

Jongkook Lee

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7534905/publications.pdf>

Version: 2024-02-01

71
papers

1,310
citations

411340

20
h-index

466096

32
g-index

78
all docs

78
docs citations

78
times ranked

2085
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of Human Kinases Involved in Hepatitis C Virus Replication by Small Interference RNA Library Screening. <i>Journal of Biological Chemistry</i> , 2008, 283, 29-36.	1.6	95
2	Elevated endoplasmic reticulum stress reinforced immunosuppression in the tumor microenvironment via myeloid-derived suppressor cells. <i>Oncotarget</i> , 2014, 5, 12331-12345.	0.8	87
3	A Small Molecule Antagonist of the Hedgehog Signaling Pathway. <i>ChemBioChem</i> , 2007, 8, 1916-1919.	1.3	71
4	Asymmetric Total Synthesis of (+)-Brefeldin A from (S)-Lactate by Triple Chirality Transfer Process and Nitrile Oxide Cycloaddition. <i>Journal of Organic Chemistry</i> , 2002, 67, 764-771.	1.7	63
5	Role of Conformational Effects on the Regioselectivity of Macrocyclic INOC Reactions: Two New Asymmetric Total Syntheses of (+)-Brefeldin A. <i>Journal of Organic Chemistry</i> , 2002, 67, 772-781.	1.7	45
6	An Approach To Enhance Specificity against RNA Targets Using Heteroconjugates of Aminoglycosides and Chloramphenicol (or Linezolid). <i>Journal of the American Chemical Society</i> , 2004, 126, 1956-1957.	6.6	44
7	Glyceollins, a novel class of soy phytoalexins, inhibit angiogenesis by blocking the VEGF and bFGF signaling pathways. <i>Molecular Nutrition and Food Research</i> , 2013, 57, 225-234.	1.5	37
8	Small molecule inhibitors of the hedgehog signaling pathway for the treatment of cancer. <i>Archives of Pharmacal Research</i> , 2012, 35, 1317-1333.	2.7	34
9	SB365, Pulsatilla saponin D, targets c-Met and exerts antiangiogenic and antitumor activities. <i>Carcinogenesis</i> , 2013, 34, 2156-2169.	1.3	34
10	Discovery of a potent small molecule SIRT1/2 inhibitor with anticancer effects. <i>International Journal of Oncology</i> , 2013, 43, 1205-1211.	1.4	34
11	Moracin M inhibits airway inflammation by interrupting the JNK/c-Jun and NF- κ B pathways in vitro and in vivo. <i>European Journal of Pharmacology</i> , 2016, 783, 64-72.	1.7	32
12	Discovery of aminopyridines substituted with benzoxazole as orally active c-Met kinase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 4223-4227.	1.0	28
13	Design and synthesis of 3-(4,5,6,7-tetrahydro-3H-imidazo[4,5-c]pyridin-2-yl)-1H-quinolin-2-ones as VEGFR-2 kinase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 2837-2842.	1.0	28
14	First Synthesis and Structural Elucidation of ($\alpha^{\prime\prime}$)-Presphaerene. <i>Journal of Organic Chemistry</i> , 2004, 69, 6433-6440.	1.7	26
15	Electrosprayed nanocomposites based on hyaluronic acid derivative and Soluplus for tumor-targeted drug delivery. <i>Colloids and Surfaces B: Biointerfaces</i> , 2016, 145, 267-274.	2.5	25
16	Novel metabolites from <i>Trichoderma atroviride</i> against human prostate cancer cells and their inhibitory effect on <i>Helicobacter pylori</i> and Shigella toxin producing <i>Escherichia coli</i> . <i>Microbial Pathogenesis</i> , 2019, 126, 19-26.	1.3	25
17	Stereoselective synthesis of (Δ^{\pm})- β^2 -elemene by a doubly diastereodifferentiating internal alkylation: a remarkable difference in the rate of enolization between syn and anti esters. <i>Tetrahedron</i> , 2001, 57, 1247-1252.	1.0	24
18	Synthesis and structure-activity relationship of aminopyridines with substituted benzoxazoles as c-Met kinase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 4044-4048.	1.0	22

#	ARTICLE	IF	CITATIONS
19	KRC-408, a novel c-Met inhibitor, suppresses cell proliferation and angiogenesis of gastric cancer. <i>Cancer Letters</i> , 2013, 332, 74-82.	3.2	22
20	Sphingosine 1-Phosphate Receptor 4 Promotes Nonalcoholic Steatohepatitis by Activating NLRP3 Inflammasome. <i>Cellular and Molecular Gastroenterology and Hepatology</i> , 2022, 13, 925-947.	2.3	22
21	Construction of the Tricyclo[5.3.1.0 ^{1,5}]undecane System by a Novel Tandem Pinacol Rearrangement-Ene Strategy: A Formal Total Synthesis of (±)-Perhydrohistrionicotoxin. <i>Journal of Organic Chemistry</i> , 2000, 65, 4864-4870.	1.7	21
22	Chemical suppression of an oncogenic splicing variant of AIMP2 induces tumour regression. <i>Biochemical Journal</i> , 2013, 454, 411-416.	1.7	21
23	Preservative effect of Chinese cabbage (<i>Brassica rapa</i> subsp. <i>pekinensis</i>) extract on their molecular docking, antioxidant and antimicrobial properties. <i>PLoS ONE</i> , 2018, 13, e0203306.	1.1	21
24	Remote Stereoinductive Intramolecular Nitrile Oxide Cycloaddition: Asymmetric Total Synthesis and Structure Revision of (±)-11 ^H -Hydroxycurvularin. <i>Journal of Organic Chemistry</i> , 2016, 81, 2612-2617.	1.7	20
25	Efficient construction of bicyclic systems by an internal SN2 ^{enolate} enolate alkylation/ring-closing metathesis (RCM) strategy: a concise synthesis of the trans-hydrindane nucleus. <i>Tetrahedron Letters</i> , 2003, 44, 7043-7044.	0.7	19
26	Synthesis and biological evaluation of 5-nitropyrimidine analogs with azabicyclic substituents as GPR119 agonists. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 1519-1521.	1.0	19
27	Total Synthesis of (+)-Pochonin D and (+)-Monocillin II via Chemo- and Regioselective Intramolecular Nitrile Oxide Cycloaddition. <i>Organic Letters</i> , 2017, 19, 6004-6007.	2.4	19
28	2-Formyl-komarovicine promotes adiponectin production in human mesenchymal stem cells through PPAR γ partial agonism. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 1069-1075.	1.4	19
29	Renal Cell Carcinoma Is Abrogated by p53 Stabilization through Transglutaminase 2 Inhibition. <i>Cancers</i> , 2018, 10, 455.	1.7	19
30	Design and synthesis of triazolopyridazines substituted with methylisoquinolinone as selective c-Met kinase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 7185-7188.	1.0	18
31	Evaluation of a multi-kinase inhibitor KRC-108 as an anti-tumor agent in vitro and in vivo. <i>Investigational New Drugs</i> , 2012, 30, 518-523.	1.2	18
32	Cross-metathesis of allyl halides with olefins bearing amide and ester groups. <i>Tetrahedron</i> , 2012, 68, 1177-1184.	1.0	17
33	Synthesis and biological evaluation of novel 2,4-disubstituted quinazoline analogues as GPR119 agonists. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 1349-1356.	1.4	16
34	Identification and Characterization of Small-Molecule Inducers of Epidermal Keratinocyte Differentiation. <i>ACS Chemical Biology</i> , 2007, 2, 171-175.	1.6	15
35	Concise Substrate-Controlled Asymmetric Total Synthesis of (+)-3-(Z)-Dihydrorhodophytin. <i>Heterocycles</i> , 2010, 82, 1113.	0.4	15
36	Stereoselective Construction of 2,6-cis-Disubstituted Tetrahydropyrans via Intramolecular Amide Enolate Alkylation: Total Synthesis of (±)-Centrolobine. <i>Journal of Organic Chemistry</i> , 2015, 80, 3315-3320.	1.7	15

#	ARTICLE	IF	CITATIONS
37	Identification of a new tadalafil analogue in an adulterated dietary supplement: Trans-Bisprehomotadalafil. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2015, 115, 352-358.	1.4	15
38	Isolation and characterisation of a novel sildenafil analogue adulterant, desmethylpiperazinyl propoxysildenafil, in a dietary supplement. <i>Science and Justice - Journal of the Forensic Science Society</i> , 2018, 58, 447-454.	1.3	15
39	Isolation and structural elucidation of a new tadalafil analogue in health supplements: bisprenortadalafil. <i>Food Additives and Contaminants - Part A Chemistry, Analysis, Control, Exposure and Risk Assessment</i> , 2016, 33, 945-952.	1.1	14
40	Identification of new synthetic cannabinoid analogue APINAC (adamantan-1-yl) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 627 Td (1-pentyl-1 Forensic Toxicology, 2017, 35, 45-55.	1.4	14
41	Novel 5-nitropyrimidine derivatives bearing endo-azabicyclic alcohols/amines as potent GPR119 agonists. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 254-260.	1.4	13
42	Resistance to the c-Met inhibitor KRC-108 induces the epithelial transition of gastric cancer cells. <i>Oncology Letters</i> , 2016, 11, 991-997.	0.8	12
43	Stereoselective Protection-Free Asymmetric Total Synthesis of (+)-Chamuvarinin, a Potent Anticancer and Antitrypanosomal Agent: Substrate-Controlled Construction of the Adjacently Linked Oxatricyclic Core by Internal Alkylation. <i>Organic Letters</i> , 2018, 20, 6398-6402.	2.4	12
44	Discovery of quinolinone derivatives as potent FLT3 inhibitors. <i>Biochemical and Biophysical Research Communications</i> , 2014, 445, 561-565.	1.0	10
45	Synthesis and Structure Revision of Dimeric Tadalafil Analogue Adulterants in Dietary Supplements. <i>Chemical and Pharmaceutical Bulletin</i> , 2017, 65, 498-503.	0.6	9
46	A small molecule inhibitor of β 4 integrin-dependent cell migration. <i>Bioorganic and Medicinal Chemistry</i> , 2009, 17, 977-980.	1.4	8
47	Identification and characterization of an indazole-3-carboxamide class synthetic cannabinoid: 2-[1-(cyclohexylmethyl)-1H-indazole-3-carboxamido]-3,3-dimethylbutanoic acid (DMBA-CHMINACA). <i>Forensic Science International</i> , 2018, 291, 167-174.	1.3	8
48	Flavonoids from <i>Scutellaria baicalensis</i> inhibit senescence-associated secretory phenotype production by interrupting $\text{I}\beta\text{B}1/\text{C}/\text{EBP}2$ pathway: Inhibition of age-related inflammation. <i>Phytomedicine</i> , 2020, 76, 153255.	2.3	8
49	Synthesis, in vitro evaluation, and computational simulations studies of 1,2,3-triazole analogues as DPP-4 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2021, 29, 115861.	1.4	8
50	Cross-metathesis of allyl halides with olefins bearing an β -alkoxy amide group. <i>Tetrahedron Letters</i> , 2011, 52, 1928-1930.	0.7	7
51	KRC-327, a selective novel inhibitor of c-Met receptor tyrosine kinase with anticancer activity. <i>Cancer Letters</i> , 2013, 331, 158-166.	3.2	7
52	Novel 1,2,3-Triazole Analogs of Sitagliptin as DPP4 Inhibitors. <i>Bulletin of the Korean Chemical Society</i> , 2016, 37, 1156-1158.	1.0	7
53	Antiviral and Anti-Inflammatory Activities of Pochonin D, a Heat Shock Protein 90 Inhibitor, against Rhinovirus Infection. <i>Biomolecules and Therapeutics</i> , 2018, 26, 576-583.	1.1	7
54	Metabolic and pharmacokinetic characterization of a new synthetic cannabinoid APINAC in rats. <i>Forensic Toxicology</i> , 2018, 36, 88-101.	1.4	6

#	ARTICLE	IF	CITATIONS
55	Development and validation of liquid chromatography-tandem mass spectrometry method for screening six selective androgen receptor modulators in dietary supplements. <i>Food Additives and Contaminants - Part A Chemistry, Analysis, Control, Exposure and Risk Assessment</i> , 2021, 38, 1075-1086.	1.1	6
56	FMS-like tyrosine kinase 3 inhibitors: a patent review. <i>Expert Opinion on Therapeutic Patents</i> , 2011, 21, 483-503.	2.4	5
57	Identification and Characterization of Small Molecule Inducers of Fetal Hemoglobin. <i>ChemMedChem</i> , 2011, 6, 777-780.	1.6	5
58	Antitumor effect of the integrin $\alpha 4$ signaling inhibitor JK273 in non-small cell lung cancer NCI-H460 cells. <i>Biochemical and Biophysical Research Communications</i> , 2017, 491, 355-360.	1.0	4
59	Highly Stereoselective Asymmetric Total Synthesis of (β)-Jimenezin via Sequential Intramolecular Amide Enolate Alkylation. <i>Organic Letters</i> , 2022, 24, 1652-1656.	2.4	4
60	Effects of KRC-108 on the Aurora A activity and growth of colorectal cancer cells. <i>Biochemical and Biophysical Research Communications</i> , 2015, 461, 605-611.	1.0	3
61	Identification of a new tadalafil analogue in commercial dietary supplements: isopropyl nortadalafil. <i>Food Additives and Contaminants - Part A Chemistry, Analysis, Control, Exposure and Risk Assessment</i> , 2017, 34, 1-8.	1.1	3
62	Collision-induced dissociation pathways of H1-antihistamines by electrospray ionization quadrupole time-of-flight mass spectrometry. <i>Archives of Pharmacol Research</i> , 2017, 40, 736-745.	2.7	3
63	Cross-Metathesis of Methallyl Halides: Concise Enantioselective Formal Total Synthesis of (β)-Presphaerene. <i>Frontiers in Chemistry</i> , 2020, 8, 494.	1.8	2
64	First Syntheses of (β)-Butesuperins A and B. <i>Bulletin of the Korean Chemical Society</i> , 2017, 38, 944-947.	1.0	1
65	Isolation and structural identification of a novel minoxidil analogue in an illegal dietary supplement: triaminodil. <i>Food Additives and Contaminants - Part A Chemistry, Analysis, Control, Exposure and Risk Assessment</i> , 2018, 35, 2-9.	1.1	1
66	KRCA-0008 suppresses ALK-positive anaplastic large-cell lymphoma growth. <i>Investigational New Drugs</i> , 2020, 38, 1282-1291.	1.2	1
67	Potential Moracin M Prodrugs Strongly Attenuate Airway Inflammation <i>in Vivo</i> . <i>Biomolecules and Therapeutics</i> , 2020, 28, 344-353.	1.1	1
68	Characterization of KRC-108 as a TrkA Kinase Inhibitor with Anti-Tumor Effects. <i>Biomolecules and Therapeutics</i> , 2022, 30, 360-367.	1.1	1
69	Efficient Construction of Bicyclic Systems by an Internal SN2 Enolate Alkylation/Ring-Closing Metathesis (RCM) Strategy: A Concise Synthesis of the trans-Hydrindane Nucleus. <i>ChemInform</i> , 2003, 34, no.	0.1	0
70	Identification of a new M-ALPHA analog and MDMA in an illegal health product. <i>Forensic Science International</i> , 2020, 313, 110332.	1.3	0
71	Development of a method for simultaneous screening of four natural-derived steroids and their analogues used as dietary supplements via liquid chromatography-quadrupole-time of flight mass spectrometry and liquid chromatography-tandem mass spectrometry. <i>Food Additives and Contaminants - Part A Chemistry, Analysis, Control, Exposure and Risk Assessment</i> , 2022, , 1-9.	1.1	0