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	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
2	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
3	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
4	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
5	When a Tumor Becomes a Legacy: A Collection of Perspectives. Journal of Palliative Medicine, 2021, 24, 1572-1574.	1.1	0
6	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
7	The Impact of COVID-19 on Cancer Screening: Challenges and Opportunities. JMIR Cancer, 2020, 6, e21697.	2.4	98
8	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
9	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
10	Surveillance Recommendations for Children with Overgrowth Syndromes and Predisposition to Wilms Tumors and Hepatoblastoma. Clinical Cancer Research, 2017, 23, e115-e122.	7.0	140
11	Cancer Screening Recommendations and Clinical Management of Inherited Gastrointestinal Cancer Syndromes in Childhood. Clinical Cancer Research, 2017, 23, e107-e114.	7.0	91
12	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
13	Hepatoblastoma modeling in mice places Nrf2 within a cancer field established by mutant Î²-catenin. JCI Insight, 2016, 1, e88549.	5.0	24
14	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
15	FGF19 functions as autocrine growth factor for hepatoblastoma. Genes and Cancer, 2016, 7, 125-135.	1.9	18
16	Clinical, histologic, and genetic features of mesothelioma in a 7â€¥earâ€old child. Pediatric Blood and Cancer, 2013, 60, 146-148.	1.5	8
17	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
18	Genetics and epigenetics of hepatoblastoma. Pediatric Blood and Cancer, 2012, 59, 785-792.	1.5	67

#	Article	IF	Citations
19	Cytogenetics of hepatoblastoma. Frontiers in Bioscience - Elite, 2012, E4, 1287.	1.8	6
20	Activated NOTCH2 is Overexpressed in Hepatoblastomas: An Immunohistochemical Study. Pediatric and Developmental Pathology, 2011, 14, 378-383.	1.0	23
21	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
22	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
23	Liver Tumors in Children. Oncologist, 2008, 13, 812-820.	3.7	200
24	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
25	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
26	The Spectrum of APC Mutations in Children with Hepatoblastoma from Familial Adenomatous Polyposis Kindreds. Journal of Pediatrics, 2005, 147, 263-266.	1.8	120
27	CCND1 polymorphism and age of onset of hepatoblastoma. Oncogene, 2004, 23, 4789-4792.	5.9	37
28	Myeloperoxidase promotor polymorphism and risk of hepatoblastoma. International Journal of Cancer, 2003, 106, 205-207.	5.1	46
29	Aberrant promoter methylation and silencing of the RASSF1A gene in pediatric tumors and cell lines. Oncogene, 2002, 21, 4345-4349.	5.9	207
30	Searching for microsatellite mutations in coding regions in lung, breast, ovarian and colorectal cancers. Oncogene, 2001, 20, 1005-1009.	5.9	17
31	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
32	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
33	The first recurring chromosome translocation in hepatoblastoma: Der(4)t(1;4)(q12;q34). , 1997, 19, 291-294.		60
34	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