

Kuo-Hsiung Lee

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5535906/publications.pdf>

Version: 2024-02-01

357
papers

14,791
citations

22099

59
h-index

33814

99
g-index

362
all docs

362
docs citations

362
times ranked

15172
citing authors

#	ARTICLE	IF	CITATIONS
1	Biology of quinoline and quinazoline alkaloids. <i>The Alkaloids Chemistry and Biology</i> , 2022, 88, 1-47.	0.8	2
2	Development of Novel Dihydrofuro[3,4- <i>d</i>]pyrimidine Derivatives as HIV-1 NNRTIs to Overcome the Highly Resistant Mutant Strains F227L/V106A and K103N/Y181C. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 2458-2470.	2.9	10
3	Molecular hybridization used to design and synthesize neo-tanshinlactone derivatives as PD-1/PD-L1 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2022, 54, 116579.	1.4	3
4	Phyto-sesquiterpene lactones DET and DETD-35 induce ferroptosis in vemurafenib sensitive and resistant melanoma via GPX4 inhibition and metabolic reprogramming. <i>Pharmacological Research</i> , 2022, 178, 106148.	3.1	16
5	Bioactivity inspired C19-diterpenoid alkaloids for overcoming multidrug-resistant cancer. <i>Journal of Natural Medicines</i> , 2022, 76, 796-802.	1.1	2
6	Fluorinated Modification of Neo-Tanshinlactone and Antiproliferative Activity Evaluation. <i>Chemistry of Natural Compounds</i> , 2022, 58, 398-403.	0.2	1
7	Biologically active indolizidine alkaloids. <i>Medicinal Research Reviews</i> , 2021, 41, 928-960.	5.0	46
8	Design, synthesis, and evaluation of â€œdual-siteâ€•binding diarylpyrimidines targeting both NNIBP and the NNRTI adjacent site of the HIV-1 reverse transcriptase. <i>European Journal of Medicinal Chemistry</i> , 2021, 211, 113063.	2.6	15
9	2,4,5-Trisubstituted Pyrimidines as Potent HIV-1 NNRTIs: Rational Design, Synthesis, Activity Evaluation, and Crystallographic Studies. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 4239-4256.	2.9	33
10	Sesquiterpene Lactone Deoxyelephantopin Isolated from <i>Elephantopus scaber</i> and Its Derivative DETD-35 Suppress BRAFV600E Mutant Melanoma Lung Metastasis in Mice. <i>International Journal of Molecular Sciences</i> , 2021, 22, 3226.	1.8	12
11	Design, synthesis, and structure activity relationship analysis of new betulinic acid derivatives as potent HIV inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021, 215, 113287.	2.6	20
12	Selectfluor-Enabled C(sp ³)â€•H Alkoxylation of 3-Methylfuranocoumarins. <i>Journal of Organic Chemistry</i> , 2021, 86, 7864-7871.	1.7	9
13	Hyperdioxanes, dibenzo-1,4-dioxane derivatives from the roots of <i>Hypericum ascyron</i> . <i>Journal of Natural Medicines</i> , 2021, 75, 907-914.	1.1	4
14	Identification of anti-HIV macrocyclic daphnane orthoesters from <i>Wikstroemia ligustrina</i> by LCâ€•MS analysis and phytochemical investigation. <i>Journal of Natural Medicines</i> , 2021, 75, 1058-1066.	1.1	9
15	Design and Synthesis of Quinolizidine Derivatives as Influenza Virus and HIV-1 Inhibitors. <i>Current Medicinal Chemistry</i> , 2021, 28, 4995-5003.	1.2	4
16	LC-MS Identification, Isolation, and Structural Elucidation of Anti-HIV Tigliane Diterpenoids from <i>Wikstroemia lamatsoensis</i> . <i>Journal of Natural Products</i> , 2021, 84, 2366-2373.	1.5	10
17	New phorbol ester derivatives as potent anti-HIV agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 50, 128319.	1.0	5
18	Discovery of potent and selective Cdc25 phosphatase inhibitors via rapid assembly and in situ screening of Quinonoid-focused libraries. <i>Bioorganic Chemistry</i> , 2021, 115, 105254.	2.0	12

#	ARTICLE	IF	CITATIONS
19	Design, synthesis, and antiviral activity of phenylalanine derivatives as HIV-1 capsid inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2021, 48, 116414.	1.4	4
20	Identification of 3, 4-disubstituted pyridine derivatives as novel CDK8 inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021, 223, 113634.	2.6	4
21	Diosgenin Derivatives as Potential Antitumor Agents: Synthesis, Cytotoxicity, and Mechanism of Action. <i>Journal of Natural Products</i> , 2021, 84, 616-629.	1.5	24
22	Synthesis and <i>in vitro</i> anticancer activities of biotinylated derivatives of glaucocalyxin A and oridonin. <i>Journal of Asian Natural Products Research</i> , 2021, 23, 703-711.	0.7	2
23	Eleven new C19-diterpenoid alkaloids from <i>Delphinium elatum</i> cv. Pacific Giant. <i>Journal of Natural Medicines</i> , 2021, , 1.	1.1	1
24	Lead Optimization: Synthesis and Biological Evaluation of PBT-1 Derivatives as Novel Antitumor Agents. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 1948-1954.	1.3	3
25	Design and synthesis of novel 7-[(<i>N</i> -substituted-thioureidopiperazinyl)-methyl]-camptothecin derivatives as potential cytotoxic agents. <i>Natural Product Research</i> , 2020, 34, 2022-2029.	1.0	5
26	Kalshiolin A, new lignan from <i>Kalimeris shimadai</i> . <i>Journal of Asian Natural Products Research</i> , 2020, 22, 489-495.	0.7	9
27	Cytotoxic diterpenoid alkaloid from <i>Aconitum japonicum</i> subsp. <i>subcuneatum</i> . <i>Journal of Natural Medicines</i> , 2020, 74, 83-89.	1.1	8
28	Daphneodorins A-C, Anti-HIV Gnidimacrin Related Macrocyclic Daphnane Orthoesters from <i>Daphne odora</i> . <i>Organic Letters</i> , 2020, 22, 11-15.	2.4	30
29	Scaffold Hopping-Driven Optimization of 4-(Quinazolin-4-yl)-3,4-dihydroquinoxalin-2(1 <i>H</i>)-ones as Novel Tubulin Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 83-89.	1.3	12
30	Rapid Recognition and Targeted Isolation of Anti-HIV Daphnane Diterpenes from <i>Daphne genkwa</i> Guided by UPLC-MS ⁿ . <i>Journal of Natural Products</i> , 2020, 83, 134-141.	1.5	18
31	Carbazole Alkaloids from <i>Clausena anisum-olens</i> : Isolation, Characterization, and Anti-HIV Evaluation. <i>Molecules</i> , 2020, 25, 99.	1.7	20
32	Design, synthesis and antineoplastic activity of novel 20(S)-acylthiourea derivatives of camptothecin. <i>European Journal of Medicinal Chemistry</i> , 2020, 187, 111971.	2.6	13
33	<i>In Silico</i> De Novo Curcuminoid Derivatives From the Compound Library of Natural Products Research Laboratories Inhibit COVID-19 3CL ^{pro} Activity. <i>Natural Product Communications</i> , 2020, 15, 1934578X2095326.	0.2	4
34	Isolation, Structural Elucidation, and Anti-HIV Activity of Daphnane Diterpenoids from <i>Daphne odora</i> . <i>Journal of Natural Products</i> , 2020, 83, 3270-3277.	1.5	16
35	Recent advances in natural anti-HIV triterpenoids and analogs. <i>Medicinal Research Reviews</i> , 2020, 40, 2339-2385.	5.0	23
36	Design and synthesis of benzylidenecyclohexenones as TrxR inhibitors displaying high anticancer activity and inducing ROS, apoptosis, and autophagy. <i>European Journal of Medicinal Chemistry</i> , 2020, 204, 112610.	2.6	19

#	ARTICLE	IF	CITATIONS
37	Development of anti-influenza agents from natural products. <i>Medicinal Research Reviews</i> , 2020, 40, 2290-2338.	5.0	48
38	New Seco-DSP derivatives as potent chemosensitizers. <i>European Journal of Medicinal Chemistry</i> , 2020, 204, 112555.	2.6	3
39	Fsp3: A new parameter for drug-likeness. <i>Drug Discovery Today</i> , 2020, 25, 1839-1845.	3.2	156
40	Biologically active isoquinoline alkaloids covering 2014–2018. <i>Medicinal Research Reviews</i> , 2020, 40, 2212-2289.	5.0	107
41	Palisanines A–E, 3,4-Methylenedioxyquinoline Alkaloids Fused with a Phenyl-14-oxabicyclo[3.2.1]octane Unit from <i>Melochia umbellata</i> var. <i>deglabrata</i> . <i>Journal of Natural Products</i> , 2020, 83, 2931-2939.	1.5	5
42	Novel Betulinic Acid–Nucleoside Hybrids with Potent Anti-HIV Activity. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 2290-2293.	1.3	18
43	Anti-HIV Tigliane Diterpenoids from <i>Wikstroemia scytophylla</i> . <i>Journal of Natural Products</i> , 2020, 83, 3584-3590.	1.5	17
44	Nanocarrier-mediated immunogenic chemotherapy for triple negative breast cancer. <i>Journal of Controlled Release</i> , 2020, 323, 431-441.	4.8	39
45	Discovery of potential dual-target prodrugs of HIV-1 reverse transcriptase and nucleocapsid protein 7. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127287.	1.0	3
46	In situ click chemistry-based rapid discovery of novel HIV-1 NNRTIs by exploiting the hydrophobic channel and tolerant regions of NNIBP. <i>European Journal of Medicinal Chemistry</i> , 2020, 193, 112237.	2.6	23
47	Design, synthesis and structure-activity relationships of 4-phenyl-1H-1,2,3-triazole phenylalanine derivatives as novel HIV-1 capsid inhibitors with promising antiviral activities. <i>European Journal of Medicinal Chemistry</i> , 2020, 190, 112085.	2.6	65
48	Synthesis, Anticancer Activity, and Preliminary Pharmacokinetic Evaluation of 4,4-Disubstituted Curcuminoid 2,2-bis(Hydroxymethyl)Propionate Derivatives. <i>Molecules</i> , 2020, 25, 479.	1.7	11
49	Cucurbitane-Type Triterpenoids from the Vines of <i>Momordica charantia</i> and Their Anti-inflammatory Activities. <i>Journal of Natural Products</i> , 2020, 83, 1400-1408.	1.5	15
50	Anti-HIV tigliane diterpenoids from <i>Reutealis trisperma</i> . <i>Phytochemistry</i> , 2020, 174, 112360.	1.4	15
51	Design, Synthesis, and Mechanism Study of Benzenesulfonamide-Containing Phenylalanine Derivatives as Novel HIV-1 Capsid Inhibitors with Improved Antiviral Activities. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 4790-4810.	2.9	41
52	Talarolactone A, an Isocoumarin Derivative Fused with Dihydrothiophene with Selective Antimigratory Activity from the Endolichenic Fungus <i>Talaromyces</i> sp.. <i>Journal of Natural Products</i> , 2020, 83, 1716-1720.	1.5	12
53	Novel potent antiplatelet thrombotic agent derived from biguanide for ischemic stroke. <i>European Journal of Medicinal Chemistry</i> , 2020, 200, 112462.	2.6	11
54	Wilforine resensitizes multidrug resistant cancer cells via competitive inhibition of P-glycoprotein. <i>Phytomedicine</i> , 2020, 71, 153239.	2.3	21

#	ARTICLE	IF	CITATIONS
55	Danazol mediates collateral sensitivity via STAT3/Myc related pathway in multidrug-resistant cancer cells. <i>Scientific Reports</i> , 2019, 9, 11628.	1.6	7
56	Spiro[3.5]nonenyl Meroterpenoid Lactones, Cryptolaevilactones Gâ€“L, an Ionone Derivative, and Total Synthesis of Cryptolaevilactone M from <i>Cryptocarya laevigata</i> . <i>Journal of Natural Products</i> , 2019, 82, 2368-2378.	1.5	9
57	Synthesis and Structureâ€“Activity Relationship Correlations of Gnidimacrin Derivatives as Potent HIV-1 Inhibitors and HIV Latency Reversing Agents. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 6958-6971.	2.9	17
58	Discovery of piperidine-substituted thiazolo[5,4-d]pyrimidine derivatives as potent and orally bioavailable HIV-1 non-nucleoside reverse transcriptase inhibitors. <i>Communications Chemistry</i> , 2019, 2, .	2.0	24
59	Isolation, Structure Elucidation, and Antiproliferative Activity of Butanolides and Lignan Glycosides from the Fruit of <i>Hernandia nymphaeifolia</i> . <i>Molecules</i> , 2019, 24, 4005.	1.7	8
60	Identification of highly potent and selective Cdc25 protein phosphatases inhibitors from miniaturization click-chemistry-based combinatorial libraries. <i>European Journal of Medicinal Chemistry</i> , 2019, 183, 111696.	2.6	26
61	Prenylated Acetophloroglucinol Dimers from <i>Acronychia trifoliolata</i> : Structure Elucidation and Total Synthesis. <i>Journal of Natural Products</i> , 2019, 82, 2852-2858.	1.5	9
62	Recent applications of click chemistry in drug discovery. <i>Expert Opinion on Drug Discovery</i> , 2019, 14, 779-789.	2.5	151
63	Structureâ€“activity relationships and evaluation of esterified diterpenoid alkaloid derivatives as antiproliferative agents. <i>Journal of Natural Medicines</i> , 2019, 73, 789-799.	1.1	13
64	Potent Anti-HIV Ingenane Diterpenoids from <i>Euphorbia ebracteolata</i> . <i>Journal of Natural Products</i> , 2019, 82, 1587-1592.	1.5	30
65	Ochrocephalamines Bâ€“D, Three Alkaloids from <i>Oxytropis ochrocephala</i> Bunge. <i>Organic Letters</i> , 2019, 21, 5051-5054.	2.4	12
66	Design, synthesis and evaluation of antiproliferative activity of fluorinated betulinic acid. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 2871-2882.	1.4	9
67	One-step templated synthesis of chiral organometallic salicyloxazoline complexes. <i>BMC Chemistry</i> , 2019, 13, 51.	1.6	4
68	New Dammarane-type Saponins from <i>Gynostemma pentaphyllum</i> . <i>Molecules</i> , 2019, 24, 1375.	1.7	9
69	Antiproliferative <i>Aspidosperma</i> -Type Monoterpenoid Indole Alkaloids from <i>Bousigonia mekongensis</i> Inhibit Tubulin Polymerization. <i>Molecules</i> , 2019, 24, 1256.	1.7	15
70	Four new triterpenoids from the bark of <i>Euonymus alatus forma ciliato-dentatus</i> . <i>Phytochemistry Letters</i> , 2019, 31, 140-146.	0.6	11
71	Discovery of an Oleanolic Acid/Hederageninâ€“Nitric Oxide Donor Hybrid as an EGFR Tyrosine Kinase Inhibitor for Non-Small-Cell Lung Cancer. <i>Journal of Natural Products</i> , 2019, 82, 3065-3073.	1.5	32
72	Kalshinoids Aâ€“F, Anti-inflammatory Sesquiterpenes from <i>Kalimeris shimadae</i> . <i>Journal of Natural Products</i> , 2019, 82, 3372-3378.	1.5	22

#	ARTICLE	IF	CITATIONS
73	Discovery of novel 1,4-disubstituted 1,2,3-triazole phenylalanine derivatives as HIV-1 capsid inhibitors. <i>RSC Advances</i> , 2019, 9, 28961-28986.	1.7	42
74	Tenulin and isotenulin inhibit P-glycoprotein function and overcome multidrug resistance in cancer cells. <i>Phytomedicine</i> , 2019, 53, 252-262.	2.3	29
75	Design, synthesis, and biologic evaluation of novel galloyl derivatives as HIV-1 RNase H inhibitors. <i>Chemical Biology and Drug Design</i> , 2019, 93, 582-589.	1.5	14
76	Contemporary medicinal-chemistry strategies for the discovery of selective butyrylcholinesterase inhibitors. <i>Drug Discovery Today</i> , 2019, 24, 629-635.	3.2	35
77	Seco-4-methyl-DCK derivatives as potent chemosensitizers. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 28-31.	1.0	3
78	New transformation pathway and cytotoxic derivatives from the acid hydrolysis of timosaponin B III. <i>Natural Product Research</i> , 2019, 33, 2755-2761.	1.0	3
79	Identification of Dihydrofuro[3,4-d]pyrimidine Derivatives as Novel HIV-1 Non-Nucleoside Reverse Transcriptase Inhibitors with Promising Antiviral Activities and Desirable Physicochemical Properties. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 1484-1501.	2.9	70
80	Discovery and synthesis of novel beesioside I derivatives with potent anti-HIV activity. <i>European Journal of Medicinal Chemistry</i> , 2019, 166, 159-166.	2.6	8
81	Discovery of potent HIV-1 non-nucleoside reverse transcriptase inhibitors by exploring the structure-activity relationship of solvent-exposed regions I. <i>Chemical Biology and Drug Design</i> , 2019, 93, 430-437.	1.5	13
82	Biologically active quinoline and quinazoline alkaloids part II. <i>Medicinal Research Reviews</i> , 2018, 38, 1614-1660.	5.0	134
83	Discovery of Novel Diarylpyrimidine Derivatives as Potent HIV-1 NNRTIs Targeting the NNRTI Adjacent Binding Site. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 334-338.	1.3	32
84	Further Exploring Solvent-Exposed Tolerant Regions of Allosteric Binding Pocket for Novel HIV-1 NNRTIs Discovery. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 370-375.	1.3	28
85	Secondary Metabolites, Monoterpene Polyketides Containing a Spiro[3.5]nonane from <i>Cryptocarya laevigata</i> . <i>Organic Letters</i> , 2018, 20, 2282-2286.	2.4	13
86	Salvisertin A, a New Hexacyclic Triterpenoid, and Other Bioactive Terpenes from <i>Salvia deserta</i> Root. <i>Chemistry and Biodiversity</i> , 2018, 15, e1800019.	1.0	9
87	Mechanism of action of cytotoxic compounds from the seeds of <i>Euphorbia lathyris</i> . <i>Phytomedicine</i> , 2018, 41, 62-66.	2.3	24
88	Elimination of HIV-1 Latently Infected Cells by Gnidimacrin and a Selective HDAC Inhibitor. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 268-273.	1.3	23
89	Evaluation of in vitro/in vivo anti-diabetic effects and identification of compounds from <i>Physalis alkekengi</i> . <i>FITOTERAPIA</i> , 2018, 127, 129-137.	1.1	27
90	Lewis acid-mediated defluorinative [3+2] cycloaddition/aromatization cascade of 2,2-difluoroethanol systems with nitriles. <i>Advanced Synthesis and Catalysis</i> , 2018, 360, 1605-1610.	2.1	19

#	ARTICLE	IF	CITATIONS
91	Structure-activity relationships of cryptopleurine analogs with E-ring modifications as anti-hepatitis C virus agents. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 630-636.	1.4	4
92	Corymbulosins A-W, Cytotoxic Clerodane Diterpenes from the Bark of <i>Laetia corymbulosa</i> . <i>Journal of Organic Chemistry</i> , 2018, 83, 951-963.	1.7	12
93	Four new C ₁₉ -diterpenoid alkaloids from <i>Delphinium elatum</i> . <i>Phytochemistry Letters</i> , 2018, 24, 6-9.	0.6	10
94	Taburnaemines A-I, Cytotoxic Vobasinyllboga-Type Bisindole Alkaloids from <i>Tabernaemontana corymbosa</i> . <i>Journal of Natural Products</i> , 2018, 81, 562-571.	1.5	39
95	SAR study on <i>N</i> ² , <i>N</i> ⁴ -disubstituted pyrimidine-2,4-diamines as effective CDK2/CDK9 inhibitors and antiproliferative agents. <i>RSC Advances</i> , 2018, 8, 11871-11885.	1.7	9
96	Targeting the entrance channel of NNIBP: Discovery of diarylnicotinamide 1,4-disubstituted 1,2,3-triazoles as novel HIV-1 NNRTIs with high potency against wild-type and E138K mutant virus. <i>European Journal of Medicinal Chemistry</i> , 2018, 151, 339-350.	2.6	68
97	Four new diterpenoid alkaloids from <i>Aconitum japonicum</i> subsp. <i>subcuneatum</i> . <i>Journal of Natural Medicines</i> , 2018, 72, 230-237.	1.1	17
98	Biologically active quinoline and quinazoline alkaloids part I. <i>Medicinal Research Reviews</i> , 2018, 38, 775-828.	5.0	262
99	Cytotoxicity, Hemolytic Toxicity, and Mechanism of Action of Pulsatilla Saponin D and Its Synthetic Derivatives. <i>Journal of Natural Products</i> , 2018, 81, 465-474.	1.5	35
100	Discovery of phenylalanine derivatives as potent HIV-1 capsid inhibitors from click chemistry-based compound library. <i>European Journal of Medicinal Chemistry</i> , 2018, 158, 478-492.	2.6	51
101	Development of novel amino-quinoline-5,8-dione derivatives as NAD(P)H:quinone oxidoreductase 1 (NQO1) inhibitors with potent antiproliferative activities. <i>European Journal of Medicinal Chemistry</i> , 2018, 154, 199-209.	2.6	27
102	Antidiabetic potential of the ethyl acetate extract of <i>Physalis alkekengi</i> and chemical constituents identified by HPLC-ESI-QTOF-MS. <i>Journal of Ethnopharmacology</i> , 2018, 225, 202-210.	2.0	22
103	Design, synthesis, and antiviral evaluation of novel hydrazone-substituted thiophene[3,2-d]pyrimidine derivatives as potent human immunodeficiency virus-1 inhibitors. <i>Chemical Biology and Drug Design</i> , 2018, 92, 2009-2021.	1.5	16
104	Identification, structural modification, and dichotomous effects on human immunodeficiency virus type 1 (HIV-1) replication of ingenane esters from <i>Euphorbia kansui</i> . <i>European Journal of Medicinal Chemistry</i> , 2018, 156, 618-627.	2.6	36
105	Chemical Structures and Biological Activities of Limonoids from the Genus <i>Swietenia</i> (Meliaceae). <i>Molecules</i> , 2018, 23, 1588.	1.7	19
106	Kleinhospitine E and Cycloartane Triterpenoids from <i>Kleinhovia hospita</i> . <i>Journal of Natural Products</i> , 2018, 81, 1619-1627.	1.5	17
107	Antiproliferative Alkaloids from <i>Alangium longiflorum</i> , an Endangered Tropical Plant Species. <i>Journal of Natural Products</i> , 2018, 81, 1884-1891.	1.5	10
108	Synthesis and antitumor activity of bis(hydroxymethyl)propionate analogs of pterostilbene in cisplatin-resistant human oral cancer cells. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 3909-3916.	1.4	10

#	ARTICLE	IF	CITATIONS
109	($\hat{\alpha}$)-Neocaryachine, an Antiproliferative Pavine Alkaloid from <i>Cryptocarya laevigata</i> , Induces DNA Double-Strand Breaks. <i>Journal of Natural Products</i> , 2017, 80, 220-224.	1.5	25
110	<i>Piper sarmentosum</i> Roxb. produces antidepressant-like effects in rodents, associated with activation of the CREB-BDNF-ERK signaling pathway and reversal of HPA axis hyperactivity. <i>Journal of Ethnopharmacology</i> , 2017, 199, 9-19.	2.0	25
111	Synthesis and biological evaluation of chalcone, dihydrochalcone, and 1,3-diarylpropane analogs as anti-inflammatory agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 1547-1550.	1.0	36
112	Design, synthesis and potent cytotoxic activity of novel 7-(N)-Tj ETQqO O O rgBT /Overlock 10 Tf 50 627 Td (-(substituted-sulfonyl)piperazine) derivatives. <i>Chemistry Letters</i> , 2017, 27, 1750-1753.	1.0	18
113	Xanthohumol isolated from <i>Humulus lupulus</i> prevents thrombosis without increased bleeding risk by inhibiting platelet activation and mtDNA release. <i>Free Radical Biology and Medicine</i> , 2017, 108, 247-257.	1.3	35
114	Vitepyrroloids A-D, 2-Cyanopyrrole-Containing Labdane Diterpenoid Alkaloids from the Leaves of <i>Vitex trifolia</i> . <i>Journal of Natural Products</i> , 2017, 80, 1679-1683.	1.5	22
115	Drug-like property-driven optimization of 4-substituted 1,5-diarylanilines as potent HIV-1 non-nucleoside reverse transcriptase inhibitors against rilpivirine-resistant mutant virus. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 2788-2792.	1.0	2
116	Design, synthesis and biological evaluation of novel indolin-2-ones as potent anticancer compounds. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 3326-3331.	1.0	17
117	New bis(hydroxymethyl) alkanolate curcuminoid derivatives exhibit activity against triple-negative breast cancer <i>in vitro</i> and <i>in vivo</i> . <i>European Journal of Medicinal Chemistry</i> , 2017, 131, 141-151.	2.6	23
118	Corymbulosins H, 2-Hydroxy- and 2-Oxo-clerodane Diterpenes from the Bark of <i>Laetia corymbulosa</i> . <i>Journal of Natural Products</i> , 2017, 80, 1065-1072.	1.5	11
119	Design, synthesis and structure-activity relationships of ($\hat{\alpha}$)-isochailulactone derivatives. <i>MedChemComm</i> , 2017, 8, 2040-2049.	3.5	4
120	Design, semisynthesis and potent cytotoxic activity of novel 10-fluorocamptothecin derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 4694-4697.	1.0	18
121	Tylophorine Analogs Allosterically Regulates Heat Shock Cognate Protein 70 And Inhibits Hepatitis C Virus Replication. <i>Scientific Reports</i> , 2017, 7, 10037.	1.6	16
122	Tabercorymines A and B, Two Vobasinyl-Ibogan-Type Bisindole Alkaloids from <i>Tabernaemontana corymbosa</i> . <i>Organic Letters</i> , 2017, 19, 4964-4967.	2.4	46
123	Design, synthesis, and cytotoxic activity of novel 7-substituted camptothecin derivatives incorporating piperazinyl-sulfonylamidine moieties. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 3959-3962.	1.0	9
124	Total Synthesis of (+)-Medicarpin. <i>Journal of Natural Products</i> , 2017, 80, 3284-3288.	1.5	23
125	In Vivo and Mechanistic Studies on Antitumor Lead 7-Methoxy-4-(2-methylquinazolin-4-yl)-3,4-dihydroquinoxalin-2(1H)-one and Its Modification as a Novel Class of Tubulin-Binding Tumor-Vascular Disrupting Agents. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 5586-5598.	2.9	26
126	Bioactive chemical constituents from the root bark of <i>Morus australis</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 309-313.	1.0	19

#	ARTICLE	IF	CITATIONS
127	Phytoagent Deoxyelephantopin and Its Derivative Inhibit Triple Negative Breast Cancer Cell Activity through ROS-Mediated Exosomal Activity and Protein Functions. <i>Frontiers in Pharmacology</i> , 2017, 8, 398.	1.6	23
128	Phytoagent deoxyelephantopin derivative inhibits triple negative breast cancer cell activity by inducing oxidative stress-mediated paraptosis-like cell death. <i>Oncotarget</i> , 2017, 8, 56942-56958.	0.8	27
129	Synthesis, biological evaluation, and physicochemical property assessment of 4-substituted 2-phenylaminoquinazolines as <i>Mer</i> tyrosine kinase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3083-3092.	1.4	5
130	Novel sesquiterpene lactone analogues as potent anti-breast cancer agents. <i>Molecular Oncology</i> , 2016, 10, 921-937.	2.1	30
131	Novel HIV-1 Non-nucleoside Reverse Transcriptase Inhibitor Agents: Optimization of Diarylanilines with High Potency against Wild-Type and Rilpivirine-Resistant E138K Mutant Virus. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 3689-3704.	2.9	31
132	A Novel Plant Sesquiterpene Lactone Derivative, DETD-35, Suppresses BRAFV600E Mutant Melanoma Growth and Overcomes Acquired Vemurafenib Resistance in Mice. <i>Molecular Cancer Therapeutics</i> , 2016, 15, 1163-1176.	1.9	19
133	Ochrocephalamine A, a new quinolizidine alkaloid from <i>Oxytropis ochrocephala</i> Bunge. <i>Tetrahedron Letters</i> , 2016, 57, 5047-5049.	0.7	11
134	Incorporation of Privileged Structures into Bevirimat Can Improve Activity against Wild-Type and Bevirimat-Resistant HIV-1. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 9262-9268.	2.9	38
135	Triethylated chromones with substituted naphthalenes as tubulin inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 6048-6057.	1.4	15
136	Strategies for the Optimization of Natural Leads to Anticancer Drugs or Drug Candidates. <i>Medicinal Research Reviews</i> , 2016, 36, 32-91.	5.0	121
137	Euphomilones A and B, ent-Rosane Diterpenoids with 7/5/6 and 5/7/6 Skeletons from <i>Euphorbia milii</i> . <i>Organic Letters</i> , 2016, 18, 6132-6135.	2.4	38
138	Clerodane diterpenes: sources, structures, and biological activities. <i>Natural Product Reports</i> , 2016, 33, 1166-1226.	5.2	175
139	Acetophenone Monomers from <i>Acronychia trifoliolata</i> . <i>Journal of Natural Products</i> , 2016, 79, 2883-2889.	1.5	15
140	Total Synthesis and in Vitro Anti-Tumor-Promoting Activities of Racemic Acetophenone Monomers from <i>Acronychia trifoliolata</i> . <i>Journal of Natural Products</i> , 2016, 79, 2890-2897.	1.5	10
141	Metformin Uniquely Prevents Thrombosis by Inhibiting Platelet Activation and mtDNA Release. <i>Scientific Reports</i> , 2016, 6, 36222.	1.6	91
142	(+)-(14 <i>E</i>)-14-Ethylmatridin-15-one, a New Quinolizidine Alkaloid from the Poisonous Plant <i>Oxytropis ochrocephala</i> . <i>Helvetica Chimica Acta</i> , 2016, 99, 225-227.	1.0	6
143	Aloperine and Its Derivatives as a New Class of HIV-1 Entry Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2016, 7, 240-244.	1.3	61
144	Carolignans from the Aerial Parts of <i>Euphorbia sikkimensis</i> and Their Anti-HIV Activity. <i>Journal of Natural Products</i> , 2016, 79, 578-583.	1.5	35

#	ARTICLE	IF	CITATIONS
145	Fluorinated betulinic acid derivatives and evaluation of their anti-HIV activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 68-71.	1.0	32
146	The application of NCTS (N-cyano-N-phenyl-p-toluenesulfonamide) in palladium-catalyzed cyanation of arenediazonium tetrafluoroborates and aryl halides. <i>Tetrahedron Letters</i> , 2016, 57, 1205-1209.	0.7	26
147	Anti-inflammatory neolignans from the roots of <i>Magnolia officinalis</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 1439-1445.	1.4	17
148	Design, synthesis, cytotoxic activity and molecular docking studies of new 20(S)-sulfonylamidine camptothecin derivatives. <i>European Journal of Medicinal Chemistry</i> , 2016, 115, 109-120.	2.6	28
149	Synthesis and structure-activity relationship studies of novel 3,9-substituted $\hat{\pm}$ -carboline derivatives with high cytotoxic activity against colorectal cancer cells. <i>European Journal of Medicinal Chemistry</i> , 2016, 110, 98-114.	2.6	16
150	Alkaloids from <i>Oxytropis ochrocephala</i> and antiproliferative activity of sophoridine derivatives against cancer cell lines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 1495-1497.	1.0	24
151	Flavonoids Isolated from Heat-Processed <i>Epimedium koreanum</i> and Their Anti-HIV Activities. <i>Helvetica Chimica Acta</i> , 2015, 98, 1177-1187.	1.0	14
152	A novel derivative of betulinic acid, SYK023, suppresses lung cancer growth and malignancy. <i>Oncotarget</i> , 2015, 6, 13671-13687.	0.8	26
153	Development of a Novel Class of Tubulin Inhibitor from Desmosdumotin B with a Hydroxylated Bicyclic B-Ring. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 2378-2389.	2.9	60
154	Design and synthesis of novel PEG-conjugated 20(S)-camptothecin sulfonylamidine derivatives with potent in vitro antitumor activity via Cu-catalyzed three-component reaction. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 2690-2693.	1.0	35
155	Stelleralides J and Anti-HIV Daphnane Diterpenes from <i>Stellera chamaejasme</i> . <i>Journal of Natural Products</i> , 2015, 78, 2712-2718.	1.5	38
156	Selective cytotoxic eremophilane-type sesquiterpenes from <i>Penicillium citreonigrum</i> . <i>Journal of Asian Natural Products Research</i> , 2015, 17, 1239-1244.	0.7	11
157	Novel curcumin analogs to overcome EGFR-TKI lung adenocarcinoma drug resistance and reduce EGFR-TKI-induced GI adverse effects. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 1507-1514.	1.4	28
158	neo-Clerodane Diterpenoids from <i>Scutellaria barbata</i> with Activity against Epstein-Barr Virus Lytic Replication. <i>Journal of Natural Products</i> , 2015, 78, 500-509.	1.5	42
159	Phenolic Diterpenoid Derivatives as Anti-Influenza A Virus Agents. <i>ACS Medicinal Chemistry Letters</i> , 2015, 6, 355-358.	1.3	19
160	Evaluation of Aconitum diterpenoid alkaloids as antiproliferative agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 1525-1531.	1.0	37
161	3,5-Diarylpyrazole Derivatives Obtained by Ammonolysis of the Total Flavonoids from <i>Chrysanthemum indicum</i> Extract Show Potential for the Treatment of Alzheimer's Disease. <i>Journal of Natural Products</i> , 2015, 78, 1593-1599.	1.5	33
162	Perspectives on Biologically Active Camptothecin Derivatives. <i>Medicinal Research Reviews</i> , 2015, 35, 753-789.	5.0	156

#	ARTICLE	IF	CITATIONS
163	Gnidimacrin, a Potent Anti-HIV Diterpene, Can Eliminate Latent HIV-1 Ex Vivo by Activation of Protein Kinase C β . <i>Journal of Medicinal Chemistry</i> , 2015, 58, 8638-8646.	2.9	35
164	Optimization of N-aryl-6-methoxy-1,2,3,4-tetrahydroquinolines as tubulin polymerization inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5740-5747.	1.4	21
165	Total synthesis of cordatanine, structural reassignment of drymaritin, and anti-inflammatory activity of synthetic precursors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 3822-3824.	1.0	7
166	Discovery of novel 5-fluoro-N ² ,N ⁴ -diphenylpyrimidine-2,4-diamines as potent inhibitors against CDK2 and CDK9. <i>MedChemComm</i> , 2015, 6, 444-454.	3.5	8
167	Design and synthesis of new 2-arylnaphthyridin-4-ones as potent antitumor agents targeting tumorigenic cell lines. <i>European Journal of Medicinal Chemistry</i> , 2015, 90, 775-787.	2.6	3
168	Synthesis and SAR studies of novel 6,7,8-substituted 4-substituted benzyloxyquinolin-2(1H)-one derivatives for anticancer activity. <i>British Journal of Pharmacology</i> , 2015, 172, 1195-1221.	2.7	17
169	Recent Progress on Modified Podophyllotoxin Analogs as Potent Antitumor Agents. <i>Medicinal Research Reviews</i> , 2015, 35, 1-62.	5.0	106
170	eIF4E binding protein 1 expression is associated with clinical survival outcomes in colorectal cancer. <i>Oncotarget</i> , 2015, 6, 24092-24104.	0.8	16
171	Cytotoxic cardiac glycosides and coumarins from <i>Antiaris toxicaria</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 1889-1898.	1.4	27
172	Discovery of novel antitumor dibenzocyclooctatetraene derivatives and related biphenyls as potent inhibitors of NF- κ B signaling pathway. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 325-333.	1.4	4
173	A-ring modified betulinic acid derivatives as potent cancer preventive agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 1005-1008.	1.0	8
174	Design, synthesis and cytotoxic activity of novel sulfonylurea derivatives of podophyllotoxin. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 204-210.	1.4	22
175	Synthesis of novel spin-labeled derivatives of 5-FU as potential antineoplastic agents. <i>Medicinal Chemistry Research</i> , 2014, 23, 3269-3273.	1.1	7
176	Design and synthesis of novel spin-labeled camptothecin derivatives as potent cytotoxic agents. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 6453-6458.	1.4	16
177	Design, synthesis, crystal structure, bioactivity, and molecular docking studies of novel sulfonylamidine-derived neonicotinoid analogs. <i>Medicinal Chemistry Research</i> , 2014, 23, 5043-5057.	1.1	13
178	Total synthesis of plagiochin G and derivatives as potential cancer chemopreventive agents. <i>Tetrahedron Letters</i> , 2014, 55, 6500-6503.	0.7	9
179	In vitro anti-inflammatory effects of diterpenoids and sesquiterpenoids from traditional Chinese medicine <i>Siegesbeckia pubescens</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 3944-3947.	1.0	18
180	Optimization of the Antiviral Potency and Lipophilicity of Halogenated 2,6-Diarylpyridinamines as a Novel Class of HIV-1 NNRTIS. <i>ChemMedChem</i> , 2014, 9, 1546-1555.	1.6	12

#	ARTICLE	IF	CITATIONS
181	Identification and Synthesis of Quinolizidines with Anti-Influenza A Virus Activity. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 942-946.	1.3	50
182	The Antitumor Agent PBT-1 Directly Targets HSP90 and hnRNP A2/B1 and Inhibits Lung Adenocarcinoma Growth and Metastasis. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 677-685.	2.9	24
183	Multidrug resistance-selective antiproliferative activity of Piper amide alkaloids and synthetic analogues. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 4818-4821.	1.0	15
184	Design, Synthesis, Mechanisms of Action, and Toxicity of Novel 20(S)-Sulfonylamidine Derivatives of Camptothecin as Potent Antitumor Agents. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 6008-6018.	2.9	66
185	Phytotoxic cis-clerodane diterpenoids from the Chinese liverwort <i>Scapania stephanii</i> . <i>Phytochemistry</i> , 2014, 105, 85-91.	1.4	10
186	MJ-66 induces malignant glioma cells G2/M phase arrest and mitotic catastrophe through regulation of cyclin B1/Cdk1 complex. <i>Neuropharmacology</i> , 2014, 86, 219-227.	2.0	21
187	Design and synthesis of new 7-(N-substituted-methyl)-camptothecin derivatives as potent cytotoxic agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 3850-3853.	1.0	14
188	Physicochemical property-driven optimization of diarylaniline compounds as potent HIV-1 non-nucleoside reverse transcriptase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 3719-3723.	1.0	6
189	Optimization of 4-(N-Cycloamino)phenylquinazolines as a Novel Class of Tubulin-Polymerization Inhibitors Targeting the Colchicine Site. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 1390-1402.	2.9	60
190	Synthesis of novel spin-labeled podophyllotoxin derivatives as potential antineoplastic agents: Part XXV. <i>Medicinal Chemistry Research</i> , 2014, 23, 4926-4931.	1.1	5
191	Toward synthesis of third-generation spin-labeled podophyllotoxin derivatives using isocyanide multicomponent reactions. <i>European Journal of Medicinal Chemistry</i> , 2014, 75, 282-288.	2.6	17
192	Anticancer Principles from Medicinal Piper (<i>Piper</i>) Plants. <i>Journal of Traditional and Complementary Medicine</i> , 2014, 4, 8-16.	1.5	52
193	NPRL-Z-1, as a New Topoisomerase II Poison, Induces Cell Apoptosis and ROS Generation in Human Renal Carcinoma Cells. <i>PLoS ONE</i> , 2014, 9, e112220.	1.1	10
194	N-Aryl-6-methoxy-1,2,3,4-tetrahydroquinolines: A novel class of antitumor agents targeting the colchicine site on tubulin. <i>European Journal of Medicinal Chemistry</i> , 2013, 67, 196-207.	2.6	49
195	Design and synthesis of 2-(3-alkylaminophenyl)-6-(pyrrolidin-1-yl)quinolin-4-ones as potent antitumor agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 699-701.	1.0	7
196	Synthesis and biological evaluation of N-alkyl-N-(4-methoxyphenyl)pyridin-2-amines as a new class of tubulin polymerization inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 632-642.	1.4	23
197	New Betulinic Acid Derivatives for Bevirimat-Resistant Human Immunodeficiency Virus Type-1. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 2029-2037.	2.9	69
198	Isolation, Structure Determination, and Anti-HIV Evaluation of Tigliane-Type Diterpenes and Biflavonoid from <i>Stellera chamaejasme</i> . <i>Journal of Natural Products</i> , 2013, 76, 852-857.	1.5	51

#	ARTICLE	IF	CITATIONS
199	Design, synthesis and potent cytotoxic activity of novel podophyllotoxin derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 2363-2369.	1.4	22
200	Design and synthesis of 6,7-methylenedioxy-4-substituted phenylquinolin-2(1H)-one derivatives as novel anticancer agents that induce apoptosis with cell cycle arrest at G2/M phase. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 5064-5075.	1.4	34
201	Cytotoxic and potential anticancer constituents from the stems of <i>Schisandra pubescens</i> . <i>Pharmaceutical Biology</i> , 2013, 51, 1204-1207.	1.3	4
202	Cancer preventive agents 11. Novel analogs of dimethyl dicarboxylate biphenyl as potent cancer chemopreventive agents. <i>Pharmaceutical Biology</i> , 2012, 50, 18-24.	1.3	2
203	Recent Progress of Research on Herbal Products Used in Traditional Chinese Medicine: the Herbs belonging to The Divine Husbandman's Herbal Foundation Canon (《神農本草經》). <i>Journal of Traditional and Complementary Medicine</i> , 2012, 2, 6-26.	1.3	4
204	Recent progress of research on medicinal mushrooms, foods, and other herbal products used in traditional Chinese medicine. <i>Journal of Traditional and Complementary Medicine</i> , 2012, 2, 1-12.	1.5	48
205	Design and one-pot synthesis of new 7-acyl camptothecin derivatives as potent cytotoxic agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 7659-7661.	1.0	15
206	Synthesis and biological evaluation of novel spin labeled 18 ² -glycyrrhetic acid derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 7530-7533.	1.0	11
207	Antitumor agents 290. Design, synthesis, and biological evaluation of new LNCaP and PC-3 cytotoxic curcumin analogs conjugated with anti-androgens. <i>Bioorganic and Medicinal Chemistry</i> , 2012, 20, 4020-4031.	1.4	31
208	Antitumor agents 294. Novel E-ring-modified camptothecin-4-anilino-4-O-demethyl-epipodophyllotoxin conjugates as DNA topoisomerase I inhibitors and cytotoxic agents. <i>Bioorganic and Medicinal Chemistry</i> , 2012, 20, 4489-4494.	1.4	9
209	Synthesis of betulinic acid derivatives as entry inhibitors against HIV-1 and bevirimat-resistant HIV-1 variants. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 5190-5194.	1.0	33
210	Anti-AIDS Agents 90. Novel C-28 Modified Bevirimat Analogues as Potent HIV Maturation Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 8128-8136.	2.9	54
211	Antitumor Agents. 293. Nontoxic Dimethyl-4,4'-dimethoxy-5,6,5'-dimethylenedioxybiphenyl-2,2'-dicarboxylate (DDB) Analogues Chemosensitize Multidrug-Resistant Cancer Cells to Clinical Anticancer Drugs. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 5413-5424.	2.9	29
212	1-(3,4,5-Trimethoxyphenyl)ethane-1,2-diyl esters, a novel compound class with potent chemoreversal activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 7726-7729.	1.0	1
213	Cryptopleurine Analogs with Modification of E Ring Exhibit Different Mechanism to Rac-Cryptopleurine and Tylophorine. <i>PLoS ONE</i> , 2012, 7, e51138.	1.1	6
214	Anti-AIDS agents 88. Anti-HIV conjugates of betulin and betulinic acid with AZT prepared via click chemistry. <i>Tetrahedron Letters</i> , 2012, 53, 1987-1989.	0.7	65
215	Design, Synthesis, and Preclinical Evaluations of Novel 4-Substituted 1,5-Diarylanilines as Potent HIV-1 Non-Nucleoside Reverse Transcriptase Inhibitor (NNRTI) Drug Candidates. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 7219-7229.	2.9	47
216	Bis-chalcone analogues as potent NO production inhibitors and as cytotoxic agents. <i>European Journal of Medicinal Chemistry</i> , 2012, 47, 97-103.	2.6	31

#	ARTICLE	IF	CITATIONS
217	Anti-AIDS agents 89. Identification of DCX derivatives as anti-HIV and chemosensitizing dual function agents to overcome P-gp-mediated drug resistance for AIDS therapy. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 3219-3222.	1.0	1
218	Recent progress of research on medicinal mushrooms, foods, and other herbal products used in traditional Chinese medicine. <i>Journal of Traditional and Complementary Medicine</i> , 2012, 2, 84-95.	1.5	16
219	Antitumor Agents 288: Design, Synthesis, SAR, and Biological Studies of Novel Heteroatom-Incorporated Antofine and Cryptopleurine Analogues as Potent and Selective Antitumor Agents. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 5097-5107.	2.9	35
220	Stelleralides A-C, Novel Potent Anti-HIV Daphnane-Type Diterpenoids from <i>Stellera chamaejasme</i> L. <i>Organic Letters</i> , 2011, 13, 2904-2907.	2.4	78
221	5-Methoxyaristololactam I, the First Natural Substituted Aristololactam from <i>Asarum ichangense</i> . <i>Natural Product Communications</i> , 2011, 6, 1934578X1100600.	0.2	3
222	Picomolar Dichotomous Activity of Gnidimacrin Against HIV-1. <i>PLoS ONE</i> , 2011, 6, e26677.	1.1	33
223	Synthesis of new 2-deoxy-2-fluoro-4-azido nucleoside analogues as potent anti-HIV agents. <i>European Journal of Medicinal Chemistry</i> , 2011, 46, 4178-4183.	2.6	40
224	Anti-AIDS agents 86. Synthesis and anti-HIV evaluation of 2,3-seco-3-nor DCP and DCK analogues. <i>European Journal of Medicinal Chemistry</i> , 2011, 46, 4924-4936.	2.6	17
225	New bichalcone analogs as NF- κ B inhibitors and as cytotoxic agents inducing Fas/CD95-dependent apoptosis. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 1895-1906.	1.4	25
226	Antitumor agents 283. Further elaboration of Desmosdumotin C analogs as potent antitumor agents: Activation of spindle assembly checkpoint as possible mode of action. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 1816-1822.	1.4	14
227	Synthesis, in vitro anti-inflammatory and cytotoxic evaluation, and mechanism of action studies of 1-benzoyl- β -carboline and 1-benzoyl-3-carboxy- β -carboline derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 1674-1682.	1.4	41
228	Antitumor agents 281. Design, synthesis, and biological activity of substituted 4-amino-7,8,9,10-tetrahydro-2H-benzo[h]chromen-2-one analogs (ATBO) as potent in vitro anticancer agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 546-549.	1.0	15
229	Efficient synthesis and biological evaluation of epiceanothoic acid and related compounds. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 338-341.	1.0	18
230	Antitumor agents 279. Structure-activity relationship and in vivo studies of novel 2-(furan-2-yl)naphthalen-1-ol (FNO) analogs as potent and selective anti-breast cancer agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 52-57.	1.0	14
231	Cytotoxic geranylflavonoids from <i>Bonannia graeca</i> . <i>Phytochemistry</i> , 2011, 72, 942-945.	1.4	14
232	Discovery and Development of Natural Product-Derived Chemotherapeutic Agents Based on a Medicinal Chemistry Approach. <i>Journal of Natural Products</i> , 2010, 73, 500-516.	1.5	236
233	Antitumor agents. 271: Total synthesis and evaluation of brazilein and analogs as anti-inflammatory and cytotoxic agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 1037-1039.	1.0	51
234	Conjugates of betulin derivatives with AZT as potent anti-HIV agents. <i>Bioorganic and Medicinal Chemistry</i> , 2010, 18, 6451-6469.	1.4	30

#	ARTICLE	IF	CITATIONS
235	Antitumor agents 278. 4-Amino-2H-benzo[h]chromen-2-one (ABO) analogs as potent in vitro anti-cancer agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 4085-4087.	1.0	41
236	Synthesis and anti-HIV activity of 2'-deoxy-2'-fluoro-4'-C-ethynyl nucleoside analogs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 4053-4056.	1.0	23
237	Cancer preventive agents 10. Prenylated dehydrozingerone analogs as potent chemopreventive agents. <i>Journal of Asian Natural Products Research</i> , 2010, 12, 227-232.	0.7	11
238	Cytotoxic Polyisoprenyl Benzophenonoids from <i>Garcinia subelliptica</i> . <i>Journal of Natural Products</i> , 2010, 73, 557-562.	1.5	57
239	Antitumor Agents. 274. A New Synthetic Strategy for E-Ring SAR Study of Antofine and Cryptopleurine Analogues. <i>Organic Letters</i> , 2010, 12, 1416-1419.	2.4	50
240	Anti-AIDS Agents 81. Design, Synthesis, and Structure-Activity Relationship Study of Betulinic Acid and Moronic Acid Derivatives as Potent HIV Maturation Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 3133-3141.	2.9	59
241	HIV entry inhibitors and their potential in HIV therapy. <i>Medicinal Research Reviews</i> , 2009, 29, 369-393.	5.0	108
242	Novel N-(3-carboxyl-9-benzyl- β -carboline-1-yl)ethylamino acids: Synthesis, anti-tumor evaluation, intercalating determination, 3D QSAR analysis and docking investigation. <i>European Journal of Medicinal Chemistry</i> , 2009, 44, 4153-4161.	2.6	85
243	Cytotoxic Phenanthrenequinones and 9,10-Dihydrophenanthrenes from <i>Calanthe arisanensis</i> . <i>Journal of Natural Products</i> , 2009, 72, 210-213.	1.5	51
244	Plant-derived triterpenoids and analogues as antitumor and anti-HIV agents. <i>Natural Product Reports</i> , 2009, 26, 1321.	5.2	154
245	Anti-AIDS Agents. 78. Design, Synthesis, Metabolic Stability Assessment, and Antiviral Evaluation of Novel Betulinic Acid Derivatives as Potent Anti-Human Immunodeficiency Virus (HIV) Agents. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 3248-3258.	2.9	72
246	Betulinic Acid Derivatives as Human Immunodeficiency Virus Type 2 (HIV-2) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 7887-7891.	2.9	49
247	Antitumor Agents 268. Design, Synthesis, and Mechanistic Studies of New 9-Substituted Phenanthrene-Based Tylophorine Analogues as Potent Cytotoxic Agents. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 5262-5268.	2.9	45
248	5-Hydroxymethyl-2-furfural, a clinical trials agent for sickle cell anemia, and its mono/di-glucosides from classically processed steamed <i>Rehmanniae Radix</i> . <i>Journal of Natural Medicines</i> , 2008, 62, 164-167.	1.1	60
249	Plant-derived natural product research aimed at new drug discovery. <i>Journal of Natural Medicines</i> , 2008, 62, 263-280.	1.1	126
250	Total synthesis of phenanthroindolizidine alkaloids (Δ^{\pm})-antofine, (Δ^{\pm})-deoxypergularinine, and their dehydro congeners and evaluation of their cytotoxic activity. <i>Bioorganic and Medicinal Chemistry</i> , 2008, 16, 6233-6241.	1.4	38
251	Structural analogs of tylophora alkaloids may not be functional analogs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 704-709.	1.0	59
252	Cytotoxic calanquinone A from <i>Calanthe arisanensis</i> and its first total synthesis. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 4275-4277.	1.0	22

#	ARTICLE	IF	CITATIONS
253	Antitumor agents 248. Chemistry and antitumor activity of tylophorinerelated alkaloids. <i>Studies in Natural Products Chemistry</i> , 2008, 34, 3-34.	0.8	12
254	Antitumor Agents 253. Design, Synthesis, and Antitumor Evaluation of Novel 9-Substituted Phenanthrene-Based Tylophorine Derivatives as Potential Anticancer Agents. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 3674-3680.	2.9	79
255	Cancer Preventive Agents. 7. Antitumor-Promoting Effects of Seven Active Flavonolignans from Milk Thistle (<i>Silybum marianum</i>) on Epstein-Barr Virus Activation. <i>Pharmaceutical Biology</i> , 2007, 45, 735-738.	1.3	9
256	New developments in natural products-based anti-AIDS research. <i>Medicinal Research Reviews</i> , 2007, 27, 108-132.	5.0	85
257	New developments in the chemistry and biology of the bioactive constituents of tanshen. <i>Medicinal Research Reviews</i> , 2007, 27, 133-148.	5.0	493
258	Anti-AIDS agents 66: Syntheses and anti-HIV activity of phenolic and aza 3 α ,4 α -di-O-(β)-camphanoyl-(+)-cis-khellactone (DCK) derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 6852-6858.	1.4	10
259	Antitumor agents. 256. Conjugation of paclitaxel with other antitumor agents: Evaluation of novel conjugates as cytotoxic agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 2894-2898.	1.0	29
260	Anti-AIDS agents 72. Bioisosteres (7-carbon-DCKs) of the potent anti-HIV lead DCK. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 4316-4319.	1.0	14
261	Anti-AIDS agents 73: Structure-activity relationship study and asymmetric synthesis of 3-O-monomethylsuccinyl-betulinic acid derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 6553-6557.	1.0	31
262	Antitumor Agents 252. Application of validated QSAR models to database mining: discovery of novel tylophorine derivatives as potential anticancer agents. <i>Journal of Computer-Aided Molecular Design</i> , 2007, 21, 97-112.	1.3	85
263	Cancer Preventive Agents. Part 5. Anti-tumor-Promoting Effects of Coumarins and Related Compounds on Epstein-Barr Virus Activation and Two-stage Mouse Skin Carcinogenesis. <i>Pharmaceutical Biology</i> , 2006, 44, 178-182.	1.3	39
264	Recent Progress and Prospects on Plant-Derived Anti-HIV Agents and Analogs. , 2006, , 357-397.		4
265	Cancer Preventive Agents. Part 6: Chemopreventive Potential of Furanocoumarins and Related Compounds. <i>Pharmaceutical Biology</i> , 2006, 44, 116-120.	1.3	20
266	Anti-AIDS Agents 69. Moronic Acid and Other Triterpene Derivatives as Novel Potent Anti-HIV Agents. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 5462-5469.	2.9	113
267	Antitumor agents 251: Synthesis, cytotoxic evaluation, and structure-activity relationship studies of phenanthrene-based tylophorine derivatives (PBTs) as a new class of antitumor agents. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 6560-6569.	1.4	79
268	Synthesis and anti-HIV activity of bi-functional betulinic acid derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 2279-2289.	1.4	76
269	Cytotoxic Alangium alkaloids from <i>Alangium longiflorum</i> . <i>Phytochemistry</i> , 2006, 67, 894-897.	1.4	12
270	Terpenoids from <i>Juniperus polycarpus</i> var. <i>seravschanica</i> . <i>Phytochemistry</i> , 2006, 67, 2635-2640.	1.4	46

#	ARTICLE	IF	CITATIONS
271	Philinopsides A and B, Two New Sulfated Triterpene Glycosides from the Sea Cucumber <i>Pentacta quadrangularis</i> . <i>Helvetica Chimica Acta</i> , 2006, 89, 54-63.	1.0	33
272	Structure-Activity Relationships of Curcumin and Its Analogs with Different Biological Activities as Antitumor Agents 241.. <i>Studies in Natural Products Chemistry</i> , 2006, 33, 785-812.	0.8	16
273	Cancer preventive agents. Part 1: Chemopreventive potential of cimigenol, cimigenol-3,15-dione, and related compounds. <i>Bioorganic and Medicinal Chemistry</i> , 2005, 13, 1403-1408.	1.4	26
274	Antitumor agents 243. Syntheses and cytotoxicity of desmosdumotin C derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2005, 13, 2325-2330.	1.4	15
275	Molecular modeling, design, synthesis, and biological evaluation of novel 3- β ,4-dicamphanoyl-(+)-cis-khellactone (DCK) analogs as potent anti-HIV agents. <i>Bioorganic and Medicinal Chemistry</i> , 2005, 13, 6435-6449.	1.4	10
276	Two new sesquiterpenoids and anti-HIV principles from the root bark of <i>Zanthoxylum ailanthoides</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2005, 13, 5915-5920.	1.4	73
277	Total synthesis and bioactivity of unique flavone desmosdumotin B and its analogs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 3016-3019.	1.0	28
278	Cancer Preventive Agents 3. Antitumor Promoting Effects of <i>Agaricus blazei</i> .. <i>Pharmaceutical Biology</i> , 2005, 43, 568-572.	1.3	6
279	The discovery of a class of novel HIV-1 maturation inhibitors and their potential in the therapy of HIV. <i>Expert Opinion on Investigational Drugs</i> , 2005, 14, 681-693.	1.9	45
280	New Dammarane-Type Saponins from the Galls of <i>Sapindus mukorossi</i> . <i>Journal of Agricultural and Food Chemistry</i> , 2005, 53, 4722-4727.	2.4	52
281	Antitumor agents. Part 230: C4-esters of GL-331 as cytotoxic agents and DNA topoisomerase II inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 2979-2982.	1.0	8
282	Anti-AIDS agents. Part 62: Anti-HIV activity of 2-substituted 4-methyl-3,4-di-O-(α^{\sim})-camphanoyl-(+)-cis-khellactone (4-methyl DCK) analogs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 5855-5857.	1.0	16
283	Cytotoxic Isoprenylated Flavonoids from the Roots of <i>Sophora flavescens</i> . <i>Helvetica Chimica Acta</i> , 2004, 87, 2574-2580.	1.0	32
284	Antitumor agents. Part 235: Novel 4-ester etoposide analogues as potent DNA topoisomerase II inhibitors with improved therapeutic potential. <i>Bioorganic and Medicinal Chemistry</i> , 2004, 12, 3363-3369.	1.4	6
285	Antitumor agents. Part 227: Studies on novel 4-O-demethyl-epipodophyllotoxins as antitumor agents targeting topoisomerase II. <i>Bioorganic and Medicinal Chemistry</i> , 2004, 12, 3339-3344.	1.4	11
286	Anti-AIDS agents. Part 56: Synthesis and anti-HIV activity of 7-thia-di-O-(α^{\sim})-camphanoyl-(+)-cis-khellactone (7-thia-DCK) analogs. <i>Bioorganic and Medicinal Chemistry</i> , 2004, 12, 6383-6387.	1.4	51
287	Anti-AIDS Agents. Part 57. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 1329-1332.	1.0	45
288	Antitumor agents. Part 232: Synthesis of cyclosulfite podophyllotoxin analogues as novel prototype antitumor agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 1581-1584.	1.0	14

#	ARTICLE	IF	CITATIONS
289	Anti-AIDS agents. Part 58: Synthesis and anti-HIV activity of 1-thia-di-O-(α^{\sim})-camphanoyl-(+)-cis-khellactone (1-thia-DCK) analogues. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 3341-3343.	1.0	10
290	3-O-Glutaryl-dihydrobetulin and related monoacyl derivatives as potent anti-HIV agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 5851-5853.	1.0	21
291	Antitumor agents. Part 227: Studies on novel 4 \prime -O-demethyl-epipodophyllotoxins as antitumor agents targeting topoisomerase II*1. <i>Bioorganic and Medicinal Chemistry</i> , 2004, 12, 3339-3344.	1.4	19
292	Antitumor agents. Part 235: Novel 4 \prime -ester etoposide analogues as potent DNA topoisomerase II inhibitors with improved therapeutic potential*1. <i>Bioorganic and Medicinal Chemistry</i> , 2004, 12, 3363-3369.	1.4	16
293	Antitumor Agents. 239. Isolation, Structure Elucidation, Total Synthesis, and Anti-Breast Cancer Activity of Neo-tanshinlactone from <i>Salvia miltiorrhiza</i> . <i>Journal of Medicinal Chemistry</i> , 2004, 47, 5816-5819.	2.9	193
294	Current Developments in the Discovery and Design of New Drug Candidates from Plant Natural Product Leads. <i>Journal of Natural Products</i> , 2004, 67, 273-283.	1.5	163
295	Lignans in treatment of cancer and other diseases. <i>Phytochemistry Reviews</i> , 2003, 2, 341-362.	3.1	90
296	Recent progress in the development of coumarin derivatives as potent anti-HIV agents. <i>Medicinal Research Reviews</i> , 2003, 23, 322-345.	5.0	455
297	Antitumor agents 222. For Part 221, see ref 1. Synthesis and anti-androgen activity of new diarylheptanoids. <i>Bioorganic and Medicinal Chemistry</i> , 2003, 11, 5083-5090.	1.4	44
298	Anti-AIDS agents 54. A potent anti-HIV chalcone and flavonoids from genus <i>Desmos</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003, 13, 1813-1815.	1.0	261
299	Antitumor agents. Part 218: Cappamensin A, a new <i>In vitro</i> anticancer principle, from <i>Capparis sikkimensis</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003, 13, 2223-2225.	1.0	34
300	Antitumor agents. Part 226: synthesis and cytotoxicity of 2-Phenyl-4-quinolone acetic acids and their esters. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003, 13, 2891-2893.	1.0	48
301	Antitumor agents 220. Antitumor-Promoting effects of cimigenol and related compounds on Epstein-Barr virus activation and two-stage mouse skin carcinogenesis. <i>Bioorganic and Medicinal Chemistry</i> , 2003, 11, 1137-1140.	1.4	23
302	Antitumor agents 216. Synthesis and evaluation of paclitaxel-camptothecin conjugates as novel cytotoxic agents. <i>Bioorganic and Medicinal Chemistry</i> , 2003, 11, 1851-1857.	1.4	32
303	Antitumor Agents. 221. Buceracidins A and B, Two New Flavanones from <i>Bucida buceras</i> . <i>Journal of Natural Products</i> , 2003, 66, 125-127.	1.5	15
304	Oriental Herbal Products: The Basis for Development of Dietary Supplements and New Medicines in the 21st Century. <i>ACS Symposium Series</i> , 2003, , 2-31.	0.5	2
305	Progress in the Recent Discovery and Development of Promising Anticancer and Anti-HIV Agents from Natural Products in the United States. <i>Journal of the Chinese Chemical Society</i> , 2003, 50, 11-22.	0.8	1
306	Antitumor agents. Part 212. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2002, 12, 345-348.	1.0	32

#	ARTICLE	IF	CITATIONS
307	Antitumor agents. Part 215: Bioorganic and Medicinal Chemistry Letters, 2002, 12, 2851-2853.	1.0	6
308	Antitumor agents. Part 209: Pheophorbide-a derivatives as photo-Independent cytotoxic agents. Bioorganic and Medicinal Chemistry, 2002, 10, 583-591.	1.4	50
309	Chemopreventive potential of cyclic diarylheptanoids. Bioorganic and Medicinal Chemistry, 2002, 10, 3361-3365.	1.4	53
310	Antitumor Agents. Part 214: For paper 213, see ref 1. Synthesis and Evaluation of Curcumin Analogues as Cytotoxic Agents. Bioorganic and Medicinal Chemistry, 2002, 10, 3481-3487.	1.4	130
311	New Cytotoxic Butanolides from <i>Litsea acutivena</i> . Journal of Natural Products, 2001, 64, 1502-1505.	1.5	40
312	Anti-AIDS Agents. 46.1 Anti-HIV Activity of Harman, an Anti-HIV Principle from <i>Symplocos setchuensis</i> , and Its Derivatives. Journal of Natural Products, 2001, 64, 958-960.	1.5	100
313	Antitumor Agents. Part 204: Synthesis and Biological Evaluation of Substituted 2-Aryl Quinazolinones. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 1193-1196.	1.0	178
314	Synthesis and anti-HIV activity of oleanolic acid derivatives. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 3115-3118.	1.0	82
315	Anti-HIV and Cytotoxic Activities of Ru(II)/Ru(III) Polypyridyl Complexes Containing 2,6-(2-Benzimidazolyl)-pyridine/chalcone as Co-Ligand. Bioorganic and Medicinal Chemistry, 2001, 9, 1667-1671.	1.4	66
316	Simple Isoquinoline and Benzylisoquinoline Alkaloids as Potential Antimicrobial, Antimalarial, Cytotoxic, and Anti-HIV Agents. Bioorganic and Medicinal Chemistry, 2001, 9, 2871-2884.	1.4	115
317	Antitumor agents 210. synthesis and evaluation of taxoid epipodophyllotoxin conjugates as novel cytotoxic agents. Bioorganic and Medicinal Chemistry, 2001, 9, 2999-3004.	1.4	43
318	Antitumor agents. Part 202: Novel 2-amino chalcones: design, synthesis and biological evaluation. Bioorganic and Medicinal Chemistry Letters, 2000, 10, 699-701.	1.0	186
319	Research and future trends in the pharmaceutical development of medicinal herbs from Chinese medicine. Public Health Nutrition, 2000, 3, 515-522.	1.1	104
320	Antitumor-promoting effects of cyclic diarylheptanoids on Epstein-Barr virus activation and two-stage mouse skin carcinogenesis. Cancer Letters, 2000, 159, 135-140.	3.2	56
321	6-Alkylamino- and 2,3-Dihydro-3-methoxy-2-phenyl-4-quinazolinones and Related Compounds: Their Synthesis, Cytotoxicity, and Inhibition of Tubulin Polymerization. Journal of Medicinal Chemistry, 2000, 43, 4479-4487.	2.9	376
322	Anti-HIV Agents 451 and Antitumor Agents 205.2 Two New Sesquiterpenes, Leitneridanins A and B, and the Cytotoxic and Anti-HIV Principles from <i>Leitneria floridana</i> . Journal of Natural Products, 2000, 63, 1712-1715.	1.5	48
323	Substituted 7H-pyrido[4,3-c]carbazoles with potent anti-HIV activity. Bioorganic and Medicinal Chemistry Letters, 1999, 9, 119-122.	1.0	44
324	Anti-AIDS agents. Part 36: 17-carboxylated steroids as potential anti-HIV agents. Bioorganic and Medicinal Chemistry, 1999, 7, 1907-1911.	1.4	12

#	ARTICLE	IF	CITATIONS
325	Anticancer drug design based on plant-derived natural products. <i>Journal of Biomedical Science</i> , 1999, 6, 236-250.	2.6	97
326	Novel antitumor agents from higher plants. , 1999, 19, 569-596.		140
327	Antitumor Agents. 196. Substituted 2-Thienyl-1,8-naphthyridin-4-ones: Their Synthesis, Cytotoxicity, and Inhibition of Tubulin Polymerization. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 4081-4087.	2.9	70
328	Asymmetric Solid-Phase Synthesis of (3 <i>R</i> ,4 <i>R</i>)-Di-O-cis-acyl 3-Carboxyl Khellactones. <i>Organic Letters</i> , 1999, 1, 2113-2115.	2.4	20
329	Anti-AIDS Agents. 37. Synthesis and Structure-Activity Relationships of (3 <i>R</i> ,4 <i>R</i>)-(+)-cis-Khellactone Derivatives as Novel Potent Anti-HIV Agents. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 2662-2672.	2.9	147
330	Recent Advances in the Discovery and Development of Quinolones and Analogs as Antitumor Agents'. <i>Current Medicinal Chemistry</i> , 1999, 6, 179-194.	1.2	29
331	Anti-AIDS Agents. 34. Synthesis and Structure-Activity Relationships of Betulin Derivatives as Anti-HIV Agents. <i>Journal of Medicinal Chemistry</i> , 1998, 41, 4648-4657.	2.9	139
332	Antitumor Agents. Part 184. Syntheses and antitubulin activity of compounds derived from reaction of thiocolchicone with amines: Lactams, alcohols, and ester analogs of allothiocolchicinoids. <i>Helvetica Chimica Acta</i> , 1998, 81, 1023-1037.	1.0	39
333	Synthesis and Cytotoxicity of 2-Acetyl-4,8-dihydrobenzodithiophene-4,8-dione Derivatives. <i>Journal of Medicinal Chemistry</i> , 1998, 41, 4658-4661.	2.9	10
334	Oxidation Products of Phenolic Thiocolchicines: Ring a Quinones and Dienones. <i>Synthetic Communications</i> , 1998, 28, 1585-1591.	1.1	10
335	Cytotoxic Principles from <i>Saussurea Lappa</i> and <i>Corydalis Turtshaninovii</i> f. <i>Yanhusuo</i> . <i>Journal of the Chinese Chemical Society</i> , 1997, 44, 357-359.	0.8	9
336	Antitumor Agents. 178. Synthesis and Biological Evaluation of Substituted 2-Aryl-1,8-naphthyridin-4(1H)-ones as Antitumor Agents That Inhibit Tubulin Polymerization. <i>Journal of Medicinal Chemistry</i> , 1997, 40, 3049-3056.	2.9	77
337	Antitumor Agents. 180. Chemical Studies and Cytotoxic Evaluation of Cumingianosides and Cumindoside A, Antileukemic Triterpene Glucosides with a 14,18-Cyclopotirucallane Skeleton. <i>Journal of Natural Products</i> , 1997, 60, 1105-1114.	1.5	11
338	Cytotoxic and Antiplatelet Aggregation Principles from <i>Aglaia elliptifolia</i> . <i>Journal of Natural Products</i> , 1997, 60, 606-608.	1.5	60
339	Anti-AIDS agents-XXVII. Synthesis and anti-HIV activity of betulinic acid and dihydrobetulinic acid derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 1997, 5, 2133-2143.	1.4	108
340	Recent advances in the discovery and development of topoisomerase inhibitors as antitumor agents. , 1997, 17, 367-425.		139
341	Synthesis and Cytotoxicity of 1,2-Disubstituted Naphth[2,3-d]imidazole-4,9-diones and Related Compounds. <i>Journal of Medicinal Chemistry</i> , 1996, 39, 1447-1451.	2.9	35
342	Antitumor Agents. 166. Synthesis and Biological Evaluation of 5,6,7,8-Substituted-2-phenylthiochromen-4-ones. <i>Journal of Medicinal Chemistry</i> , 1996, 39, 1975-1980.	2.9	70

#	ARTICLE	IF	CITATIONS
343	Betulinic Acid and Dihydrobetulinic Acid Derivatives as Potent Anti-HIV Agents ¹ . <i>Journal of Medicinal Chemistry</i> , 1996, 39, 1016-1017.	2.9	262
344	Identification of ent-16 β ,17-Dihydroxykauran-19-oic Acid as an Anti-HIV Principle and Isolation of the New Diterpenoids Annosquamosins A and B from <i>Annona squamosa</i> . <i>Journal of Natural Products</i> , 1996, 59, 635-637.	1.5	131
345	Napalolides A-D, Four New Sesquiterpene Lactones from <i>Carpesium nepalense</i> . <i>Journal of Natural Products</i> , 1996, 59, 991-993.	1.5	23
346	Anti-AIDS Agents, 11. Betulinic Acid and Platanic Acid as Anti-HIV Principles from <i>Syzygium claviflorum</i> , and the Anti-HIV Activity of Structurally Related Triterpenoids. <i>Journal of Natural Products</i> , 1994, 57, 243-247.	1.5	424
347	Antineoplastic Alkaloids From Chinese Medicinal Plants and Their Analogs. <i>Journal of the Chinese Chemical Society</i> , 1994, 41, 371-384.	0.8	8
348	Antitumor agents. 126. Novel 4 beta-substituted anilino derivatives of 3',4'-O,O-didemethylpodophyllotoxin as potent inhibitors of human DNA topoisomerase II. <i>Pharmaceutical Research</i> , 1993, 10, 343-350.	1.7	7
349	Antitumor agents. 125. New 4 beta-benzoylamino derivatives of 4'-O-demethyl-4-desoxypodophyllotoxin and 4 beta-benzoyl derivatives of 4'-O-demethylpodophyllotoxin as potent inhibitors of human DNA topoisomerase II. <i>Pharmaceutical Research</i> , 1993, 10, 214-219.	1.7	17
350	Antineoplastic Agents and Their Analogues from Chinese Traditional Medicine. <i>ACS Symposium Series</i> , 1993, , 170-190.	0.5	18
351	Plant Phenolic Compounds as Cytotoxic Antitumor Agents. <i>ACS Symposium Series</i> , 1992, , 367-379.	0.5	8
352	Anti-AIDS Agents, 4. Tripterifordin, a Novel Anti-HIV Principle from <i>Tripterium wilfordii</i> : Isolation and Structural Elucidation. <i>Journal of Natural Products</i> , 1992, 55, 88-92.	1.5	72
353	Cytotoxic Principles of <i>Securinega virosa</i> : Virosecurinine and Viroallosecurinine and Related Derivatives. <i>Journal of Pharmaceutical Sciences</i> , 1991, 80, 325-327.	1.6	41
354	X-Ray Crystal Structure of Acrovestone, a Cytotoxic Principle from <i>Acronychia pedunculata</i> . <i>Journal of Natural Products</i> , 1989, 52, 1284-1289.	1.5	36
355	Antimalarial agents. III. Mechanism of action of artesunate against <i>Plasmodium berghei</i> infection.. <i>Chemical and Pharmaceutical Bulletin</i> , 1987, 35, 2052-2061.	0.6	27
356	Structure-Activity Relationships for Binding and Inactivation of Rabbit Reticulocyte Ribosomes by Quassinoid Antineoplastic Agents. <i>FEBS Journal</i> , 1983, 132, 157-163.	0.2	7
357	Genkwadaphnin, a potent antileukemic diterpene from <i>Daphne genkwa</i> . <i>Phytochemistry</i> , 1981, 20, 2592-2594.	1.4	50