

Cheng-Wei Yang

List of Publications by Year in descending order

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papers

968
citations

394286

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#	ARTICLE	IF	CITATIONS
1	Tyrphostin AG1024 Suppresses Coronaviral Replication by Downregulating JAK1 via an IR/IGF-1R Independent Proteolysis Mediated by Ndfip1/2_NEDD4-like E3 Ligase Itch. <i>Pharmaceuticals</i> , 2022, 15, 241.	1.7	1
2	Remdesivir and Cyclosporine Synergistically Inhibit the Human Coronaviruses OC43 and SARS-CoV-2. <i>Frontiers in Pharmacology</i> , 2021, 12, 706901.	1.6	16
3	The cardenolides ouabain and reevesioside A promote FGF2 secretion and subsequent FGFR1 phosphorylation via converged ERK1/2 activation. <i>Biochemical Pharmacology</i> , 2020, 172, 113741.	2.0	2
4	Tylophorine-based compounds are therapeutic in rheumatoid arthritis by targeting the caprin-1 ribonucleoprotein complex and inhibiting expression of associated c-Myc and HIF-1 α . <i>Pharmacological Research</i> , 2020, 152, 104581.	3.1	21
5	Inhibition of SARS-CoV-2 by Highly Potent Broad-Spectrum Anti-Coronaviral Tylophorine-Based Derivatives. <i>Frontiers in Pharmacology</i> , 2020, 11, 606097.	1.6	17
6	Natural cardenolides suppress coronaviral replication by downregulating JAK1 via a Na ⁺ /K ⁺ -ATPase independent proteolysis. <i>Biochemical Pharmacology</i> , 2020, 180, 114122.	2.0	10
7	Repurposing old drugs as antiviral agents for coronaviruses. <i>Biomedical Journal</i> , 2020, 43, 368-374.	1.4	54
8	The cardenolide ouabain suppresses coronaviral replication via augmenting a Na ⁺ /K ⁺ -ATPase-dependent PI3K_PDK1 axis signaling. <i>Toxicology and Applied Pharmacology</i> , 2018, 356, 90-97.	1.3	27
9	Identification of anti-viral activity of the cardenolides, Na ⁺ / K ⁺ -ATPase inhibitors, against porcine transmissible gastroenteritis virus. <i>Toxicology and Applied Pharmacology</i> , 2017, 332, 129-137.	1.3	24
10	Design, Synthesis, and Evaluation of Thiazolidine-2,4-dione Derivatives as a Novel Class of Glutaminase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 5599-5612.	2.9	30
11	Targeting Coronaviral Replication and Cellular JAK2 Mediated Dominant NF- κ B Activation for Comprehensive and Ultimate Inhibition of Coronaviral Activity. <i>Scientific Reports</i> , 2017, 7, 4105.	1.6	57
12	Targeting a ribonucleoprotein complex containing the caprin-1 protein and the c-Myc mRNA suppresses tumor growth in mice: an identification of a novel oncotarget. <i>Oncotarget</i> , 2015, 6, 2148-2163.	0.8	24
13	Discovery of selective inhibitors of Glutaminase-2, which inhibit mTORC1, activate autophagy and inhibit proliferation in cancer cells. <i>Oncotarget</i> , 2014, 5, 6087-6101.	0.8	63
14	c-Jun-mediated anticancer mechanisms of tylophorine. <i>Carcinogenesis</i> , 2013, 34, 1304-1314.	1.3	46
15	Cytotoxic cardenolide glycosides from the root of <i>Reevesia formosana</i> . <i>Phytochemistry</i> , 2013, 87, 86-95.	1.4	16
16	Exploration of the role of tylophorine E ring in anti-coronavirus activity: Tylophorine derived dibenzoquinolines impart multi-biological activities as orally active agents. <i>FASEB Journal</i> , 2013, 27, lb71.	0.2	0
17	Synthesis and Biological Evaluation of Tylophorine-Derived Dibenzoquinolines as Orally Active Agents: Exploration of the Role of Tylophorine E Ring on Biological Activity. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 10363-10377.	2.9	47
18	Isolation and Biological Activities of Phenanthroindolizidine and Septicine Alkaloids from the Formosan <i>Tylophora ovata</i> . <i>Planta Medica</i> , 2011, 77, 1932-1938.	0.7	32

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19	Discovery of novel benzoisoquinoline compounds for anti- β -coronaviruses. <i>FASEB Journal</i> , 2011, 25, 1b149.	0.2	0
20	Cytotoxic Flavonoids from the Leaves of <i>Cryptocarya chinensis</i> . <i>Journal of Natural Products</i> , 2010, 73, 1470-1475.	1.5	44
21	Identification of phenanthroindolizines and phenanthroquinolizidines as novel potent anti-coronaviral agents for porcine enteropathogenic coronavirus transmissible gastroenteritis virus and human severe acute respiratory syndrome coronavirus. <i>Antiviral Research</i> , 2010, 88, 160-168.	1.9	86
22	Cytotoxic Sesquiterpenes from <i>Magnolia kachirachirai</i> . <i>Chemistry and Biodiversity</i> , 2010, 7, 2737-2747.	1.0	29
23	Cytotoxic alkyl benzoquinones and alkyl phenols from <i>Ardisia virens</i> . <i>Phytochemistry</i> , 2009, 70, 2064-2071.	1.4	42
24	Tylophorine arrests carcinoma cells at G1 phase by downregulating cyclin A2 expression. <i>Biochemical and Biophysical Research Communications</i> , 2009, 386, 140-145.	1.0	51
25	Secondary Metabolites from the Leaves of <i>Litsea liliifolia</i> var. <i>nunkaoensis</i> . <i>Helvetica Chimica Acta</i> , 2008, 91, 1036-1044.	1.0	11
26	Analogues of 2-phenyl-ethanesulfonic acid phenyl ester have dual functions of inhibiting expression of inducible nitric oxide synthase and activating peroxisome proliferator-activated receptor β . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 5676-5679.	1.0	6
27	Novel Small-Molecule Inhibitors of Transmissible Gastroenteritis Virus. <i>Antimicrobial Agents and Chemotherapy</i> , 2007, 51, 3924-3931.	1.4	16
28	Anti-inflammatory effects of 7-methoxycryptopleurine and structure-activity relations of phenanthroindolizidines and phenanthroquinolizidines. <i>Biochemical and Biophysical Research Communications</i> , 2007, 354, 942-948.	1.0	48
29	Expedient synthesis and structure-activity relationships of phenanthroindolizidine and phenanthroquinolizidine alkaloids. <i>Organic and Biomolecular Chemistry</i> , 2006, 4, 860.	1.5	67
30	Anti-Inflammatory Mechanisms of Phenanthroindolizidine Alkaloids. <i>Molecular Pharmacology</i> , 2006, 69, 749-758.	1.0	80
31	Analysis of the structure-activity relationships between phenanthroindolizidines and phenanthroquinolizidines. <i>FASEB Journal</i> , 2006, 20, LB108.	0.2	0