(i) catalyzed carboxylation of the C α -F bond with CO ₂ : a DFT study. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 9065-9071.	2.8	7
30	Non-reducible disulfide bond replacement implies that disulfide exchange is not required for hepcidin α -ferroportin interaction. <i>Chemical Communications</i> , 2019, 55, 2821-2824.	4.1	6
31	One-Pot Synthesis of a Bis-Thio-Acetone Linked Ubiquitinated Histones Using 1,3-Dibromoacetone. <i>Journal of Organic Chemistry</i> , 2020, 85, 15631-15637.	3.2	6
32	Chemical Synthesis of Six α -Atom Thioether Bridged Diaminiodiacid for Solid α -Phase Synthesis of Peptide Disulfide Bond Mimics. <i>ChemistrySelect</i> , 2020, 5, 1359-1363.	1.5	6
33	Semisynthesis of Ubiquitin and SUMO-Rhodamine 110-Glycine through Aminolysis of Boc-Protected Thioester Counterparts. <i>Journal of Organic Chemistry</i> , 2019, 84, 14861-14867.	3.2	5
34	Efficient Semi α -Synthesis of Atypical Ubiquitin Chains and Ubiquitin α -Based Probes Forged by Thioether Isopeptide Bonds. <i>Chemistry - A European Journal</i> , 2019, 25, 16668-16675.	3.3	5
35	Chemical synthesis of disulfide surrogate peptides by using beta-carbon dimethyl modified diaminiodiacids. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 9021-9025.	2.8	5
36	Chemical Synthesis of diSUMO Photoaffinity Probes for the Identification of PolySUMO Chain-Specific Interacting Proteins. <i>CCS Chemistry</i> , 2021, 3, 1157-1168.	7.8	4

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37	Photocaging of Activity-Based Ubiquitin Probes via a C-Terminal Backbone Modification Strategy. <i>Angewandte Chemie</i> , 2022, 134, .	2.0	4
38	An E1-Catalyzed Chemoenzymatic Strategy to Isopeptide-N-Ethylated Deubiquitylase-Resistant Ubiquitin Probes. <i>Angewandte Chemie</i> , 2020, 132, 13598-13603.	2.0	3
39	Desulfurization Mechanism of Cysteine in Synthesis of Polypeptides. <i>Chinese Journal of Chemical Physics</i> , 2015, 28, 269-276.	1.3	2
40	Acid-sensitive auxiliary assisted atypical diubiquitin synthesis exploiting thiol-ene coupling. <i>Tetrahedron Letters</i> , 2019, 60, 151123.	1.4	2
41	Density Functional Theory Calculations on Ni-Ligand Bond Dissociation Enthalpies. <i>Chinese Journal of Chemical Physics</i> , 2014, 27, 640-646.	1.3	1
42	QUANTUM-CHEMICAL PREDICTION OF FORMATION ENTHALPY OF CYCLOALKANE. <i>Journal of Theoretical and Computational Chemistry</i> , 2010, 09, 155-166.	1.8	0
43	Efficient synthesis of terminal-diazirine-based histone peptide probes. <i>Tetrahedron Letters</i> , 2022, , 153878.	1.4	0