

Ichiro Hayakawa

List of Publications by Year in descending order

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papers

920
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#	ARTICLE	IF	CITATIONS
1	Structure-activity relationship studies on an antitumor marine macrolide using aplyronine A-swinholide A hybrid. <i>Organic and Biomolecular Chemistry</i> , 2022, 20, 2922-2938.	1.5	1
2	Development of Gatastatin G2, a β -Tubulin-specific Inhibitor. <i>Yuki Gosei Kagaku Kyokaiishi/Journal of Synthetic Organic Chemistry</i> , 2022, 80, 563-573.	0.0	0
3	Toward the Synthesis of Paspaline-Type Indole-Terpenes: Stereoselective Construction of Core Scaffold with Contiguous Asymmetric Quaternary Carbon Centers. <i>Journal of Organic Chemistry</i> , 2021, 86, 9802-9810.	1.7	8
4	Quantitative analysis of the <i>Tricholoma ustale</i> -derived toxin, ustalic acid, in mushroom and food samples by LC-MS/MS. <i>Forensic Science International</i> , 2020, 317, 110554.	1.3	3
5	Kinetic Resolution of \pm -Nitrolactones by Catalytic Asymmetric Hydrolysis or Ester-Amide Exchange Reaction. <i>Synlett</i> , 2020, 31, 2018-2022.	1.0	3
6	Structure Optimization of Gatastatin for the Development of β -Tubulin-Specific Inhibitor. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 1125-1129.	1.3	5
7	Dual Inhibition of β -Tubulin and Plk1 Induces Mitotic Cell Death. <i>Frontiers in Pharmacology</i> , 2020, 11, 620185.	1.6	4
8	Enantioselective Diels-Alder Reaction of 3-Nitrocoumarins Promoted by Chiral Organoammonium Salt Catalysts. <i>Synlett</i> , 2020, 31, 2013-2017.	1.0	4
9	Toward the Synthesis of Yuzurimine-Type Alkaloids: Stereoselective Construction of the Heterocyclic Portions of Deoxyyuzurimine and Macrodaphnine. <i>Organic Letters</i> , 2019, 21, 6337-6341.	2.4	10
10	Toward the Synthesis of SB-203207: Construction of Four Contiguous Nitrogen-Containing Stereogenic Centers. <i>Journal of Organic Chemistry</i> , 2019, 84, 15614-15623.	1.7	4
11	Catalytic enantioselective Hosomi-Sakurai reaction of \pm -ketoesters promoted by chiral copper complexes. <i>Chemical Communications</i> , 2019, 55, 3923-3926.	2.2	6
12	Structure-Activity Relationship Study of Gatastatin Based on the Topliss Tree Approach. <i>Heterocycles</i> , 2019, 99, 238.	0.4	3
13	Development of a novel inducer of protein-protein interactions based on aplyronine A. <i>Chemical Communications</i> , 2018, 54, 9537-9540.	2.2	8
14	Stimulation of microtubule-based transport by nucleation of microtubules on pigment granules. <i>Molecular Biology of the Cell</i> , 2017, 28, 1418-1425.	0.9	0
15	Reinvestigation of the Biomimetic Cyclization of 3,5-Diketo Esters: Application to the Total Synthesis of Cyercene A, an \pm -Methoxy- β -Pyrone-Containing Polypropionate. <i>Synlett</i> , 2017, 28, 1596-1600.	1.0	4
16	Total Synthesis of Biselide A, A Cytotoxic Macrolide of Marine Origin. <i>Synthesis</i> , 2017, 49, 2958-2970.	1.2	7
17	Total Synthesis of Biselide E, a Marine Polyketide. <i>Organic Letters</i> , 2017, 19, 5713-5716.	2.4	4
18	Second-generation total synthesis of aplyronine A featuring Ni/Cr-mediated coupling reactions. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 124-131.	1.5	11

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19	Regioselective DMAD-Insertion Reaction of Silyl Dienol Ether of $\hat{1}^3$ -Pyrone under Catalyst- and Heating-Free Conditions. <i>Heterocycles</i> , 2017, 94, 2299.	0.4	0
20	Discovery of O6-benzyl glaziovianin A, a potent cytotoxic substance and a potent inhibitor of $\hat{1}^{\pm}, \hat{1}^2$ -tubulin polymerization. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 5639-5645.	1.4	10
21	Total synthesis of natural derivatives and artificial analogs of 13-oxyingenol and their biological evaluation. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 11426-11437.	1.5	4
22	Recent Progress in The Synthetic Study of an Antitumor Marine Macrolide Aplyronine A and Related Molecules. <i>Heterocycles</i> , 2015, 91, 1137.	0.4	9
23	Construction of the [6-7-5-5] tetracyclic core of all the carbocyclic frameworks of yuzurimine-type alkaloids. <i>Chemical Communications</i> , 2015, 51, 11568-11571.	2.2	19
24	The $\hat{1}^3$ -tubulin-specific inhibitor gatastatin reveals temporal requirements of microtubule nucleation during the cell cycle. <i>Nature Communications</i> , 2015, 6, 8722.	5.8	47
25	Synthesis and structure-activity relationships for cytotoxicity and apoptosis-inducing activity of (+)-halichonine B. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 9969-9976.	1.5	3
26	Practical Synthesis of Glaziovianin A, a Cytotoxic Isoflavone, and Its 7-Propargyl Analogue. <i>Bulletin of the Chemical Society of Japan</i> , 2014, 87, 544-549.	2.0	6
27	Development of Biofunctional Molecules Based on Total Synthesis of Natural Products. <i>Yuki Gosei Kagaku Kyokaiishi/Journal of Synthetic Organic Chemistry</i> , 2014, 72, 126-136.	0.0	1
28	Glaziovianin A Prevents Endosome Maturation via Inhibiting Microtubule Dynamics. <i>ACS Chemical Biology</i> , 2013, 8, 884-889.	1.6	18
29	Total Synthesis and Biological Evaluation of Auripyrones A and B. <i>Bulletin of the Chemical Society of Japan</i> , 2012, 85, 1077-1092.	2.0	5
30	Design, Synthesis, and Biological Evaluations of Aplyronine-Mycolalide B Hybrid Compound. <i>Organic Letters</i> , 2012, 14, 1290-1293.	2.4	31
31	Toward the synthesis of $\hat{1}^3$ -pyrone-containing natural products: diastereoselective aldol-type reaction of a $\hat{1}^3$ -pyrone. <i>Tetrahedron</i> , 2012, 68, 6477-6484.	1.0	9
32	Design, synthesis, and biological evaluation of the analogues of glaziovianin A, a potent antitumor isoflavone. <i>Bioorganic and Medicinal Chemistry</i> , 2012, 20, 5745-5756.	1.4	20
33	Total Synthesis of ($\hat{1}^3$)-oxyingenol and its Natural Derivative. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 4972-4975.	7.2	42
34	Synthetic studies toward biselides. Part 1: synthesis of the core carbon framework of biselides A, B, and E using Stille coupling. <i>Tetrahedron Letters</i> , 2012, 53, 1390-1392.	0.7	5
35	Synthetic studies toward biselides. Part 2: synthesis of the macrolactone part of biselides A and B using allylic oxidation. <i>Tetrahedron Letters</i> , 2012, 53, 1393-1396.	0.7	5
36	Synthetic Studies toward 13-Oxyingenol: Construction of the Fully Substituted Tetracyclic Compound. <i>Organic Letters</i> , 2011, 13, 2160-2163.	2.4	14

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37	Toward the Second Generation Synthesis of Aplyronine A: Stereocontrolled Assembly of the C1~C19 Segment by Using an Asymmetric Nozaki~Hiyama~Kishi Coupling. <i>Organic Letters</i> , 2011, 13, 900-903.	2.4	29
38	Synthetic Studies towards Optically Active 13-Oxyingenol via Asymmetric ~Cyclopropanation. <i>Synthesis</i> , 2011, 2011, 769-777.	1.2	2
39	Total Synthesis of Auripyrones~A and B and Determination of the Absolute Configuration of Auripyrone~B. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 2401-2405.	7.2	22
40	Structure~activity relationship study of glaziovianin A against cell cycle progression and spindle formation of HeLa S3 cells. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 5402-5404.	1.0	16
41	Total Synthesis and Cytotoxicity of Haterumalides NA and B and Their Artificial Analogues. Yuki Gosei Kagaku Kyokaiishi/Journal of Synthetic Organic Chemistry, 2010, 68, 814-823.	0.0	4
42	Practical Syntheses of Enantiomerically Pure Key Intermediates of Opioid Receptor-like 1 (ORL1) Antagonists. <i>Synthesis</i> , 2009, 2009, 1153-1162.	1.2	4
43	Aldol-type reaction of a 4-pyrone: a straightforward approach to 4-pyrone-containing natural products. <i>Tetrahedron Letters</i> , 2009, 50, 325-328.	0.7	19
44	Total Synthesis and Cytotoxicity of Haterumalides NA and B and Their Artificial Analogues. <i>Journal of Organic Chemistry</i> , 2009, 74, 3370-3377.	1.7	34
45	Structure of zamamistatin~a correction. <i>Tetrahedron Letters</i> , 2008, 49, 5383-5384.	0.7	11
46	Synthesis of ustalic acid, an inhibitor of Na+,K+-ATPase. <i>Tetrahedron</i> , 2008, 64, 5873-5877.	1.0	16
47	Second-Generation Total Synthesis of Haterumalide NA Using B-Alkyl Suzuki~Miyaura Coupling. <i>Organic Letters</i> , 2008, 10, 1859-1862.	2.4	28
48	Synthesis of Glaziovianin A: A Potent Antitumor Isoflavone. <i>Chemistry Letters</i> , 2007, 36, 1382-1383.	0.7	21
49	Synthetic Studies on Mycalolide B: Synthesis of the C7~C35 Fragment. <i>Chemistry Letters</i> , 2007, 36, 1490-1491.	0.7	11
50	Development of Practical Synthesis of Halichondrins. Yuki Gosei Kagaku Kyokaiishi/Journal of Synthetic Organic Chemistry, 2007, 65, 1225-1226.	0.0	0
51	Synthetic study on 13-oxyingenol: construction of the full carbon framework. <i>Tetrahedron Letters</i> , 2007, 48, 6221-6224.	0.7	18
52	Marine cytotoxic macrolides haterumalides and biselides, and related natural products. <i>Chemical Record</i> , 2007, 7, 254-264.	2.9	32
53	Enantioselective synthesis of aurisides A and B, cytotoxic macrolide glycosides of marine origin. <i>Tetrahedron</i> , 2006, 62, 7687-7698.	1.0	23
54	Revised structure of zamamistatin. <i>Tetrahedron Letters</i> , 2006, 47, 155-158.	0.7	21

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55	Synthesis of the tricyclic core of halichlorine. <i>Chemical Communications</i> , 2004, , 1222.	2.2	22
56	Synthesis of (±)-Pinnaic Acid. <i>Heterocycles</i> , 2003, 59, 441.	0.4	32
57	Three New Alkaloids, Convolutamines F and G, and Convolutamydine E, from the Floridian Marine Bryozoan <i>Amathia convoluta</i> . <i>Collection of Czechoslovak Chemical Communications</i> , 1999, 64, 1147-1153.	1.0	35
58	Multi-valent polymer of vancomycin: enhanced antibacterial activity against VRE. <i>Chemical Communications</i> , 1999, , 1361-1362.	2.2	87
59	Absolute stereochemistry of halichlorine; A potent inhibitor of VCAM-1 induction. <i>Tetrahedron Letters</i> , 1998, 39, 861-862.	0.7	86