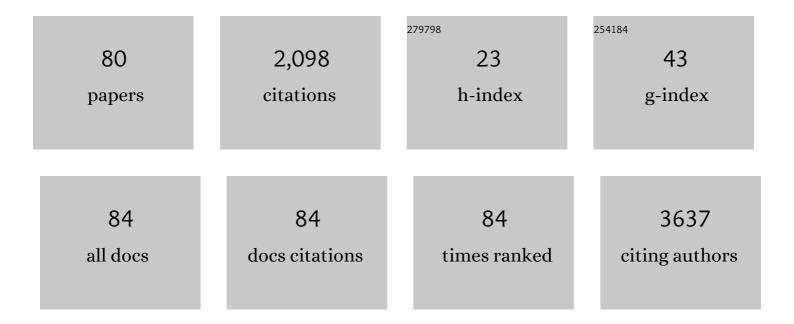
List of Publications by Year in descending order

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Version: 2024-02-01



#	Article	IF	CITATIONS
1	Methuosis Contributes to Jaspine-B-Induced Cell Death. International Journal of Molecular Sciences, 2022, 23, 7257.	4.1	4
2	The anti-cancer drug ABTL0812 induces ER stress-mediated cytotoxic autophagy by increasing dihydroceramide levels in cancer cells. Autophagy, 2021, 17, 1349-1366.	9.1	72
3	Synthesis and characterization of bichromophoric 1-deoxyceramides as FRET probes. Organic and Biomolecular Chemistry, 2021, 19, 2456-2467.	2.8	4
4	Discovery of deoxyceramide analogs as highly selective ACER3 inhibitors in live cells. European Journal of Medicinal Chemistry, 2021, 216, 113296.	5.5	9
5	Chirality-Puckering correlation and intermolecular interactions in Sphingosines: Rotational spectroscopy of jaspine B3 and its monohydrate. Spectrochimica Acta - Part A: Molecular and Biomolecular Spectroscopy, 2021, 267, 120531.	3.9	1
6	A Mechanism-Based Sphingosine-1-phosphate Lyase Inhibitor. Journal of Organic Chemistry, 2020, 85, 419-429.	3.2	5
7	Ceramide Analogue SACLAC Modulates Sphingolipid Levels and <i>MCL-1</i> Splicing to Induce Apoptosis in Acute Myeloid Leukemia. Molecular Cancer Research, 2020, 18, 352-363.	3.4	22
8	Click and count: specific detection of acid ceramidase activity in live cells. Chemical Science, 2020, 11, 13044-13051.	7.4	9
9	Rotational spectroscopy of organophosphorous chemical agents: cresyl and phenyl saligenin phosphates. Physical Chemistry Chemical Physics, 2019, 21, 16418-16422.	2.8	0
10	Activity-Based Imaging of Acid Ceramidase in Living Cells. Journal of the American Chemical Society, 2019, 141, 7736-7742.	13.7	17
11	SCOTfluors: Small, Conjugatable, Orthogonal, and Tunable Fluorophores for Inâ€Vivo Imaging of Cell Metabolism. Angewandte Chemie - International Edition, 2019, 58, 6911-6915.	13.8	100
12	New fluorogenic probes for neutral and alkaline ceramidases. Journal of Lipid Research, 2019, 60, 1174-1181.	4.2	5
13	Dihydroceramide Desaturase 1 Inhibitors Reduce Amyloid-β Levels in Primary Neurons from an Alzheimer's Disease Transgenic Model. Pharmaceutical Research, 2018, 35, 49.	3.5	14
14	Analysis of the neurotoxic effects of neuropathic organophosphorus compounds in adult zebrafish. Scientific Reports, 2018, 8, 4844.	3.3	11
15	Azide-tagged sphingolipids for the proteome-wide identification of C16-ceramide-binding proteins. Chemical Communications, 2018, 54, 13742-13745.	4.1	7
16	Clearly Detectable, Kinetically Restricted Solid–Solid Phase Transition in cis-Ceramide Monolayers. Langmuir, 2018, 34, 11749-11758.	3.5	6
17	Inhibitors of ceramide de novo biosynthesis rescue damages induced by cigarette smoke in airways epithelia. Naunyn-Schmiedeberg's Archives of Pharmacology, 2017, 390, 753-759.	3.0	17
18	The first fluorogenic sensor for sphingosine-1-phosphate lyase activity in intact cells. Chemical Communications, 2017, 53, 5441-5444.	4.1	12

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19	Rotational spectra of tetracyclic quinolizidine alkaloids: does a water molecule flip sparteine?. Physical Chemistry Chemical Physics, 2017, 19, 17553-17559.	2.8	4
20	Stereoselective preparation of quaternary 2-vinyl sphingosines and ceramides and their effect on basal sphingolipid metabolism. Chemistry and Physics of Lipids, 2017, 205, 34-41.	3.2	0
21	Jaspine B induces nonapoptotic cell death in gastric cancer cells independently of its inhibition of ceramide synthase. Journal of Lipid Research, 2017, 58, 1500-1513.	4.2	18
22	From the configurational preference of dihydroceramide desaturase-1 towards Δ ⁶ -unsaturated substrates to the discovery of a new inhibitor. Chemical Communications, 2017, 53, 4394-4397.	4.1	7
23	Abiotic amidine and guanidine hydrolysis of lamotrigine-N2-glucuronide and related compounds in wastewater: The role of pH and N2-substitution on reaction kinetics. Water Research, 2016, 100, 466-475.	11.3	14
24	3-Ketosphinganine provokes the accumulation of dihydroshingolipids and induces autophagy in cancer cells. Molecular BioSystems, 2016, 12, 1166-1173.	2.9	12
25	Studies on the inhibition of sphingosine-1-phosphate lyase by stabilized reaction intermediates and stereodefined azido phosphates. European Journal of Medicinal Chemistry, 2016, 123, 905-915.	5.5	2
26	Dihydroceramide accumulation mediates cytotoxic autophagy of cancer cells via autolysosome destabilization. Autophagy, 2016, 12, 2213-2229.	9.1	118
27	Bacterial versus human sphingosine-1-phosphate lyase (S1PL) in the design of potential S1PL inhibitors. Bioorganic and Medicinal Chemistry, 2016, 24, 4381-4389.	3.0	3
28	Investigating the formation and toxicity of nitrogen transformation products of diclofenac and sulfamethoxazole in wastewater treatment plants. Journal of Hazardous Materials, 2016, 309, 157-164.	12.4	72
29	Approaches to polyunsaturated sphingolipids: new conformationally restrained analogs with minimal structural modifications. Tetrahedron, 2016, 72, 605-612.	1.9	2
30	Inhibitors of sphingosine-1-phosphate metabolism (sphingosine kinases and sphingosine-1-phosphate) Tj ETQqO	0	Overlock 101
31	Fluorescent Polyene Ceramide Analogues as Membrane Probes. Langmuir, 2015, 31, 2484-2492.	3.5	8
32	Azideâ€Tagged Sphingolipids: New Tools for Metabolic Flux Analysis. ChemBioChem, 2015, 16, 641-650.	2.6	24
33	A straightforward synthesis of the CERT inhibitor (1′R,3′S)-HPA-12. Tetrahedron Letters, 2015, 56, 1706-1708.	1.4	8
34	Activity of neutral and alkaline ceramidases on fluorogenic N-acylated coumarin-containing aminodiols. Journal of Lipid Research, 2015, 56, 2019-2028.	4.2	13
35	Chemical Probes of Sphingolipid Metabolizing Enzymes. , 2015, , 437-469.		1
36	Structure elucidation of phototransformation products of unapproved analogs of the erectile dysfunction drug sildenafil in artificial freshwater with UPLCâ€Q Exactiveâ€MS. Journal of Mass Spectrometry, 2014, 49, 1279-1289.	1.6	10

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37	Simultaneous determination of diclofenac, its human metabolites and microbial nitration/nitrosation transformation products in wastewaters by liquid chromatography/quadrupole-linear ion trap mass spectrometry. Journal of Chromatography A, 2014, 1347, 63-71.	3.7	59
38	In Situ Synthesis of Fluorescent Membrane Lipids (Ceramides) using Click Chemistry. Biophysical Journal, 2013, 104, 373a.	0.5	0
39	Natural Products as Platforms for the Design of Sphingolipid-Related Anticancer Agents. Advances in Cancer Research, 2013, 117, 237-281.	5.0	20
40	Straightforward Access to Spisulosine and 4,5-Dehydrospisulosine Stereoisomers: Probes for Profiling Ceramide Synthase Activities in Intact Cells. Journal of Organic Chemistry, 2013, 78, 5858-5866.	3.2	43
41	Acid ceramidase as a therapeutic target in metastatic prostate cancer. Journal of Lipid Research, 2013, 54, 1207-1220.	4.2	61
42	Cellular Changes that Accompany Shedding of Human Corneocytes. Journal of Investigative Dermatology, 2012, 132, 2430-2439.	0.7	48
43	Dihydroceramide delays cell cycle G1/S transition via activation of ER stress and induction of autophagy. International Journal of Biochemistry and Cell Biology, 2012, 44, 2135-2143.	2.8	66
44	ldentification of phototransformation products of sildenafil (Viagra) and its Nâ€demethylated human metabolite under simulated sunlight. Journal of Mass Spectrometry, 2012, 47, 701-711.	1.6	19
45	In situ synthesis of fluorescent membrane lipids (ceramides) using click chemistry. Journal of Chemical Biology, 2012, 5, 119-123.	2.2	8
46	C6-Ceramide and targeted inhibition of acid ceramidase induce synergistic decreases in breast cancer cell growth. Breast Cancer Research and Treatment, 2012, 133, 447-458.	2.5	69
47	3-Deoxy-3,4-dehydro analogs of XM462. Preparation and activity on sphingolipid metabolism and cell fate. Bioorganic and Medicinal Chemistry, 2012, 20, 3173-3179.	3.0	11
48	Sphingolipid Modulation: A Strategy for Cancer Therapy. Anti-Cancer Agents in Medicinal Chemistry, 2012, 12, 285-302.	1.7	22
49	Ceramidases in Hematological Malignancies: Senseless or Neglected Target?. Anti-Cancer Agents in Medicinal Chemistry, 2011, 11, 830-843.	1.7	12
50	Dihydrosphingomyelin Impairs HIV-1 Infection by Rigidifying Liquid-Ordered Membrane Domains. Chemistry and Biology, 2010, 17, 766-775.	6.0	76
51	A simple fluorogenic method for determination of acid ceramidase activity and diagnosis of Farber disease. Journal of Lipid Research, 2010, 51, 3542-3547.	4.2	53
52	An Unexpected Access to a New Sphingoid Base Containing a Vinyl Sulfide Unit. Synlett, 2010, 2010, 2950-2952.	1.8	1
53	Control of metabolism and signaling of simple bioactive sphingolipids: Implications in disease. Progress in Lipid Research, 2010, 49, 316-334.	11.6	124
54	Synthesis of a Fluorogenic Analogue of Sphingosineâ€1â€Phosphate and Its Use to Determine Sphingosineâ€1â€Phosphate Lyase Activity. ChemBioChem, 2009, 10, 820-822.	2.6	30

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55	A multifunctional desaturase involved in the biosynthesis of the processionary moth sex pheromone. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 16444-16449.	7.1	46
56	Substrate-Dependent Stereochemical Course of the (Z)-13-Desaturation Catalyzed by the Processionary Moth Multifunctional Desaturase. Journal of the American Chemical Society, 2007, 129, 15007-15012.	13.7	20
57	Synthesis and Use of Deuterated Palmitic Acids to Decipher the Cryptoregiochemistry of a Δ13Desaturation. Journal of Organic Chemistry, 2007, 72, 760-764.	3.2	4
58	Chemical Tools to Investigate Sphingolipid Metabolism and Functions. ChemMedChem, 2007, 2, 580-606.	3.2	50
59	Synthesis and Use of Probes to Investigate the Cryptoregiochemistry of the First Animal Acetylenase. Journal of Organic Chemistry, 2006, 71, 7558-7564.	3.2	6
60	Detection of DNA Adducts Derived from the Reactive Metabolite of Furan, cis-2-Butene-1,4-dial. Chemical Research in Toxicology, 2006, 19, 414-420.	3.3	87
61	Inhibitors of sphingolipid metabolism enzymes. Biochimica Et Biophysica Acta - Biomembranes, 2006, 1758, 1957-1977.	2.6	156
62	Synthesis of deuterated fatty acids to investigate the biosynthetic pathway of disparlure, the sex pheromone of the Gypsy Moth, Lymantria dispar. Lipids, 2004, 39, 397-401.	1.7	9
63	Arylacetic acid derivatization of 2,3- and internal erythro-squalene diols. Separation and absolute configuration determination. Tetrahedron, 2004, 60, 11519-11525.	1.9	2
64	Active Site Contacts in the Purine Nucleoside Phosphorylaseâ^'Hypoxanthine Complex by NMR andab InitioCalculationsâ€. Biochemistry, 2004, 43, 15966-15974.	2.5	19
65	Synthesis and Use of Stereospecifically Deuterated Analogues of Palmitic Acid To Investigate the Stereochemical Course of the Δ11 Desaturase of the Processionary Moth. Journal of Organic Chemistry, 2004, 69, 7108-7113.	3.2	18
66	Synthesis of fluorinated analogs of myristic acid as potential inhibitors of egyptian armyworm (Spodoptera littorialis) Δ11 desaturasedesaturase. Lipids, 2003, 38, 865-871.	1.7	7
67	Novel Chemoenzymatic Strategy for the Synthesis of Enantiomerically Pure Secondary Alcohols with Sterically Similar Substituents. Journal of Organic Chemistry, 2003, 68, 5351-5356.	3.2	10
68	Sex pheromone biosynthetic pathway for disparlure in the gypsy moth, Lymantria dispar. Proceedings of the United States of America, 2003, 100, 809-814.	7.1	53
69	Stereospecificity of the (Z)-9 desaturase that converts (E)-11-tetradecenoic acid into (Z,E)-9,11-tetradecadienoic acid in the biosynthesis of Spodoptera littoralis sex pheromone. Insect Biochemistry and Molecular Biology, 2001, 31, 799-803.	2.7	15
70	ls Hydrogen Tunneling Involved in AcylCoA Desaturase Reactions? The Case of a Δ9 Desaturase That Transforms (E)-11-Tetradecenoic Acid into (Z,E)-9,11-Tetradecadienoic Acid. Angewandte Chemie - International Edition, 2000, 39, 3279-3281.	13.8	51
71	Synthesis of Dideuterated and Enantiomers of Monodeuterated Tridecanoic Acids at C-9 and C-10 Positions. Journal of Organic Chemistry, 2000, 65, 8582-8588.	3.2	21
72	15N-Multilabeled Adenine and Guanine Nucleosides. Syntheses of [1,3,NH2-15N3]- and [2-13C-1,3,NH2-15N3]-Labeled Adenosine, Guanosine, 2â€~-Deoxyadenosine, and 2â€~-Deoxyguanosine. Journal c Organic Chemistry, 1999, 64, 6575-6582.	of 3.2	27

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73	Use of13C tags with specifically15N-labeled DNA and RNA. Biopolymers, 1998, 48, 57-63.	2.4	5
74	High yield protection of purine ribonucleosides for H-phosphonate RNA synthesis. Tetrahedron Letters, 1997, 38, 7135-7138.	1.4	13
75	2,3,18,19-Dioxidosqualene Stereoisomers:Â Characterization and Activity as Inhibitors of Purified Pig Liver 2,3-Oxidosqualeneâ^'Lanosterol Cyclase. Journal of Organic Chemistry, 1996, 61, 7603-7607.	3.2	8
76	Internal Oxidosqualenes: Determination of Absolute Configuration and Activity as Inhibitors of Purified Pig Liver Squalene Epoxidase. Journal of Organic Chemistry, 1995, 60, 3648-3656.	3.2	12
77	Unequivocal Identification of Compounds Formed in the Photodegradation of Fenitrothion in Water/Methanol and Proposal of Selected Transformation Pathways. Journal of Agricultural and Food Chemistry, 1994, 42, 814-821.	5.2	38
78	Epoxidation of 6,7- and 10,11-oxidosqualenes by the squalene epoxidase present in rat liver microsomes. Bioorganic and Medicinal Chemistry Letters, 1993, 3, 2581-2586.	2.2	4
79	Dioxidosqualenes: characterization and activity as inhibitors of 2,3-oxidosqualene-lanosterol cyclase. Journal of Organic Chemistry, 1993, 58, 3991-3997.	3.2	15
80	2,3:18,19-dioxidosqualene: synthesis and activity as a potent inhibitor of 2,3-oxidosqualene-lanosterol cyclase in rat liver microsomes. Bioorganic and Medicinal Chemistry Letters, 1992, 2, 1239-1242.	2.2	15