

Seth B Herzon

List of Publications by Year in descending order

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83
papers

2,859
citations

126858

33
h-index

197736

49
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99
all docs

99
docs citations

99
times ranked

2465
citing authors

#	ARTICLE	IF	CITATIONS
1	Fragmentverknüpfungen in der Totalsynthese – Bildung von C–C-Bindungen über intermediäre Carbanionen oder freie Radikale. <i>Angewandte Chemie</i> , 2021, 133, 1132-1167.	1.6	5
2	Fragment Coupling Reactions in Total Synthesis That Form Carbon–Carbon Bonds via Carbanionic or Free Radical Intermediates. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 1116-1150.	7.2	32
3	Synthesis and Biological Evaluation of (2S,2'-S)-Lomaiviticin A. <i>Journal of the American Chemical Society</i> , 2021, 143, 1126-1132.	6.6	8
4	Enantioselective Synthesis of Euonyminol. <i>Journal of the American Chemical Society</i> , 2021, 143, 699-704.	6.6	15
5	General Method for the Synthesis of 1- or 2-Deoxyaminoglycosides Bearing Basic Nitrogen. <i>Journal of the American Chemical Society</i> , 2021, 143, 2777-2783.	6.6	20
6	Metric-Based Analysis of Convergence in Complex Molecule Synthesis. <i>Accounts of Chemical Research</i> , 2021, 54, 903-916.	7.6	10
7	Structure Revision of the Lomaiviticins. <i>Journal of the American Chemical Society</i> , 2021, 143, 6578-6585.	6.6	36
8	On the Stability and Spectroscopic Properties of 5-Hydroxyoxazole-4-carboxylic Acid Derivatives. <i>Organic Letters</i> , 2021, 23, 5457-5460.	2.4	2
9	Natural Products: An Era of Discovery in Organic Chemistry. <i>Journal of Organic Chemistry</i> , 2021, 86, 10943-10945.	1.7	3
10	Probing Microbiome Genotoxicity: A Stable Colibactin Provides Insight into Structure–Activity Relationships and Facilitates Mechanism of Action Studies. <i>Journal of the American Chemical Society</i> , 2021, 143, 15824-15833.	6.6	8
11	Chemoproteomic Profiling by Cysteine Fluoroalkylation Reveals Myrocin G as an Inhibitor of the Nonhomologous End Joining DNA Repair Pathway. <i>Journal of the American Chemical Society</i> , 2021, 143, 20332-20342.	6.6	22
12	Development of an Enantioselective Synthesis of (R)-Euonyminol. <i>Journal of Organic Chemistry</i> , 2021, 86, 17011-17035.	1.7	6
13	Employing chemical synthesis to study the structure and function of colibactin, a “dark matter” metabolite. <i>Natural Product Reports</i> , 2020, 37, 1532-1548.	5.2	12
14	Macrocyclic colibactins. <i>Nature Chemistry</i> , 2020, 12, 1005-1006.	6.6	6
15	Structure and bioactivity of colibactin. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127280.	1.0	44
16	New Leads for the Treatment of Multidrug Resistant Mycobacterium tuberculosis. <i>ACS Central Science</i> , 2020, 6, 833-835.	5.3	3
17	Development of a Convergent Enantioselective Synthetic Route to (R)-Myrocin G. <i>Journal of Organic Chemistry</i> , 2020, 85, 8952-8989.	1.7	5
18	Synthesis of the bis(cyclohexenone) core of (R)-lomaiviticin A. <i>Chemical Science</i> , 2020, 11, 7462-7467.	3.7	6

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19	Depurination of Colibactin-Derived Interstrand Cross-Links. <i>Biochemistry</i> , 2020, 59, 892-900.	1.2	25
20	Synthesis of (â€“)â€“)-Myrocin G via a Cascade Coupling. <i>Trends in Chemistry</i> , 2020, 2, 776-777.	4.4	0
21	Antibacterial properties and clinical potential of pleuromutilins. <i>Natural Product Reports</i> , 2019, 36, 220-247.	5.2	64
22	Structure elucidation of colibactin and its DNA cross-links. <i>Science</i> , 2019, 365, .	6.0	158
23	Synthesis and reactivity of precolibactin 886. <i>Nature Chemistry</i> , 2019, 11, 890-898.	6.6	31
24	Programmable Synthesis of 2-Deoxyglycosides. <i>Journal of the American Chemical Society</i> , 2019, 141, 8098-8103.	6.6	51
25	A convergent approach to batzelladine alkaloids. Total syntheses of (+)-batzelladine E, (â€“)â€“)-dehydrobatzelladine C, and (+)-batzelladine K. <i>Tetrahedron</i> , 2018, 74, 3188-3197.	1.0	11
26	Directed Câ€“H Bond Oxidation of (+)-Pleuromutilin. <i>Journal of Organic Chemistry</i> , 2018, 83, 6843-6892.	1.7	23
27	DNA Repair: Unconventional Lesions Require Unconventional Repair. <i>Biochemistry</i> , 2018, 57, 1057-1058.	1.2	0
28	Synthesis of Myrocin G, the Putative Active Form of the Myrocin Antitumor Antibiotics. <i>Journal of the American Chemical Society</i> , 2018, 140, 16058-16061.	6.6	16
29	Model Colibactins Exhibit Human Cell Genotoxicity in the Absence of Host Bacteria. <i>ACS Chemical Biology</i> , 2018, 13, 3286-3293.	1.6	23
30	Cobalt bis(acetylacetonate)â€“tert-butyl hydroperoxideâ€“triethylsilane: a general reagent combination for the Markovnikov-selective hydrofunctionalization of alkenes by hydrogen atom transfer. <i>Beilstein Journal of Organic Chemistry</i> , 2018, 14, 2259-2265.	1.3	16
31	Characterization of Natural Colibactinâ€“Nucleobase Adducts by Tandem Mass Spectrometry and Isotopic Labeling. Support for DNA Alkylation by Cyclopropane Ring Opening. <i>Biochemistry</i> , 2018, 57, 6391-6394.	1.2	39
32	Emergent Properties of Natural Products. <i>Synlett</i> , 2018, 29, 1823-1835.	1.0	3
33	Domain-Targeted Metabolomics Delineates the Heterocycle Assembly Steps of Colibactin Biosynthesis. <i>Journal of the American Chemical Society</i> , 2017, 139, 4195-4201.	6.6	48
34	A complex stereochemical relay approach to the antimalarial alkaloid ocimicide A ₁ . Evidence for a structural revision. <i>Chemical Science</i> , 2017, 8, 4867-4871.	3.7	3
35	A modular and enantioselective synthesis of the pleuromutilin antibiotics. <i>Science</i> , 2017, 356, 956-959.	6.0	57
36	Hydroheteroarylation of Unactivated Alkenes Using <i>N</i> -Methoxyheteroarenium Salts. <i>Journal of the American Chemical Society</i> , 2017, 139, 5998-6007.	6.6	133

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37	Molecular Basis of Gut Microbiome-Associated Colorectal Cancer: A Synthetic Perspective. <i>Journal of the American Chemical Society</i> , 2017, 139, 14817-14824.	6.6	39
38	The Mechanism of Action of (âˆ²)-Lomaiviticin A. <i>Accounts of Chemical Research</i> , 2017, 50, 2577-2588.	7.6	20
39	Introduction: Natural Product Synthesis. <i>Chemical Reviews</i> , 2017, 117, 11649-11650.	23.0	7
40	Development of a Modular Synthetic Route to (+)-Pleuromutilin, (+)-12- <i>epi</i> -Mutilins, and Related Structures. <i>Journal of the American Chemical Society</i> , 2017, 139, 16377-16388.	6.6	46
41	Structure and Functional Analysis of ClbQ, an Unusual Intermediate-Releasing Thioesterase from the Colibactin Biosynthetic Pathway. <i>ACS Chemical Biology</i> , 2017, 12, 2598-2608.	1.6	32
42	Scalable Synthesis of a Key Intermediate for the Production of Pleuromutilin-Based Antibiotics. <i>Organic Letters</i> , 2017, 19, 4980-4983.	2.4	10
43	ClbS Is a Cyclopropane Hydrolase That Confers Colibactin Resistance. <i>Journal of the American Chemical Society</i> , 2017, 139, 17719-17722.	6.6	52
44	Intermolecular Hydropyridylation of Unactivated Alkenes. <i>Journal of the American Chemical Society</i> , 2016, 138, 8718-8721.	6.6	153
45	Convergent and Modular Synthesis of Candidate Precolibactins. Structural Revision of Precolibactin A. <i>Journal of the American Chemical Society</i> , 2016, 138, 5426-5432.	6.6	49
46	Synthesis of Ketones and Esters from Heteroatom-Functionalized Alkenes by Cobalt-Mediated Hydrogen Atom Transfer. <i>Journal of Organic Chemistry</i> , 2016, 81, 8673-8695.	1.7	37
47	Stereoselective Multicomponent Reactions Using Zincate Nucleophiles: Î²-Dicarbonyl Synthesis and Functionalization. <i>Organic Letters</i> , 2016, 18, 4880-4883.	2.4	12
48	A Mechanistic Model for Colibactin-Induced Genotoxicity. <i>Journal of the American Chemical Society</i> , 2016, 138, 15563-15570.	6.6	66
49	Mechanism of Nucleophilic Activation of (âˆ²)-Lomaiviticin A. <i>Journal of the American Chemical Society</i> , 2016, 138, 15559-15562.	6.6	12
50	Synergistic potentiation of (âˆ²)-lomaiviticin A cytotoxicity by the ATR inhibitor VE-821. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 3122-3126.	1.0	4
51	Structural basis for DNA cleavage by the potent antiproliferative agent (â€“)lomaiviticin A. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, 2851-2856.	3.3	29
52	Characterization of Cardiac Glycoside Natural Products as Potent Inhibitors of DNA Double-Strand Break Repair by a Whole-Cell Double Immunofluorescence Assay. <i>Journal of the American Chemical Society</i> , 2016, 138, 3844-3855.	6.6	43
53	Synthesis of 1,3-Amino Alcohols, 1,3-Diols, Amines, and Carboxylic Acids from Terminal Alkynes. <i>Journal of Organic Chemistry</i> , 2015, 80, 8604-8618.	1.7	28
54	Mechanism of Action Studies of Lomaiviticin A and the Monomeric Lomaiviticin Aglycon. Selective and Potent Activity Toward DNA Double-Strand Break Repair-Deficient Cell Lines. <i>Journal of the American Chemical Society</i> , 2015, 137, 5741-5747.	6.6	17

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55	Multigram synthesis of 1- O -acetyl-3- O -(4-methoxybenzyl)-4- N -(9-fluorenylmethoxycarbonyl)-4- N -methyl-1 -pyrrolidine. <i>Tetrahedron Letters</i> , 2015, 56, 3231-3234.	0.7	1
56	The Discovery of a Novel Route to Highly Substituted $\hat{\pm}$ -Tropolones Enables Expedient Entry to the Core of the Gukulenins. <i>Organic Letters</i> , 2015, 17, 2030-2033.	2.4	24
57	Non-classical selectivities in the reduction of alkenes by cobalt-mediated hydrogen atom transfer. <i>Chemical Science</i> , 2015, 6, 6250-6255.	3.7	74
58	A concise synthesis of (+)-batzelladine B from simple pyrrole-based starting materials. <i>Nature</i> , 2015, 525, 507-510.	13.7	54
59	The Hasubanan and Acutumine Alkaloids. <i>The Alkaloids Chemistry and Biology</i> , 2014, 73, 161-222.	0.8	18
60	Analysis of Diazofluorene DNA Binding and Damaging Activity: DNA Cleavage by a Synthetic Monomeric Diazofluorene. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 9325-9328.	7.2	22
61	The cytotoxicity of ($\hat{\pm}$)-lomaiviticin A arises from induction of double-strand breaks in DNA. <i>Nature Chemistry</i> , 2014, 6, 504-510.	6.6	73
62	Scope and Limitations of 2-Deoxy- and 2,6-Dideoxyglycosyl Bromides as Donors for the Synthesis of $\hat{\pm}$ -2-Deoxy- and $\hat{\pm}$ -2,6-Dideoxyglycosides. <i>Organic Letters</i> , 2014, 16, 2776-2779.	2.4	53
63	Temporal separation of catalytic activities allows anti-Markovnikov reductive functionalization of terminal alkynes. <i>Nature Chemistry</i> , 2014, 6, 22-27.	6.6	51
64	Substrate-Modified Functional Group Reactivity: Hasubanan and Acutumine Alkaloid Syntheses. <i>Journal of Organic Chemistry</i> , 2014, 79, 8937-8947.	1.7	18
65	Broad $\hat{\pm}$ -Spectrum Catalysts for the Ambient Temperature Anti-Markovnikov Hydration of Alkynes. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 7892-7895.	7.2	50
66	A Method for the Selective Hydrogenation of Alkenyl Halides to Alkyl Halides. <i>Journal of the American Chemical Society</i> , 2014, 136, 6884-6887.	6.6	134
67	A practical method for regiocontrolled one-carbon ring contraction. <i>Tetrahedron</i> , 2013, 69, 5634-5639.	1.0	6
68	Development of Enantioselective Synthetic Routes to the Hasubanan and Acutumine Alkaloids. <i>Journal of Organic Chemistry</i> , 2013, 78, 10031-10057.	1.7	44
69	Total Syntheses of ($\hat{\pm}$)-Acutumine and ($\hat{\pm}$)-Dechloroacutumine. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 3642-3645.	7.2	61
70	Direct Synthesis of $\hat{\pm}$ -Glycosides by the Reductive Glycosylation of Azides with Protected and Native Carbohydrate Donors. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 6068-6071.	7.2	17
71	Characterization of a reductively-activated elimination pathway relevant to the biological chemistry of the kinamycins and lomaiviticins. <i>Chemical Science</i> , 2012, 3, 1070-1074.	3.7	16
72	The diazofluorene antitumor antibiotics: Structural elucidation, biosynthetic, synthetic, and chemical biological studies. <i>Natural Product Reports</i> , 2012, 29, 87-118.	5.2	70

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73	Isolation of Lomaiviticins Câ€“E, Transformation of Lomaiviticin C to Lomaiviticin A, Complete Structure Elucidation of Lomaiviticin A, and Structureâ€“Activity Analyses. Journal of the American Chemical Society, 2012, 134, 15285-15288.	6.6	63
74	Synthesis of (<i>R</i>)-(+)-4-Methylcyclohex-2-ene-1-one. Journal of Organic Chemistry, 2012, 77, 9422-9425.	1.7	5
75	The Kinamycins. , 2012, , 39-65.		7
76	Development of Enantioselective Synthetic Routes to (âˆ™)-Kinamycin F and (âˆ™)-Lomaiviticin Aglycon. Journal of the American Chemical Society, 2012, 134, 17262-17273.	6.6	37
77	A robust and scalable synthesis of the potent neuroprotective agent (âˆ™)-huperzine A. Chemical Science, 2011, 2, 2251.	3.7	51
78	11-Step Enantioselective Synthesis of (âˆ™)-Lomaiviticin Aglycon. Journal of the American Chemical Society, 2011, 133, 7260-7263.	6.6	68
79	Efficient Entry to the Hasubanan Alkaloids: First Enantioselective Total Syntheses of (âˆ™)â€“Hasubanonine, (âˆ™)â€“Runanine, (âˆ™)â€“Delavayine, and (+)â€“Periglaucineâ€“B. Angewandte Chemie - International Edition, 2011, 50, 8863-8866.		60
80	Cover Picture: Efficient Entry to the Hasubanan Alkaloids: First Enantioselective Total Syntheses of (âˆ™)â€“Hasubanonine, (âˆ™)â€“Runanine, (âˆ™)â€“Delavayine, and (+)â€“Periglaucineâ€“B (Angew. Chem. Int. Ed. 38/2011). Angewandte Chemie - International Edition, 2011, 50, 8761-8761.		0
81	Single-Step Synthesis of Secondary Phosphine Oxides. Organometallics, 2010, 29, 4193-4195.	1.1	22
82	Development of a Convergent Entry to the Diazofluorene Antitumor Antibiotics: Enantioselective Synthesis of Kinamycin F. Journal of the American Chemical Society, 2010, 132, 2540-2541.	6.6	46
83	Synthesis of the Fully Glycosylated Cyclohexenone Core of Lomaiviticin A. Organic Letters, 2009, 11, 4322-4325.	2.4	39