

# Masaki Ohtawa

## List of Publications by Year in descending order

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33  
papers

512  
citations

567144

15  
h-index

677027

22  
g-index

38  
all docs

38  
docs citations

38  
times ranked

535  
citing authors

#	ARTICLE	IF	CITATIONS
1	Synthesis of (âˆ™)-11-O-Debenzoyltashironin: Neurotrophic Sesquiterpenes Cause Hyperexcitation. <i>Journal of the American Chemical Society</i> , 2017, 139, 9637-9644.	6.6	54
2	Synthetic, Mechanistic, and Biological Interrogation of <i>Ginkgo biloba</i> Chemical Space En Route to (âˆ™)-Bilobalide. <i>Journal of the American Chemical Society</i> , 2020, 142, 18599-18618.	6.6	40
3	Concise asymmetric synthesis of (âˆ™)-bilobalide. <i>Nature</i> , 2019, 575, 643-646.	13.7	39
4	Enantioselective total synthesis of atpenin A5. <i>Journal of Antibiotics</i> , 2009, 62, 289-294.	1.0	33
5	Total synthesis of pyripyropene A. <i>Tetrahedron</i> , 2011, 67, 8195-8203.	1.0	30
6	Synthesis and structure-activity relationship of pyripyropene A derivatives as potent and selective acyl-CoA:cholesterol acyltransferase 2 (ACAT2) inhibitors: Part 1. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 1285-1287.	1.0	26
7	Synthesis and biological activity of 5-(4-methoxyphenyl)-oxazole derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 313-316.	1.0	23
8	The total synthesis and biological evaluation of nafuredin-Î³ and its analogues. <i>Tetrahedron</i> , 2008, 64, 8117-8127.	1.0	21
9	Synthesis and structure-activity relationship of pyripyropene A derivatives as potent and selective acyl-CoA:cholesterol acyltransferase 2 (ACAT2) inhibitors: Part 3. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 3798-3801.	1.0	21
10	Selectivity of Pyripyropene Derivatives in Inhibition toward Acyl-CoA:cholesterol Acyltransferase 2 Isozyme. <i>Journal of Antibiotics</i> , 2008, 61, 503-508.	1.0	20
11	Potential of Bleomycin in Jurkat Cells by Fungal Pycnidione. <i>Biological and Pharmaceutical Bulletin</i> , 2012, 35, 18-28.	0.6	20
12	Synthesis and structure-activity relationship of pyripyropene A derivatives as potent and selective acyl-CoA:cholesterol acyltransferase 2 (ACAT2) inhibitors: Part 2. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 2659-2662.	1.0	20
13	Simplifungin and Valsafungins, Antifungal Antibiotics of Fungal Origin. <i>Journal of Organic Chemistry</i> , 2016, 81, 7373-7383.	1.7	18
14	New Pyripyropene A Derivatives, Highly SOAT2-Selective Inhibitors, Improve Hypercholesterolemia and Atherosclerosis in Atherogenic Mouse Models. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2015, 355, 297-307.	1.3	17
15	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.	2.0	17
16	Scopranones with Two Atypical Scooplike Moieties Produced by <i>Streptomyces</i> sp. BYK-11038. <i>Organic Letters</i> , 2017, 19, 5980-5983.	2.4	16
17	9-(2-C-Cyano-2-deoxy-Î²-d-arabino-pentofuranosyl)guanine, a Potential Antitumor Agent against B-Lymphoma Infected with Kaposi's Sarcoma-Associated Herpesvirus. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 2007-2010.	2.9	12
18	Stereoselective Total Synthesis of Atpenins A4 and B, Harzianopyridone, and NBRI23477 B. <i>Chemical and Pharmaceutical Bulletin</i> , 2012, 60, 898-906.	0.6	10

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19	In vitro metabolism of pyripyropene A and ACAT inhibitory activity of its metabolites. <i>Journal of Antibiotics</i> , 2015, 68, 27-34.	1.0	9
20	One-Pot $\hat{I}^3$ -Lactonization of Homopropargyl Alcohols via Intramolecular Ketene Trapping. <i>Organic Letters</i> , 2021, 23, 2831-2835.	2.4	9
21	Design, synthesis, and biological evaluation of air-stable nafuredin- $\hat{I}^3$ analogs as complex I inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 932-943.	1.4	8
22	Design and Synthesis of A Ring Simplified Pyripyropene A Analogues as Potent and Selective Synthetic SOAT2 Inhibitors. <i>ChemMedChem</i> , 2018, 13, 411-421.	1.6	8
23	Regioselective Mono-Deprotection of Di- <i>tert</i> -butylsilylene Acetal Derived from 1,3-Diol with Ammonium Fluoride. <i>Bulletin of the Chemical Society of Japan</i> , 2014, 87, 113-118.	2.0	6
24	Synthesis and Structural Revision of Cyslabdan. <i>Chemical and Pharmaceutical Bulletin</i> , 2016, 64, 1370-1377.	0.6	6
25	Total Synthesis and Absolute Configuration of Simpotentin, a Potentiator of Amphotericin B Activity. <i>Organic Letters</i> , 2019, 21, 5596-5599.	2.4	6
26	Total synthesis of 4-epi-atpenin A5 as a potent nematode complex II inhibitor. <i>Tetrahedron</i> , 2019, 75, 3178-3185.	1.0	6
27	Deprotection of silyl ethers by using SO <sub>3</sub> H silica gel: Application to sugar, nucleoside, and alkaloid derivatives. <i>Tetrahedron</i> , 2017, 73, 5425-5429.	1.0	5
28	Structure-Activity Relationship Study and Total Synthesis of Pyripyropene A as a Potent ACAT2-Selective Inhibitor. <i>Yuki Gosei Kagaku Kyokaiishi/Journal of Synthetic Organic Chemistry</i> , 2013, 71, 830-843.	0.0	3
29	Synthesis and Evaluation of Habiterpenol Analogs. <i>Chemical and Pharmaceutical Bulletin</i> , 2022, 70, 261-268.	0.6	3
30	Structure-activity relationship studies of atpenin A5 analogs with chemical modification of the side chain moiety. <i>Tetrahedron Letters</i> , 2019, 60, 1037-1042.	0.7	2
31	Repair of DNA damage induced by the novel nucleoside analogue CNDAG through homologous recombination. <i>Cancer Chemotherapy and Pharmacology</i> , 2020, 85, 661-672.	1.1	2
32	Development of a new air-stable structure-simplified nafuredin- $\hat{I}^3$ analog as a potent and selective nematode complex I inhibitor. <i>Journal of Antibiotics</i> , 2017, 70, 647-654.	1.0	1
33	Mixture of clopidogrel bisulfate and magnesium oxide tablets reduces clopidogrel dose administered through a feeding tube. <i>Journal of Pharmaceutical Health Care and Sciences</i> , 2021, 7, 18.	0.4	1