

Qiang Yu

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8201996/publications.pdf>

Version: 2024-02-01

72
papers

7,342
citations

94269

37
h-index

88477

70
g-index

72
all docs

72
docs citations

72
times ranked

13076
citing authors

#	ARTICLE	IF	CITATIONS
1	Colorectal cancer-associated fibroblasts promote metastasis by up-regulating LRG1 through stromal IL-6/STAT3 signaling. <i>Cell Death and Disease</i> , 2022, 13, 16.	2.7	36
2	Hypoxia induces HIF1 α -dependent epigenetic vulnerability in triple negative breast cancer to confer immune effector dysfunction and resistance to anti-PD-1 immunotherapy. <i>Nature Communications</i> , 2022, 13, .	5.8	48
3	Targeting the IRAK1 α -S100A9 Axis Overcomes Resistance to Paclitaxel in Nasopharyngeal Carcinoma. <i>Cancer Research</i> , 2021, 81, 1413-1425.	0.4	19
4	Stromal induction of BRD4 phosphorylation Results in Chromatin Remodeling and BET inhibitor Resistance in Colorectal Cancer. <i>Nature Communications</i> , 2021, 12, 4441.	5.8	49
5	Targeting enhancer reprogramming to mitigate MEK inhibitor resistance in preclinical models of advanced ovarian cancer. <i>Journal of Clinical Investigation</i> , 2021, 131, .	3.9	6
6	CREBBP cooperates with the cell cycle machinery to attenuate chidamide sensitivity in relapsed/refractory diffuse large B-cell lymphoma. <i>Cancer Letters</i> , 2021, 521, 268-280.	3.2	10
7	Interleukin enhancer-binding factor 2 promotes cell proliferation and DNA damage response in metastatic melanoma. <i>Clinical and Translational Medicine</i> , 2021, 11, e608.	1.7	8
8	Inhibition of the PLK1 α -Coupled Cell Cycle Machinery Overcomes Resistance to Oxaliplatin in Colorectal Cancer. <i>Advanced Science</i> , 2021, 8, e2100759.	5.6	29
9	EZH2-mediated PP2A inactivation confers resistance to HER2-targeted breast cancer therapy. <i>Nature Communications</i> , 2020, 11, 5878.	5.8	29
10	HER2-L755S mutation induces hyperactive MAPK and PI3K-mTOR signaling, leading to resistance to HER2 tyrosine kinase inhibitor treatment. <i>Cell Cycle</i> , 2019, 18, 1513-1522.	1.3	15
11	Methionine is a metabolic dependency of tumor-initiating cells. <i>Nature Medicine</i> , 2019, 25, 825-837.	15.2	226
12	Inhibition of interleukin-1 receptor-associated kinase 1 (IRAK1) as a therapeutic strategy. <i>Oncotarget</i> , 2018, 9, 33416-33439.	0.8	107
13	KDM4B-regulated unfolded protein response as a therapeutic vulnerability in <i>PTEN</i> -deficient breast cancer. <i>Journal of Experimental Medicine</i> , 2018, 215, 2833-2849.	4.2	33
14	Hypoxic tumor microenvironment activates GLI2 via HIF-1 α and TGF- β 2 to promote chemoresistance in colorectal cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E5990-E5999.	3.3	203
15	KDM6B Counteracts EZH2-Mediated Suppression of <i>IGFBP5</i> to Confer Resistance to PI3K/AKT Inhibitor Treatment in Breast Cancer. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 1973-1983.	1.9	35
16	MiR-200a Regulates CDK4/6 Inhibitor Effect by Targeting CDK6 in Metastatic Melanoma. <i>Journal of Investigative Dermatology</i> , 2017, 137, 1955-1964.	0.3	32
17	Chromosome 1q21.3 amplification is a trackable biomarker and actionable target for breast cancer recurrence. <i>Nature Medicine</i> , 2017, 23, 1319-1330.	15.2	116
18	<i>VHL</i> Deficiency Drives Enhancer Activation of Oncogenes in Clear Cell Renal Cell Carcinoma. <i>Cancer Discovery</i> , 2017, 7, 1284-1305.	7.7	111

#	ARTICLE	IF	CITATIONS
19	Pericyte-targeting prodrug overcomes tumor resistance to vascular disrupting agents. <i>Journal of Clinical Investigation</i> , 2017, 127, 3689-3701.	3.9	71
20	EZH2 phosphorylation by JAK3 mediates a switch to noncanonical function in natural killer/T-cell lymphoma. <i>Blood</i> , 2016, 128, 948-958.	0.6	110
21	Molecular switch of EZH2 in hypoxia. <i>Cell Cycle</i> , 2016, 15, 3007-3008.	1.3	6
22	HIF1 α activation underlies a functional switch in the paradoxical role of Ezh2/PRC2 in breast cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E3735-44.	3.3	62
23	Preclinical pharmacokinetic studies of 3-deazaneplanocin A, a potent epigenetic anticancer agent, and its human pharmacokinetic prediction using GastroPlus \textregistered . <i>European Journal of Pharmaceutical Sciences</i> , 2015, 77, 290-302.	1.9	29
24	IRAK1 is a therapeutic target that drives breast cancer metastasis and resistance to paclitaxel. <i>Nature Communications</i> , 2015, 6, 8746.	5.8	125
25	3-Deazaneplanocin A and Neplanocin A Analogues and Their Effects on Apoptotic Cell Death. <i>ChemMedChem</i> , 2015, 10, 173-182.	1.6	24
26	Elevated expression of long intergenic non-coding RNA HOTAIR in a basal-like variant of MCF7 breast cancer cells. <i>Molecular Carcinogenesis</i> , 2015, 54, 1656-1667.	1.3	35
27	Functional Characterization of D9, a Novel Deazaneplanocin A (DZNep) Analog, in Targeting Acute Myeloid Leukemia (AML). <i>PLoS ONE</i> , 2015, 10, e0122983.	1.1	18
28	Herbal compound Naoshuantong capsule attenuates retinal injury in ischemia/reperfusion rat model by inhibiting apoptosis. <i>International Journal of Clinical and Experimental Medicine</i> , 2015, 8, 12252-63.	1.3	3
29	The KDM2B-Let-7b-EZH2 Axis in Myelodysplastic Syndromes as a Target for Combined Epigenetic Therapy. <i>PLoS ONE</i> , 2014, 9, e107817.	1.1	27
30	Tumor Necrosis Factor- α and Apoptosis Induction in Melanoma Cells through Histone Modification by 3-Deazaneplanocin A. <i>Journal of Investigative Dermatology</i> , 2014, 134, 1470-1473.	0.3	3
31	EZH2-Mediated Inactivation of IFN- γ -JAK-STAT1 Signaling Is an Effective Therapeutic Target in MYC-Driven Prostate Cancer. <i>Cell Reports</i> , 2014, 8, 204-216.	2.9	87
32	RASAL2 activates RAC1 to promote triple-negative breast cancer progression. <i>Journal of Clinical Investigation</i> , 2014, 124, 5291-5304.	3.9	72
33	PDK1 Signaling Toward PLK1 \rightarrow MYC Activation Confers Oncogenic Transformation, Tumor-Initiating Cell Activation, and Resistance to mTOR-Targeted Therapy. <i>Cancer Discovery</i> , 2013, 3, 1156-1171.	7.7	119
34	EZH2 overexpression in natural killer/T-cell lymphoma confers growth advantage independently of histone methyltransferase activity. <i>Blood</i> , 2013, 121, 4512-4520.	0.6	131
35	Protein tyrosine phosphatase <i>UBASH3B</i> is overexpressed in triple-negative breast cancer and promotes invasion and metastasis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 11121-11126.	3.3	57
36	Molecular mechanisms of tumor resistance to PI3K-mTOR-targeted therapy. <i>Chinese Journal of Cancer</i> , 2013, 32, 376-379.	4.9	32

#	ARTICLE	IF	CITATIONS
37	Heterogeneous epigenetic regulation of <i>TIMP3</i> in prostate cancer. <i>Epigenetics</i> , 2012, 7, 1279-1289.	1.3	34
38	<i>TP53</i> Genomic Status Regulates Sensitivity of Gastric Cancer Cells to the Histone Methylation Inhibitor 3-Deazaneplanocin A (DZNep). <i>Clinical Cancer Research</i> , 2012, 18, 4201-4212.	3.2	65
39	Glycine Decarboxylase Activity Drives Non-Small Cell Lung Cancer Tumor-Initiating Cells and Tumorigenesis. <i>Cell</i> , 2012, 148, 259-272.	13.5	593
40	Loading 3-deazaneplanocin A into pegylated unilamellar liposomes by forming transient phenylboronic acid-drug complex and its pharmacokinetic features in Sprague-Dawley rats. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2012, 80, 323-331.	2.0	15
41	TXNIP (VDUP-1, TBP-2): A major redox regulator commonly suppressed in cancer by epigenetic mechanisms. <i>International Journal of Biochemistry and Cell Biology</i> , 2011, 43, 1668-1673.	1.2	94
42	Context-Specific Regulation of NF- κ B Target Gene Expression by EZH2 in Breast Cancers. <i>Molecular Cell</i> , 2011, 43, 798-810.	4.5	338
43	Determinants of Sensitivity to DZNep Induced Apoptosis in Multiple Myeloma Cells. <i>PLoS ONE</i> , 2011, 6, e21583.	1.1	29
44	The histone methyltransferase inhibitor, DZNep, up-regulates TXNIP, increases ROS production, and targets leukemia cells in AML. <i>Blood</i> , 2011, 118, 2830-2839.	0.6	205
45	Quantification of 3-deazaneplanocin A, a novel epigenetic anticancer agent, in rat biosamples by hydrophilic interaction liquid chromatography-tandem mass spectrometric detection. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2011, 879, 285-290.	1.2	7
46	PDK1-driven Myc signaling regulates cellular response to mTOR inhibitors. <i>Cell Cycle</i> , 2011, 10, 1019-1020.	1.3	1
47	B55 β -Associated PP2A Complex Controls PDK1-Directed Myc Signaling and Modulates Rapamycin Sensitivity in Colorectal Cancer. <i>Cancer Cell</i> , 2010, 18, 459-471.	7.7	104
48	Systems Pharmacology in Cancer. , 2010, , 377-397.		0
49	miR-449 regulates CDK-Rb-E2F1 through an auto-regulatory feedback circuit. <i>Cell Cycle</i> , 2010, 9, 213-214.	1.3	35
50	Dual Regulation of Cdc25A by Chk1 and p53-ATF3 in DNA Replication Checkpoint Control. <i>Journal of Biological Chemistry</i> , 2009, 284, 4132-4139.	1.6	38
51	<i>miR-449a</i> and <i>miR-449b</i> are direct transcriptional targets of E2F1 and negatively regulate pRb-E2F1 activity through a feedback loop by targeting <i>CDK6</i> and <i>CDC25A</i> . <i>Genes and Development</i> , 2009, 23, 2388-2393.	2.7	242
52	Combinatorial pharmacologic approaches target EZH2-mediated gene repression in breast cancer cells. <i>Molecular Cancer Therapeutics</i> , 2009, 8, 3191-3202.	1.9	65
53	The E2F family and the role of E2F1 in apoptosis. <i>International Journal of Biochemistry and Cell Biology</i> , 2009, 41, 2389-2397.	1.2	57
54	BRCA1-deficient mammary tumor cells are dependent on EZH2 expression and sensitive to Polycomb Repressive Complex 2-inhibitor 3-deazaneplanocin A. <i>Breast Cancer Research</i> , 2009, 11, R63.	2.2	98

#	ARTICLE	IF	CITATIONS
55	CDKN1C (p57KIP2) Is a Direct Target of EZH2 and Suppressed by Multiple Epigenetic Mechanisms in Breast Cancer Cells. <i>PLoS ONE</i> , 2009, 4, e5011.	1.1	155
56	Intrahepatic biliary cystadenocarcinoma: clinical analysis of 4 cases. <i>Hepatobiliary and Pancreatic Diseases International</i> , 2009, 8, 71-4.	0.6	12
57	DACT3 Is an Epigenetic Regulator of Wnt/ β ² -Catenin Signaling in Colorectal Cancer and Is a Therapeutic Target of Histone Modifications. <i>Cancer Cell</i> , 2008, 13, 529-541.	7.7	216
58	Cancer gene silencing without DNA hypermethylation. <i>Epigenetics</i> , 2008, 3, 315-317.	1.3	8
59	Pharmacologic disruption of Polycomb-repressive complex 2-mediated gene repression selectively induces apoptosis in cancer cells. <i>Genes and Development</i> , 2007, 21, 1050-1063.	2.7	804
60	Ribosomal Protein S27-like, a p53-Inducible Modulator of Cell Fate in Response to Genotoxic Stress. <i>Cancer Research</i> , 2007, 67, 11317-11326.	0.4	56
61	c-Myc overexpression sensitizes Bim-mediated Bax activation for apoptosis induced by histone deacetylase inhibitor suberoylanilide hydroxamic acid (SAHA) through regulating Bcl-2/Bcl-xL expression. <i>International Journal of Biochemistry and Cell Biology</i> , 2007, 39, 1016-1025.	1.2	21
62	A Global Map of p53 Transcription-Factor Binding Sites in the Human Genome. <i>Cell</i> , 2006, 124, 207-219.	13.5	1,060
63	Restoring p53-mediated apoptosis in cancer cells: New opportunities for cancer therapy. <i>Drug Resistance Updates</i> , 2006, 9, 19-25.	6.5	71
64	Apoptosis Signal-regulating Kinase 1 Is a Direct Target of E2F1 and Contributes to Histone Deacetylase Inhibitor-induced Apoptosis through Positive Feedback Regulation of E2F1 Apoptotic Activity. <i>Journal of Biological Chemistry</i> , 2006, 281, 10508-10515.	1.6	60
65	Pharmacologic Modulation of Glycogen Synthase Kinase-3 β Promotes p53-Dependent Apoptosis through a Direct Bax-Mediated Mitochondrial Pathway in Colorectal Cancer Cells. <i>Cancer Research</i> , 2005, 65, 9012-9020.	0.4	115
66	Inhibitors of histone deacetylases target the Rb-E2F1 pathway for apoptosis induction through activation of proapoptotic protein Bim. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 16090-16095.	3.3	234
67	p53-regulated Transcriptional Program Associated with Genotoxic Stress-induced Apoptosis. <i>Journal of Biological Chemistry</i> , 2004, 279, 21183-21192.	1.6	133
68	Identification of Myc-mediated Death Response Pathways by Microarray Analysis. <i>Journal of Biological Chemistry</i> , 2002, 277, 13059-13066.	1.6	27
69	UCN-01 inhibits p53 up-regulation and abrogates gamma-radiation-induced G(2)-M checkpoint independently of p53 by targeting both of the checkpoint kinases, Chk2 and Chk1. <i>Cancer Research</i> , 2002, 62, 5743-8.	0.4	115
70	Antisense inhibition of Chk2/hCds1 expression attenuates DNA damage-induced S and G2 checkpoints and enhances apoptotic activity in HEK-293 cells. <i>FEBS Letters</i> , 2001, 505, 7-12.	1.3	62
71	A truncated cytoplasmic topoisomerase II α in a drug-resistant lung cancer cell line is encoded by a TOP2A allele with a partial deletion of exon 34. , 2000, 85, 534-539.		20
72	Two COOH-Terminal Truncated Cytoplasmic Forms of Topoisomerase II α in a VP-16-Selected Lung Cancer Cell Line Result from Partial Gene Deletion and Alternative Splicing. <i>Biochemistry</i> , 1997, 36, 5868-5877.	1.2	30