

Karl-Heinz Altmann

List of Publications by Year in descending order

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134
papers

6,103
citations

71004

43
h-index

90395

73
g-index

160
all docs

160
docs citations

160
times ranked

7154
citing authors

#	ARTICLE	IF	CITATIONS
1	Synthesis and Biological Evaluation of Endocannabinoid Uptake Inhibitors Derived from WOBE437. <i>ChemMedChem</i> , 2021, 16, 145-154.	1.6	6
2	Synthesis of Morpholine-Based Analogues of (Δ^9)-Zampanolide and Their Biological Activity. <i>Chemistry - A European Journal</i> , 2021, 27, 5936-5943.	1.7	7
3	Studies toward the Synthesis of an Oxazole-Based Analog of (Δ^9)-Zampanolide. <i>Organic Letters</i> , 2021, 23, 2238-2242.	2.4	5
4	Mechanism of substrate transport and inhibition of the human LAT1-4F2hc amino acid transporter. <i>Cell Discovery</i> , 2021, 7, 16.	3.1	40
5	An Antigen Capture Assay for the Detection of Mycolactone, the Polyketide Toxin of <i>Mycobacterium ulcerans</i> . <i>Journal of Immunology</i> , 2021, 206, 2753-2762.	0.4	3
6	Synthesis and Structure-Activity Relationship Studies of C2-Modified Analogs of the Antimycobacterial Natural Product Pyridomycin. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 1105-1131.	2.9	11
7	Taxanes convert regions of perturbed microtubule growth into rescue sites. <i>Nature Materials</i> , 2020, 19, 355-365.	13.3	44
8	Small molecule inhibitors provide insights into the relevance of LAT1 and LAT2 in maternal-fetal amino acid transport. <i>Journal of Cellular and Molecular Medicine</i> , 2020, 24, 12681-12693.	1.6	12
9	A Method for the Stereoselective Construction of the Hemiaminal Center in Zampanolides. <i>Organic Letters</i> , 2020, 22, 8345-8348.	2.4	5
10	Ring-Closing Metathesis Approaches towards the Total Synthesis of Rhizoxins. <i>Molecules</i> , 2020, 25, 4527.	1.7	6
11	Morphing of Amphipathic Helices to Explore the Activity and Selectivity of Membranolytic Antimicrobial Peptides. <i>Biochemistry</i> , 2020, 59, 3772-3781.	1.2	4
12	An RCM-Based Total Synthesis of the Antibiotic Disciformycin...B. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 17393-17397.	7.2	12
13	Die Totalsynthese des Antibiotikums Disciformycin...B durch Ringschlussmetathese. <i>Angewandte Chemie</i> , 2020, 132, 17546-17550.	1.6	2
14	On the Importance of the Thiazole Nitrogen in Epothilones: Semisynthesis and Microtubule-Binding Affinity of Deaza-Epothilone C. <i>Chemistry</i> , 2020, 2, 499-509.	0.9	0
15	Development of an ELISA for the quantification of mycolactone, the cytotoxic macrolide toxin of <i>Mycobacterium ulcerans</i> . <i>PLoS Neglected Tropical Diseases</i> , 2020, 14, e0008357.	1.3	9
16	Ammonia uptake by transmembrane pH gradient poly(isoprene)-block-poly(ethylene glycol) polymersomes. <i>Soft Matter</i> , 2020, 16, 2725-2735.	1.2	2
17	Total Synthesis of the Endocannabinoid Uptake Inhibitor Guineensine and SAR Studies. <i>ChemMedChem</i> , 2019, 14, 1590-1596.	1.6	6
18	Configurally Stabilized Analogs of <i>M. ulcerans</i> Exotoxins Mycolactones A and B Reveal the Importance of Side Chain Geometry for Mycolactone Virulence. <i>Organic Letters</i> , 2019, 21, 5853-5857.	2.4	11

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19	Synthesis, Microtubule-Binding Affinity, and Antiproliferative Activity of New Epothilone Analogs and of an EGFR-Targeted Epothilone-Peptide Conjugate. <i>International Journal of Molecular Sciences</i> , 2019, 20, 1113.	1.8	4
20	Synthesis, Profiling, and Bioactive Conformation of trans ϵ -Cyclopropyl Epothilones. <i>Helvetica Chimica Acta</i> , 2019, 102, e1900078.	1.0	3
21	Crystal Structure of the Cyclostreptin-Tubulin Adduct: Implications for Tubulin Activation by Taxane-Site Ligands. <i>International Journal of Molecular Sciences</i> , 2019, 20, 1392.	1.8	24
22	Primary resistance mechanism of the canine distemper virus fusion protein against a small-molecule membrane fusion inhibitor. <i>Virus Research</i> , 2019, 259, 28-37.	1.1	10
23	Structural basis of small-molecule inhibition of human multidrug transporter ABCG2. <i>Nature Structural and Molecular Biology</i> , 2018, 25, 333-340.	3.6	258
24	Total Synthesis of Ripostatin B and Structure-Activity Relationship Studies on Ripostatin Analogs. <i>Journal of Organic Chemistry</i> , 2018, 83, 7150-7172.	1.7	22
25	Zampanolide Binding to Tubulin Indicates Cross-Talk of Taxane Site with Colchicine and Nucleotide Sites. <i>Journal of Natural Products</i> , 2018, 81, 494-505.	1.5	15
26	Studies toward the Total Synthesis of the Marine Macrolide Salarin A. <i>Organic Letters</i> , 2018, 20, 7679-7683.	2.4	11
27	The LAT1 inhibitor JPH203 reduces growth of thyroid carcinoma in a fully immunocompetent mouse model. <i>Journal of Experimental and Clinical Cancer Research</i> , 2018, 37, 234.	3.5	72
28	Arylvinylpiperazine Amides, a New Class of Potent Inhibitors Targeting QcrB of <i>Mycobacterium tuberculosis</i> . <i>MBio</i> , 2018, 9, .	1.8	52
29	A fluorescence anisotropy assay to discover and characterize ligands targeting the maytansine site of tubulin. <i>Nature Communications</i> , 2018, 9, 2106.	5.8	41
30	Tumor Targeting with Small Molecule-Drug Conjugates (SMDCs) – Can They be Better than ADCs?. <i>Chimia</i> , 2018, 72, 154.	0.3	5
31	Recent developments in natural product-based drug discovery for tuberculosis. <i>Drug Discovery Today</i> , 2017, 22, 585-591.	3.2	31
32	RNA polymerase motions during promoter melting. <i>Science</i> , 2017, 356, 863-866.	6.0	85
33	The Macrolide Toxin Mycolactone Promotes Bim-Dependent Apoptosis in Buruli Ulcer through Inhibition of mTOR. <i>ACS Chemical Biology</i> , 2017, 12, 1297-1307.	1.6	62
34	Peptide-Membrane Interaction between Targeting and Lysis. <i>ACS Chemical Biology</i> , 2017, 12, 2254-2259.	1.6	12
35	Rational Design of Membrane-Pore-Forming Peptides. <i>Small</i> , 2017, 13, 1701316.	5.2	24
36	Drugs from the Oceans: Marine Natural Products as Leads for Drug Discovery. <i>Chimia</i> , 2017, 71, 646.	0.3	74

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37	The chemistry and biology of mycolactones. <i>Beilstein Journal of Organic Chemistry</i> , 2017, 13, 1596-1660.	1.3	35
38	Zampanolide, a Microtubule-Stabilizing Agent, Is Active in Resistant Cancer Cells and Inhibits Cell Migration. <i>International Journal of Molecular Sciences</i> , 2017, 18, 971.	1.8	24
39	Chemical probes to potently and selectively inhibit endocannabinoid cellular reuptake. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E5006-E5015.	3.3	72
40	On the Chemistry and Biology of the Marine Macrolides Zampanolide and Dactylolide. , 2017, , 555-599.		0
41	Synthesis, Biological Profiling and Determination of the Tubulin-Bound Conformation of 12-Aza-Epothilones (Azathilones). <i>Molecules</i> , 2016, 21, 1010.	1.7	6
42	Total Synthesis and Biological Assessment of Mandelalide. <i>Chemistry - A European Journal</i> , 2016, 22, 1292-1300.	1.7	27
43	Deorphaning the Macromolecular Targets of the Natural Anticancer Compound Dolicolide. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 12408-12411.	7.2	31
44	Deorphaning the Macromolecular Targets of the Natural Anticancer Compound Dolicolide. <i>Angewandte Chemie</i> , 2016, 128, 12596-12599.	1.6	3
45	Actin-binding dolicolide causes premature senescence in p53 wild type cells. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 123-129.	1.4	9
46	Antibody-Mediated Neutralization of the Exotoxin Mycolactone, the Main Virulence Factor Produced by <i>Mycobacterium ulcerans</i> . <i>PLoS Neglected Tropical Diseases</i> , 2016, 10, e0004808.	1.3	38
47	Directed Hydrogenations and an Ireland-Claisen Rearrangement Linked to Evans-Tishchenko Chemistry: The Highly Efficient Total Synthesis of the Marine Cyclodepsipeptide Dolicolide. <i>Chemistry - A European Journal</i> , 2015, 21, 8403-8407.	1.7	11
48	High affinity and covalent-binding microtubule stabilizing agents show activity in chemotherapy-resistant acute myeloid leukemia cells. <i>Cancer Letters</i> , 2015, 368, 97-104.	3.2	12
49	Total Synthesis of the Tiacumicin (Lipiamycin A3/Fidaxomicin) Aglycone. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 1937-1940.	7.2	52
50	Xenicane Natural Products: Biological Activity and Total Synthesis. <i>Current Pharmaceutical Design</i> , 2015, 21, 5467-5488.	0.9	16
51	The Impact of Cyclopropane Configuration on the Biological Activity of Cyclopropyl-Epothilones. <i>ChemMedChem</i> , 2014, 9, 2227-2232.	1.6	13
52	Pyridomycin bridges the NADH- and substrate-binding pockets of the enoyl reductase InhA. <i>Nature Chemical Biology</i> , 2014, 10, 96-98.	3.9	63
53	Structural Basis of Microtubule Stabilization by Laulimalide and Peloruside. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 1621-1625.	7.2	154
54	The cytoskeleton and its interactions with small molecules. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 5038-5039.	1.4	5

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55	A new tubulin-binding site and pharmacophore for microtubule-destabilizing anticancer drugs. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 13817-13821.	3.3	229
56	Molecular Recognition of Epothilones by Microtubules and Tubulin Dimers Revealed by Biochemical and NMR Approaches. ACS Chemical Biology, 2014, 9, 1033-1043.	1.6	30
57	Pharmacological characterization of actin-binding (α)-Doliculide. Bioorganic and Medicinal Chemistry, 2014, 22, 5117-5122.	1.4	14
58	Mycobacterial Toxin Induces Analgesia in Buruli Ulcer by Targeting the Angiotensin Pathways. Cell, 2014, 157, 1565-1576.	13.5	160
59	Methods for Studying Microtubule Binding Site Interactions. Methods in Cell Biology, 2013, 115, 303-325.	0.5	4
60	Synthesis and Biological Activity of 7,8,9-trideoxy- and 7 <i>R</i> -DesTHP-Peloruside A. Chemistry - A European Journal, 2013, 19, 13105-13111.	1.7	14
61	Synthesis and Antimycobacterial Activity of 2,1-dihydropyridomycins. ACS Medicinal Chemistry Letters, 2013, 4, 264-268.	1.3	24
62	Molecular Mechanism of Action of Microtubule-Stabilizing Anticancer Agents. Science, 2013, 339, 587-590.	6.0	436
63	Total Synthesis of the Tubulin Inhibitor WF1360F Based on Macrocyclic Formation through Ring-Closing Alkyne Metathesis. Angewandte Chemie - International Edition, 2013, 52, 5866-5870.	7.2	67
64	Total Synthesis of the Mycobacterial Macrolide Ripostatin B. Chimia, 2013, 67, 227.	0.3	8
65	Structure-Activity Relationship Studies on the Macrolide Exotoxin Mycolactone of Mycobacterium ulcerans. PLoS Neglected Tropical Diseases, 2013, 7, e2143.	1.3	53
66	Total Synthesis of (α)-Zampanolide and Structure-Activity Relationship Studies on (α)-Dactylolide Derivatives. Chemistry - A European Journal, 2012, 18, 16868-16883.	1.7	47
67	Zampanolide, a Potent New Microtubule-Stabilizing Agent, Covalently Reacts with the Taxane Luminal Site in Tubulin α , β -Heterodimers and Microtubules. Chemistry and Biology, 2012, 19, 686-698.	6.2	81
68	Towards a new tuberculosis drug: pyridomycin - nature's isoniazid. EMBO Molecular Medicine, 2012, 4, 1032-1042.	3.3	175
69	Total Synthesis of the Bacterial RNA Polymerase Inhibitor Ripostatin B. Angewandte Chemie - International Edition, 2012, 51, 3405-3409.	7.2	39
70	Kinase Inhibition by Deoxy Analogues of the Resorcylic Lactone L-783277. ACS Medicinal Chemistry Letters, 2011, 2, 22-27.	1.3	24
71	Stereoselective Synthesis of 12,13-Cyclopropyl-Epothilone B and Side-Chain-Modified Variants. Organic Letters, 2011, 13, 1436-1439.	2.4	18
72	Diversity through semisynthesis: the chemistry and biological activity of semisynthetic epothilone derivatives. Molecular Diversity, 2011, 15, 383-399.	2.1	34

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73	A Ring-Closing Metathesis (RCM)-Based Approach to Mycolactones...A/B. Chemistry - A European Journal, 2011, 17, 13017-13031.	1.7	45
74	Natural Products as Leads for Anticancer Drug Discovery: Discovery of New Chemotypes of Microtubule Stabilizers through Reengineering of the Epothilone Scaffold. Chimia, 2010, 64, 8.	0.3	6
75	Synthesis and Biological Activity of New Functionalized Epothilones for Prodrug Design and Tumor Targeting. Chimia, 2010, 64, 136-139.	0.3	1
76	Evaluation of Novel Epothilone Analogues by means of a Common Pharmacophore and a QSAR Pseudoreceptor Model for Taxanes and Epothilones. ChemMedChem, 2010, 5, 35-40.	1.6	26
77	The Binding Mode of Side Chain- and C3-Modified Epothilones to Tubulin. ChemMedChem, 2010, 5, 911-920.	1.6	14
78	Highly Potent Modulation of GABA _A Receptors by Valerenic Acid Derivatives. ChemMedChem, 2010, 5, 678-681.	1.6	12
79	Treasures from the Sea: Discovery and Total Synthesis of Ammosamides. Angewandte Chemie - International Edition, 2010, 49, 6936-6938.	7.2	11
80	Stereoselective Synthesis of a Monocyclic Peloruside A Analogue. Organic Letters, 2010, 12, 1120-1123.	2.4	52
81	Synthesis of (â)-Dactyloide and 13-Desmethylene-(â)-dactyloide and Their Effects on Tubulin. Organic Letters, 2010, 12, 2302-2305.	2.4	60
82	The Laulimalide Family: Total Synthesis and Biological Evaluation of Neolaulimalide, Isolaulimalide, Laulimalide and a Nonnatural Analogue. Chemistry - A European Journal, 2009, 15, 5979-5997.	1.7	57
83	Epothilone Analogues with Benzimidazole and Quinoline Side Chains: Chemical Synthesis, Antiproliferative Activity, and Interactions with Tubulin. Chemistry - A European Journal, 2009, 15, 10144-10157.	1.7	27
84	Differential Effects of Natural Product Microtubule Stabilizers on Microtubule Assembly: Single Agent and Combination Studies with Taxol, Epothilone B, and Discodermolide. ChemBioChem, 2009, 10, 166-175.	1.3	12
85	Making Epothilones Fluoresce: Design, Synthesis, and Biological Characterization of a Fluorescent N12-Aza-Epothilone (Azathilone). ChemBioChem, 2009, 10, 2513-2521.	1.3	11
86	Synthesis and biological evaluation of epothilone A dimeric compounds. Bioorganic and Medicinal Chemistry, 2009, 17, 7435-7440.	1.4	13
87	Synthesis and SAR of C12-C13-oxazoline derivatives of epothilone A. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 3760-3763.	1.0	22
88	GABAA receptors as in vivo substrate for the anxiolytic action of valerenic acid, a major constituent of valerian root extracts. Neuropharmacology, 2009, 56, 174-181.	2.0	173
89	Semisynthetic Derivatives of Epothilones. Progress in the Chemistry of Organic Natural Products, 2009, 90, 135-156.	0.8	3
90	Preclinical Pharmacology and Structure-Activity Studies of Epothilones. Progress in the Chemistry of Organic Natural Products, 2009, 90, 157-220.	0.8	4

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91	Clinical Studies with Epothilones. Progress in the Chemistry of Organic Natural Products, 2009, 90, 221-237.	0.8	1
92	Resorcylic acid lactones as new lead structures for kinase inhibition. Comptes Rendus Chimie, 2008, 11, 1318-1335.	0.2	60
93	Total Synthesis of the Marine Diterpenoid Blumiolide...C. Angewandte Chemie - International Edition, 2008, 47, 10081-10085.	7.2	38
94	Epothilones – A fascinating family of microtubule stabilizing antitumor agents. Comptes Rendus Chimie, 2008, 11, 1336-1368.	0.2	35
95	Synthesis of 12-aza analogs of epothilones and (E)-9,10-dehydroepothilones. Tetrahedron, 2008, 64, 7920-7928.	1.0	10
96	Epothilones as lead structures for new anticancer drugs – pharmacology, fermentation, and structure-activity-relationships. , 2008, 66, 273-334.		5
97	Unraveling a molecular target of macrolides. Nature Chemical Biology, 2008, 4, 388-389.	3.9	7
98	Conformational Preferences of Natural and C3-Modified Epothilones in Aqueous Solution. Journal of Medicinal Chemistry, 2008, 51, 1469-1473.	2.9	49
99	Epothilones as Lead Structures for the Synthesis-Based Discovery of New Chemotypes for Microtubule Stabilization. Accounts of Chemical Research, 2008, 41, 21-31.	7.6	89
100	Total Synthesis of Hypermodified Epothilone Analogs with Potent in Vitro Antitumor Activity. Organic Letters, 2008, 10, 1183-1186.	2.4	29
101	Synthesis and Biological Activity of 12-Aza-Epothilones (Azathilones) – Non-Natural Natural Products with Potent Antiproliferative Activity. Chimia, 2007, 61, 143-146.	0.3	4
102	Synthetic Studies on Mycolactones: Synthesis of the Mycolactone Core – Structure through Ring-Closing Olefin Metathesis. Synlett, 2007, 2007, 0415-0418.	1.0	4
103	The Chemistry and Biology of Epothilones – The Wheel Keeps Turning. ChemMedChem, 2007, 2, 396-423.	1.6	119
104	Chemical Tools from Biology-Oriented Synthesis. Chemistry and Biology, 2007, 14, 347-349.	6.2	4
105	Anticancer drugs from nature – natural products as a unique source of new microtubule-stabilizing agents. Natural Product Reports, 2007, 24, 327-357.	5.2	230
106	Total Synthesis and Biological Assessment of Benzimidazole-Based Analogues of Epothilone A: Ambivalent Effects on Cancer Cell Growth Inhibition. ChemBioChem, 2006, 7, 54-57.	1.3	31
107	Design and Synthesis of 12-Aza-Epothilones (Azathilones) – Non-Natural – Natural Products with Potent Anticancer Activity. Angewandte Chemie - International Edition, 2006, 45, 5880-5885.	7.2	28
108	Structure-Activity Relationships in Side-Chain-Modified Epothilone Analogues – How Important is the Position of the Nitrogen Atom?. ChemMedChem, 2006, 1, 37-40.	1.6	29

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109	Total Synthesis and Biological Assessment of Cyclopropane-Based Epothilone Analogues - Modulation of Drug Efflux through Polarity Adjustments. <i>Synlett</i> , 2006, 2006, 1384-1388.	1.0	6
110	Scaffolds for Microtubule Inhibition through Extensive Modification of the Epothilone Template. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 7469-7473.	7.2	44
111	Patupilone (epothilone B, EPO906) inhibits growth and metastasis of experimental prostate tumors in vivo. <i>Prostate</i> , 2005, 65, 231-240.	1.2	46
112	Recent Developments in the Chemical Biology of Epothilones. <i>Current Pharmaceutical Design</i> , 2005, 11, 1595-1613.	0.9	99
113	Total Synthesis of 26-Fluoro-epothilone B. <i>Synlett</i> , 2004, 2004, 693-697.	1.0	11
114	Synthesis and Biological Evaluation of Furano-Epothilone C. <i>Synlett</i> , 2004, 2004, 1375-1378.	1.0	21
115	Total Synthesis and Biological Evaluation of a C(10)/C(12)-Phenylene-Bridged Analog of Epothilone?D. <i>Chemistry and Biodiversity</i> , 2004, 1, 1771-1784.	1.0	8
116	The merger of natural product synthesis and medicinal chemistry: on the chemistry and chemical biology of epothilones. <i>Organic and Biomolecular Chemistry</i> , 2004, 2, 2137.	1.5	95
117	Natural Product-Based Drug Discovery â€œ Epothilones as Lead Structures for the Discovery of New Anticancer Agents. <i>Chimia</i> , 2004, 58, 686-690.	0.3	20
118	Title is missing!. <i>Angewandte Chemie</i> , 2003, 115, 2615-2619.	1.6	24
119	Design, Synthesis, and Biological Properties of Highly Potent Epothilone B Analogues. <i>Angewandte Chemie</i> , 2003, 115, 3639-3644.	1.6	13
120	The High-Resolution Solution Structure of Epothilone A Bound to Tubulin: An Understanding of the Structureâ€œActivity Relationships for a Powerful Class of Antitumor Agents. <i>Angewandte Chemie - International Edition</i> , 2003, 42, 2511-2515.	7.2	103
121	Design, Synthesis, and Biological Properties of Highly Potent Epothilone B Analogues. <i>Angewandte Chemie - International Edition</i> , 2003, 42, 3515-3520.	7.2	66
122	Epothilone B and its Analogs - A New Family of Anticancer Agents. <i>Mini-Reviews in Medicinal Chemistry</i> , 2003, 3, 149-158.	1.1	96
123	The Total Synthesis and Biological Assessment of trans-Epothilone A. <i>Helvetica Chimica Acta</i> , 2002, 85, 4086-4110.	1.0	76
124	Chemical synthesis and biological evaluation of novel epothilone B and trans-12,13-cyclopropyl epothilone B analogues. <i>Tetrahedron</i> , 2002, 58, 6413-6432.	1.0	57
125	Synthetic and Semisynthetic Analogs of Epothilones: Chemistry and Biological Activity. <i>ACS Symposium Series</i> , 2001, , 112-130.	0.5	8
126	Microtubule-stabilizing agents: a growing class of important anticancer drugs. <i>Current Opinion in Chemical Biology</i> , 2001, 5, 424-431.	2.8	134

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127	Chemical Synthesis and Biological Evaluation of cis- and trans-12,13-Cyclopropyl and 12,13-Cyclobutyl Epothilones and Related Pyridine Side Chain Analogues. <i>Journal of the American Chemical Society</i> , 2001, 123, 9313-9323.	6.6	205
128	Epothilones and their analogues - a new class of promising microtubule inhibitors. <i>Expert Opinion on Therapeutic Patents</i> , 2001, 11, 951-968.	2.4	23
129	Synthesis and Biological Evaluation of Aza-Epothilones. <i>ChemBioChem</i> , 2000, 1, 67-70.	1.3	26
130	Synthesis and biological evaluation of highly potent analogues of epothilones B and D. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2000, 10, 2765-2768.	1.0	64
131	Epothilones and related structures – a new class of microtubule inhibitors with potent in vivo antitumor activity. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2000, 1470, M79-M91.	3.3	120
132	New Anilinophthalazines as Potent and Orally Well Absorbed Inhibitors of the VEGF Receptor Tyrosine Kinases Useful as Antagonists of Tumor-Driven Angiogenesis. <i>Journal of Medicinal Chemistry</i> , 2000, 43, 2310-2323.	2.9	224
133	The Solution Conformation of a Carbocyclic Analog of the Dickerson-Drew Dodecamer: Comparison with its own X-ray Structure and that of the NMR Structure of the Native Counterpart. <i>Journal of Biomolecular Structure and Dynamics</i> , 1998, 16, 547-568.	2.0	23
134	Correlating Structure and Stability of DNA Duplexes with Incorporated 2'-O-Modified RNA Analogues. <i>Biochemistry</i> , 1998, 37, 10626-10634.	1.2	57