

# Hongliang Zong

## List of Publications by Year in descending order

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43  
papers

2,244  
citations

304743

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docs citations

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times ranked

3952  
citing authors

#	ARTICLE	IF	CITATIONS
1	FDA-approved ferumoxytol displays anti-leukaemia efficacy against cells with low ferroportin levels. <i>Nature Nanotechnology</i> , 2019, 14, 616-622.	31.5	199
2	Cranberry A-type proanthocyanidins selectively target acute myeloid leukemia cells. <i>Blood Advances</i> , 2019, 3, 3261-3265.	5.2	3
3	The epichaperome is an integrated chaperome network that facilitates tumour survival. <i>Nature</i> , 2016, 538, 397-401.	27.8	233
4	Small Molecule Inhibitor of CBF $\beta$ -RUNX Binding for RUNX Transcription Factor Driven Cancers. <i>EBioMedicine</i> , 2016, 8, 117-131.	6.1	84
5	A Hyperactive Signalosome in Acute Myeloid Leukemia Drives Addiction to a Tumor-Specific Hsp90 Species. <i>Cell Reports</i> , 2015, 13, 2159-2173.	6.4	51
6	Synthesis, anticancer activity and molecular docking studies on a series of heterocyclic trans-cyanocombretastatin analogues as antitubulin agents. <i>European Journal of Medicinal Chemistry</i> , 2015, 92, 212-220.	5.5	18
7	A small-molecule inhibitor of the aberrant transcription factor CBF $\beta$ -SMMHC delays leukemia in mice. <i>Science</i> , 2015, 347, 779-784.	12.6	104
8	Synthesis and evaluation of a series of resveratrol analogues as potent anti-cancer agents that target tubulin. <i>MedChemComm</i> , 2015, 6, 788-794.	3.4	31
9	Pharmacologic Inhibition of the Menin-MLL Interaction Blocks Progression of MLL Leukemia In Vivo. <i>Cancer Cell</i> , 2015, 27, 589-602.	16.8	290
10	A Hyperactive Signalosome Results in High Sensitivity to HSP90 Inhibitors in AML. <i>Blood</i> , 2015, 126, 2567-2567.	1.4	0
11	A-Type Proanthocyanidins Prevent Engraftment Of Primary Acute Myelogenous Leukemia Cells In Mice and Exhibit Potentially Novel Anti-Leukemia Mechanisms. <i>Blood</i> , 2013, 122, 3962-3962.	1.4	0
12	Novel Multistage Nanoparticle Drug Delivery to Ablate Leukemia Stem Cells in Their Niche.. <i>Blood</i> , 2012, 120, 2631-2631.	1.4	7
13	Synthesis of purine-scaffold fluorescent probes for heat shock protein 90 with use in flow cytometry and fluorescence microscopy. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 5347-5352.	2.2	20
14	Design, synthesis, and evaluation of small molecule Hsp90 probes. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 2603-2614.	3.0	43
15	AGEs, RAGE, and Diabetic Retinopathy. <i>Current Diabetes Reports</i> , 2011, 11, 244-252.	4.2	190
16	Thr-370 Is Responsible for CDK11p58 Autophosphorylation, Dimerization, and Kinase Activity. <i>Journal of Biological Chemistry</i> , 2011, 286, 1748-1757.	3.4	17
17	Affinity-based proteomics reveal cancer-specific networks coordinated by Hsp90. <i>Nature Chemical Biology</i> , 2011, 7, 818-826.	8.0	240
18	The Acrylonitrile Analog, VJ-289 Ablates Acute Myelogenous Leukemia Blast, Progenitor and Stem Cell Populations by Inducing Tubulin Acetylation and Caspase Activation. <i>Blood</i> , 2011, 118, 2496-2496.	1.4	0

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19	FLT3-ITD+ AML Blast, Progenitor and Stem Cell Populations Demonstrate Higher Sensitivity to the Hsp90 Inhibitor PU-H71. <i>Blood</i> , 2011, 118, 3500-3500.	1.4	0
20	HSP70 Inhibitor, YK5, Ablates Blast, Progenitor and Stem Cell Populations in Primary Acute Myelogenous Leukemia Cells. <i>Blood</i> , 2011, 118, 2493-2493.	1.4	0
21	Trihydrophobin 1 attenuates androgen signal transduction through promoting androgen receptor degradation. <i>Journal of Cellular Biochemistry</i> , 2010, 109, 1013-1024.	2.6	3
22	Homodimerization Is Essential for the Receptor for Advanced Glycation End Products (RAGE)-mediated Signal Transduction. <i>Journal of Biological Chemistry</i> , 2010, 285, 23137-23146.	3.4	115
23	Repression of Estrogen Receptor alpha by CDK11p58 Through Promoting its Ubiquitin-Proteasome Degradation. <i>Journal of Biochemistry</i> , 2009, 145, 331-343.	1.7	18
24	HSP70 protects BCL2L12 and BCL2L12A from N-terminal ubiquitination-mediated proteasomal degradation. <i>FEBS Letters</i> , 2009, 583, 1409-1414.	2.8	22
25	CDK11p58 represses vitamin D receptor-mediated transcriptional activation through promoting its ubiquitin-proteasome degradation. <i>Biochemical and Biophysical Research Communications</i> , 2009, 386, 493-498.	2.1	22
26	Knockdown of BCL2L12 leads to cisplatin resistance in MDA-MB-231 breast cancer cells. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2008, 1782, 649-657.	3.8	28
27	Cdc34-mediated Degradation of ATF5 Is Blocked by Cisplatin. <i>Journal of Biological Chemistry</i> , 2008, 283, 18773-18781.	3.4	24
28	Cyclin D3/CDK11 <sup>p58</sup> Complex Is Involved in the Repression of Androgen Receptor. <i>Molecular and Cellular Biology</i> , 2007, 27, 7125-7142.	2.3	71
29	Functional Interaction of E1AF and Sp1 in Glioma Invasion. <i>Molecular and Cellular Biology</i> , 2007, 27, 8770-8782.	2.3	24
30	CDK11p58 protein kinase activity is associated with Bcl-2 down-regulation in pro-apoptosis pathway. <i>Molecular and Cellular Biochemistry</i> , 2007, 304, 213-218.	3.1	13
31	Ubiquitin-dependent proteolysis of trihydrophobin 1 (TH1) by the human papilloma virus E6-associated protein (E6-AP). <i>Journal of Cellular Biochemistry</i> , 2007, 101, 167-180.	2.6	10
32	SGT, a Hsp90 <sup>β</sup> binding partner, is accumulated in the nucleus during cell apoptosis. <i>Biochemical and Biophysical Research Communications</i> , 2006, 343, 1153-1158.	2.1	30
33	Cell Surface Beta 1, 4-galactosyltransferase 1 promotes apoptosis by inhibiting epidermal growth factor receptor pathway. <i>Molecular and Cellular Biochemistry</i> , 2006, 291, 69-76.	3.1	9
34	β1,4-Galactosyltransferase V Functions as a Positive Growth Regulator in Glioma. <i>Journal of Biological Chemistry</i> , 2006, 281, 9482-9489.	3.4	31
35	Down-regulation of β1,4-galactosyltransferase V is a critical part of etoposide-induced apoptotic process and could be mediated by decreasing Sp1 levels in human glioma cells. <i>Glycobiology</i> , 2006, 16, 1045-1051.	2.5	20
36	Protein expression pattern of CDK11p58 during testicular development in the mouse. <i>Molecular and Cellular Biochemistry</i> , 2005, 270, 99-106.	3.1	5

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37	Elevated $\beta$ 1,4-Galactosyltransferase I in Highly Metastatic Human Lung Cancer Cells. <i>Journal of Biological Chemistry</i> , 2005, 280, 12503-12516.	3.4	88
38	Downregulation of $\beta$ 1,4-galactosyltransferase 1 inhibits CDK1/p58-mediated apoptosis induced by cycloheximide. <i>Biochemical and Biophysical Research Communications</i> , 2005, 327, 628-636.	2.1	29
39	Overexpression of small glutamine-rich TPR-containing protein promotes apoptosis in 7721 cells. <i>FEBS Letters</i> , 2005, 579, 1279-1284.	2.8	32
40	Cyclin-dependent kinase 1/p58 interacts with HBO1 and enhances its histone acetyltransferase activity. <i>FEBS Letters</i> , 2005, 579, 3579-3588.	2.8	21
41	Identification of the p28 subunit of eukaryotic initiation factor 3 (eIF3k) as a new interaction partner of cyclin D3. <i>FEBS Letters</i> , 2004, 573, 139-146.	2.8	23
42	The $\beta$ -(1 $\rightarrow$ 6)-branched $\beta$ -(1 $\rightarrow$ 3) glucohexaose and its analogues containing an $\alpha$ -(1 $\rightarrow$ 3)-linked bond have similar stimulatory effects on the mouse spleen as Lentinan. <i>International Immunopharmacology</i> , 2003, 3, 1861-1871.	3.8	33
43	The C-terminal Kinase Domain of the p34cdc2-related PITSLRE Protein Kinase (p110C) Associates with p21-activated Kinase 1 and Inhibits Its Activity during Anoikis. <i>Journal of Biological Chemistry</i> , 2003, 278, 20029-20036.	3.4	43