

Jonathan Sperry

List of Publications by Year in descending order

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141
papers

3,403
citations

136950

32
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197818

49
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159
all docs

159
docs citations

159
times ranked

3818
citing authors

#	ARTICLE	IF	CITATIONS
1	Catalytic deep eutectic solvent for levoglucosenone production by pyrolysis of cellulose. <i>Bioresource Technology</i> , 2022, 344, 126323.	9.6	10
2	Progress toward a biomimetic synthesis of pegaharmaline A. <i>Organic and Biomolecular Chemistry</i> , 2022, 20, 1275-1283.	2.8	1
3	Isolation and biological activity of azocine and azocane alkaloids. <i>Bioorganic and Medicinal Chemistry</i> , 2022, 54, 116560.	3.0	13
4	Production of biomass-based composite from reed pretreated by ball-milling combined with p-toluenesulfonic acid. <i>Industrial Crops and Products</i> , 2022, 180, 114712.	5.2	1
5	Itaconate is a covalent inhibitor of the <i>Mycobacterium tuberculosis</i> isocitrate lyase. <i>RSC Medicinal Chemistry</i> , 2021, 12, 57-61.	3.9	28
6	Synthetic Studies toward Bisindigotin: Polyheteroaromatic Scaffolds via Skeletal Rearrangements of a Diacetoxytetraindole. <i>Journal of Organic Chemistry</i> , 2021, 86, 74-78.	3.2	5
7	Remediation of poly- and perfluoroalkyl substances (PFAS) contaminated soils – To mobilize or to immobilize or to degrade?. <i>Journal of Hazardous Materials</i> , 2021, 401, 123892.	12.4	169
8	Synthetic studies toward inducamide C. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 416-420.	2.8	5
9	Synthesis of 3-nitroindoles by sequential paired electrolysis. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 7903-7913.	2.8	9
10	Bioinspired Synthesis of the Europyrazine Alkaloid Hyrtioseragamine A. <i>Journal of Organic Chemistry</i> , 2021, 86, 4779-4785.	3.2	4
11	Tetrahydrocarbazoles by mechanochemical Fischer indolisation. <i>Tetrahedron Letters</i> , 2021, 72, 153068.	1.4	6
12	The Mechanism of Surface-Radical Generation and Amorphization of Crystalline Quartz Sand upon Mechanochemical Grinding. <i>Journal of Physical Chemistry C</i> , 2021, 125, 20877-20886.	3.1	18
13	Synthesis of bio-based 2-thiothiophenes. <i>Philosophical Transactions Series A, Mathematical, Physical, and Engineering Sciences</i> , 2021, 379, 20200350.	3.4	0
14	Octacycles and Nonacycles from 3-Hydroxy-2,2-bisindole. <i>Journal of Organic Chemistry</i> , 2021, , .	3.2	2
15	Influence of ionic liquid type on porous carbon formation during the ionothermal pyrolysis of cellulose. <i>Journal of Analytical and Applied Pyrolysis</i> , 2020, 145, 104728.	5.5	19
16	Insights into the active sites and catalytic mechanism of oxidative esterification of 5-hydroxymethylfurfural by metal-organic frameworks-derived N-doped carbon. <i>Journal of Catalysis</i> , 2020, 381, 570-578.	6.2	56
17	Pyridine alkaloids with activity in the central nervous system. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115820.	3.0	50
18	The curious yellow colouring matter of the Iceland poppy. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 5278-5286.	2.8	2

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19	Jiangrine-like scaffolds from biorenewable platforms. <i>Tetrahedron Letters</i> , 2020, 61, 152538.	1.4	6
20	Manganese catalyzed transfer hydrogenation of biomass-derived aldehydes: Insights to the catalytic performance and mechanism. <i>Journal of Catalysis</i> , 2020, 389, 157-165.	6.2	28
21	Haber-independent, diversity-oriented synthesis of nitrogen compounds from biorenewable chitin. <i>Green Chemistry</i> , 2020, 22, 1978-1984.	9.0	53
22	One-pot oxidative hydrolysis-oxidative cleavage of 7-borylindoles enables access to <i>o</i> -amidophenols and 4-acylbenzoxazoles. <i>Chemical Communications</i> , 2020, 56, 3559-3562.	4.1	3
23	Structural Revision of Pseudocerosine and Validation of a Biosynthetic Proposal for E-ring Formation in Pyridoacridine Alkaloids. <i>Organic Letters</i> , 2020, 22, 3495-3498.	4.6	12
24	Synthesis of the 1,2,4-Thiadiazole Alkaloid Polyaurine B. <i>Journal of Natural Products</i> , 2020, 83, 1721-1724.	3.0	12
25	Impact of the alkaloid colletotrichumine A on the pathogenicity of <i>Colletotrichum capsici</i> in <i>Capsicum annum</i> L. <i>Rhizosphere</i> , 2020, 16, 100247.	3.0	7
26	Synthesis of the Tetracyclic Cores of the Integrastatins, Epicoccolide A and Epicocconigrone A. <i>Journal of Organic Chemistry</i> , 2019, 84, 11935-11944.	3.2	9
27	Cleavage of lignin model compounds and lignin using aqueous oxalic acid. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 7408-7415.	2.8	11
28	Cu ¹ -Cu ⁰ bicomponent CuNPs@ZIF-8 for highly selective hydrogenation of biomass derived 5-hydroxymethylfurfural. <i>Green Chemistry</i> , 2019, 21, 4319-4323.	9.0	52
29	Acetyl-CoA-mediated activation of <i>Mycobacterium tuberculosis</i> isocitrate lyase 2. <i>Nature Communications</i> , 2019, 10, 4639.	12.8	23
30	Two-Step Preparation of Diverse 3-Amidofurans from Chitin. <i>ChemistrySelect</i> , 2019, 4, 10097-10099.	1.5	25
31	Synthetic Studies Toward the Flatworm-Derived Alkaloid Pseudocerosine. <i>ChemistrySelect</i> , 2019, 4, 11367-11369.	1.5	2
32	Synthesis of MCM-41-Supported Metal Catalysts in Deep Eutectic Solvent for the Conversion of Carbohydrates into 5-Hydroxymethylfurfural. <i>ChemSusChem</i> , 2019, 12, 978-982.	6.8	42
33	Transferring the biorenewable nitrogen present in chitin to several N-functional groups. <i>Sustainable Chemistry and Pharmacy</i> , 2019, 13, 100143.	3.3	18
34	Clean Synthesis of 5-Hydroxymethylfurfural and Levulinic Acid by Aqueous Phase Conversion of Levoglucosenone over Solid Acid Catalysts. <i>ACS Sustainable Chemistry and Engineering</i> , 2019, 7, 5892-5899.	6.7	34
35	Biomimetic synthesis of nudicaulins I and II, yellow pigments from the Iceland poppy <i>Papaver nudicaule</i> . <i>Chemical Communications</i> , 2019, 55, 13594-13597.	4.1	8
36	Oxidative Ring-Expansion of a Chitin-Derived Platform Enables Access to Unexplored Amino Sugar Chemical Space. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 1355-1360.	2.4	33

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37	Bio-Based Chiral Amines via Aza-Michael Additions to Levoglucosenone Under Aqueous Conditions. <i>European Journal of Organic Chemistry</i> , 2018, 2018, 2028-2038.	2.4	9
38	C4-H alkoxylation of 6-bromoindole and its application to the synthesis of breifussin B. <i>Tetrahedron</i> , 2018, 74, 1199-1202.	1.9	9
39	A novel dihydrodifuropyridine scaffold derived from ketones and the chitin-derived heterocycle 3-acetamido-5-acetylfuran. <i>Monatshefte für Chemie</i> , 2018, 149, 857-861.	1.8	24
40	Towards the Shell Biorefinery: Sustainable Synthesis of the Anticancer Alkaloid Proximicin A from Chitin. <i>ChemSusChem</i> , 2018, 11, 532-535.	6.8	79
41	Synthesis of three <i>Tricholoma</i> -derived indoles via an ortho-quinone methide. <i>Arkivoc</i> , 2018, 2018, 6-12.	0.5	3
42	Biomimetic synthesis of the bisindole framework present in sciadole, an alkaloid from <i>Tricholoma sciodes</i> . <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 6882-6885.	2.8	1
43	Total Synthesis of an <i>Isatis indigotica</i> -Derived Alkaloid Using a Biomimetic Thio-Diels-Alder Reaction. <i>Organic Letters</i> , 2018, 20, 3545-3548.	4.6	25
44	Photosensitized Cross-Linking of Tryptophan and Tyrosine Derivatives by Rose Bengal in Aqueous Solutions. <i>Journal of Organic Chemistry</i> , 2018, 83, 10835-10844.	3.2	12
45	Non-monoterpenoid azepinoindole alkaloids. <i>Natural Product Reports</i> , 2018, 35, 1347-1382.	10.3	40
46	Flavoalkaloids Isolation, Biological Activity, and Total Synthesis. <i>The Alkaloids Chemistry and Biology</i> , 2017, 77, 85-115.	2.0	15
47	Synthetic Access to 3,5,7-Trisubstituted Indoles Enabled by Iridium-Catalyzed C-H Borylation. <i>Synthesis</i> , 2017, 49, 4731-4737.	2.3	9
48	Targeting isocitrate lyase for the treatment of latent tuberculosis. <i>Drug Discovery Today</i> , 2017, 22, 1008-1016.	6.4	40
49	Bioinspired Total Synthesis and Stereochemical Revision of the Fungal Metabolite Pestalospirane B. <i>Organic Letters</i> , 2017, 19, 3414-3417.	4.6	8
50	Synthesis of putative clausenal from carbazole using sequential C-H borylations. <i>Tetrahedron Letters</i> , 2017, 58, 1699-1701.	1.4	6
51	Production of Levoglucosenone and Dihydrolevoglucosenone by Catalytic Reforming of Volatiles from Cellulose Pyrolysis Using Supported Ionic Liquid Phase. <i>ACS Sustainable Chemistry and Engineering</i> , 2017, 5, 1132-1140.	6.7	78
52	Development of NMR and thermal shift assays for the evaluation of <i>Mycobacterium tuberculosis</i> isocitrate lyase inhibitors. <i>MedChemComm</i> , 2017, 8, 2155-2163.	3.4	11
53	Mushroom-Derived Indole Alkaloids. <i>Journal of Natural Products</i> , 2017, 80, 2178-2187.	3.0	116
54	Natural Products with Heteroatom-Rich Ring Systems. <i>Journal of Natural Products</i> , 2017, 80, 3060-3079.	3.0	69

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55	Observations arising from a Beckmann rearrangement-Mannich cyclization approach to the azepinobisindole alkaloid lheyamine A. <i>Tetrahedron</i> , 2017, 73, 4355-4362.	1.9	17
56	Pyranonaphthoquinones – isolation, biology and synthesis: an update. <i>Natural Product Reports</i> , 2017, 34, 25-61.	10.3	45
57	Synthesis of the Azepinobisindole Alkaloid lheyamine A Enabled by a Cross-Mannich Reaction. <i>Organic Letters</i> , 2016, 18, 5404-5407.	4.6	33
58	4,4- ϵ^2 -Bismoschamine: biomimetic synthesis and evidence to support structural equivalency to montamine. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 8838-8847.	2.8	1
59	Synthesis and electrochemical detection of a thiazolyl-indole natural product isolated from the nosocomial pathogen <i>Pseudomonas aeruginosa</i> . <i>Analytical and Bioanalytical Chemistry</i> , 2016, 408, 6361-6367.	3.7	13
60	Flexible synthesis of diverse N-heterocycles from substrates attainable from biomass. <i>Green Chemistry</i> , 2016, 18, 2453-2459.	9.0	21
61	Total syntheses of (Δ^{\pm})-spiroindimicins B and C enabled by a late-stage Sch \ddot{A} llkopf \ddot{A} Magnus \ddot{A} Barton \ddot{A} Zard (SMBZ) reaction. <i>Chemical Communications</i> , 2016, 52, 800-802.	4.1	36
62	Synthetic studies towards putative yuremamine using an iterative C(sp ³) \ddot{A} H arylation strategy. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 5728-5743.	2.8	12
63	Synthesis of Inducamides A and B. <i>Journal of Natural Products</i> , 2016, 79, 519-522.	3.0	4
64	Synthesis of the 1,2,4-thiadiazole alkaloids polycarpathiamines A and B. <i>Organic Chemistry Frontiers</i> , 2016, 3, 38-42.	4.5	25
65	Bioinspired total synthesis and structural revision of yuremamine, an alkaloid from the entheogenic plant <i>Mimosa tenuiflora</i> . <i>Chemical Communications</i> , 2015, 51, 6202-6205.	4.1	34
66	A Procedure for Transforming Indoles into Indolequinones. <i>Journal of Organic Chemistry</i> , 2015, 80, 1006-1017.	3.2	20
67	Synthesis of 2-(3- ϵ^2 -Indolyl)tetrahydrofurans by Oxidative Cycloetherification. <i>Journal of Organic Chemistry</i> , 2015, 80, 2900-2906.	3.2	9
68	Biomimetic Synthesis of Dendridine A. <i>Organic Letters</i> , 2015, 17, 1344-1346.	4.6	27
69	Formal synthesis of nanaomycin D via a Hauser \ddot{A} Kraus annulation using a chiral enone-lactone. <i>Tetrahedron</i> , 2015, 71, 7137-7143.	1.9	23
70	Alkaloids from the traditional chinese medicine ChanSu: synthesis-enabled structural reassignment of bufopyramide to bufoserotonin C. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 7911-7914.	2.8	10
71	Synthesis of colletotrichumine A. <i>Heterocyclic Communications</i> , 2015, 21, 335-336.	1.2	1
72	Iridium-Catalyzed Triborylation of 3-Substituted Indoles. <i>Australian Journal of Chemistry</i> , 2015, 68, 1810.	0.9	9

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73	Synthesis of scalaridine A. <i>Tetrahedron Letters</i> , 2015, 56, 5914-5915.	1.4	14
74	Synthesis of Alocasin A. <i>Journal of Natural Products</i> , 2015, 78, 3080-3082.	3.0	11
75	Iridium-Catalysed C-H Borylation Facilitates a Total Synthesis of the HRV 3C Protease Inhibitor (±)-Thysanone. <i>Synlett</i> , 2014, 25, 556-558.	1.8	7
76	A Complementary Synthetic Approach to Fluorazone. <i>Journal of Heterocyclic Chemistry</i> , 2014, 51, 282-284.	2.6	12
77	Heterocycle construction using the biomass-derived building block itaconic acid. <i>Green Chemistry</i> , 2014, 16, 2084-2101.	9.0	38
78	Towards a biomimetic synthesis of schischkiniin: assembling the bis-dihydropyrazinone cycloaddition precursor. <i>Tetrahedron</i> , 2014, 70, 3430-3439.	1.9	12
79	Toward an Asymmetric Synthesis of the Dimeric Pyranonaphthoquinone Antibiotic Crisamicin A. <i>Journal of Organic Chemistry</i> , 2014, 79, 7169-7178.	3.2	28
80	Synthesis of the 2-methylene analogue of the HRV 3C protease inhibitor thysanone (2-carbathysanone). <i>Organic and Biomolecular Chemistry</i> , 2014, 12, 905-912.	2.8	17
81	Total Synthesis of (±)-Aspergilazine A. <i>Organic Letters</i> , 2014, 16, 5056-5059.	4.6	24
82	Total synthesis of putative montamine and a proposed structural reassignment. <i>Organic and Biomolecular Chemistry</i> , 2014, 12, 6878-6884.	2.8	8
83	A short synthesis of the endogenous plant metabolite 7-hydroxyoxindole-3-acetic acid (7-OH-OxIAA) using simultaneous C-H borylations. <i>Tetrahedron Letters</i> , 2014, 55, 5798-5800.	1.4	20
84	Discovery of a 1,2-bis(3-indolyl)ethane that selectively inhibits the pyruvate kinase of methicillin-resistant <i>Staphylococcus aureus</i> over human isoforms. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 5059-5062.	2.2	15
85	Synthesis and evaluation of 9-deoxy analogues of (±)-thysanone, an inhibitor of HRV 3C protease. <i>European Journal of Medicinal Chemistry</i> , 2014, 87, 220-227.	5.5	12
86	Synthesis and Biological Evaluation of 7-Deoxy Analogues of the Human Rhinovirus 3C Protease Inhibitor Thysanone. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 122-128.	2.4	9
87	Schischkiniin support studies: synthetic access to 1,1-bisindoles. <i>Chemical Communications</i> , 2013, 49, 4349-4351.	4.1	18
88	Synthesis and cytotoxicity of pyranonaphthoquinone natural product analogues under bioreductive conditions. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 7971-7980.	3.0	18
89	Studies towards the synthesis of montamine: synthesis of the 1,2-bis(indolyl)ethylhydrazine fragment. <i>Tetrahedron Letters</i> , 2013, 54, 1980-1982.	1.4	6
90	Telomerase Inhibition Studies of Novel Spiroketal-Containing Rubromycin Derivatives. <i>Australian Journal of Chemistry</i> , 2013, 66, 530.	0.9	11

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91	Natural Product-Inspired Pyranonaphthoquinone Inhibitors of Indoleamine 2,3-Dioxygenase-1 (IDO-1). <i>Australian Journal of Chemistry</i> , 2013, 66, 40.	0.9	20
92	Natural Products Containing a Nitrogen–Nitrogen Bond. <i>Journal of Natural Products</i> , 2013, 76, 794-812.	3.0	299
93	Total syntheses of the dipyrrolobenzoquinone (+)-terreusinone enabled by an evaluation of 4-methylpent-1-yn-3-ols in the Larock indole synthesis. <i>Tetrahedron</i> , 2013, 69, 4563-4577.	1.9	25
94	Palladium-Catalyzed Heteroannulation Approach to 1,2-Bis(3-indolyl)ethanes. <i>Synlett</i> , 2013, 24, 1931-1936.	1.8	5
95	Extending the Utility of the Bartoli Indolization: Synthesis of Marinoquinolines C and E. <i>Synlett</i> , 2013, 24, 461-464.	1.8	17
96	A Bidirectional Synthesis of (+)-Terreusinone. <i>Synlett</i> , 2012, 23, 1824-1828.	1.8	14
97	Biomimetic Synthesis of Phenazine-1,6-dicarboxylic Acid (PDC). <i>Synlett</i> , 2012, 23, 2827-2829.	1.8	5
98	Total Synthesis of Danshenspiroketalactone. <i>Synlett</i> , 2012, 2012, 128-130.	1.8	5
99	Pyrazine alkaloids via dimerization of amino acid-derived α -amino aldehydes: biomimetic synthesis of 2,5-diisopropylpyrazine, 2,5-bis(3-indolylmethyl)pyrazine and actinopolymorphol C. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 2126.	2.8	19
100	A furoindoline synthesis by remote radical functionalization. <i>Tetrahedron Letters</i> , 2012, 53, 5426-5429.	1.4	7
101	A simple solid phase, peptide-based fluorescent assay for the efficient and universal screening of HRV 3C protease inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 5018-5024.	2.2	5
102	Synthesis of the Tetracyclic Core of Berkelic Acid Using Gold(I)-Catalyzed Hydroarylation and Oxidative Radical Cyclizations. <i>Organic Letters</i> , 2012, 14, 5820-5823.	4.6	20
103	Total Synthesis of the Initially Reported and Revised Structures of the Neuroprotective Agent Palmyrolide A. <i>Organic Letters</i> , 2012, 14, 5374-5377.	4.6	34
104	Iridium-Catalyzed C–H Borylation-Based Synthesis of Natural Indolequinones. <i>Journal of Organic Chemistry</i> , 2012, 77, 2584-2587.	3.2	25
105	Synthetic studies towards dendridine A: synthesis of hemi-dendridine A acetate by Fischer indolization. <i>Tetrahedron Letters</i> , 2012, 53, 3623-3626.	1.4	9
106	Enantioselective synthesis of pyranonaphthoquinone antibiotics using a CBS reduction/cross-metathesis/oxa-Michael strategy. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 5423.	2.8	24
107	Natural products targeting telomere maintenance. <i>MedChemComm</i> , 2011, 2, 229.	3.4	37
108	Total Synthesis of the Photoprotecting Dipyrrolobenzoquinone (+)-Terreusinone. <i>Organic Letters</i> , 2011, 13, 6444-6447.	4.6	36

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109	Concise syntheses of 5,6-dibromotryptamine and 5,6-dibromo-N,N-dimethyltryptamine en route to the antibiotic alternatamide D. <i>Tetrahedron Letters</i> , 2011, 52, 4042-4044.	1.4	13
110	A concise synthesis of meridianin F. <i>Tetrahedron Letters</i> , 2011, 52, 4537-4538.	1.4	17
111	The Oxidation of Amides to Imides: A Powerful Synthetic Transformation. <i>Synthesis</i> , 2011, 2011, 3569-3580.	2.3	48
112	Efficient Synthesis of the Spiroacetal Core of Paecilospirone via Oxidative Radical Cyclisation. <i>Synlett</i> , 2011, 2011, 1395-1398.	1.8	6
113	Synthesis of Benzannulated Spiroketal Using an Oxidative Radical Cyclization. <i>Synthesis</i> , 2011, 2011, 1383-1398.	2.3	6
114	Biomimetic Synthesis of 2,5-Bis(indol-3-ylmethyl)pyrazine via Intermolecular Amino Aldehyde Cyclization. <i>Synlett</i> , 2011, 2011, 2339-2342.	1.8	2
115	Synthesis of the Selective Neuronal Nitric Oxide Synthase (nNOS) Inhibitor 5,6-Dibromo-2- ϵ^2 -demethylaplysinopsin. <i>Synlett</i> , 2011, 2011, 826-830.	1.8	3
116	Synthesis of Benzotriazole Analogues of the Helicobactericidal Agents CJ-13,015, CJ-13,102, CJ-13,108, and CJ-13,104 Using a Regioselective 1,3-Dipolar Cycloaddition. <i>Synlett</i> , 2011, 2011, 99-103.	1.8	2
117	Total Synthesis and Absolute Configuration of (α^+)-Berkeleyamide A. <i>Organic Letters</i> , 2010, 12, 420-423.	4.6	16
118	Diazonamide studies. A direct synthesis of the indole bis-oxazole fragment from tri- and tetra-peptides using biomimetic oxidative cyclizations. <i>Tetrahedron</i> , 2010, 66, 6483-6495.	1.9	18
119	Synthesis of triazole analogues of the nanaomycin antibiotics using ϵ^{click} chemistry ϵ^{TM} . <i>Tetrahedron</i> , 2010, 66, 4002-4009.	1.9	22
120	Synthesis of the Pyranonaphthoquinones Dehydroherbarin, (+)-Astropaquinone B and (+)-Astropaquinone C en Route to Ascomycones A and B. <i>Synthesis</i> , 2010, 2010, 2604-2608.	2.3	3
121	Improved Synthesis of the Benzyne Precursor 2-(Trimethylsilyl)phenyl Trifluoromethanesulfonate. <i>Synthesis</i> , 2010, 2010, 911-913.	2.3	12
122	Heteroatom-Directed Reverse Wacker Oxidations. Synthesis of the Reported Structure of (α^+)-Herbaric Acid. <i>Journal of Organic Chemistry</i> , 2010, 75, 7388-7392.	3.2	42
123	Isolation, biological activity and synthesis of benzannulated spiroketal natural products. <i>Natural Product Reports</i> , 2010, 27, 1117.	10.3	138
124	Synthesis of natural products containing spiroketals via intramolecular hydrogen abstraction. <i>Organic and Biomolecular Chemistry</i> , 2010, 8, 29-38.	2.8	61
125	Enantioselective Synthesis of the 3C-Protease Inhibitor (-)-Thysanone by a Staunton-Weinreb Annulation Strategy. <i>Synthesis</i> , 2009, 2009, 2561-2569.	2.3	4
126	A Facile Cross-Metathesis-Radical-Cyclisation Approach to Monobenzannulated Spiroketal. <i>Synlett</i> , 2009, 2009, 793-797.	1.8	6

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127	Pyranonaphthoquinone derivatives of eleutherin, ventiloquinone L, thysanone and nanaomycin A possessing a diverse topoisomerase II inhibition and cytotoxicity spectrum. <i>Bioorganic and Medicinal Chemistry</i> , 2009, 17, 7131-7137.	3.0	33
128	Biomimetic studies towards the cardinalins: synthesis of (+)-ventiloquinone L and an unusual dimerisation. <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 2599.	2.8	26
129	Chemoenzymatic synthesis of deoxy analogues of the DNA topoisomerase II inhibitor eleutherin and the 3C-protease inhibitor thysanone. <i>Tetrahedron</i> , 2008, 64, 4827-4834.	1.9	23
130	Synthesis of a C8 oxygenated pyranonaphthoquinone: a useful precursor to dimeric pyranonaphthoquinones. <i>Tetrahedron</i> , 2008, 64, 3343-3350.	1.9	13
131	An approach to an enantioselective synthesis of crisamicin A via a novel double Hauser-Kraus annulation strategy. <i>Tetrahedron</i> , 2008, 64, 3912-3927.	1.9	27
132	Enantioselective synthesis of the dimeric pyranonaphthoquinone core of the cardinalins using a late-stage homocoupling strategy. <i>Organic and Biomolecular Chemistry</i> , 2008, 6, 4261.	2.8	27
133	Pyranonaphthoquinones— isolation, biological activity and synthesis. <i>Natural Product Reports</i> , 2008, 25, 376-400.	10.3	87
134	An Efficient Enantioselective Synthesis of the 3C Protease Inhibitor (-)-Thysanone. <i>Synlett</i> , 2008, 2008, 1910-1912.	1.8	1
135	A Facile Enantioselective Synthesis of the Dimeric Pyranonaphthoquinone Core of the Cardinalins. <i>Synlett</i> , 2008, 2008, 867-870.	1.8	20
136	(1R,1aR,3S,3aS)-5,5,10,10-tetramethoxy-1,3,3,4-tetramethyl-3,4-tetrahydro-1H,8-bi[benzo]g Acta Crystallographica Section E: Structure Reports Online, 2008, 64, o758-o758.	0.2	1
137	Enantioselective Synthesis of an Analogue of Nanaomycin A. <i>Synthesis</i> , 2007, 2007, 2887-2893.	2.3	21
138	Synthesis of the calothrixins, pentacyclic indolo[3,2-j]phenanthridine alkaloids, using a biomimetic approach. <i>Tetrahedron</i> , 2007, 63, 10963-10970.	1.9	44
139	The influence of microwave irradiation on lipase-catalyzed kinetic resolution of racemic secondary alcohols. <i>Tetrahedron: Asymmetry</i> , 2007, 18, 1618-1624.	1.8	33
140	A biomimetic synthesis of calothrixin B. <i>Tetrahedron Letters</i> , 2007, 48, 231-234.	1.4	37
141	Biomimetic approaches to diazonamide A. Direct synthesis of the indole bis-oxazole fragment by oxidation of a TyrValTrpTrp tetrapeptide. <i>Chemical Communications</i> , 2006, , 2397.	4.1	17