

# Kuo-Hsiung Lee

## List of Publications by Year in descending order

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357  
papers

14,791  
citations

22132

59  
h-index

33869

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 dual-site-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 <sup>3</sup> )-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 LC-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

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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 <sup>n</sup> . <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 <sup>pro</sup> 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

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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

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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 Structure-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	Structure-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

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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>FÄ-toterapÄ-Äç</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

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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 16–18, 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 <sub>19</sub> -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 2,4-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

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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)pipe Chemistry Letters, 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 Aâ€“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 in vitro and in vivo. <i>European Journal of Medicinal Chemistry</i> , 2017, 131, 141-151.	2.6	23
118	Corymbulosins Dâ€“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 structureâ€“activity relationships of ( $\hat{\Delta}$ )-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 Vobasinylâ€“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(1 <i>H</i> )-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



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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
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