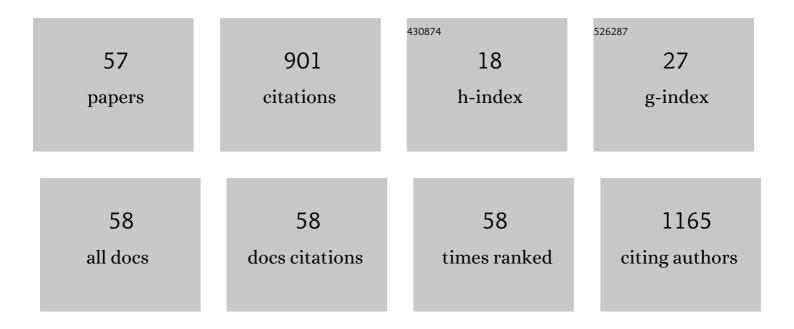
Jingwei Tian

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	H6, a novel hederagenin derivative, reverses multidrug resistance in vitro and in vivo. Toxicology and Applied Pharmacology, 2018, 341, 98-105.	2.8	82
2	Lx2-32c, a novel semi-synthetic taxane, exerts antitumor activity against prostate cancer cells in vitro and in vivo. Acta Pharmaceutica Sinica B, 2017, 7, 52-58.	12.0	62
3	Pirfenidone controls the feedback loop of the AT1R/p38 MAPK/renin-angiotensin system axis by regulating liver X receptor-α in myocardial infarction-induced cardiac fibrosis. Scientific Reports, 2017, 7, 40523.	3.3	57
4	PCC0208017, a novel small-molecule inhibitor of MARK3/MARK4, suppresses glioma progression inÂvitro and inÂvivo. Acta Pharmaceutica Sinica B, 2020, 10, 289-300.	12.0	39
5	Reciprocating friction and wear performances of nanometer <scp>sizedâ€₹iO₂</scp> filled epoxy composites. Polymer Composites, 2021, 42, 2061-2072.	4.6	38
6	Protective Effect of RA on Myocardial Infarction-Induced Cardiac Fibrosis via AT1R/p38 MAPK Pathway Signaling and Modulation of the ACE2/ACE Ratio. Journal of Agricultural and Food Chemistry, 2016, 64, 6716-6722.	5.2	36
7	Combinatorial antitumor effects of indoleamine 2,3-dioxygenase inhibitor NLG919 and paclitaxel in a murine B16-F10 melanoma model. International Journal of Immunopathology and Pharmacology, 2017, 30, 215-226.	2.1	33
8	Toll-like receptor 3 modulates the behavioral effects of cocaine in mice. Journal of Neuroinflammation, 2018, 15, 93.	7.2	32
9	Preclinical safety of ginsenoside compound K: Acute, and 26-week oral toxicity studies in mice and rats. Food and Chemical Toxicology, 2019, 131, 110578.	3.6	29
10	Acute and repeated dose 26-week oral toxicity study of 20(S)-ginsenoside Rg3 in Kunming mice and Sprague–Dawley rats. Journal of Ginseng Research, 2020, 44, 222-228.	5.7	28
11	Design, synthesis, and discovery of ocotillol-type amide derivatives as orally available modulators of P-glycoprotein-mediated multidrug resistance. European Journal of Medicinal Chemistry, 2019, 161, 118-130.	5.5	27
12	PCC0208009 enhances the anti-tumor effects of temozolomide through direct inhibition and transcriptional regulation of indoleamine 2,3-dioxygenase in glioma models. International Journal of Immunopathology and Pharmacology, 2018, 32, 205873841878799.	2.1	23
13	Cocaine induces differential circular RNA expression in striatum. Translational Psychiatry, 2019, 9, 199.	4.8	23
14	LPM580098, a Novel Triple Reuptake Inhibitor of Serotonin, Noradrenaline, and Dopamine, Attenuates Neuropathic Pain. Frontiers in Pharmacology, 2019, 10, 53.	3.5	23
15	PCC0208025 (BMS202), a small molecule inhibitor of PD-L1, produces an antitumor effect in B16-F10 melanoma-bearing mice. PLoS ONE, 2020, 15, e0228339.	2.5	22
16	Nonclinical safety of astilbin: A 4-week oral toxicity study in rats with genotoxicity, chromosomal aberration, and mammalian micronucleus tests. Food and Chemical Toxicology, 2017, 107, 1-9.	3.6	21
17	The discovery of tetrahydropyridine analogs as h Nav1.7 selective inhibitors for analgesia. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2210-2215.	2.2	19
18	Discovery of the Next-Generation Pan-TRK Kinase Inhibitors for the Treatment of Cancer. Journal of Medicinal Chemistry, 2021, 64, 10286-10296.	6.4	19

Jingwei Tian

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19	A small-molecule inhibitor of MDMX suppresses cervical cancer cells via the inhibition of E6-E6AP-p53 axis. Pharmacological Research, 2022, 177, 106128.	7.1	18
20	PCC0208027, a novel tyrosine kinase inhibitor, inhibits tumor growth of NSCLC by targeting EGFR and HER2 aberrations. Scientific Reports, 2019, 9, 5692.	3.3	17
21	Role of <scp>BRD4</scp> phosphorylation in the nucleus accumbens in relapse to cocaineâ€seeking behavior in mice. Addiction Biology, 2020, 25, e12808.	2.6	17
22	Acute, subchronic oral toxicity, and genotoxicity evaluations of LPM570065, a new potent triple reuptake inhibitor. Regulatory Toxicology and Pharmacology, 2018, 98, 129-139.	2.7	15
23	Endocannabinoid signaling regulates the reinforcing and psychostimulant effects of ketamine in mice. Nature Communications, 2020, 11, 5962.	12.8	15
24	Acute and a 28-day repeated-dose toxicity study of total flavonoids from Clinopodium chinense (Benth.) O. Ktze in mice and rats. Regulatory Toxicology and Pharmacology, 2017, 91, 117-123.	2.7	14
25	Synthesis, biological, and structural explorations of a series of μ-opioid receptor (MOR) agonists with high G protein signaling bias. European Journal of Medicinal Chemistry, 2022, 228, 113986.	5.5	14
26	Antidepressant-like Effects of LPM580153, A Novel Potent Triple Reuptake Inhibitor. Scientific Reports, 2016, 6, 24233.	3.3	12
27	A 26-week 20(S)-ginsenoside Rg3 oral toxicity study in Beagle dogs. Regulatory Toxicology and Pharmacology, 2020, 110, 104522.	2.7	12
28	PCC0208023, a potent SHP2 allosteric inhibitor, imparts an antitumor effect against KRAS mutant colorectal cancer. Toxicology and Applied Pharmacology, 2020, 398, 115019.	2.8	12
29	FNDC3B is associated with ER stress and poor prognosis in cervical cancer. Oncology Letters, 2020, 19, 406-414.	1.8	12
30	Toxicity effects of a novel potent triple reuptake inhibitor, LPM570065, on the fertility and early embryonic development in Sprague-Dawley rats. Regulatory Toxicology and Pharmacology, 2018, 100, 45-51.	2.7	11
31	Discovery and synthesis of 3- and 21-substituted fusidic acid derivatives as reversal agents of P-glycoprotein-mediated multidrug resistance. European Journal of Medicinal Chemistry, 2019, 182, 111668.	5.5	11
32	Pharmacological Characterization of Toludesvenlafaxine as a Triple Reuptake Inhibitor. Frontiers in Pharmacology, 2021, 12, 741794.	3.5	10
33	Epigenetic Mechanism of 5-HT/NE/DA Triple Reuptake Inhibitor on Adult Depression Susceptibility in Early Stress Mice. Frontiers in Pharmacology, 2022, 13, 848251.	3.5	10
34	(-)-SCR1693 Protects against Memory Impairment and Hippocampal Damage in a Chronic Cerebral Hypoperfusion Rat Model. Scientific Reports, 2016, 6, 28908.	3.3	9
35	Repeated-dose 26-week oral toxicity study of ginsenoside compound K in Beagle dogs. Journal of Ethnopharmacology, 2020, 248, 112323.	4.1	9
36	Discovery of aminocyclohexene analogues as selective and orally bioavailable h Nav1.7 inhibitors for analgesia. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 4979-4984.	2.2	7

JINGWEI TIAN

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37	Toxicological evaluation of, red rice yeast extract, Xuezhikang: Acute, 26-week chronic and genotoxicity studies. Regulatory Toxicology and Pharmacology, 2020, 114, 104654.	2.7	7
38	Mechanical, bonding and tribological performances of epoxyâ€based nanocomposite coatings with multiple fillers. Journal of Applied Polymer Science, 2022, 139, .	2.6	7
39	4R Tau Modulates Cocaine-Associated Memory through Adult Dorsal Hippocampal Neurogenesis. Journal of Neuroscience, 2021, 41, 6753-6774.	3.6	6
40	Comparison study of different indoleamine-2,3 dioxygenase inhibitors from the perspective of pharmacodynamic effects. International Journal of Immunopathology and Pharmacology, 2020, 34, 205873842095058.	2.1	5
41	Improving the treatment of Parkinson's disease: Structure-based development of novel 5-HT2A receptor antagonists/inverse agonists. European Journal of Medicinal Chemistry, 2022, 234, 114246.	5.5	5
42	A study of D-allulose-associated reproductive toxicity in rats. Food and Chemical Toxicology, 2019, 131, 110548.	3.6	4
43	PCC0208018 exerts antitumor effects by activating effector T cells. International Journal of Immunopathology and Pharmacology, 2019, 33, 205873841984336.	2.1	4
44	PCC-0105002, a novel small molecule inhibitor of PSD95-nNOS protein-protein interactions, attenuates neuropathic pain and corrects motor disorder associated with neuropathic pain model. Toxicology and Applied Pharmacology, 2021, 429, 115698.	2.8	4
45	LPM3770277, a Potent Novel CDK4/6 Degrader, Exerts Antitumor Effect Against Triple-Negative Breast Cancer. Frontiers in Pharmacology, 2022, 13, 853993.	3.5	4
46	Synthesis and analysis of dihydrotetrabenazine derivatives as novel vesicular monoamine transporter 2 inhibitors. European Journal of Medicinal Chemistry, 2021, 224, 113718.	5.5	3
47	Lipidomics Reveals Dysregulated Glycerophospholipid Metabolism in the Corpus Striatum of Mice Treated with Cefepime. ACS Chemical Neuroscience, 2021, 12, 4449-4464.	3.5	3
48	Pharmacokinetics of S-EPA, and its inhibition on indoleamine 2,3-dioxgenase: a case of sulfur-substitution affecting distributions in blood cells. Xenobiotica, 2019, 49, 1338-1343.	1.1	2
49	Assessment of the toxicity and toxicokinetics of the novel potent tropomyosin receptor kinase (Trk) inhibitor LPM4870108 in rhesus monkeys. Regulatory Toxicology and Pharmacology, 2021, 122, 104886.	2.7	2
50	Integrated lipidomic and transcriptomic analysis reveals clarithromycin-induced alteration of glycerophospholipid metabolism in the cerebral cortex of mice. Cell Biology and Toxicology, 2023, 39, 771-793.	5.3	2
51	Neonatal exposure to sevoflurane induces adolescent neurobehavioral dysfunction by interfering with hippocampal glycerophoslipid metabolism in rats. Cerebral Cortex, 2023, 33, 1955-1971.	2.9	2
52	Pharmacokinetics of Sâ€epacadostat, an indoleamine 2,3â€dioxygenase 1 inhibitor, in dog plasma and identification of its metabolites in vivo and in vitro. Biomedical Chromatography, 2021, 35, e5226.	1.7	1
53	The tyrosine kinase inhibitor LPM4870108 impairs learning and memory and induces transcriptomic and gene‑specific DNA methylation changes in rats. Archives of Toxicology, 2022, 96, 845-857.	4.2	1
54	(+)-9-Trifluoroethoxy-α-Dihydrotetrabenazine as a Highly Potent Vesicular Monoamine Transporter 2 Inhibitor for Tardive Dyskinesia. Frontiers in Pharmacology, 2021, 12, 770377.	3.5	1

JINGWEI TIAN

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55	In Vitro and In Vivo Characterization of PCC0104005, a Novel Modulator of Serotonin-Dopamine Activity, as an Atypical Antipsychotic Drug. Scientific Reports, 2018, 8, 6892.	3.3	0
56	A 26-week toxicological study of Xuezhikang (XZK), red yeast rice extract, in Beagle dogs with a 4-week recovery period. Regulatory Toxicology and Pharmacology, 2020, 117, 104781.	2.7	0
57	9-Cyclopropylmethoxy-dihydrotetrabenazine and its stereoisomers as vesicular monoamine transporter-2 inhibitors. Future Medicinal Chemistry, 2022, 14, 991-1003.	2.3	Ο