

Yuyan Chen

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

1,832
citations

777949

13
h-index

799663

21
g-index

26
all docs

26
docs citations

26
times ranked

3717
citing authors

#	ARTICLE	IF	CITATIONS
1	Rare germline variants in childhood cancer patients suspected of genetic predisposition to cancer. <i>Genes Chromosomes and Cancer</i> , 2022, 61, 81-93.	1.5	2
2	Delayed recruiting of TPD52 to lipid droplets “ evidence for a “second wave” of lipid droplet-associated proteins that respond to altered lipid storage induced by Brefeldin A treatment. <i>Scientific Reports</i> , 2019, 9, 9790.	1.6	5
3	Investigation of clinically relevant germline variants detected by next-generation sequencing in patients with childhood cancer: a review of the literature. <i>Journal of Medical Genetics</i> , 2018, 55, 785-793.	1.5	17
4	Tumor Protein D52 (TPD52). , 2018, , 5779-5786.		0
5	Dropping in on the lipid droplet- tumor protein D52 (TPD52) as a new regulator and resident protein. <i>Adipocyte</i> , 2016, 5, 326-332.	1.3	5
6	Tumor Protein D52 (TPD52). , 2016, , 1-8.		0
7	TPD52 expression increases neutral lipid storage within cultured cells. <i>Journal of Cell Science</i> , 2015, 128, 3223-38.	1.2	31
8	Genome-wide approach to identify second gene targets for malignant rhabdoid tumors using high-density oligonucleotide microarrays. <i>Cancer Science</i> , 2014, 105, 258-264.	1.7	12
9	TPD52 represents a survival factor in <i>ERBB2</i> -amplified breast cancer cells. <i>Molecular Carcinogenesis</i> , 2014, 53, 807-819.	1.3	31
10	Tumor protein D52 (TPD52) and cancer oncogene understudy or understudied oncogene?. <i>Tumor Biology</i> , 2014, 35, 7369-7382.	0.8	51
11	Identification of PLP2 and RAB5C as novel TPD52 binding partners through yeast two-hybrid screening. <i>Molecular Biology Reports</i> , 2014, 41, 4565-4572.	1.0	11
12	Tumor protein D52 represents a negative regulator of ATM protein levels. <i>Cell Cycle</i> , 2013, 12, 3083-3097.	1.3	26
13	Aberrant activation of ALK kinase by a novel truncated form ALK protein in neuroblastoma. <i>Oncogene</i> , 2012, 31, 4667-4676.	2.6	49
14	Challenges in Identifying Candidate Amplification Targets in Human Cancers: Chromosome 8q21 as a Case Study. <i>Genes and Cancer</i> , 2012, 3, 87-101.	0.6	9
15	Aberrations of <i>NEGR1</i> on 1p31 and <i>MYEOV</i> on 11q13 in neuroblastoma. <i>Cancer Science</i> , 2011, 102, 1645-1650.	1.7	37
16	Hepatoblastoma in a Patient with Sotos Syndrome. <i>Journal of Pediatrics</i> , 2009, 155, 937-939.	0.9	23
17	Frequent inactivation of A20 in B-cell lymphomas. <i>Nature</i> , 2009, 459, 712-716.	13.7	520
18	Aberrations of Genes Regulating NF Kappa B Pathway in B-Cell Malignant Lymphoma.. <i>Blood</i> , 2009, 114, 971-971.	0.6	0

#	ARTICLE	IF	CITATIONS
19	Oncogenic mutations of ALK kinase in neuroblastoma. <i>Nature</i> , 2008, 455, 971-974.	13.7	795
20	Genome-Wide Analysis of B Cell Non-Hodgkin's Lymphoma Disclosed Frequent Involvement of Genes in NFκB Pathway. <i>Blood</i> , 2008, 112, 807-807.	0.6	3
21	AML1 Mutation and FLT3-internal Tandem Duplication in Leukemia Transformed From Myelodysplastic Syndrome. <i>Journal of Pediatric Hematology/Oncology</i> , 2007, 29, 666-667.	0.3	1
22	Mutation and expression analyses of the MET and CDKN2A genes in rhabdomyosarcoma with emphasis on MET overexpression. <i>Genes Chromosomes and Cancer</i> , 2007, 46, 348-358.	1.5	38
23	High-Resolution Analyses of Genetic and Epigenetic Aberrations in Infant Leukemia with MLL Rearrangement. <i>Blood</i> , 2007, 110, 4238-4238.	0.6	0
24	Mutations of the PTPN11 and RAS genes in rhabdomyosarcoma and pediatric hematological malignancies. <i>Genes Chromosomes and Cancer</i> , 2006, 45, 583-591.	1.5	70
25	Allelic imbalance on chromosome 2q and alterations of the caspase 8 gene in neuroblastoma. <i>Oncogene</i> , 2001, 20, 4424-4432.	2.6	92