

Hiroyuki Osada

List of Publications by Year in descending order

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375
papers

11,617
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22153

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all docs

390
docs citations

390
times ranked

12468
citing authors

#	ARTICLE	IF	CITATIONS
1	Involvement of p38 Mitogen-activated Protein Kinase Signaling Pathway in Osteoclastogenesis Mediated by Receptor Activator of NF- κ B Ligand (RANKL). <i>Journal of Biological Chemistry</i> , 2000, 275, 31155-31161.	3.4	482
2	Novel mammalian cell cycle inhibitors, spirotryprostatins A and B, produced by <i>Aspergillus fumigatus</i> , which inhibit mammalian cell cycle at G2/M phase. <i>Tetrahedron</i> , 1996, 52, 12651-12666.	1.9	478
3	Tryprostatin A, a specific and novel inhibitor of microtubule assembly. <i>Biochemical Journal</i> , 1998, 333, 543-548.	3.7	211
4	Novel Mammalian Cell Cycle Inhibitors, Tryprostatins A, B and Other Diketopiperazines Produced by <i>Aspergillus fumigatus</i> . I. Taxonomy, Fermentation, Isolation and Biological Properties.. <i>Journal of Antibiotics</i> , 1996, 49, 527-533.	2.0	187
5	Immobilization of Natural Products on Glass Slides by Using a Photoaffinity Reaction and the Detection of Protein-Small-Molecule Interactions. <i>Angewandte Chemie - International Edition</i> , 2003, 42, 5584-5587.	13.8	153
6	Novel mammalian cell cycle inhibitors, cycloprostatins A-D, produced by <i>Aspergillus fumigatus</i> , which inhibit mammalian cell cycle at G2/M phase. <i>Tetrahedron</i> , 1997, 53, 59-72.	1.9	145
7	Epolactaene, a Novel Neuritogenic Compound in Human Neuroblastoma Cells, Produced by a Marine Fungus.. <i>Journal of Antibiotics</i> , 1995, 48, 733-735.	2.0	140
8	Novel Mammalian Cell Cycle Inhibitors, Tryprostatins A, B and Other Diketopiperazines Produced by <i>Aspergillus fumigatus</i> . II. Physico-chemical Properties and Structures.. <i>Journal of Antibiotics</i> , 1996, 49, 534-540.	2.0	140
9	Azaspirene: A Novel Angiogenesis Inhibitor Containing a 1-Oxa-7-azaspiro[4.4]non-2-ene-4,6-dione Skeleton Produced by the Fungus <i>Neosartorya</i> sp.. <i>Organic Letters</i> , 2002, 4, 2845-2848.	4.6	128
10	The identification of an osteoclastogenesis inhibitor through the inhibition of glyoxalase I. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 11691-11696.	7.1	125
11	Mutations in the Plk gene lead to instability of Plk protein in human tumour cell lines. <i>Nature Cell Biology</i> , 2000, 2, 852-854.	10.3	122
12	Inhibition of bone and muscle metastases of lung cancer cells by a decrease in the number of monocytes/macrophages. <i>Cancer Science</i> , 2008, 99, 1595-1602.	3.9	116
13	SPR Imaging of Photo-Cross-Linked Small-Molecule Arrays on Gold. <i>Analytical Chemistry</i> , 2006, 78, 2226-2230.	6.5	111
14	Biosynthesis of the mycotoxin tenuazonic acid by a fungal NRPS-PKS hybrid enzyme. <i>Nature Communications</i> , 2015, 6, 8758.	12.8	108
15	Photo-Cross-Linked Small-Molecule Affinity Matrix for Facilitating Forward and Reverse Chemical Genetics. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 3559-3562.	13.8	107
16	A Novel Action of Terpendole E on the Motor Activity of Mitotic Kinesin Eg5. <i>Chemistry and Biology</i> , 2003, 10, 131-137.	6.0	105
17	The phosphorylation status and anti-apoptotic activity of Bcl-2 are regulated by ERK and protein phosphatase 2A on the mitochondria. <i>FEBS Letters</i> , 2004, 569, 249-255.	2.8	105
18	Characterization of Giant Modular PKSs Provides Insight into Genetic Mechanism for Structural Diversification of Aminopolyol Polyketides. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 1740-1745.	13.8	103

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19	Deficiency in Chromosome Congression by the Inhibition of Plk1 Polo Box Domain-dependent Recognition. <i>Journal of Biological Chemistry</i> , 2009, 284, 2344-2353.	3.4	102
20	Reversal of breast cancer resistance protein-mediated drug resistance by tryprostatin A. <i>International Journal of Cancer</i> , 2003, 107, 721-728.	5.1	99
21	The Anticancer Natural Product Pironetin Selectively Targets Lys352 of β -Tubulin. <i>Chemistry and Biology</i> , 2004, 11, 799-806.	6.0	95
22	Cell cycle arrest and antitumor activity of pironetin and its derivatives. <i>Cancer Letters</i> , 1998, 126, 29-32.	7.2	94
23	Epolactaene binds human Hsp60 Cys442 resulting in the inhibition of chaperone activity. <i>Biochemical Journal</i> , 2005, 387, 835-840.	3.7	94
24	Toxic tau oligomer formation blocked by capping of cysteine residues with 1,2-dihydroxybenzene groups. <i>Nature Communications</i> , 2015, 6, 10216.	12.8	94
25	Epoxyquinol A, a Highly Functionalized Pentaketide Dimer with Antiangiogenic Activity Isolated from Fungal Metabolites. <i>Journal of the American Chemical Society</i> , 2002, 124, 3496-3497.	13.7	93
26	A small-molecule inhibitor shows that pirin regulates migration of melanoma cells. <i>Nature Chemical Biology</i> , 2010, 6, 667-673.	8.0	91
27	Neuritogenic Effect of Epolactaene Derivatives on Human Neuroblastoma Cells Which Lack High-Affinity Nerve Growth Factor Receptors. <i>Journal of Medicinal Chemistry</i> , 1997, 40, 391-394.	6.4	90
28	Morphobase, an Encyclopedic Cell Morphology Database, and Its Use for Drug Target Identification. <i>Chemistry and Biology</i> , 2012, 19, 1620-1630.	6.0	89
29	Proteomic profiling of small-molecule inhibitors reveals dispensability of MTH1 for cancer cell survival. <i>Scientific Reports</i> , 2016, 6, 26521.	3.3	87
30	RECK Negatively Regulates Matrix Metalloproteinase-9 Transcription. <i>Cancer Research</i> , 2009, 69, 1502-1508.	0.9	86
31	Fusarisetin A, an Acinar Morphogenesis Inhibitor from a Soil Fungus, <i>Fusarium</i> sp. FN080326. <i>Journal of the American Chemical Society</i> , 2011, 133, 6865-6867.	13.7	84
32	Vipirinin, a Coumarin-based HIV-1 Vpr Inhibitor, Interacts with a Hydrophobic Region of VPR. <i>Journal of Biological Chemistry</i> , 2011, 286, 14049-14056.	3.4	83
33	Inhibition of Hsp90 activates osteoclast c-Src signaling and promotes growth of prostate carcinoma cells in bone. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 15541-15546.	7.1	82
34	Application of Proteomic Profiling Based on 2D-DIGE for Classification of Compounds According to the Mechanism of Action. <i>Chemistry and Biology</i> , 2010, 17, 460-470.	6.0	82
35	Reveromycin A biosynthesis uses RevG and RevJ for stereospecific spiroacetal formation. <i>Nature Chemical Biology</i> , 2011, 7, 461-468.	8.0	80
36	Discovery of a Small Molecule PDI Inhibitor That Inhibits Reduction of HIV-1 Envelope Glycoprotein gp120. <i>ACS Chemical Biology</i> , 2011, 6, 245-251.	3.4	80

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37	Spectomycin B1 as a Novel SUMOylation Inhibitor That Directly Binds to SUMO E2. ACS Chemical Biology, 2013, 8, 2635-2642.	3.4	80
38	Reveromycin a, a new antibiotic which inhibits the mitogenic activity of epidermal growth factor.. Journal of Antibiotics, 1991, 44, 259-261.	2.0	79
39	Reveromycin A, an agent for osteoporosis, inhibits bone resorption by inducing apoptosis specifically in osteoclasts. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 4729-4734.	7.1	79
40	In vitro reconstruction of tetronate RK-682 biosynthesis. Nature Chemical Biology, 2010, 6, 99-101.	8.0	79
41	Functional annotation of chemical libraries across diverse biological processes. Nature Chemical Biology, 2017, 13, 982-993.	8.0	76
42	Target identification of small molecules based on chemical biology approaches. Molecular BioSystems, 2013, 9, 897.	2.9	74
43	Synthesis and structure-activity relationship studies on tryprostatin A, an inhibitor of breast cancer resistance protein. Bioorganic and Medicinal Chemistry, 2008, 16, 4626-4651.	3.0	73
44	Cobtorin target analysis reveals that pectin functions in the deposition of cellulose microfibrils in parallel with cortical microtubules. Plant Journal, 2010, 64, 657-667.	5.7	73
45	Construction of a microbial natural product library for chemical biology studies. Current Opinion in Chemical Biology, 2012, 16, 101-108.	6.1	72
46	Selective Inhibition of the Bacterial Peptidoglycan Biosynthesis by the New Types of Liposidomycins.. Journal of Antibiotics, 1998, 51, 1099-1104.	2.0	71
47	Amphidinolide H, a Potent Cytotoxic Macrolide, Covalently Binds on Actin Subdomain 4 and Stabilizes Actin Filament. Chemistry and Biology, 2004, 11, 1269-1277.	6.0	70
48	Xanthohumol Impairs Autophagosome Maturation through Direct Inhibition of Valosin-Containing Protein. ACS Chemical Biology, 2012, 7, 892-900.	3.4	70
49	Identification of Cytochrome P450s Required for Fumitremorgin Biosynthesis in <i>Aspergillus fumigatus</i> . ChemBioChem, 2009, 10, 920-928.	2.6	69
50	Sulfonamides identified as plant immune-priming compounds in high-throughput chemical screening increase disease resistance in <i>Arabidopsis thaliana</i> . Frontiers in Plant Science, 2012, 3, 245.	3.6	68
51	Identification of <i>Saccharomyces cerevisiae</i> Isoleucyl-tRNA Synthetase as a Target of the G1-specific Inhibitor Reveromycin A. Journal of Biological Chemistry, 2002, 277, 28810-28814.	3.4	67
52	A new enzyme involved in the control of the stereochemistry in the decalin formation during equisetin biosynthesis. Biochemical and Biophysical Research Communications, 2015, 460, 210-215.	2.1	65
53	Biochemical Characterization of a Novel Indole Prenyltransferase from <i>Streptomyces</i> sp. SN-593. Journal of Bacteriology, 2010, 192, 2839-2851.	2.2	64
54	Discovery of novel antiviral agents directed against the influenza A virus nucleoprotein using photo-cross-linked chemical arrays. Biochemical and Biophysical Research Communications, 2010, 394, 721-727.	2.1	64

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55	Affinity-based target identification for bioactive small molecules. <i>MedChemComm</i> , 2014, 5, 277-287.	3.4	64
56	Rapid screening method for inhibitors of protein kinase C.. <i>Journal of Antibiotics</i> , 1988, 41, 925-931.	2.0	63
57	Enhancement and Selective Production of Phoslactomycin B, a Protein Phosphatase IIa Inhibitor, through Identification and Engineering of the Corresponding Biosynthetic Gene Cluster. <i>Journal of Biological Chemistry</i> , 2003, 278, 35552-35557.	3.4	63
58	Secretion of Heparanase Protein Is Regulated by Glycosylation in Human Tumor Cell Lines. <i>Journal of Biological Chemistry</i> , 2004, 279, 2697-2703.	3.4	62
59	Photo-Cross-Linked Small-Molecule Microarrays as Chemical Genomic Tools for Dissecting Protein-Ligand Interactions. <i>Chemistry - an Asian Journal</i> , 2006, 1, 789-797.	3.3	62
60	Heparanase as a molecular target of cancer chemotherapy. <i>Cancer Science</i> , 2004, 95, 553-558.	3.9	61
61	Terpendole E, a Kinesin Eg5 Inhibitor, Is a Key Biosynthetic Intermediate of Indole-Diterpenes in the Producing Fungus <i>Chaetomium alba</i> . <i>Chemistry and Biology</i> , 2012, 19, 1611-1619.	6.0	61
62	Reveromycins, new inhibitors of eukaryotic cell growth. III. Structures of reveromycins A, B, C and D.. <i>Journal of Antibiotics</i> , 1992, 45, 1420-1427.	2.0	59
63	Apoptosis induction via microtubule disassembly by an antitumour compound, pironetin. <i>Biochemical Journal</i> , 1999, 340, 411-416.	3.7	58
64	RECK-Mediated Suppression of Tumor Cell Invasion Is Regulated by Glycosylation in Human Tumor Cell Lines. <i>Cancer Research</i> , 2005, 65, 7455-7461.	0.9	58
65	Epoxyquinol B, a Fungal Metabolite with a Potent Antiangiogenic Activity.. <i>Journal of Antibiotics</i> , 2002, 55, 829-831.	2.0	55
66	Structure-activity relationships of epolactaene derivatives: structural requirements for inhibition of Hsp60 chaperone activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 4425-4429.	2.2	55
67	Reveromycins, new inhibitors of eukaryotic cell growth. II. Biological activities.. <i>Journal of Antibiotics</i> , 1992, 45, 1414-1419.	2.0	53
68	Lucilactaene, a New Cell Cycle Inhibitor in p53-Transfected Cancer Cells, Produced by a <i>Fusarium</i> sp.. <i>Journal of Antibiotics</i> , 2001, 54, 850-854.	2.0	53
69	Cytotrienin A, a Novel Apoptosis Inducer in Human Leukemia HL-60 Cells.. <i>Journal of Antibiotics</i> , 1997, 50, 370-372.	2.0	51
70	Crystal structure of the predicted phospholipase LYPLAL1 reveals unexpected functional plasticity despite close relationship to acyl protein thioesterases. <i>Journal of Lipid Research</i> , 2012, 53, 43-50.	4.2	50
71	Control of the Stereochemical Course of [4+2] Cycloaddition during <i>trans</i> -Decalin Formation by Fsa Family Enzymes. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 9754-9758.	13.8	49
72	Phoslactomycin targets cysteine-269 of the protein phosphatase 2A catalytic subunit in cells. <i>FEBS Letters</i> , 2005, 579, 2463-2468.	2.8	48

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73	Reveromycins, new inhibitors of eukaryotic cell growth. I. Producing organism, fermentation, isolation and physico-chemical properties.. Journal of Antibiotics, 1992, 45, 1409-1413.	2.0	47
74	Screening of cell cycle inhibitors from microbial metabolites by a bioassay using a mouse cdc2 mutant cell line, tsFT210. Bioorganic and Medicinal Chemistry, 1997, 5, 193-203.	3.0	47
75	Reveromycin A Inhibits Osteolytic Bone Metastasis of Small-Cell Lung Cancer Cells, SBC-5, through an Antiosteoclastic Activity. Clinical Cancer Research, 2005, 11, 8822-8828.	7.0	47
76	Involvement of Disulfide Bond Formation in the Activation of Heparanase. Cancer Research, 2007, 67, 7841-7849.	0.9	47
77	Identification of a Small-Molecule Inhibitor of DNA Topoisomerase II by Proteomic Profiling. Chemistry and Biology, 2011, 18, 743-751.	6.0	47
78	A new biological role of sangivamycin; Inhibition of protein kinases.. Journal of Antibiotics, 1989, 42, 102-106.	2.0	46
79	Phenotypic Identification of a Novel Autophagy Inhibitor Chemotype Targeting Lipid Kinase VPS34. Angewandte Chemie - International Edition, 2017, 56, 8153-8157.	13.8	45
80	Design and synthesis of a dimeric derivative of RK-682 with increased inhibitory activity against VHR, a dual-specificity ERK phosphatase: implications for the molecular mechanism of the inhibition. Chemistry and Biology, 2001, 8, 1209-1220.	6.0	44
81	Induction of morphological change of human myeloid leukemia and activation of protein kinase C by a novel antibiotic, tautomycin.. Journal of Antibiotics, 1988, 41, 932-937.	2.0	42
82	A new inhibitor of protein kinase C, RK-286C (4'-demethylamino-4'-hydroxystaurosporine) I. Screening, taxonomy, fermentation and biological activity.. Journal of Antibiotics, 1990, 43, 163-167.	2.0	42
83	Introduction of New Tools for Chemical Biology Research on Microbial Metabolites. Bioscience, Biotechnology and Biochemistry, 2010, 74, 1135-1140.	1.3	42
84	A new inhibitor of protein kinase C, RK-1409 (7-oxostaurosporine). I. Taxonomy and biological activity.. Journal of Antibiotics, 1992, 45, 189-194.	2.0	41
85	p62 Functions as a p38 MAP Kinase Regulator. Biochemical and Biophysical Research Communications, 2000, 269, 521-525.	2.1	41
86	lejimalides Show Anti-Osteoclast Activity via V-ATPase Inhibition. Bioscience, Biotechnology and Biochemistry, 2006, 70, 1364-1370.	1.3	41
87	An overproduction of astellolides induced by genetic disruption of chromatin-remodeling factors in Aspergillus oryzae. Journal of Antibiotics, 2016, 69, 4-8.	2.0	41
88	Expression of heparanase in human tumor cell lines and human head and neck tumors. Cancer Letters, 2003, 193, 83-89.	7.2	40
89	Dephosphorylation of Bcl-2 by protein phosphatase 2A results in apoptosis resistance. Cancer Science, 2004, 95, 266-270.	3.9	40
90	Phosphorylation-Dependent Protein-Protein Interaction Modules As Potential Molecular Targets for Cancer Therapy. Current Drug Targets, 2012, 13, 1654-1658.	2.1	40

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91	A new inhibitor of protein kinase C, RK-286C (4'-demethylamino-4'-hydroxystaurosporine) II. Isolation, physico-chemical properties and structure.. <i>Journal of Antibiotics</i> , 1990, 43, 168-173.	2.0	39
92	Robust and Systematic Drug Screening Method Using Chemical Arrays and the Protein Library: Identification of Novel Inhibitors of Carbonic Anhydrase II. <i>Bioscience, Biotechnology and Biochemistry</i> , 2008, 72, 2739-2749.	1.3	39
93	Verticilactam, a New Macrolactam Isolated from a Microbial Metabolite Fraction Library. <i>Organic Letters</i> , 2010, 12, 4564-4567.	4.6	39
94	Methyl 3-((6-Methoxy-1,4-dihydroindeno[1,2- <i>c</i>]pyrazol-3-yl)amino)benzoate (GN39482) as a Tubulin Polymerization Inhibitor Identified by MorphoBase and ChemProteoBase Profiling Methods. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 4230-4241.	6.4	38
95	New Types of Liposidomycins that Inhibit Bacterial Peptidoglycan Synthesis and are Produced by <i>Streptomyces</i> . II. Isolation and Structure Elucidation.. <i>Journal of Antibiotics</i> , 1998, 51, 647-654.	2.0	37
96	New Types of Liposidomycins that Inhibit Bacterial Peptidoglycan Synthesis and are Produced by <i>Streptomyces</i> . I. Producing Organism and Medium Components.. <i>Journal of Antibiotics</i> , 1998, 51, 640-646.	2.0	37
97	Osteoclast-targeting small molecules for the treatment of neoplastic bone metastases. <i>Cancer Science</i> , 2009, 100, 1999-2005.	3.9	37
98	p38 mitogen-activated protein kinase plays a key role in regulating MAPKAPK2 expression. <i>Biochemical and Biophysical Research Communications</i> , 2005, 337, 415-421.	2.1	36
99	Azaspirene, a fungal product, inhibits angiogenesis by blocking Raf-1 activation. <i>Cancer Science</i> , 2008, 99, 1853-1858.	3.9	36
100	Novel Heparan Sulfate Mimetic Compounds as Antitumor Agents. <i>Chemistry and Biology</i> , 2004, 11, 367-377.	6.0	35
101	Discovery of the novel autophagy inhibitor aumitin that targets mitochondrial complex I. <i>Chemical Science</i> , 2018, 9, 3014-3022.	7.4	34
102	A Novel Antiviral Target Structure Involved in the RNA Binding, Dimerization, and Nuclear Export Functions of the Influenza A Virus Nucleoprotein. <i>PLoS Pathogens</i> , 2015, 11, e1005062.	4.7	34
103	New Cyclic Lipopeptides of the Iturin Class Produced by Saltern-Derived <i>Bacillus</i> sp. KCB14S006. <i>Marine Drugs</i> , 2016, 14, 72.	4.6	33
104	Identification of a Small Compound Targeting PKM2-Regulated Signaling Using 2D Gel Electrophoresis-Based Proteome-wide CETSA. <i>Cell Chemical Biology</i> , 2020, 27, 186-196.e4.	5.2	33
105	Pyrrrolizilactone, a new pyrrolizidinone metabolite produced by a fungus. <i>Journal of Antibiotics</i> , 2013, 66, 621-623.	2.0	32
106	Identification of a novel sesquiterpene biosynthetic machinery involved in astellolide biosynthesis. <i>Scientific Reports</i> , 2016, 6, 32865.	3.3	32
107	Distribution of photo-cross-linked products from 3-aryl-3-trifluoromethyl-diazirines and alcohols. <i>Tetrahedron</i> , 2008, 64, 5692-5698.	1.9	31
108	Gene Disruption and Biochemical Characterization of Verruculogen Synthase of <i>Aspergillus fumigatus</i> . <i>ChemBioChem</i> , 2011, 12, 711-714.	2.6	31

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109	Structures and biological activities of azaphilones produced by <i>Penicillium</i> sp. KCB11A109 from a ginseng field. <i>Phytochemistry</i> , 2016, 122, 154-164.	2.9	31
110	Stereocomplementary Chemoenzymatic Pictet-Spengler Reactions for Formation of Rare Azepino-indole Frameworks: Discovery of Antimalarial Compounds. <i>ACS Catalysis</i> , 2019, 9, 7443-7448.	11.2	31
111	Identification of a novel Vpr-binding compound that inhibits HIV-1 multiplication in macrophages by chemical array. <i>Biochemical and Biophysical Research Communications</i> , 2010, 403, 40-45.	2.1	30
112	Boron-Based Inhibitors of Acyl Protein Thioesterases 1 and 2. <i>ChemBioChem</i> , 2013, 14, 115-122.	2.6	30
113	RK-1355A and B, novel quinomycin derivatives isolated from a microbial metabolites fraction library based on NPPlot screening. <i>Journal of Antibiotics</i> , 2014, 67, 323-329.	2.0	30
114	Apoptosis induction via microtubule disassembly by an antitumour compound, pironetin. <i>Biochemical Journal</i> , 1999, 340, 411.	3.7	29
115	Determination by Asymmetric Total Synthesis of the Absolute Configuration of Lucilactaene, a Cell-Cycle Inhibitor in p53-Transfected Cancer Cells. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 3110-3115.	13.8	29
116	Genetic Safeguard against Mycotoxin Cyclopiazonic Acid Production in <i>Aspergillus oryzae</i> . <i>ChemBioChem</i> , 2011, 12, 1376-1382.	2.6	29
117	Identification of small molecule inhibitors of p27 ^{Kip1} ubiquitination by high-throughput screening. <i>Cancer Science</i> , 2013, 104, 1461-1467.	3.9	29
118	Proteomic profiling reveals that collismycin A is an iron chelator. <i>Scientific Reports</i> , 2016, 6, 38385.	3.3	29
119	High-throughput screening identifies artesunate as selective inhibitor of cancer stemness: Involvement of mitochondrial metabolism. <i>Biochemical and Biophysical Research Communications</i> , 2016, 477, 737-742.	2.1	29
120	Biosynthesis and Structure-Activity Relationship Studies of Okaramines That Target Insect Glutamate-Gated Chloride Channels. <i>ACS Chemical Biology</i> , 2018, 13, 561-566.	3.4	29
121	A new inhibitor of protein kinase C, RK-1409 (7-oxostaurosporine). II. Fermentation, isolation, physico-chemical properties and structure.. <i>Journal of Antibiotics</i> , 1992, 45, 195-198.	2.0	28
122	Chemical modification of reveromycin A and its biological activities. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2002, 12, 3363-3366.	2.2	28
123	A Point Mutation in <i>ftmD</i> Blocks the Fumitremorgin Biosynthetic Pathway in <i>Aspergillus fumigatus</i> Strain Af293. <i>Bioscience, Biotechnology and Biochemistry</i> , 2013, 77, 1061-1067.	1.3	28
124	Amino-group carrier-protein-mediated secondary metabolite biosynthesis in <i>Streptomyces</i> . <i>Nature Chemical Biology</i> , 2016, 12, 967-972.	8.0	28
125	Stachybotrysin, an Osteoclast Differentiation Inhibitor from the Marine-Derived Fungus <i>Stachybotrys</i> sp. KCB13F013. <i>Journal of Natural Products</i> , 2016, 79, 2703-2708.	3.0	28
126	Evolutionarily conserved BIL4 suppresses the degradation of brassinosteroid receptor BRI1 and regulates cell elongation. <i>Scientific Reports</i> , 2017, 7, 5739.	3.3	28

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127	Anti-allergy effect of mojabanchromanol isolated from <i>Sargassum horneri</i> in bone marrow-derived cultured mast cells. <i>Algal Research</i> , 2020, 48, 101898.	4.6	28
128	Cleavable Linker for Photo-Cross-Linked Small-Molecule Affinity Matrix. <i>Bioconjugate Chemistry</i> , 2010, 21, 182-186.	3.6	27
129	Furaquinocins I and J: novel polyketide isoprenoid hybrid compounds from <i>Streptomyces reveromyceticus</i> SN-593. <i>Journal of Antibiotics</i> , 2011, 64, 509-513.	2.0	27
130	Development of a Terpenoid-Production Platform in <i>Streptomyces reveromyceticus</i> SN-593. <i>ACS Synthetic Biology</i> , 2017, 6, 2339-2349.	3.8	27
131	ATM blocks tunicamycin-induced endoplasmic reticulum stress. <i>FEBS Letters</i> , 2009, 583, 903-908.	2.8	26
132	Identification of a Molecular Target of a Novel Fungal Metabolite, Pyrrolizilactone, by Phenotypic Profiling Systems. <i>ChemBioChem</i> , 2013, 14, 2456-2463.	2.6	26
133	Unantimycin A, a new neoantimycin analog isolated from a microbial metabolite fraction library. <i>Journal of Antibiotics</i> , 2016, 69, 456-458.	2.0	26
134	FPX is a Novel Chemical Inducer that Promotes Callus Formation and Shoot Regeneration in Plants. <i>Plant and Cell Physiology</i> , 2018, 59, 1555-1567.	3.1	26
135	Discovery and applications of nucleoside antibiotics beyond polyoxin. <i>Journal of Antibiotics</i> , 2019, 72, 855-864.	2.0	26
136	Biosynthesis of phoslactomycins: cyclohexanecarboxylic acid as the starter unit. <i>Tetrahedron</i> , 2003, 59, 7465-7471.	1.9	25
137	Haenamindole, an unusual diketopiperazine derivative from a marine-derived <i>Penicillium</i> sp. KCB12F005. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 5398-5401.	2.2	25
138	Epiblastin A Induces Reprogramming of Epiblast Stem Cells Into Embryonic Stem Cells by Inhibition of Casein Kinase 1. <i>Cell Chemical Biology</i> , 2016, 23, 494-507.	5.2	25
139	Small molecules that target phosphorylation dependent protein-protein interaction. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3246-3254.	3.0	25
140	Structure-based design of a selective heparanase inhibitor as an antimetastatic agent. <i>Molecular Cancer Therapeutics</i> , 2004, 3, 1069-77.	4.1	25
141	Indomethacin Induction of Metamorphosis from the Asexual Stage to Sexual Stage in the Moon Jellyfish, <i>Aurelia aurita</i> . <i>Bioscience, Biotechnology and Biochemistry</i> , 2012, 76, 1397-1400.	1.3	24
142	Affinity-based screening of MDM2/MDMX-p53 interaction inhibitors by chemical array: Identification of novel peptidic inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 3802-3805.	2.2	24
143	Antiangiogenic Potential of Microbial Metabolite Elaiophylin for Targeting Tumor Angiogenesis. <i>Molecules</i> , 2018, 23, 563.	3.8	24
144	A new inhibitor of protein kinase C, RK-1409B (4'-demethylamino-4'-hydroxy-3'-epistaurosporine).. <i>Journal of Antibiotics</i> , 1992, 45, 1428-1432.	2.0	23

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