Xuanxuan Dai

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	α-Klotho is a non-enzymatic molecular scaffold for FGF23 hormone signalling. Nature, 2018, 553, 461-466.	27.8	348
2	Exploration and synthesis of curcumin analogues with improved structural stability both in vitro and in vivo as cytotoxic agents. Bioorganic and Medicinal Chemistry, 2009, 17, 2623-2631.	3.0	288
3	Dihydroartemisinin (DHA) induces ferroptosis and causes cell cycle arrest in head and neck carcinoma cells. Cancer Letters, 2016, 381, 165-175.	7.2	227
4	Feedback Activation of STAT3 as a Cancer Drug-Resistance Mechanism. Trends in Pharmacological Sciences, 2016, 37, 47-61.	8.7	190
5	Evaluation and Discovery of Novel Synthetic Chalcone Derivatives as Anti-Inflammatory Agents. Journal of Medicinal Chemistry, 2011, 54, 8110-8123.	6.4	182
6	Saturated palmitic acid induces myocardial inflammatory injuries through direct binding to TLR4 accessory protein MD2. Nature Communications, 2017, 8, 13997.	12.8	166
7	Inhibition of JNK Phosphorylation by a Novel Curcumin Analog Prevents High Glucose–Induced Inflammation and Apoptosis in Cardiomyocytes and the Development of Diabetic Cardiomyopathy. Diabetes, 2014, 63, 3497-3511.	0.6	160
8	Luteolin protects against diabetic cardiomyopathy by inhibiting NF-κB-mediated inflammation and activating the Nrf2-mediated antioxidant responses. Phytomedicine, 2019, 59, 152774.	5.3	157
9	The role of sphingosine 1â€phosphate receptor 2 in bileâ€acid–induced cholangiocyte proliferation and cholestasisâ€induced liver injury in mice. Hepatology, 2017, 65, 2005-2018.	7.3	153
10	Cholangiocyteâ€Đerived Exosomal Long Noncoding RNA H19 Promotes Hepatic Stellate Cell Activation and Cholestatic Liver Fibrosis. Hepatology, 2019, 70, 1317-1335.	7.3	150
11	Inhibition of high glucoseâ€induced inflammatory response and macrophage infiltration by a novel curcumin derivative prevents renal injury in diabetic rats. British Journal of Pharmacology, 2012, 166, 1169-1182.	5.4	142
12	Curcumin protects hearts from FFA-induced injury by activating Nrf2 and inactivating NF-κB both in vitro and in vivo. Journal of Molecular and Cellular Cardiology, 2015, 79, 1-12.	1.9	141
13	Celastrol Attenuates Angiotensin Il–Induced Cardiac Remodeling by Targeting STAT3. Circulation Research, 2020, 126, 1007-1023.	4.5	127
14	ROS generation mediates the anti-cancer effects of WZ35 via activating JNK and ER stress apoptotic pathways in gastric cancer. Oncotarget, 2015, 6, 5860-5876.	1.8	126
15	Synthesis and anti-inflammatory activities of mono-carbonyl analogues of curcumin. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 1525-1529.	2.2	123
16	Synthesis, crystal structure and anti-inflammatory properties of curcumin analogues. European Journal of Medicinal Chemistry, 2009, 44, 915-919.	5.5	117
17	Fibroblast growth factor 1 ameliorates diabetic nephropathy by an anti-inflammatory mechanism. Kidney International, 2018, 93, 95-109.	5.2	117
18	Piperlongumine as a direct TrxR1 inhibitor with suppressive activity against gastric cancer. Cancer Letters, 2016, 375, 114-126.	7.2	115

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19	Celastrol induces ROS-mediated apoptosis via directly targeting peroxiredoxin-2 in gastric cancer cells. Theranostics, 2020, 10, 10290-10308.	10.0	113
20	Auranofin induces apoptosis by ROS-mediated ER stress and mitochondrial dysfunction and displayed synergistic lethality with piperlongumine in gastric cancer. Oncotarget, 2015, 6, 36505-36521.	1.8	111
21	Anticancer molecules targeting fibroblast growth factor receptors. Trends in Pharmacological Sciences, 2012, 33, 531-541.	8.7	110
22	Recent Progress of Small-Molecule Epidermal Growth Factor Receptor (EGFR) Inhibitors against C797S Resistance in Non-Small-Cell Lung Cancer. Journal of Medicinal Chemistry, 2018, 61, 4290-4300.	6.4	102
23	miR-196b-5p–mediated downregulation of TSPAN12 and GATA6 promotes tumor progression in non-small cell lung cancer. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 4347-4357.	7.1	95
24	FGF21 improves cognition by restored synaptic plasticity, dendritic spine density, brain mitochondrial function and cell apoptosis in obese-insulin resistant male rats. Hormones and Behavior, 2016, 85, 86-95.	2.1	92
25	Synthesis and Anti-inflammatory Evaluation of Novel Benzimidazole and Imidazopyridine Derivatives. ACS Medicinal Chemistry Letters, 2013, 4, 69-74.	2.8	91
26	MD2 activation by direct AGE interaction drives inflammatory diabetic cardiomyopathy. Nature Communications, 2020, 11, 2148.	12.8	90
27	Synthesis and biological evaluation of allylated and prenylated mono-carbonyl analogs of curcumin as anti-inflammatory agents. European Journal of Medicinal Chemistry, 2014, 74, 671-682.	5.5	89
28	Novel curcumin analog C66 prevents diabetic nephropathy via JNK pathway with the involvement of p300/CBP-mediated histone acetylation. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2015, 1852, 34-46.	3.8	86
29	iRGD decorated lipid-polymer hybrid nanoparticles for targeted co-delivery of doxorubicin and sorafenib to enhance anti-hepatocellular carcinoma efficacy. Nanomedicine: Nanotechnology, Biology, and Medicine, 2016, 12, 1303-1311.	3.3	86
30	MD2 mediates angiotensin II-induced cardiac inflammation and remodeling via directly binding to Ang II and activating TLR4/NF-κB signaling pathway. Basic Research in Cardiology, 2017, 112, 9.	5.9	84
31	C66 ameliorates diabetic nephropathy in mice by both upregulating NRF2 function via increase in miR-200a and inhibiting miR-21. Diabetologia, 2016, 59, 1558-1568.	6.3	81
32	Curcumin derivative WZ35 inhibits tumor cell growth via ROS-YAP-JNK signaling pathway in breast cancer. Journal of Experimental and Clinical Cancer Research, 2019, 38, 460.	8.6	75
33	Inhibition of STAT3 in tubular epithelial cells prevents kidney fibrosis and nephropathy in STZ-induced diabetic mice. Cell Death and Disease, 2019, 10, 848.	6.3	75
34	Anti-inflammatory effects of novel curcumin analogs in experimental acute lung injury. Respiratory Research, 2015, 16, 43.	3.6	73
35	Angiotensin II induces kidney inflammatory injury and fibrosis through binding to myeloid differentiation protein-2 (MD2). Scientific Reports, 2017, 7, 44911.	3.3	73
36	Kaempferol attenuates hyperglycemia-induced cardiac injuries by inhibiting inflammatory responses and oxidative stress. Endocrine, 2018, 60, 83-94.	2.3	72

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37	Synergistic antitumor activity of rapamycin and EF24 via increasing ROS for the treatment of gastric cancer. Redox Biology, 2016, 10, 78-89.	9.0	70
38	<scp>MD</scp> â€2 as the target of a novel small molecule, <scp>L6H</scp> 21, in the attenuation of <scp>LPS</scp> â€induced inflammatory response and sepsis. British Journal of Pharmacology, 2015, 172, 4391-4405.	5.4	69
39	Novel curcumin analogue 14p protects against myocardial ischemia reperfusion injury through Nrf2-activating anti-oxidative activity. Toxicology and Applied Pharmacology, 2015, 282, 175-183.	2.8	69
40	Licochalcone A Inhibits the Proliferation of Human Lung Cancer Cell Lines A549 and H460 by Inducing G2/M Cell Cycle Arrest and ER Stress. International Journal of Molecular Sciences, 2017, 18, 1761.	4.1	69
41	Inhibition of LPSâ€induced production of inflammatory factors in the macrophages by mono arbonyl analogues of curcumin. Journal of Cellular and Molecular Medicine, 2009, 13, 3370-3379.	3.6	68
42	Curcumin analog EF24 induces apoptosis via ROS-dependent mitochondrial dysfunction in human colorectal cancer cells. Cancer Chemotherapy and Pharmacology, 2016, 78, 1151-1161.	2.3	65
43	Inhibition of epidermal growth factor receptor attenuates atherosclerosis via decreasing inflammation and oxidative stress. Scientific Reports, 2017, 7, 45917.	3.3	65
44	Curcumin Analog L48H37 Prevents Lipopolysaccharide-Induced TLR4 Signaling Pathway Activation and Sepsis via Targeting MD2. Journal of Pharmacology and Experimental Therapeutics, 2015, 353, 539-550.	2.5	64
45	MicroRNA-224 is implicated in lung cancer pathogenesis through targeting caspase-3 and caspase-7. Oncotarget, 2015, 6, 21802-21815.	1.8	63
46	Rhein sensitizes human pancreatic cancer cells to EGFR inhibitors by inhibiting STAT3 pathway. Journal of Experimental and Clinical Cancer Research, 2019, 38, 31.	8.6	63
47	Design, Synthesis, and Structure–Activity Relationship Study of Novel Indole-2-carboxamide Derivatives as Anti-inflammatory Agents for the Treatment of Sepsis. Journal of Medicinal Chemistry, 2016, 59, 4637-4650.	6.4	61
48	Curcumin suppresses gastric tumor cell growth via ROS-mediated DNA polymerase γ depletion disrupting cellular bioenergetics. Journal of Experimental and Clinical Cancer Research, 2017, 36, 47.	8.6	61
49	Blockage of ROS and NF-l [®] B-mediated inflammation by a new chalcone L6H9 protects cardiomyocytes from hyperglycemia-induced injuries. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2015, 1852, 1230-1241.	3.8	60
50	Curcuminoid B63 induces ROS-mediated paraptosis-like cell death by targeting TrxR1 in gastric cells. Redox Biology, 2019, 21, 101061.	9.0	60
51	Shikonin inhibits myeloid differentiation protein 2 to prevent LPSâ€induced acute lung injury. British Journal of Pharmacology, 2018, 175, 840-854.	5.4	59
52	Small molecule inhibition of fibroblast growth factor receptors in cancer. Cytokine and Growth Factor Reviews, 2013, 24, 467-475.	7.2	58
53	Discovery and evaluation of piperid-4-one-containing mono-carbonyl analogs of curcumin as anti-inflammatory agents. Bioorganic and Medicinal Chemistry, 2013, 21, 3058-3065.	3.0	58
54	Novel <scp>EGFR</scp> inhibitors attenuate cardiac hypertrophy induced by angiotensin II. Journal of Cellular and Molecular Medicine, 2016, 20, 482-494.	3.6	58

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55	EGFR Inhibition Blocks Palmitic Acid-induced inflammation in cardiomyocytes and Prevents Hyperlipidemia-induced Cardiac Injury in Mice. Scientific Reports, 2016, 6, 24580.	3.3	58
56	Oncogenic KRAS Reduces Expression of FGF21 in Acinar Cells to Promote Pancreatic Tumorigenesis in Mice on a High-Fat Diet. Gastroenterology, 2019, 157, 1413-1428.e11.	1.3	57
57	Inhibition of JNK by novel curcumin analog C66 prevents diabetic cardiomyopathy with a preservation of cardiac metallothionein expression. American Journal of Physiology - Endocrinology and Metabolism, 2014, 306, E1239-E1247.	3.5	54
58	Piperlongumine, a Novel TrxR1 Inhibitor, Induces Apoptosis in Hepatocellular Carcinoma Cells by ROS-Mediated ER Stress. Frontiers in Pharmacology, 2019, 10, 1180.	3.5	54
59	Discovery of a New Inhibitor of Myeloid Differentiation 2 from Cinnamamide Derivatives with Anti-Inflammatory Activity in Sepsis and Acute Lung Injury. Journal of Medicinal Chemistry, 2016, 59, 2436-2451.	6.4	52
60	Inhibition of high glucose-induced inflammation and fibrosis by a novel curcumin derivative prevents renal and heart injury in diabetic mice. Toxicology Letters, 2017, 278, 48-58.	0.8	52
61	New MD2 inhibitors derived from curcumin with improved anti-inflammatory activity. European Journal of Medicinal Chemistry, 2018, 148, 291-305.	5.5	52
62	Myeloid Differentiation Primary Response Protein 88 (MyD88): The Central Hub of TLR/IL-1R Signaling. Journal of Medicinal Chemistry, 2020, 63, 13316-13329.	6.4	52
63	EGFR inhibition protects cardiac damage and remodeling through attenuating oxidative stress in STZ-induced diabetic mouse model. Journal of Molecular and Cellular Cardiology, 2015, 82, 63-74.	1.9	51
64	Synthesis and optimization of novel allylated mono-carbonyl analogs of curcumin (MACs) act as potent anti-inflammatory agents against LPS-induced acute lung injury (ALI) in rats. European Journal of Medicinal Chemistry, 2016, 121, 181-193.	5.5	51
65	Enhancement of oxaliplatin-induced colon cancer cell apoptosis by alantolactone, a natural product inducer of ROS. International Journal of Biological Sciences, 2019, 15, 1676-1684.	6.4	51
66	miR-130a Deregulates PTEN and Stimulates Tumor Growth. Cancer Research, 2017, 77, 6168-6178.	0.9	50
67	Osthole inhibits triple negative breast cancer cells by suppressing STAT3. Journal of Experimental and Clinical Cancer Research, 2018, 37, 322.	8.6	50
68	Inhibition of EGFR attenuates fibrosis and stellate cell activation in diet-induced model of nonalcoholic fatty liver disease. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2018, 1864, 133-142.	3.8	48
69	Rational combination of MEK inhibitor and the STAT3 pathway modulator for the therapy in K-Ras mutated pancreatic and colon cancer cells. Oncotarget, 2015, 6, 14472-14487.	1.8	46
70	Selective killing of gastric cancer cells by a small molecule via targeting TrxR1 and ROS-mediated ER stress activation. Oncotarget, 2016, 7, 16593-16609.	1.8	46
71	Discovery of new MD2 inhibitor from chalcone derivatives with anti-inflammatory effects in LPS-induced acute lung injury. Scientific Reports, 2016, 6, 25130.	3.3	45
72	Metal-free regioselective C–H chalcogenylation of coumarins/(hetero)arenes at ambient temperature. Chemical Communications, 2020, 56, 1847-1850.	4.1	45

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73	EF24 induces ROS-mediated apoptosis <i>via</i> targeting <i>thioredoxin reductase 1</i> in gastric cancer cells. Oncotarget, 2016, 7, 18050-18064.	1.8	45
74	The Prevention of Diabetic Cardiomyopathy by Non-Mitogenic Acidic Fibroblast Growth Factor Is Probably Mediated by the Suppression of Oxidative Stress and Damage. PLoS ONE, 2013, 8, e82287.	2.5	44
75	Metabolism-Associated Molecular Patterns (MAMPs). Trends in Endocrinology and Metabolism, 2020, 31, 712-724.	7.1	44
76	Angiotensin II Causes Biphasic STAT3 Activation Through TLR4 to Initiate Cardiac Remodeling. Hypertension, 2018, 72, 1301-1311.	2.7	43
77	Escherichia coli adhesion portion FimH functions as an adjuvant for cancer immunotherapy. Nature Communications, 2020, 11, 1187.	12.8	43
78	FGF1ΔHBS prevents diabetic cardiomyopathy by maintaining mitochondrial homeostasis and reducing oxidative stress via AMPK/Nur77 suppression. Signal Transduction and Targeted Therapy, 2021, 6, 133.	17.1	43
79	Curcuminoid EF24 enhances the antiâ€ŧumour activity of Akt inhibitor MKâ€⊋206 through ROSâ€mediated endoplasmic reticulum stress and mitochondrial dysfunction in gastric cancer. British Journal of Pharmacology, 2017, 174, 1131-1146.	5.4	42
80	Alantolactone sensitizes human pancreatic cancer cells to EGFR inhibitors through the inhibition of STAT3 signaling. Molecular Carcinogenesis, 2019, 58, 565-576.	2.7	42
81	Selective Targeting of Cancer Cells by Oxidative Vulnerabilities with Novel Curcumin Analogs. Scientific Reports, 2017, 7, 1105.	3.3	41
82	Schisandrin B alleviates diabetic nephropathy through suppressing excessive inflammation and oxidative stress. Biochemical and Biophysical Research Communications, 2019, 508, 243-249.	2.1	41
83	Macrophage-derived myeloid differentiation protein 2 plays an essential role in ox-LDL-induced inflammation and atherosclerosis. EBioMedicine, 2020, 53, 102706.	6.1	41
84	Isoliquiritigenin attenuates diabetic cardiomyopathy via inhibition of hyperglycemia-induced inflammatory response and oxidative stress. Phytomedicine, 2020, 78, 153319.	5.3	40
85	Base promoted synthesis of novel indole-dithiocarbamate compounds as potential anti-inflammatory therapeutic agents for treatment of acute lung injury. European Journal of Medicinal Chemistry, 2019, 171, 54-65.	5.5	39
86	Development of 2-amino-4-phenylthiazole analogues to disrupt myeloid differentiation factor 88 and prevent inflammatory responses in acute lung injury. European Journal of Medicinal Chemistry, 2019, 161, 22-38.	5.5	39
87	A Newly Designed Curcumin Analog Y20 Mitigates Cardiac Injury via Anti-Inflammatory and Anti-Oxidant Actions in Obese Rats. PLoS ONE, 2015, 10, e0120215.	2.5	38
88	Curcumin analog L48H37 induces apoptosis through ROSâ€mediated endoplasmic reticulum stress and STAT3 pathways in human lung cancer cells. Molecular Carcinogenesis, 2017, 56, 1765-1777.	2.7	38
89	Costunolide specifically binds and inhibits thioredoxin reductase 1 to induce apoptosis in colon cancer. Cancer Letters, 2018, 412, 46-58.	7.2	38
90	miRNA-mediated TUSC3 deficiency enhances UPR and ERAD to promote metastatic potential of NSCLC. Nature Communications, 2018, 9, 5110.	12.8	38

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91	Kaempferol reduces K63-linked polyubiquitination to inhibit nuclear factor-κB and inflammatory responses in acute lung injury in mice. Toxicology Letters, 2019, 306, 53-60.	0.8	38
92	Ruthenium(II)-Catalyzed C–H Activation of Chromones with Maleimides to Synthesize Succinimide/Maleimide-Containing Chromones. Journal of Organic Chemistry, 2020, 85, 9230-9243.	3.2	38
93	Design, Synthesis, and Biological Evaluation of Novel Quinazoline Derivatives as Antiâ€inflammatory Agents against Lipopolysaccharideâ€induced Acute Lung Injury in Rats. Chemical Biology and Drug Design, 2015, 85, 672-684.	3.2	37
94	A novel chalcone derivative attenuates the diabetes-induced renal injury via inhibition of high glucose-mediated inflammatory response and macrophage infiltration. Toxicology and Applied Pharmacology, 2015, 282, 129-138.	2.8	37
95	Discovery of 3-(Indol-5-yl)-indazole Derivatives as Novel Myeloid Differentiation Protein 2/Toll-like Receptor 4 Antagonists for Treatment of Acute Lung Injury. Journal of Medicinal Chemistry, 2019, 62, 5453-5469.	6.4	37
96	FGF21 and DPP-4 inhibitor equally prevents cognitive decline in obese rats. Biomedicine and Pharmacotherapy, 2018, 97, 1663-1672.	5.6	36
97	CHK2-FOXK axis promotes transcriptional control of autophagy programs. Science Advances, 2020, 6, eaax5819.	10.3	36
98	Pattern recognition receptorâ€mediated inflammation in diabetic vascular complications. Medicinal Research Reviews, 2020, 40, 2466-2484.	10.5	36
99	Exercise-Induced Irisin Decreases Inflammation and Improves NAFLD by Competitive Binding with MD2. Cells, 2021, 10, 3306.	4.1	36
100	Recent progress in the discovery of myeloid differentiation 2 (MD2) modulators for inflammatory diseases. Drug Discovery Today, 2018, 23, 1187-1202.	6.4	35
101	An Aza resveratrol–chalcone derivative 6b protects mice against diabetic cardiomyopathy by alleviating inflammation and oxidative stress. Journal of Cellular and Molecular Medicine, 2018, 22, 1931-1943.	3.6	35
102	Schisandrin A inhibits triple negative breast cancer cells by regulating Wnt/ER stress signaling pathway. Biomedicine and Pharmacotherapy, 2019, 115, 108922.	5.6	35
103	Design and synthesis novel di-carbonyl analogs of curcumin (DACs) act as potent anti-inflammatory agents against LPS-induced acute lung injury (ALI). European Journal of Medicinal Chemistry, 2019, 167, 414-425.	5.5	35
104	EGFR mediates hyperlipidemia-induced renal injury via regulating inflammation and oxidative stress: the detrimental role and mechanism of EGFR activation. Oncotarget, 2016, 7, 24361-24373.	1.8	34
105	Arachidonic acid inhibits inflammatory responses by binding to myeloid differentiation factor-2 (MD2) and preventing MD2/toll-like receptor 4 signaling activation. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2020, 1866, 165683.	3.8	34
106	Evaluation of a curcumin analog as an anti-cancer agent inducing ER stress-mediated apoptosis in non-small cell lung cancer cells. BMC Cancer, 2013, 13, 494.	2.6	33
107	Selective killing of gastric cancer cells by a small molecule targeting ROSâ€mediated ER stress activation. Molecular Carcinogenesis, 2016, 55, 1073-1086.	2.7	33
108	Inhibition of myeloid differentiation factor 2 by baicalein protects against acute lung injury. Phytomedicine, 2019, 63, 152997.	5.3	33

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109	Discovery and identification of new non-ATP competitive FGFR1 inhibitors with therapeutic potential on non-small-cell lung cancer. Cancer Letters, 2014, 344, 82-89.	7.2	32
110	EGFR inhibition attenuates diabetic nephropathy through decreasing ROS and endoplasmic reticulum stress. Oncotarget, 2017, 8, 32655-32667.	1.8	32
111	Discovery of caffeic acid phenethyl ester derivatives as novel myeloid differentiation protein 2 inhibitors for treatment of acute lung injury. European Journal of Medicinal Chemistry, 2018, 143, 361-375.	5.5	32
112	<p>Rhein shows potent efficacy against non-small-cell lung cancer through inhibiting the STAT3 pathway</p> . Cancer Management and Research, 2019, Volume 11, 1167-1176.	1.9	32
113	Kaempferol attenuates streptozotocin-induced diabetic nephropathy by downregulating TRAF6 expression: The role of TRAF6 in diabetic nephropathy. Journal of Ethnopharmacology, 2021, 268, 113553.	4.1	32
114	Tetrahydroisoquinoline-7-carboxamide Derivatives as New Selective Discoidin Domain Receptor 1 (DDR1) Inhibitors. ACS Medicinal Chemistry Letters, 2017, 8, 327-332.	2.8	31
115	Synthesis and evaluation of asymmetric curcuminoid analogs as potential anticancer agents that downregulate NF-ήB activation and enhance the sensitivity of gastric cancer cell lines to irinotecan chemotherapy. European Journal of Medicinal Chemistry, 2017, 139, 917-925.	5.5	31
116	Schisandrin B exhibits potent anticancer activity in triple negative breast cancer by inhibiting STAT3. Toxicology and Applied Pharmacology, 2018, 358, 110-119.	2.8	31
117	Curcuminoid WZ35 synergize with cisplatin by inducing ROS production and inhibiting TrxR1 activity in gastric cancer cells. Journal of Experimental and Clinical Cancer Research, 2019, 38, 207.	8.6	31
118	Blockade of myeloid differentiation 2 attenuates diabetic nephropathy by reducing activation of the reninâ€angiotensin system in mouse kidneys. British Journal of Pharmacology, 2019, 176, 2642-2657.	5.4	31
119	Molecular basis for receptor tyrosine kinase A-loop tyrosine transphosphorylation. Nature Chemical Biology, 2020, 16, 267-277.	8.0	31
120	A novel fibroblast growth factor receptor 1 inhibitor protects against cartilage degradation in a murine model of osteoarthritis. Scientific Reports, 2016, 6, 24042.	3.3	30
121	Osthole Protects against Acute Lung Injury by Suppressing NF- <i>κ</i> B-Dependent Inflammation. Mediators of Inflammation, 2018, 2018, 1-12.	3.0	30
122	Acetyl-bufalin shows potent efficacy against non-small-cell lung cancer by targeting the CDK9/STAT3 signalling pathway. British Journal of Cancer, 2021, 124, 645-657.	6.4	30
123	Chemopreventive effect of chalcone derivative, L2H17, in colon cancer development. BMC Cancer, 2015, 15, 870.	2.6	29
124	Niclosamide inhibition of STAT3 synergizes with erlotinib in human colon cancer. OncoTargets and Therapy, 2017, Volume 10, 1767-1776.	2.0	29
125	Fibroblast growth factor-21 restores insulin sensitivity but induces aberrant bone microstructure in obese insulin-resistant rats. Journal of Bone and Mineral Metabolism, 2017, 35, 142-149.	2.7	28
126	Increased Intracellular Reactive Oxygen Species Mediates the Anti ancer Effects of WZ35 via Activating Mitochondrial Apoptosis Pathway in Prostate Cancer Cells. Prostate, 2017, 77, 489-504.	2.3	28

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127	Alantolactone promotes ER stressâ€mediated apoptosis by inhibition of TrxR1 in tripleâ€negative breast cancer cell lines and in a mouse model. Journal of Cellular and Molecular Medicine, 2019, 23, 2194-2206.	3.6	28
128	Baicalein attenuates OVA-induced allergic airway inflammation through the inhibition of the NF-κB signaling pathway. Aging, 2019, 11, 9310-9327.	3.1	28
129	Discovery and evaluation of asymmetrical monocarbonyl analogs of curcumin as anti-inflammatory agents. Drug Design, Development and Therapy, 2014, 8, 373.	4.3	27
130	Synthesis and biological evaluation of a novel class of curcumin analogs as anti-inflammatory agents for prevention and treatment of sepsis in mouse model. Drug Design, Development and Therapy, 2015, 9, 1663.	4.3	27
131	(S)-crizotinib induces apoptosis in human non-small cell lung cancer cells by activating ROS independent of MTH1. Journal of Experimental and Clinical Cancer Research, 2017, 36, 120.	8.6	27
132	Novel allylated monocarbonyl analogs of curcumin induce mitotic arrest and apoptosis by reactive oxygen species-mediated endoplasmic reticulum stress and inhibition of STAT3. Oncotarget, 2017, 8, 101112-101129.	1.8	27
133	Schizandrin B attenuates angiotensin II induced endothelial to mesenchymal transition in vascular endothelium by suppressing NF-κB activation. Phytomedicine, 2019, 62, 152955.	5.3	27
134	Blockage of ROS and MAPKs-mediated inflammation via restoring SIRT1 by a new compound LF10 prevents type 1 diabetic cardiomyopathy. Toxicology and Applied Pharmacology, 2019, 370, 24-35.	2.8	27
135	Synthesis and biological evaluation of novel oxindole-based RTK inhibitors as anti-cancer agents. Bioorganic and Medicinal Chemistry, 2014, 22, 6953-6960.	3.0	26
136	Design, synthesis and biological evaluation of paralleled Aza resveratrol–chalcone compounds as potential anti-inflammatory agents for the treatment of acute lung injury. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 2998-3004.	2.2	26
137	Inhibition of inflammation and oxidative stress by an imidazopyridine derivative X22 prevents heart injury from obesity. Journal of Cellular and Molecular Medicine, 2016, 20, 1427-1442.	3.6	26
138	Design, Synthesis, and Structure–Activity Relationship Analysis of Thiazolo[3,2â€ <i>a</i>]pyrimidine Derivatives with Antiâ€inflammatory Activity in Acute Lung Injury. ChemMedChem, 2017, 12, 1022-1032.	3.2	26
139	A self-assembling luminescent lanthanide molecular nanoparticle with potential for liveÂcell imaging. Chemical Science, 2018, 9, 4630-4637.	7.4	26
140	An anti-inflammatory chalcone derivative prevents heart and kidney from hyperlipidemia-induced injuries by attenuating inflammation. Toxicology and Applied Pharmacology, 2018, 338, 43-53.	2.8	26
141	Selective targeting of the TLR4 co-receptor, MD2, prevents colon cancer growth and lung metastasis. International Journal of Biological Sciences, 2020, 16, 1288-1301.	6.4	26
142	Feedback activation of EGFR is the main cause for STAT3 inhibition-irresponsiveness in pancreatic cancer cells. Oncogene, 2020, 39, 3997-4013.	5.9	26
143	Attenuation of inflammatory response by a novel chalcone protects kidney and heart from hyperglycemia-induced injuries in type 1 diabetic mice. Toxicology and Applied Pharmacology, 2015, 288, 179-191.	2.8	25
144	11β-Hydroxysteroid Dehydrogenase Type 1(11β-HSD1) mediates insulin resistance through JNK activation in adipocytes. Scientific Reports, 2016, 6, 37160.	3.3	25

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145	Small-molecule RL71-triggered excessive autophagic cell death as a potential therapeutic strategy in triple-negative breast cancer. Cell Death and Disease, 2017, 8, e3049-e3049.	6.3	25
146	Biomimetic synthesis of the natural product salviadione and its hybrids: discovery of tissue-specific anti-inflammatory agents for acute lung injury. Chemical Science, 2019, 10, 4667-4672.	7.4	25
147	Novel Epidermal Growth Factor Receptor Inhibitor Attenuates Angiotensin II-Induced Kidney Fibrosis. Journal of Pharmacology and Experimental Therapeutics, 2015, 356, 32-42.	2.5	24
148	Insights into the binding mode of curcumin to MD-2: studies from molecular docking, molecular dynamics simulations and experimental assessments. Molecular BioSystems, 2015, 11, 1933-1938.	2.9	24
149	A novel STAT3 inhibitor HO-3867 induces cell apoptosis by reactive oxygen species-dependent endoplasmic reticulum stress in human pancreatic cancer cells. Anti-Cancer Drugs, 2017, 28, 392-400.	1.4	24
150	Development of resveratrol-curcumin hybrids as potential therapeutic agents for inflammatory lung diseases. European Journal of Medicinal Chemistry, 2017, 125, 478-491.	5.5	24
151	<p>Acid-responsive nanoparticles as a novel oxidative stress-inducing anticancer therapeutic agent for colon cancer</p> . International Journal of Nanomedicine, 2019, Volume 14, 1597-1618.	6.7	24
152	Synthesis and Evaluation of a Series of Novel Asymmetrical Curcumin Analogs for the Treatment of Inflammation. Molecules, 2014, 19, 7287-7307.	3.8	23
153	Design, synthesis, and anticancer evaluation of long-chain alkoxylated mono-carbonyl analogues of curcumin. European Journal of Medicinal Chemistry, 2015, 103, 44-55.	5.5	23
154	A novel MyD88 inhibitor LM9 prevents atherosclerosis by regulating inflammatory responses and oxidative stress in macrophages. Toxicology and Applied Pharmacology, 2019, 370, 44-55.	2.8	23
155	Direct cardio-protection of Dapagliflozin against obesity-related cardiomyopathy via NHE1/MAPK signaling. Acta Pharmacologica Sinica, 2022, 43, 2624-2635.	6.1	23
156	Inhibition of MD2â€dependent inflammation attenuates the progression of nonâ€alcoholic fatty liver disease. Journal of Cellular and Molecular Medicine, 2018, 22, 936-947.	3.6	22
157	MD2 Blockage Protects Obesityâ€Induced Vascular Remodeling via Activating AMPK/Nrf2. Obesity, 2017, 25, 1532-1539.	3.0	22
158	Inhibition of epidermal growth factor receptor attenuates LPS-induced inflammation and acute lung injury in rats. Oncotarget, 2017, 8, 26648-26661.	1.8	22
159	Design, synthesis and biological activity of novel asymmetric C66 analogs as anti-inflammatory agents for the treatment of acute lung injury. European Journal of Medicinal Chemistry, 2015, 94, 436-446.	5.5	21
160	Da0324, an inhibitor of nuclear factor-κB activation, demonstrates selective antitumor activity on human gastric cancer cells. Drug Design, Development and Therapy, 2016, 10, 979.	4.3	21
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