

Ryan A Shenvi

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

73
papers

3,661
citations

32
h-index

60
g-index

110
ext. papers

4,355
ext. citations

15.5
avg, IF

6.07
L-index

#	Paper	IF	Citations
73	Stereodivergent Attached-Ring Synthesis via Non-Covalent Interactions: A Short Formal Synthesis of Merrilactone A. <i>Angewandte Chemie</i> , 2022 , 134, e202114514	3.6	
72	Concise syntheses of GB22, GB13, and himgaline by cross-coupling and complete reduction.. <i>Science</i> , 2022 , 375, 1270-1274	33.3	3
71	Change the channel: CysLoop receptor antagonists from nature. <i>Pest Management Science</i> , 2021 , 77, 3650-3662	4.6	3
70	Cobalt-catalyzed alkene hydrogenation by reductive turnover. <i>Tetrahedron Letters</i> , 2021 , 72, 153047	2	2
69	Revision of the Unstable Picrotoxinin Hydrolysis Product. <i>Angewandte Chemie</i> , 2021 , 133, 19261-19264	3.6	
68	Natural Product Synthesis through the Lens of Informatics. <i>Accounts of Chemical Research</i> , 2021 , 54, 1157-1167	24.3	2
67	Revision of the Unstable Picrotoxinin Hydrolysis Product. <i>Angewandte Chemie - International Edition</i> , 2021 , 60, 19113-19116	16.4	0
66	Cycloisomerization of Olefins in Water. <i>Angewandte Chemie</i> , 2020 , 132, 13098-13103	3.6	6
65	Synthesis of (-)-Picrotoxinin by Late-Stage Strong Bond Activation. <i>Journal of the American Chemical Society</i> , 2020 , 142, 11376-11381	16.4	15
64	Electronic complementarity permits hindered butenolide heterodimerization and discovery of novel cGAS/STING pathway antagonists. <i>Nature Chemistry</i> , 2020 , 12, 310-317	17.6	16
63	Cycloisomerization of Olefins in Water. <i>Angewandte Chemie - International Edition</i> , 2020 , 59, 12998-13006	16.4	17
62	Catalytic hydrogen atom transfer to alkenes: a roadmap for metal hydrides and radicals. <i>Chemical Science</i> , 2020 , 11, 12401-12422	9.4	55
61	Synthetic, Mechanistic, and Biological Interrogation of Chemical Space En Route to (-)-Bilobalide. <i>Journal of the American Chemical Society</i> , 2020 , 142, 18599-18618	16.4	20
60	Chemical syntheses of the salvinorin chemotype of KOR agonist. <i>Natural Product Reports</i> , 2020 , 37, 1478-1496	14.96	4
59	Natural Products in the "Marketplace": Interfacing Synthesis and Biology. <i>Journal of the American Chemical Society</i> , 2019 , 141, 3332-3346	16.4	29
58	Expanding Reactivity in DNA-Encoded Library Synthesis via Reversible Binding of DNA to an Inert Quaternary Ammonium Support. <i>Journal of the American Chemical Society</i> , 2019 , 141, 9998-10006	16.4	86
57	Hydroalkylation of Olefins To Form Quaternary Carbons. <i>Journal of the American Chemical Society</i> , 2019 , 141, 7709-7714	16.4	81

56	Reanalysis of Lindenatriene, a Building Block for the Synthesis of Lindenane Oligomers. <i>Tetrahedron</i> , 2019 , 75, 3140-3144	2.4	
55	Intermolecular Heck Coupling with Hindered Alkenes Directed by Potassium Carboxylates. <i>Angewandte Chemie</i> , 2019 , 131, 2393-2398	3.6	4
54	Olefin Hydroarylation via Ni/Co Dual Catalysis. <i>Trends in Chemistry</i> , 2019 , 1, 540-541	14.8	1
53	Hydrofunctionalization of Alkenes by Hydrogen-Atom Transfer 2019 , 383-470		10
52	Concise asymmetric synthesis of (-)-bilobalide. <i>Nature</i> , 2019 , 575, 643-646	50.4	20
51	Intermolecular Heck Coupling with Hindered Alkenes Directed by Potassium Carboxylates. <i>Angewandte Chemie - International Edition</i> , 2019 , 58, 2371-2376	16.4	13
50	Mechanism of Action of the Cytotoxic Asmarine Alkaloids. <i>ACS Chemical Biology</i> , 2018 , 13, 1299-1306	4.9	5
49	O6C-20-nor-salvinorin A is a stable and potent KOR agonist. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018 , 28, 2770-2772	2.9	8
48	A review of salvinorin analogs and their kappa-opioid receptor activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018 , 28, 1436-1445	2.9	29
47	Pharmacological characterization of the neurotrophic sesquiterpene jiadifenolide reveals a non-convulsant signature and potential for progression in neurodegenerative disease studies. <i>Biochemical Pharmacology</i> , 2018 , 155, 61-70	6	12
46	Iron-Nickel Dual-Catalysis: A New Engine for Olefin Functionalization and the Formation of Quaternary Centers. <i>Journal of the American Chemical Society</i> , 2018 , 140, 11317-11324	16.4	102
45	Mechanistic Interrogation of Co/Ni-Dual Catalyzed Hydroarylation. <i>Journal of the American Chemical Society</i> , 2018 , 140, 12056-12068	16.4	108
44	Branch-Selective Addition of Unactivated Olefins into Imines and Aldehydes. <i>Journal of the American Chemical Society</i> , 2018 , 140, 16976-16981	16.4	62
43	The High Chemofidelity of Metal-Catalyzed Hydrogen Atom Transfer. <i>Accounts of Chemical Research</i> , 2018 , 51, 2628-2640	24.3	119
42	Stereocontrolled Synthesis of Kalihinol C. <i>Journal of the American Chemical Society</i> , 2017 , 139, 3647-3650	6.4	18
41	Synthesis of (-)-11-O-Debenzoyltashironin: Neurotrophic Sesquiterpenes Cause Hyperexcitation. <i>Journal of the American Chemical Society</i> , 2017 , 139, 9637-9644	16.4	38
40	Dynamic Strategic Bond Analysis Yields a Ten-Step Synthesis of 20-nor-Salvinorin A, a Potent MOR Agonist. <i>ACS Central Science</i> , 2017 , 3, 1329-1336	16.8	23
39	Branch-Selective Hydroarylation: Iodoarene-Olefin Cross-Coupling. <i>Journal of the American Chemical Society</i> , 2016 , 138, 12779-12782	16.4	170

38	Mn-, Fe-, and Co-Catalyzed Radical Hydrofunctionalizations of Olefins. <i>Chemical Reviews</i> , 2016 , 116, 8912-8900	16.2	490
37	Reaction: And You, of Tender Years 1. <i>CheM</i> , 2016 , 1, 334-335	16.2	
36	Synthesis of (+)-7,20-Diisocyanoadociane and Liver-Stage Antiplasmodial Activity of the Isocyanoterpene Class. <i>Journal of the American Chemical Society</i> , 2016 , 138, 7268-71	16.4	50
35	Synthesis and Sulfur Electrophilicity of the Nuphar Thiaspirane Pharmacophore. <i>ACS Central Science</i> , 2016 , 2, 401-8	16.8	17
34	Ph(i-PrO)SiH ₂ : An Exceptional Reductant for Metal-Catalyzed Hydrogen Atom Transfers. <i>Journal of the American Chemical Society</i> , 2016 , 138, 4962-71	16.4	156
33	Neurite outgrowth enhancement by jiadifenolide: possible targets. <i>Natural Product Reports</i> , 2016 , 33, 535-9	15.1	20
32	Conjuring a Supernatural Product [DelMarine]. <i>Synlett</i> , 2016 , 27, 1145-1164	2.2	14
31	Cluster Preface: Reinventing Radical Reactions. <i>Synlett</i> , 2016 , 27, 678-679	2.2	1
30	Synthesis of the Privileged 8-Arylmenthol Class by Radical Arylation of Isopulegol. <i>Organic Letters</i> , 2016 , 18, 2620-3	6.2	56
29	An eight-step gram-scale synthesis of (-)-jiadifenolide. <i>Nature Chemistry</i> , 2015 , 7, 604-7	17.6	67
28	A Longitudinal Study of Alkaloid Synthesis Reveals Functional Group Interconversions as Bad Actors. <i>Chemical Reviews</i> , 2015 , 115, 9465-531	68.1	39
27	Synthesis of Lepadiformine Using a Hydroamination Transform. <i>Organic Letters</i> , 2015 , 17, 5776-9	6.2	11
26	Syntheses and biological studies of marine terpenoids derived from inorganic cyanide. <i>Natural Product Reports</i> , 2015 , 32, 543-77	15.1	53
25	Nitrosipurines en route to potently cytotoxic asmarines. <i>Angewandte Chemie - International Edition</i> , 2015 , 54, 2410-5	16.4	17
24	Supramolecular catalysis: Terpenes in tight spaces. <i>Nature Chemistry</i> , 2015 , 7, 187-9	17.6	4
23	Nitrosipurines En Route to Potently Cytotoxic Asmarines. <i>Angewandte Chemie</i> , 2015 , 127, 2440-2445	3.6	3
22	Simple, chemoselective hydrogenation with thermodynamic stereocontrol. <i>Journal of the American Chemical Society</i> , 2014 , 136, 1300-3	16.4	194
21	Simple, chemoselective, catalytic olefin isomerization. <i>Journal of the American Chemical Society</i> , 2014 , 136, 16788-91	16.4	210

20	Synthesis of medicinally relevant terpenes: reducing the cost and time of drug discovery. <i>Future Medicinal Chemistry</i> , 2014 , 6, 1127-48	4.1	51
19	Synthesis of (-)-neothiobinupharidine. <i>Journal of the American Chemical Society</i> , 2013 , 135, 1209-12	16.4	49
18	Stereoinversion of tertiary alcohols to tertiary-alkyl isonitriles and amines. <i>Nature</i> , 2013 , 501, 195-9	50.4	109
17	Synthesis of a potent antimalarial amphilectene. <i>Journal of the American Chemical Society</i> , 2012 , 134, 19604-6	16.4	72
16	Synthesis of highly strained terpenes by non-stop tail-to-head polycyclization. <i>Nature Chemistry</i> , 2012 , 4, 915-20	17.6	72
15	A stereoselective hydroamination transform to access polysubstituted indolizidines. <i>Journal of the American Chemical Society</i> , 2012 , 134, 2012-5	16.4	45
14	Scalable synthesis of cortistatin A and related structures. <i>Journal of the American Chemical Society</i> , 2011 , 133, 8014-27	16.4	99
13	Synthetic access to bent polycycles by cation-pi cyclization. <i>Organic Letters</i> , 2010 , 12, 3548-51	6.2	34
12	Stereodivergent synthesis of 17-alpha and 17-beta-alkaryl steroids: application and biological evaluation of D-ring cortistatin analogues. <i>Angewandte Chemie - International Edition</i> , 2009 , 48, 4328-31	16.4	52
11	A short and efficient synthesis of (-)-7-methylomuralide, a potent proteasome inhibitor. <i>Journal of the American Chemical Society</i> , 2009 , 131, 5746-7	16.4	25
10	Chemoselectivity: the mother of invention in total synthesis. <i>Accounts of Chemical Research</i> , 2009 , 42, 530-41	24.3	221
9	Synthesis of (+)-cortistatin A. <i>Journal of the American Chemical Society</i> , 2008 , 130, 7241-3	16.4	143
8	One-Step Synthesis of 4,5-Disubstituted Pyrimidines Using Commercially Available and Inexpensive Reagents. <i>Heterocycles</i> , 2006 , 70, 581	0.8	9
7	Total synthesis of (+/-)-chartelline C. <i>Journal of the American Chemical Society</i> , 2006 , 128, 14028-9	16.4	76
6	A remarkable ring contraction en route to the chartelline alkaloids. <i>Angewandte Chemie - International Edition</i> , 2005 , 44, 3714-7	16.4	59
5	A Remarkable Ring Contraction En Route to the Chartelline Alkaloids. <i>Angewandte Chemie</i> , 2005 , 117, 3780-3783	3.6	13
4	Long-range effects on calcium binding and conformational change in the N-domain of calmodulin. <i>Biochemistry</i> , 2001 , 40, 12719-26	3.2	33
3	Synthesis and Mechanistic Interrogation of Ginkgo biloba Chemical Space en route to (-)-Bilobalide		2

2	Concise Synthesis of GB22 by Endo-Selective Siloxycyclopropane Arylation	7
1	Concise Synthesis of GB22 by Endo-Selective Siloxycyclopropane Arylation	2