Gail E Tomlinson

List of Publications by Year
in descending order

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34 papers	2,251 citations	304743 22 h-index	454955 30 g-index
35 all docs	35 docs citations	35 times ranked	2808 citing authors

#	Article	IF	Citations
1	Rhabdoid Tumor of the Kidney in The National Wilms' Tumor Study: Age at Diagnosis As a Prognostic Factor. Journal of Clinical Oncology, 2005, 23, 7641-7645.	1.6	227
2	Aberrant promoter methylation and silencing of the RASSF1A gene in pediatric tumors and cell lines. Oncogene, 2002, 21, 4345-4349.	5.9	207
3	Von Hippelâ€“Lindau and Hereditary Pheochromocytoma/Paraganglioma Syndromes: Clinical Features, Genetics, and Surveillance Recommendations in Childhood. Clinical Cancer Research, 2017, 23, e68-e75.	7.0	205
4	Liver Tumors in Children. Oncologist, 2008, 13, 812-820.	3.7	200
5	Small cell undifferentiated variant of hepatoblastoma: Adverse clinical and molecular features similar to rhabdoid tumors. Pediatric Blood and Cancer, 2009, 52, 328-334.	1.5	155
6	Surveillance Recommendations for Children with Overgrowth Syndromes and Predisposition to Wilms Tumors and Hepatoblastoma. Clinical Cancer Research, 2017, 23, e115-e122.	7.0	140
7	Grading of late effects in young adult survivors of childhood cancer followed in an ambulatory adult setting. Cancer, 2000, 88, 1687-1695.	4.1	137
8	The Spectrum of APC Mutations in Children with Hepatoblastoma from Familial Adenomatous Polyposis Kindreds. Journal of Pediatrics, 2005, 147, 263-266.	1.8	120
9	The Impact of COVID-19 on Cancer Screening: Challenges and Opportunities. JMIR Cancer, 2020, 6, e21697.	2.4	98
10	Cancer Screening Recommendations and Clinical Management of Inherited Gastrointestinal Cancer Syndromes in Childhood. Clinical Cancer Research, 2017, 23, e107-e114.	7.0	91
11	Cytogenetic evaluation of a large series of hepatoblastomas: Numerical abnormalities with recurring aberrations involving 1q12â€“q21. Genes Chromosomes and Cancer, 2005, 44, 177-184.	2.8	84
12	Genetics and epigenetics of hepatoblastoma. Pediatric Blood and Cancer, 2012, 59, 785-792.	1.5	67
13	The first recurring chromosome translocation in hepatoblastoma: Der(4)t(1;4)(q12;q34). , 1997, 19, 291-294.		60
14	Multiple Endocrine Neoplasia and Hyperparathyroid-Jaw Tumor Syndromes: Clinical Features, Genetics, and Surveillance Recommendations in Childhood. Clinical Cancer Research, 2017, 23, e123-e132.	7.0	55
15	De-Regulated MicroRNAs in Pediatric Cancer Stem Cells Target Pathways Involved in Cell Proliferation, Cell Cycle and Development. PLoS ONE, 2013, 8, e61622.	2.5	48
16	Myeloperoxidase promotor polymorphism and risk of hepatoblastoma. International Journal of Cancer, 2003, 106, 205-207.	5.1	46
17	Rhabdoid tumor of the kidney with primitive neuroectodermal tumor of the central nervous system: Associated tumors with different histologic, cytogenetic, and molecular findings. Genes Chromosomes and Cancer, 1994, 11, 146-152.	2.8	42
18	CCND1 polymorphism and age of onset of hepatoblastoma. Oncogene, 2004, 23, 4789-4792.	5.9	37

#	Article	IF	Citations
19	Reaching high-risk underserved individuals for cancer genetic counseling by video-teleconferencing. Journal of Community and Supportive Oncology, 2016, 14, 162-168.	0.1	37
20	Establishment and characterization of a cancer cell line derived from an aggressive childhood liver tumor. Pediatric Blood and Cancer, 2009, 53, 1040-1047.	1.5	27
21	Hepatoblastoma modeling in mice places Nrf2 within a cancer field established by mutant Î²-catenin. JCI Insight, 2016, 1, e88549.	5.0	24
22	Activated NOTCH2 is Overexpressed in Hepatoblastomas: An Immunohistochemical Study. Pediatric and Developmental Pathology, 2011, 14, 378-383.	1.0	23
23	Two identical triplet sisters carrying a germlineBRCA1 gene mutation acquire very similar breast cancer somatic mutations at multiple other sites throughout the genome. Genes Chromosomes and Cancer, 2000, 28, 359-369.	2.8	22
24	Examining access to care in clinical genomic research and medicine: Experiences from the CSER Consortium. Journal of Clinical and Translational Science, 2021, 5, e193.	0.6	21
25	FGF19 functions as autocrine growth factor for hepatoblastoma. Genes and Cancer, 2016, 7, 125-135.	1.9	18
26	Searching for microsatellite mutations in coding regions in lung, breast, ovarian and colorectal cancers. Oncogene, 2001, 20, 1005-1009.	5.9	17
27	Maternal and paternal occupational exposures and hepatoblastoma: results from the HOPE study through the Childrenâ€™s Oncology Group. Journal of Exposure Science and Environmental Epidemiology, 2017, 27, 359-364.	3.9	13
28	Genetic markers for treatment-related pancreatitis in a cohort of Hispanic children with acute lymphoblastic leukemia. Supportive Care in Cancer, 2021, 29, 725-731.	2.2	10
29	Clinical, histologic, and genetic features of mesothelioma in a 7â€Yearâ€old child. Pediatric Blood and Cancer, 2013, 60, 146-148.	1.5	8
30	Cytogenetics of hepatoblastoma. Frontiers in Bioscience - Elite, 2012, E4, 1287.	1.8	6
31	Disparities in Cancer Genetic Testing and Variants of Uncertain Significance in the Hispanic Population of South Texas. JCO Oncology Practice, 2022, 18, e805-e813.	2.9	3
32	Highly aggressive thoracic desmoid tumors in adolescent siblings with fatal outcomes in an FAP kindred: a need for increased vigilance and intervention in at-risk AYAs. Familial Cancer, 2020, 19, 311-314.	1.9	0
33	When a Tumor Becomes a Legacy: A Collection of Perspectives. Journal of Palliative Medicine, 2021, 24, 1572-1574.	1.1	0
34	Prevalence of pathogenic germline cancer risk variants in testicular cancer patients: Identifying high risk groups. Urologic Oncology: Seminars and Original Investigations, 2022, , .	1.6	0