

# Erik J Sorensen

## List of Publications by Year in descending order

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70  
papers

4,419  
citations

117571

34  
h-index

110317

64  
g-index

83  
all docs

83  
docs citations

83  
times ranked

4329  
citing authors

#	ARTICLE	IF	CITATIONS
1	Recent applications of C-H functionalization in complex natural product synthesis. <i>Chemical Society Reviews</i> , 2018, 47, 8925-8967.	18.7	470
2	Proteomic profiling of mechanistically distinct enzyme classes using a common chemotype. <i>Nature Biotechnology</i> , 2002, 20, 805-809.	9.4	229
3	Protein-Reactive Natural Products. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 5788-5809.	7.2	228
4	Target discovery in small-molecule cell-based screens by in situ proteome reactivity profiling. <i>Nature Biotechnology</i> , 2005, 23, 1303-1307.	9.4	210
5	Chemical Strategies for Functional Proteomics. <i>Molecular and Cellular Proteomics</i> , 2002, 1, 781-790.	2.5	162
6	Acceptorless dehydrogenation of small molecules through cooperative base metal catalysis. <i>Nature Communications</i> , 2015, 6, 10093.	5.8	162
7	The Uranyl Cation as a Visible-Light Photocatalyst for C(sp <sup>3</sup> )-H Fluorination. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 8923-8927.	7.2	142
8	An Enantioselective Synthesis of FR182877 Provides a Chemical Rationalization of Its Structure and Affords Multigram Quantities of Its Direct Precursor. <i>Journal of the American Chemical Society</i> , 2003, 125, 5393-5407.	6.6	141
9	Profiling the specific reactivity of the proteome with non-directed activity-based probes. <i>Chemistry and Biology</i> , 2001, 8, 81-95.	6.2	136
10	Cyclostreptin binds covalently to microtubule pores and luminal taxoid binding sites. , 2007, 3, 117-125.		130
11	Pd-Catalyzed, <i>ortho</i> -C-H Methylation and Fluorination of Benzaldehydes Using Orthoanilic Acids as Transient Directing Groups. <i>Journal of the American Chemical Society</i> , 2018, 140, 2789-2792.	6.6	129
12	Synthesis of the Furanosteroidal Antibiotic Viridin. <i>Angewandte Chemie - International Edition</i> , 2004, 43, 1998-2001.	7.2	115
13	A Synthesis of (+)-FR182877, Featuring Tandem Transannular Diels-Alder Reactions Inspired by a Postulated Biogenesis. <i>Journal of the American Chemical Society</i> , 2002, 124, 4552-4553.	6.6	108
14	Convergent, Enantioselective Syntheses of Guanacastepenes A and E Featuring a Selective Cyclobutane Fragmentation. <i>Journal of the American Chemical Society</i> , 2006, 128, 7025-7035.	6.6	104
15	Abyssomicins G and H and atrop-Abyssomicin C from the Marine <i>Verrucospora</i> Strain AB-18-032. <i>Journal of Antibiotics</i> , 2007, 60, 391-394.	1.0	102
16	Synthesis of Fluorenones from Benzaldehydes and Aryl Iodides: Dual C-H Functionalizations Using a Transient Directing Group. <i>Organic Letters</i> , 2017, 19, 1140-1143.	2.4	100
17	A Diels-Alder Macrocyclization Enables an Efficient Asymmetric Synthesis of the Antibacterial Natural Product Abyssomicin C. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 6533-6537.	7.2	95
18	Bond formations by intermolecular and intramolecular trappings of acylketenes and their applications in natural product synthesis. <i>Chemical Society Reviews</i> , 2009, 38, 3022.	18.7	95

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19	Pd-Catalyzed <i>ortho</i> -C-H Hydroxylation of Benzaldehydes Using a Transient Directing Group. <i>Organic Letters</i> , 2017, 19, 6280-6283.	2.4	83
20	An Enantiospecific Synthesis of Jiadifenolide. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 5332-5335.	7.2	81
21	A Convergent Synthesis of the Tricyclic Architecture of the Guanacastepenes Featuring a Selective Ring Fragmentation. <i>Organic Letters</i> , 2002, 4, 2063-2066.	2.4	71
22	( <i>â</i> )-FR182877 Is a Potent and Selective Inhibitor of Carboxylesterase-1. <i>Angewandte Chemie - International Edition</i> , 2003, 42, 5480-5484.	7.2	57
23	Cyclostreptin (FR182877), an Antitumor Tubulin-Polymerizing Agent Deficient in Enhancing Tubulin Assembly Despite Its High Affinity for the Taxoid Site. <i>Biochemistry</i> , 2005, 44, 11525-11538.	1.2	55
24	Intramolecular Allenolate Acylations in Studies toward a Synthesis of FR182877. <i>Organic Letters</i> , 2001, 3, 4307-4310.	2.4	51
25	An interrupted Ugi reaction enables the preparation of substituted indoxyls and aminoindoles. <i>Tetrahedron</i> , 2009, 65, 3096-3101.	1.0	51
26	A Rapid, Asymmetric Synthesis of the Decahydrofluorene Core of the Hirsutellones. <i>Organic Letters</i> , 2009, 11, 701-703.	2.4	50
27	Postulated Biogenesis of WS9885B and Progress toward an Enantioselective Synthesis. <i>Organic Letters</i> , 1999, 1, 645-648.	2.4	48
28	The Abyssomicin C family as in vitro inhibitors of <i>Mycobacterium tuberculosis</i> . <i>Tuberculosis</i> , 2010, 90, 298-300.	0.8	47
29	Mechanistic and structural requirements for active site labeling of phosphoglycerate mutase by spiroepoxides. <i>Molecular BioSystems</i> , 2007, 3, 495.	2.9	46
30	A concise synthesis of the molecular framework of pleuromutilin. <i>Chemical Communications</i> , 2011, 47, 1500-1502.	2.2	43
31	A chemical synthesis of 11-methoxy mitragynine pseudoindoxyl featuring the interrupted Ugi reaction. <i>Chemical Science</i> , 2012, 3, 2849.	3.7	42
32	Design, Synthesis, and Reactivity of 1-Hydrazinodienes for Use in Organic Synthesis. <i>Journal of the American Chemical Society</i> , 2005, 127, 8612-8613.	6.6	40
33	Efficient substrate screening and inhibitor testing of human CYP4Z1 using permeabilized recombinant fission yeast. <i>Biochemical Pharmacology</i> , 2017, 146, 174-187.	2.0	40
34	Pd(II)-Catalyzed Synthesis of Benzocyclobutenes by $\beta$ -Methylene-Selective C(sp <sup>3</sup> )-H Arylation with a Transient Directing Group. <i>Journal of the American Chemical Society</i> , 2021, 143, 20035-20041.	6.6	37
35	A Concise Synthesis of Fumagillol. <i>Angewandte Chemie - International Edition</i> , 1999, 38, 971-974.	7.2	35
36	Toward a mild dehydroformylation using base-metal catalysis. <i>Chemical Science</i> , 2017, 8, 1954-1959.	3.7	35

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37	Rapid Construction of a Benzo-fused Indoxamycin Core Enabled by Site-selective C-H Functionalizations. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 8270-8274.	7.2	34
38	The Uranyl Cation as a Visible-Light Photocatalyst for C(sp <sup>3</sup> )-H Fluorination. <i>Angewandte Chemie</i> , 2016, 128, 9069-9073.	1.6	33
39	Ir(III)-catalyzed ortho-C-H alkylations of (hetero)aromatic aldehydes using alkyl boron reagents. <i>Chemical Science</i> , 2018, 9, 8951-8956.	3.7	33
40	Architectural self-construction in nature and chemical synthesis. <i>Bioorganic and Medicinal Chemistry</i> , 2003, 11, 3225-3228.	1.4	32
41	Expanding the pleuromutilin class of antibiotics by de novo chemical synthesis. <i>Chemical Science</i> , 2011, 2, 1258.	3.7	26
42	CHEMISTRY: Enhanced: A Dash of Proline Makes Things Sweet. <i>Science</i> , 2004, 305, 1725-1726.	6.0	24
43	Synthesis enables a structural revision of the Mycobacterium tuberculosis-produced diterpene, edaxadiene. <i>Chemical Science</i> , 2010, 1, 202.	3.7	24
44	Toward a Synthesis of Hirsutellone B by the Concept of Double Cyclization. <i>Journal of Organic Chemistry</i> , 2013, 78, 9584-9607.	1.7	24
45	Crystal Structure of the Cyclostreptin-Tubulin Adduct: Implications for Tubulin Activation by Taxane-Site Ligands. <i>International Journal of Molecular Sciences</i> , 2019, 20, 1392.	1.8	24
46	Cyclization by C(sp <sup>3</sup> )-H Arylation with a Transient Directing Group for the Diastereoselective Preparation of Indanes. <i>ACS Catalysis</i> , 2021, 11, 3115-3127.	5.5	22
47	Transition States of the Retro-Ene Reactions of Allylic Diazenes. <i>Organic Letters</i> , 2006, 8, 3105-3107.	2.4	21
48	Tandem Diels-Alder and Retro-Ene Reactions of 1-Sulfonyl- and 1-Sulfinyl-1,3-dienes as a Traceless Route to Cyclohexenes. <i>Journal of the American Chemical Society</i> , 2014, 136, 9918-9921.	6.6	21
49	Development of a Bio-inspired Dual Catalytic System for Alkane Dehydrogenation. <i>Israel Journal of Chemistry</i> , 2017, 57, 259-269.	1.0	21
50	A Concise and Convergent Synthesis of PA-824. <i>Journal of Organic Chemistry</i> , 2010, 75, 7479-7482.	1.7	19
51	Importance of asparagine-381 and arginine-487 for substrate recognition in CYP4Z1. <i>Biochemical Pharmacology</i> , 2020, 174, 113850.	2.0	17
52	Seebach's Conjugative Reagent Enables Double Cyclizations. <i>Organic Letters</i> , 2010, 12, 2746-2749.	2.4	16
53	The Catalytic Asymmetric Diels-Alder Reactions and Post-cycloaddition Reductive Transpositions of 1-Hydrazinodienes. <i>Chemistry - A European Journal</i> , 2011, 17, 11131-11134.	1.7	16
54	In Vitro Reconstitution of OxyA Enzymatic Activity Clarifies Late Steps in Vancomycin Biosynthesis. <i>ACS Chemical Biology</i> , 2017, 12, 2248-2253.	1.6	13

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55	Modulating OxyB-Catalyzed Cross-Coupling Reactions in Vancomycin Biosynthesis by Incorporation of Diverse $\alpha$ -Tyr Analogues. <i>Journal of Organic Chemistry</i> , 2018, 83, 7309-7317.	1.7	12
56	Installation of multiple aryl ether crosslinks onto non-native substrate peptides by the vancomycin OxyB. <i>Tetrahedron</i> , 2018, 74, 3231-3237.	1.0	9
57	New ProLuciferin Substrates for Human CYP4 Family Enzymes. <i>Applied Biochemistry and Biotechnology</i> , 2021, 193, 218-237.	1.4	9
58	A C-H Functionalization Strategy Enables an Enantioselective Formal Synthesis of ( $\alpha$ )-Aflatoxin B <sub>2</sub> . <i>Organic Letters</i> , 2021, 23, 9393-9397.	2.4	9
59	An expedient synthesis of maraviroc (UK-427,857) via C-H functionalization. <i>Tetrahedron Letters</i> , 2015, 56, 3620-3623.	0.7	8
60	A Stereocontrolled Annulation of the Taccalonolide Epoxy Lactone onto the Molecular Framework of <i>trans</i> -Androsterone. <i>Organic Letters</i> , 2017, 19, 4892-4895.	2.4	7
61	Rhodium-Catalyzed C-H Amination: A Case Study of Selectivity in C-H Functionalization Reactions. <i>Journal of Chemical Education</i> , 2018, 95, 2243-2248.	1.1	7
62	Synthesis of (+)-Lineariifolianone and Related Cyclopropenone-Containing Sesquiterpenoids. <i>Journal of Organic Chemistry</i> , 2019, 84, 5524-5534.	1.7	7
63	Donor-Acceptor-Acceptor 1,3-Bisdiazo Compounds: An Exploration of Synthesis and Stepwise Reactivity. <i>Organic Letters</i> , 2020, 22, 1791-1795.	2.4	6
64	Rapid Construction of a Benzo-fused Indoxamycin Core Enabled by Site-selective C-H Functionalizations. <i>Angewandte Chemie</i> , 2016, 128, 8410-8414.	1.6	4
65	Design and synthesis of molecular scaffolds with anti-infective activity. <i>Tetrahedron</i> , 2016, 72, 3579-3592.	1.0	4
66	Diastereoselective syntheses of substituted cis-hydrindanones featuring sequential inter- and intramolecular Michael reactions. <i>Tetrahedron</i> , 2016, 72, 3713-3717.	1.0	3
67	New luciferin-based probe substrates for human CYP26A1. <i>Biochemistry and Biophysics Reports</i> , 2020, 24, 100861.	0.7	3
68	Conversion of five proLuciferin esters by human cytochrome P450 enzymes. <i>Biotechnology Journal</i> , 2021, 16, 2100007.	1.8	3
69	A Dash of Proline Makes Things Sweet. <i>ChemInform</i> , 2005, 36, no.	0.1	0
70	Protein-Reactive Natural Products. <i>ChemInform</i> , 2005, 36, no.	0.1	0