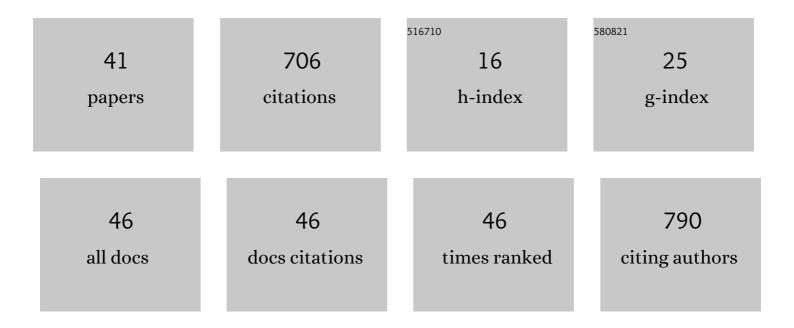
Joanne E Harvey

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/328793/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Mechanistic Studies on the Base-Promoted Ring Opening of Glycal-Derived <i>gem</i> -Dibromocyclopropanes. Journal of Organic Chemistry, 2022, 87, 301-315.	3.2	2
2	Unraveling the binding mode of a methamphetamine aptamer: A spectroscopic and calorimetric study. Biophysical Journal, 2022, 121, 2193-2205.	0.5	2
3	Gold(I)-catalyzed, one-pot, oxidative formation of 2,4-disubstituted thiazoles: Application to the synthesis of a pateamine-related macrodiolide. Tetrahedron, 2021, 88, 132109.	1.9	2
4	Total Synthesis and Bioactivity Studies of Fungal Metabolite (â^')-TAN-2483B. Organic Letters, 2020, 22, 9427-9432.	4.6	6
5	Kinase-Inhibitory Nucleoside Derivatives from the Pacific Bryozoan <i>Nelliella nelliiformis</i> . Journal of Natural Products, 2020, 83, 547-551.	3.0	7
6	Synthesis of Bioactive Sideâ€Chain Analogues of TANâ€2483B. Chemistry - an Asian Journal, 2019, 14, 1230-1237.	3.3	7
7	A colourful azulene-based protecting group for carboxylic acids. Tetrahedron, 2018, 74, 2942-2955.	1.9	6
8	Genome mining, isolation, chemical synthesis and biological evaluation of a novel lanthipeptide, tikitericin, from the extremophilic microorganism <i>Thermogemmatispora</i> strain T81. Chemical Science, 2018, 9, 7311-7317.	7.4	23
9	Synthesis of a simplified triazole analogue of pateamine A. Organic and Biomolecular Chemistry, 2016, 14, 5117-5127.	2.8	5
10	Preparation of conjugated dienoates with Bestmann ylide: Towards the synthesis of zampanolide and dactylolide using a facile linchpin approach. Beilstein Journal of Organic Chemistry, 2015, 11, 1815-1822.	2.2	2
11	Synthetic, semisynthetic and natural analogues of peloruside A. Chemical Communications, 2015, 51, 4750-4765.	4.1	18
12	Synthesis of mycothiol conjugate analogues and evaluation of their antimycobacterial activity. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 2152-2155.	2.2	3
13	Functional, water-dispersible gold nanoparticles produced with N,N′-bis(acryloyl)-(<scp>l</scp>)-cystine. RSC Advances, 2015, 5, 104079-104086.	3.6	2
14	Reactions of 1,2-cyclopropyl carbohydrates. Pure and Applied Chemistry, 2014, 86, 1377-1399.	1.9	18
15	Synthesis and Solution Aggregation Studies of a Suite of Mixed Neutral and Zwitterionic Chromophores for Second-Order Nonlinear Optics. Journal of Organic Chemistry, 2014, 79, 10153-10169.	3.2	10
16	Divergent synthesis of 2-C-branched pyranosides and oxepines from 1,2-gem-dibromocyclopropyl carbohydrates. Tetrahedron, 2014, 70, 7032-7043.	1.9	13
17	13C NMR Analysis of 3,6-Dihydro-2H-pyrans: Assignment of Remote Stereochemistry Using Axial Shielding Effects. Journal of Organic Chemistry, 2014, 79, 5521-5532.	3.2	11
18	Analytical Profile of Moxidectin. Profiles of Drug Substances, Excipients and Related Methodology, 2013, 38, 315-366.	8.0	11

JOANNE E HARVEY

#	Article	IF	CITATIONS
19	Synthesis of diastereomeric, deoxy and ring-expanded sulfone analogues of aigialomycin D. Tetrahedron, 2013, 69, 10581-10592.	1.9	8
20	Pd-Catalyzed Allylic Alkylation Cascade with Dihydropyrans: Regioselective Synthesis of Furo[3,2- <i>c</i>]pyrans. Organic Letters, 2013, 15, 2430-2433.	4.6	43
21	Evaluation of degradation kinetics for abamectin in formulations using a stability indicating method. Acta Pharmaceutica, 2013, 63, 59-69.	2.0	3
22	Separation and Identification of Degradation Products in Abamectin Formulation Using LC, LTQ FT-MS, H/D Exchange and NMR. Current Pharmaceutical Analysis, 2012, 8, 415-430.	0.6	4
23	An Overview on Chemical Derivatization and Stability Aspects of Selected Avermectin Derivatives. Chemical and Pharmaceutical Bulletin, 2012, 60, 931-944.	1.3	33
24	Isolation and characterization of degradation products of moxidectin using LC, LTQ FT-MS, H/D exchange and NMR. Analytical and Bioanalytical Chemistry, 2012, 404, 2203-2222.	3.7	15
25	Separation and identification of degradation products in eprinomectin formulation using LC, LTQ FT-MS, H/D exchange, and NMR. Journal of Pharmaceutical and Biomedical Analysis, 2012, 63, 62-73.	2.8	18
26	Synthesis of C-furanosides from a <scp>d</scp> -glucal-derived cyclopropane through a ring-expansion/ring-contraction sequence. Chemical Communications, 2011, 47, 421-423.	4.1	21
27	Alkenylphosphonates: unexpected products from reactions of methyl 2-[(diethoxyphosphoryl)methyl]benzoate under Horner–Wadsworth–Emmons conditions. Organic and Biomolecular Chemistry, 2011, 9, 4432.	2.8	9
28	Towards a simplified peloruside A: synthesis of C1–C11 of a dihydropyran analogue. Tetrahedron, 2011, 67, 9376-9381.	1.9	4
29	Synthesis of the (â^')-TAN-2483B ring system via a d-mannose-derived cyclopropane. Organic and Biomolecular Chemistry, 2011, 9, 998-1000.	2.8	10
30	Synthesis of Oxepines and 2-Branched Pyranosides from a <scp>d</scp> -Glucal-Derived <i>gem</i> -Dibromo-1,2-cyclopropanated Sugar. Journal of Organic Chemistry, 2010, 75, 955-958.	3.2	51
31	Mechanistic studies of rearrangements during the ring expansions of cyclopropanated carbohydrates. Tetrahedron Letters, 2009, 50, 7283-7285.	1.4	7
32	Heptanosides from Galactose-Derived Oxepenes via Stereoselective Addition Reactions. Journal of Organic Chemistry, 2009, 74, 7627-7632.	3.2	28
33	Total Synthesis of Aigialomycin D Using a Rambergâ^BĂœklund/RCM Strategy. Journal of Organic Chemistry, 2009, 74, 2271-2277.	3.2	66
34	Synthesis of the C12–C24 fragment of peloruside A by silyl-tethered diastereomer-discriminating RCM. Tetrahedron Letters, 2008, 49, 7021-7023.	1.4	26
35	Stereochemical Control in Carbohydrate Chemistry. Journal of Chemical Education, 2008, 85, 689.	2.3	6
36	Electrocyclic Ring-Opening Reactions of gem-Dibromocyclopropanes in the Synthesis of Natural Products and Related Compounds. Synlett, 2006, 2006, 1975-2000.	1.8	11

JOANNE E HARVEY

#	Article	IF	CITATIONS
37	Highly functionalised organolithium and organoboron reagents for the preparation of enantiomerically pure α-amino acids. Tetrahedron, 2005, 61, 3403-3417.	1.9	48
38	Synthesis of non-proteinogenic phenylalanine analogues by Suzuki cross-coupling of a serine-derived alkyl boronic acid. Tetrahedron Letters, 2004, 45, 2467-2471.	1.4	23
39	A Versatile and Stereocontrolled Route to Pyranose and FuranoseC-Glycosides. Organic Letters, 2004, 6, 2611-2614.	4.6	33
40	The first synthesis of the epoxide-containing macrolactone nucleus of oximidine I. Tetrahedron Letters, 2003, 44, 7209-7212.	1.4	18
41	Electrocyclic Ring-Opening/Ĩ€-Allyl Cation Cyclization Reaction Sequences Involvinggem-Dihalocyclopropanes as Substrates: Application to Syntheses of (±)-, (+)-, and (â^')-γ-Lycorane. Journal of Organic Chemistry, 2000, 65, 4241-4250.	3.2	71