

# Jian Li

## List of Publications by Year in descending order

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105  
papers

3,818  
citations

186265

28  
h-index

138484

58  
g-index

107  
all docs

107  
docs citations

107  
times ranked

5662  
citing authors

#	ARTICLE	IF	CITATIONS
1	Colistin: the re-emerging antibiotic for multidrug-resistant Gram-negative bacterial infections. <i>Lancet Infectious Diseases</i> , 2006, 6, 589-601.	9.1	1,170
2	Scaffold Diversity Inspired by the Natural Product Evodiamine: Discovery of Highly Potent and Multitargeting Antitumor Agents. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 6678-6696.	6.4	156
3	Small-molecule targeting of a diapophytoene desaturase inhibits <i>S. aureus</i> virulence. <i>Nature Chemical Biology</i> , 2016, 12, 174-179.	8.0	121
4	Discovering Potent Small Molecule Inhibitors of Cyclophilin A Using de Novo Drug Design Approach. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 5295-5298.	6.4	92
5	Organocatalytic Enantioselective Direct Additions of Aldehydes to 4-Vinylpyridines and Electron-Deficient Vinylarenes and Their Synthetic Applications. <i>Journal of the American Chemical Society</i> , 2015, 137, 2303-2310.	13.7	89
6	Development of Multifunctional Pyrimidinylthiourea Derivatives as Potential Anti-Alzheimer Agents. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 8326-8344.	6.4	69
7	Identification of Novel Falcipain-2 Inhibitors as Potential Antimalarial Agents through Structure-Based Virtual Screening. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 4936-4940.	6.4	68
8	Discovery of Novel Small Molecule Inhibitors of Dengue Viral NS2B-NS3 Protease Using Virtual Screening and Scaffold Hopping. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 6278-6293.	6.4	67
9	Two-Dimensional Design Strategy to Construct Smart Fluorescent Probes for the Precise Tracking of Senescence. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 10756-10765.	13.8	65
10	Discovery and Optimization of 1,3,4-Trisubstituted-pyrazolone Derivatives as Novel, Potent, and Nonsteroidal Farnesoid X Receptor (FXR) Selective Antagonists. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 7037-7053.	6.4	61
11	JX06 Selectively Inhibits Pyruvate Dehydrogenase Kinase PDK1 by a Covalent Cysteine Modification. <i>Cancer Research</i> , 2015, 75, 4923-4936.	0.9	61
12	Chemical Structure-Related Drug-Like Criteria of Global Approved Drugs. <i>Molecules</i> , 2016, 21, 75.	3.8	61
13	Discovery of novel propargylamine-modified 4-aminoalkyl imidazole substituted pyrimidinylthiourea derivatives as multifunctional agents for the treatment of Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2018, 143, 33-47.	5.5	60
14	Organocatalytic enantioselective conjugate addition of ketones to isatylidene malononitriles. <i>Chemical Communications</i> , 2012, 48, 1692-1694.	4.1	59
15	Development of the First Generation of Disulfide-Based Subtype-Selective and Potent Covalent Pyruvate Dehydrogenase Kinase 1 (PDK1) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 2227-2244.	6.4	55
16	First-generation species-selective chemical probes for fluorescence imaging of human senescence-associated $\beta$ -galactosidase. <i>Chemical Science</i> , 2020, 11, 7292-7301.	7.4	55
17	Novel Carboline Derivatives as Potent Antifungal Lead Compounds: Design, Synthesis, and Biological Evaluation. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 506-511.	2.8	49
18	Design, Synthesis, and Biological Evaluation of Orally Available First-Generation Dual-Target Selective Inhibitors of Acetylcholinesterase (AChE) and Phosphodiesterase 5 (PDE5) for the Treatment of Alzheimer's Disease. <i>ACS Chemical Neuroscience</i> , 2018, 9, 328-345.	3.5	46

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19	Ruthenium(II)-Catalyzed Redox-Neutral [3+2] Annulation of Indoles with Internal Alkynes via C-H Bond Activation: Accessing a Pyrroloindolone Scaffold. <i>Journal of Organic Chemistry</i> , 2017, 82, 5263-5273.	3.2	45
20	Extra Sugar on Vancomycin: New Analogues for Combating Multidrug-Resistant <i>Staphylococcus aureus</i> and Vancomycin-Resistant <i>Enterococci</i> . <i>Journal of Medicinal Chemistry</i> , 2018, 61, 286-304.	6.4	45
21	Design and Synthesis of Small Molecule RhoA Inhibitors: A New Promising Therapy for Cardiovascular Diseases?. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 4508-4522.	6.4	43
22	Discovery of Potent Benzofuran-Derived Diapophytoene Desaturase (CrtN) Inhibitors with Enhanced Oral Bioavailability for the Treatment of Methicillin-Resistant <i>Staphylococcus aureus</i> (MRSA) Infections. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 3215-3230.	6.4	40
23	Spiropyran <i>in Situ</i> Switching: A Real-Time Fluorescence Strategy for Tracking DNA G-Quadruplexes in Live Cells. <i>Analytical Chemistry</i> , 2019, 91, 5354-5361.	6.5	35
24	Drug Repurposing of Haloperidol: Discovery of New Benzocyclane Derivatives as Potent Antifungal Agents against Cryptococcosis and Candidiasis. <i>ACS Infectious Diseases</i> , 2020, 6, 768-786.	3.8	35
25	Protective effects of BAY 73-6691, a selective inhibitor of phosphodiesterase 9, on amyloid- $\beta^2$ peptides-induced oxidative stress in in-vivo and in-vitro models of Alzheimer's disease. <i>Brain Research</i> , 2016, 1642, 327-335.	2.2	32
26	Novel Vilazodone-Tacrine Hybrids as Potential Multitarget-Directed Ligands for the Treatment of Alzheimer's Disease Accompanied with Depression: Design, Synthesis, and Biological Evaluation. <i>ACS Chemical Neuroscience</i> , 2017, 8, 2708-2721.	3.5	32
27	Kinetics-Driven Drug Design Strategy for Next-Generation Acetylcholinesterase Inhibitors to Clinical Candidate. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 1844-1855.	6.4	32
28	Drug Repurposing of Histone Deacetylase Inhibitors That Alleviate Neutrophilic Inflammation in Acute Lung Injury and Idiopathic Pulmonary Fibrosis via Inhibiting Leukotriene A4 Hydrolase and Blocking LTB4 Biosynthesis. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 1817-1828.	6.4	30
29	Synthesis of Highly Functionalized Chiral 3,3-Disubstituted Oxindoles via an Organocatalytic Enantioselective Michael Addition of Nitroalkanes to Indolylideneacyanoacetates. <i>Organic Letters</i> , 2012, 14, 134-137.	4.6	29
30	Policresulen, a novel NS2B/NS3 protease inhibitor, effectively inhibits the replication of DENV2 virus in BHK-21 cells. <i>Acta Pharmacologica Sinica</i> , 2015, 36, 1126-1136.	6.1	28
31	Rational Design of Novel Selective Dual-Target Inhibitors of Acetylcholinesterase and Monoamine Oxidase B as Potential Anti-Alzheimer's Disease Agents. <i>ACS Chemical Neuroscience</i> , 2019, 10, 482-496.	3.5	28
32	Selective visualization of live-cell mitochondrial thiophenols and their induced oxidative stress process by a rationally designed rhodol-based fluorescent probe. <i>Sensors and Actuators B: Chemical</i> , 2019, 283, 820-830.	7.8	28
33	Discovery of a Novel CCR5 Antagonist Lead Compound Through Fragment Assembly. <i>Molecules</i> , 2008, 13, 2426-2441.	3.8	27
34	$\text{I}^{\text{sup}}\text{â}^{\text{~}}\text{C}^{\text{sub}}\text{sp}^3\text{â}^{\text{~}}\text{N/C}^{\text{sub}}\text{sp}^2\text{â}^{\text{~}}\text{N}$ bond formation via oxidative coupling with benzophenone imine in water. <i>Green Chemistry</i> , 2015, 17, 4715-4719.	9.0	26
35	A mild and regioselective Ullmann reaction of indazoles with aryl iodides in water. <i>Tetrahedron</i> , 2017, 73, 172-178.	1.9	26
36	Structural Biology-Inspired Discovery of Novel KRAS PDE1 Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 9400-9406.	6.4	26

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37	Stress response decay with aging visualized using a dual-channel logic-based fluorescent probe. <i>Chemical Science</i> , 2021, 12, 13483-13491.	7.4	24
38	Discovery of Benzocycloalkane Derivatives Efficiently Blocking Bacterial Virulence for the Treatment of Methicillin-Resistant <i>S. aureus</i> (MRSA) Infections by Targeting Diapophytoene Desaturase (CrtN). <i>Journal of Medicinal Chemistry</i> , 2016, 59, 4831-4848.	6.4	23
39	Repurposing antimycotic ciclopirox olamine as a promising anti-ischemic stroke agent. <i>Acta Pharmaceutica Sinica B</i> , 2020, 10, 434-446.	12.0	23
40	A novel multistage antiplasmodial inhibitor targeting <i>Plasmodium falciparum</i> histone deacetylase 1. <i>Cell Discovery</i> , 2020, 6, 93.	6.7	23
41	Novel Terminal Biphenyl-Based Diapophytoene Desaturases (CrtN) Inhibitors as Anti-MRSA/VISR/LRSA Agents with Reduced hERG Activity. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 224-250.	6.4	22
42	Novel Inhibitors of Staphyloxanthin Virulence Factor in Comparison with Linezolid and Vancomycin versus Methicillin-Resistant, Linezolid-Resistant, and Vancomycin-Intermediate <i>Staphylococcus aureus</i> Infections in Vivo. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 8145-8159.	6.4	21
43	TRPA1 channel mediates organophosphate-induced delayed neuropathy. <i>Cell Discovery</i> , 2017, 3, 17024.	6.7	21
44	Discovery of Novel <i>Plasmodium falciparum</i> HDAC1 Inhibitors with Dual-Stage Antimalarial Potency and Improved Safety Based on the Clinical Anticancer Drug Candidate Quisinostat. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 2254-2271.	6.4	21
45	Lysosomal polarity increases with aging as revealed by a lysosome-targetable near-infrared fluorescent probe. <i>Sensors and Actuators B: Chemical</i> , 2020, 319, 128302.	7.8	21
46	Verapamil extends lifespan in <i>Caenorhabditis elegans</i> by inhibiting calcineurin activity and promoting autophagy. <i>Aging</i> , 2020, 12, 5300-5317.	3.1	21
47	Total synthesis of the 2-arylbenzo[b]furan-containing natural products from <i>Artocarpus</i> . <i>Tetrahedron Letters</i> , 2015, 56, 4383-4387.	1.4	20
48	Fragment-based drug discovery of triazole inhibitors to block PDE1-RAS protein-protein interaction. <i>European Journal of Medicinal Chemistry</i> , 2019, 163, 597-609.	5.5	20
49	Design, synthesis and evaluation of vilazodone-tacrine hybrids as multitarget-directed ligands against depression with cognitive impairment. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 3117-3125.	3.0	19
50	Discovery, mechanism and metabolism studies of 2,3-difluorophenyl-linker-containing PARP1 inhibitors with enhanced in vivo efficacy for cancer therapy. <i>European Journal of Medicinal Chemistry</i> , 2017, 138, 514-531.	5.5	18
51	Novel Tadalafil Derivatives Ameliorates Scopolamine-Induced Cognitive Impairment in Mice via Inhibition of Acetylcholinesterase (AChE) and Phosphodiesterase 5 (PDE5). <i>ACS Chemical Neuroscience</i> , 2018, 9, 1625-1636.	3.5	18
52	Discovery of simplified sampangine derivatives as novel fungal biofilm inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2018, 143, 1510-1523.	5.5	18
53	Discovery of nitazoxanide-based derivatives as autophagy activators for the treatment of Alzheimer's disease. <i>Acta Pharmaceutica Sinica B</i> , 2020, 10, 646-666.	12.0	18
54	Targeting virulence factors as an antimicrobial approach: Pigment inhibitors. <i>Medicinal Research Reviews</i> , 2020, 40, 293-338.	10.5	18

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55	Melting Point Distribution Analysis of Globally Approved and Discontinued Drugs: A Research for Improving the Chance of Success of Drug Design and Discovery. <i>ChemistryOpen</i> , 2016, 5, 357-368.	1.9	17
56	Design, synthesis and biological evaluation of novel antitumor spirotetrahydrothiopyran-oxindole derivatives as potent p53-MDM2 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 5268-5277.	3.0	17
57	Identification of the New Covalent Allosteric Binding Site of Fructose-1,6-bisphosphatase with Disulfiram Derivatives toward Glucose Reduction. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 6238-6247.	6.4	17
58	Design, Synthesis, and Biological Evaluation of Novel Nonsteroidal Farnesoid X Receptor (FXR) Antagonists: Molecular Basis of FXR Antagonism. <i>ChemMedChem</i> , 2015, 10, 1184-1199.	3.2	16
59	Discovery of Novel Small Molecule Anti-HCV Agents via the CypA Inhibitory Mechanism Using O-Acylation-Directed Lead Optimization. <i>Molecules</i> , 2015, 20, 10342-10359.	3.8	16
60	Design and Synthesis of Pyrophosphate-Targeting Vancomycin Derivatives for Combating Vancomycin-Resistant <i>Enterococci</i> . <i>ChemMedChem</i> , 2018, 13, 1644-1657.	3.2	16
61	Development of Novel N-hydroxypyridone Derivatives as Potential Anti-Ischemic Stroke Agents. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 1051-1067.	6.4	14
62	Direct oxidative conversion of 3-aryl propionaldehydes to 3-aryl acroleins promoted by SOMO catalysis. <i>Tetrahedron Letters</i> , 2012, 53, 1207-1209.	1.4	13
63	2-Arylbenzofuran derivatives as potent human lipoxygenase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 98-105.	5.2	13
64	Discovery of Potent, Selective Stem Cell Factor Receptor/Platelet Derived Growth Factor Receptor Alpha (c-KIT/PDGFR $\alpha$ ) Dual Inhibitor for the Treatment of Imatinib-Resistant Gastrointestinal Stromal Tumors (GISTs). <i>Journal of Medicinal Chemistry</i> , 2017, 60, 5099-5119.	6.4	13
65	Sulfoximines-Assisted Rh(III)-Catalyzed C-H Activation and Intramolecular Annulation for the Synthesis of Fused Isochromeno-1,2-Benzothiazines Scaffolds under Room Temperature. <i>Molecules</i> , 2020, 25, 2515.	3.8	13
66	Discovery of Piperidol Derivatives for Combinational Treatment of Azole-Resistant Candidiasis. <i>ACS Infectious Diseases</i> , 2021, 7, 650-660.	3.8	13
67	Synthesis of 3-substituted 1,5-aldehyde esters via an organocatalytic highly enantioselective conjugate addition of new carbonylmethyl 2-pyridinylsulfone to enals. <i>Chemical Communications</i> , 2012, 48, 148-150.	4.1	12
68	Discovery of new antimalarial agents: Second-generation dual inhibitors against FP-2 and PfDHFR via fragments assembly. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 6467-6478.	3.0	12
69	Discovery of novel piperonyl derivatives as diapophytoene desaturase inhibitors for the treatment of methicillin-, vancomycin- and linezolid-resistant <i>Staphylococcus aureus</i> infections. <i>European Journal of Medicinal Chemistry</i> , 2018, 145, 235-251.	5.5	12
70	Identification, synthesis and pharmacological evaluation of novel anti-EV71 agents via cyclophilin A inhibition. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 5682-5686.	2.2	11
71	Discovery of new Syk inhibitors through structure-based virtual screening. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 1776-1779.	2.2	11
72	Discovery of novel purine nucleoside derivatives as phosphodiesterase 2 (PDE2) inhibitors: Structure-based virtual screening, optimization and biological evaluation. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 119-133.	3.0	11

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73	Chlorpromazine Sensitizes Progesterin-Resistant Endometrial Cancer Cells to MPA by Upregulating PRB. <i>Frontiers in Oncology</i> , 2021, 11, 665832.	2.8	11
74	Specific tracking of monoamine oxidase A in heart failure models by a far-red fluorescent probe with an ultra large Stokes shift. <i>Chinese Chemical Letters</i> , 2022, 33, 1572-1576.	9.0	11
75	Small-molecule fluorescence-based probes for aging diagnosis. , 2022, 1, .		11
76	Cucurbitacin B-induced G2/M cell cycle arrest of conjunctival melanoma cells mediated by GRP78â€‘FOXM1â€‘KIF20A pathway. <i>Acta Pharmaceutica Sinica B</i> , 2022, 12, 3861-3876.	12.0	11
77	2-Amido-3-(1H-Indol-3-yl)-N-Substitued-Propanamides as a New Class of Falcipain-2 Inhibitors. 1. Design, Synthesis, Biological Evaluation and Binding Model Studies. <i>Molecules</i> , 2009, 14, 494-508.	3.8	10
78	2-(3,4-Dihydro-4-Oxothieno[2,3-d]pyrimidin-2-ylthio) Acetamides as a New Class of Falcipain-2 Inhibitors. 3. Design, Synthesis and Biological Evaluation. <i>Molecules</i> , 2009, 14, 785-797.	3.8	10
79	Luciferase Reporter Gene Assay on Human 5-HT Receptor: Which Response Element Should Be Chosen?. <i>Scientific Reports</i> , 2015, 5, 8060.	3.3	9
80	Fangchinoline suppresses conjunctival melanoma by directly binding FUBP2 and inhibiting the homologous recombination pathway. <i>Cell Death and Disease</i> , 2021, 12, 380.	6.3	9
81	Repurposing of antipsychotics perphenazine for the treatment of endometrial cancer. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127239.	2.2	9
82	Drug Repurposing of Quisinostat to Discover Novel <i>Plasmodium falciparum</i> HDAC1 Inhibitors with Enhanced Triple-Stage Antimalarial Activity and Improved Safety. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 4156-4181.	6.4	9
83	Facile construction of pyrrolo[1,2-b]isoquinolin-10(5H)-ones via a redox-aminationâ€‘aromatizationâ€‘Friedelâ€‘Crafts acylation cascade reaction and discovery of novel topoisomerase inhibitors. <i>Chemical Communications</i> , 2016, 52, 9593-9596.	4.1	8
84	Discovery of potent 2,4-difluoro-linker poly(ADP-ribose) polymerase 1 inhibitors with enhanced water solubility and in vivo anticancer efficacy. <i>Acta Pharmacologica Sinica</i> , 2017, 38, 1521-1532.	6.1	8
85	Novel Staphyloxanthin Inhibitors with Improved Potency against Multidrug Resistant <i>Staphylococcus aureus</i> . <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 233-237.	2.8	8
86	Discovery of synergistic activity of fluoroquinolones in combination with antimicrobial peptides against clinical polymyxin-resistant <i>Pseudomonas aeruginosa</i> DK2. <i>Chinese Chemical Letters</i> , 2020, 31, 413-417.	9.0	8
87	Development of disulfide-derived fructose-1,6-bisphosphatase (FBPase) covalent inhibitors for the treatment of type 2 diabetes. <i>European Journal of Medicinal Chemistry</i> , 2020, 203, 112500.	5.5	8
88	Repurposing of antitumor drug candidate Quisinostat lead to novel spirocyclic antimalarial agents. <i>Chinese Chemical Letters</i> , 2021, 32, 1660-1664.	9.0	8
89	Discovery of Novel Sertraline Derivatives as Potent Anti- <i>Cryptococcus</i> Agents. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 6541-6554.	6.4	8
90	Anti-aging effects of chlorpropamide depend on mitochondrial complex-II and the production of mitochondrial reactive oxygen species. <i>Acta Pharmaceutica Sinica B</i> , 2022, 12, 665-677.	12.0	7

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91	Discovery of novel dual RAGE/SERT inhibitors for the potential treatment of the comorbidity of Alzheimer's disease and depression. <i>European Journal of Medicinal Chemistry</i> , 2022, 236, 114347.	5.5	7
92	Two-Dimensional Design Strategy to Construct Smart Fluorescent Probes for the Precise Tracking of Senescence. <i>Angewandte Chemie</i> , 2021, 133, 10851-10860.	2.0	6
93	Rapid Repurposing of Novel Combination Drugs for the Treatment of Heart Failure via a Computationally Guided Network Screening Approach. <i>Journal of Chemical Information and Modeling</i> , 2022, 62, 5223-5232.	5.4	6
94	The novel therapeutic strategy of vilazodone-donepezil chimeras as potent triple-target ligands for the potential treatment of Alzheimer's disease with comorbid depression. <i>European Journal of Medicinal Chemistry</i> , 2022, 229, 114045.	5.5	5
95	Crotamiton derivative JM03 extends lifespan and improves oxidative and hypertonic stress resistance in <i>Caenorhabditis elegans</i> via inhibiting OSM-9. <i>ELife</i> , 2022, 11, .	6.0	5
96	Discovery of Potent Benzocycloalkane Derived Diapophytoene Desaturase Inhibitors with an Enhanced Safety Profile for the Treatment of MRSA, VISA, and LRSA Infections. <i>ACS Infectious Diseases</i> , 2018, 4, 208-217.	3.8	4
97	TSPA as a novel ATF6 translocation inducer efficiently ameliorates insulin sensitivity restoration and glucose homeostasis in db/db mice. <i>Biochemical and Biophysical Research Communications</i> , 2018, 499, 948-953.	2.1	3
98	A facile method for vancomycin C-terminus functionalization and derivatization through hydrazide. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 42, 128027.	2.2	3
99	Novel niclosamide-derived adjuvants elevating the efficacy of polymyxin B against MDR <i>Pseudomonas aeruginosa</i> DK2. <i>European Journal of Medicinal Chemistry</i> , 2022, 236, 114318.	5.5	3
100	Novel chlorpromazine derivatives as anti-endometrial carcinoma agents with reduced extrapyramidal side effects. <i>Bioorganic Chemistry</i> , 2022, 127, 106008.	4.1	3
101	Antiaging Effects of <i>Vicatia thibetica</i> de Boiss Root Extract on <i>Caenorhabditis elegans</i> and Doxorubicin-Induced Premature Aging in Adult Mice. <i>Oxidative Medicine and Cellular Longevity</i> , 2021, 2021, 1-13.	4.0	1
102	Impurity Identification and Scale-Up of a Novel Glycopeptide Antibiotic. <i>Organic Process Research and Development</i> , 2021, 25, 2390-2402.	2.7	1
103	Identification of Guaifenesin-Andrographolide as a Novel Combinatorial Drug Therapy for Epilepsy Using Network Virtual Screening and Experimental Validation. <i>ACS Chemical Neuroscience</i> , 2022, 13, 978-986.	3.5	1
104	The structure of a small GTPase RhoA in complex with PDZ RhoGEF and the inhibitor HL47. <i>Biotechnology Letters</i> , 2017, 39, 745-750.	2.2	0
105	Design and synthesis of novel hydroxamic acid derivatives based on quisinostat as promising antimalarial agents with improved safety. , 2022, 1, .		0