

Lih-Ching Hsu

List of Publications by Year in descending order

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44
papers

882
citations

430874

18
h-index

552781

26
g-index

44
all docs

44
docs citations

44
times ranked

1660
citing authors

#	ARTICLE	IF	CITATIONS
1	Targeting Colorectal Cancer with Conjugates of a Glucose Transporter Inhibitor and 5-Fluorouracil. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 4450-4461.	6.4	13
2	Phosphodiesterase Type 5 Inhibitors Synergize Vincristine in Killing Castration-Resistant Prostate Cancer Through Amplifying Mitotic Arrest Signaling. <i>Frontiers in Oncology</i> , 2020, 10, 1274.	2.8	14
3	Discovery of Novel Agents on Spindle Assembly Checkpoint to Sensitize Vinorelbine-Induced Mitotic Cell Death against Human Non-Small Cell Lung Cancers. <i>International Journal of Molecular Sciences</i> , 2020, 21, 5608.	4.1	4
4	Evaluation of the Anticancer Activity of a Bile Acid-Dihydroartemisinin Hybrid Ursodeoxycholic-Dihydroartemisinin in Hepatocellular Carcinoma Cells. <i>Frontiers in Pharmacology</i> , 2020, 11, 599067.	3.5	24
5	Chalcones Display Anti-NLRP3 Inflammasome Activity in Macrophages through Inhibition of Both Priming and Activation Steps—Structure-Activity-Relationship and Mechanism Studies. <i>Molecules</i> , 2020, 25, 5960.	3.8	7
6	The (+)-Brevipolide H Displays Anticancer Activity against Human Castration-Resistant Prostate Cancer: The Role of Oxidative Stress and Akt/mTOR/p70S6K-Dependent Pathways in G1 Checkpoint Arrest and Apoptosis. <i>Molecules</i> , 2020, 25, 2929.	3.8	5
7	Dihydroartemisinin—Bile Acid Hybridization as an Effective Approach to Enhance Dihydroartemisinin Anticancer Activity. <i>ChemMedChem</i> , 2019, 14, 779-787.	3.2	18
8	The Combination of MK-2206 and WZB117 Exerts a Synergistic Cytotoxic Effect Against Breast Cancer Cells. <i>Frontiers in Pharmacology</i> , 2019, 10, 1311.	3.5	26
9	Para-Toluenesulfonamide Induces Anti-tumor Activity Through Akt-Dependent and -Independent mTOR/p70S6K Pathway: Roles of Lipid Raft and Cholesterol Contents. <i>Frontiers in Pharmacology</i> , 2018, 9, 1223.	3.5	11
10	Traversal of the Blood—Brain Barrier by Cleavable Lysine Conjugates of Apigenin. <i>Journal of Agricultural and Food Chemistry</i> , 2018, 66, 8124-8131.	5.2	8
11	Phosphodiesterase Type 5 (PDE5) Inhibitors Sensitize Topoisomerase II Inhibitors in Killing Prostate Cancer Through PDE5-Independent Impairment of HR and NHEJ DNA Repair Systems. <i>Frontiers in Oncology</i> , 2018, 8, 681.	2.8	19
12	Using precursor ion scan of 184 with liquid chromatography-electrospray ionization-tandem mass spectrometry for concentration normalization in cellular lipidomic studies. <i>Analytica Chimica Acta</i> , 2017, 971, 68-77.	5.4	24
13	Enantiomerically pure Î²-dipeptide derivative induces anticancer activity against human hormone-refractory prostate cancer through both PI3K/Akt-dependent and -independent pathways. <i>Oncotarget</i> , 2017, 8, 96668-96683.	1.8	9
14	Hemiasterlin derivative (R)(S)(S)-BF65 and Akt inhibitor MK-2206 synergistically inhibit SKOV3 ovarian cancer cell growth. <i>Biochemical Pharmacology</i> , 2016, 113, 12-23.	4.4	8
15	A novel hybrid drug between two potent anti-tubulin agents as a potential prolonged anticancer approach. <i>European Journal of Pharmaceutical Sciences</i> , 2016, 91, 50-63.	4.0	8
16	Gain-of-function p53 mutant with 21-bp deletion confers susceptibility to multidrug resistance in MCF-7 cells. <i>International Journal of Molecular Medicine</i> , 2016, 37, 233-242.	4.0	13
17	Non-immunosuppressive triazole-based small molecule induces anticancer activity against human hormone-refractory prostate cancers: the role in inhibition of PI3K/AKT/mTOR and c-Myc signaling pathways. <i>Oncotarget</i> , 2016, 7, 76995-77009.	1.8	14
18	Effects of lipopolysaccharide on the expression of plasma membrane monoamine transporter (PMAT) at the blood—brain barrier and its implications to the transport of neurotoxins. <i>Journal of Neurochemistry</i> , 2015, 135, 1178-1188.	3.9	25

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19	Repurposing of phentolamine as a potential anticancer agent against human castration-resistant prostate cancer: A central role on microtubule stabilization and mitochondrial apoptosis pathway. <i>Prostate</i> , 2015, 75, 1454-1466.	2.3	16
20	Zerumbone, a ginger sesquiterpene, induces apoptosis and autophagy in human hormone-refractory prostate cancers through tubulin binding and crosstalk between endoplasmic reticulum stress and mitochondrial insult. <i>Naunyn-Schmiedeberg's Archives of Pharmacology</i> , 2015, 388, 1223-1236.	3.0	35
21	The Akt inhibitor MK-2206 enhances the cytotoxicity of paclitaxel (Taxol) and cisplatin in ovarian cancer cells. <i>Naunyn-Schmiedeberg's Archives of Pharmacology</i> , 2015, 388, 19-31.	3.0	37
22	Reevesioside F inhibits Na ⁺ /K ⁺ -ATPase, causing cytosolic acidification, Bak activation and apoptosis in glioblastoma. <i>Oncotarget</i> , 2015, 6, 24032-24046.	1.8	7
23	Repurposing of nitroxoline as a potential anticancer agent against human prostate cancer - a crucial role on AMPK/mTOR signaling pathway and the interplay with Chk2 activation. <i>Oncotarget</i> , 2015, 6, 39806-39820.	1.8	40
24	Reevesioside A, a Cardenolide Glycoside, Induces Anticancer Activity against Human Hormone-Refractory Prostate Cancers through Suppression of c-myc Expression and Induction of G1 Arrest of the Cell Cycle. <i>PLoS ONE</i> , 2014, 9, e87323.	2.5	25
25	A unique amidoanthraquinone derivative displays antiproliferative activity against human hormone-refractory metastatic prostate cancers through activation of LKB1-AMPK-mTOR signaling pathway. <i>Naunyn-Schmiedeberg's Archives of Pharmacology</i> , 2014, 387, 979-990.	3.0	11
26	Terfenadine induces anti-proliferative and apoptotic activities in human hormone-refractory prostate cancer through histamine receptor-independent Mcl-1 cleavage and Bak up-regulation. <i>Naunyn-Schmiedeberg's Archives of Pharmacology</i> , 2014, 387, 33-45.	3.0	27
27	The K898E germline variant in the PP1-binding motif of BRCA1 causes defects in DNA Repair. <i>Scientific Reports</i> , 2014, 4, 5812.	3.3	9
28	Ardisianone, a natural benzoquinone, efficiently induces apoptosis in human hormone-refractory prostate cancers through mitochondrial damage stress and survivin downregulation. <i>Prostate</i> , 2013, 73, 133-145.	2.3	22
29	Synthesis and biological evaluation of novel C-aryl d-glucofuranosides as sodium-dependent glucose co-transporter 2 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 6282-6291.	3.0	13
30	Reevesioside F induces potent and efficient anti-proliferative and apoptotic activities through Na ⁺ /K ⁺ -ATPase β 3 subunit-involved mitochondrial stress and amplification of caspase cascades. <i>Biochemical Pharmacology</i> , 2013, 86, 1564-1575.	4.4	17
31	Development of a novel non-radioactive cell-based method for the screening of SGLT1 and SGLT2 inhibitors using 1-NBDG. <i>Molecular BioSystems</i> , 2013, 9, 2010.	2.9	17
32	Pim-1 knockdown potentiates paclitaxel-induced apoptosis in human hormone-refractory prostate cancers through inhibition of NHEJ DNA repair. <i>Cancer Letters</i> , 2012, 319, 214-222.	7.2	21
33	Development of hemiassterlin derivatives as potential anticancer agents that inhibit tubulin polymerization and synergize with a stilbene tubulin inhibitor. <i>Investigational New Drugs</i> , 2012, 30, 1379-1388.	2.6	13
34	Rottlerin potentiates camptothecin-induced cytotoxicity in human hormone refractory prostate cancers through increased formation and stabilization of topoisomerase I-DNA cleavage complexes in a PKC δ -independent pathway. <i>Biochemical Pharmacology</i> , 2012, 84, 59-67.	4.4	18
35	A PP1-binding motif present in BRCA1 plays a role in its DNA repair function. <i>International Journal of Biological Sciences</i> , 2008, 4, 352-361.	6.4	22
36	Identification and functional characterization of a PP1-binding site in BRCA1. <i>Biochemical and Biophysical Research Communications</i> , 2007, 360, 507-512.	2.1	17

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37	Loss of distal 11q is associated with DNA repair deficiency and reduced sensitivity to ionizing radiation in head and neck squamous cell carcinoma. <i>Genes Chromosomes and Cancer</i> , 2007, 46, 761-775.	2.8	79
38	Coordinate up-regulation of TMEM97 and cholesterol biosynthesis genes in normal ovarian surface epithelial cells treated with progesterone: implications for pathogenesis of ovarian cancer. <i>BMC Cancer</i> , 2007, 7, 223.	2.6	42
39	Centrosome abnormalities in ovarian cancer. <i>International Journal of Cancer</i> , 2005, 113, 746-751.	5.1	46
40	The HPV16 E6/E7 oncogene sensitizes human ovarian surface epithelial cells to low-dose but not high-dose 5-FU and 5-FUdR. <i>Biochemical and Biophysical Research Communications</i> , 2004, 320, 249-255.	2.1	6
41	Regulation of BRCA1 phosphorylation by interaction with protein phosphatase 1alpha. <i>Cancer Research</i> , 2002, 62, 6357-61.	0.9	24
42	Mapping of 55 new rat microsatellite markers from chromosome-specific libraries. <i>Mammalian Genome</i> , 1998, 9, 622-628.	2.2	20
43	Allelic imbalance in mammary carcinomas induced by either 7,12-dimethylbenz[a]anthracene or ionizing radiation in rats carrying genes conferring differential susceptibilities to mammary carcinogenesis. , 1996, 17, 134-143.		29
44	Bivariate Flow Karyotyping, Sorting, and Peak Assignment of All Rat Chromosomes. <i>Genomics</i> , 1994, 19, 75-85.	2.9	19