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List of Publications by Year
in descending order

24 papers	494 citations	840776 11 h-index	752698 20 g-index
24 all docs	24 docs citations	24 times ranked	490 citing authors

#	Article	IF	Citations
1	Characteristics of Liâ€Fraumeni syndrome in Japan: A review study by the special committee of JSHT. Cancer Science, 2021, 112, 2821-2834.	3.9	6
2	Virilizing Adrenocortical Carcinoma Invading the Right Atrium with Histological High-Grade Malignancy and p53 Mutation in a 3-Year-Old Child: Indication of Post Operative Adjuvant Chemotherapy. Clinical Pediatric Endocrinology, 2004, 13, 25-32.	0.8	4
3	Mutations/deletions of theWT1 gene, loss of heterozygosity on chromosome arms 11p and 11q, chromosome ploidy and histology in Wilms' tumors in Japan. International Journal of Cancer, 2001, 94, 396-400.	5.1	22
4	Frequent Increase of DNA Copy Number in the 2q24 Chromosomal Region and Its Association with a Poor Clinical Outcome in Hepatoblastoma: Cytogenetic and Comparative Genomic Hybridization Analysis. Japanese Journal of Cancer Research, 2001, 92, 854-862.	1.7	30
5	Hemophagocytosis by leukemic blasts in 7 acute myeloid leukemia cases with t(16;21)(p11;q22). , 2000, 88, 1970-1975.		39
6	Hemophagocytosis by leukemic blasts in 7 acute myeloid leukemia cases with t(16;21)(p11;q22). Cancer, 2000, 88, 1970-1975.	4.1	6
7	Germline p53 Mutation in a Case of Li-Fraumeni Syndrome Presenting Gastric Cancer. Japanese Journal of Clinical Oncology, 1999, 29, 513-516.	1.3	22
8	Abnormalities of the p53 gene in juvenile myelomonocytic leukaemia. British Journal of Haematology, 1999, 106, 980-986.	2.5	9
9	TheFANCA gene in Japanese Fanconi anemia: Reports of eight novel mutations and analysis of sequence variability. Human Mutation, 1999, 13, 237-244.	2.5	30
10	Correlation of chromosome abnormalities with presence or absence ofWT1 deletions/mutations in Wilms tumor. , 1999, 25, 26-32.		15
11	11p15 translocations involving theNUP98 gene in childhood therapy-related acute myeloid leukemia/myelodysplastic syndrome. Genes Chromosomes and Cancer, 1999, 26, 215-220.	2.8	69
12	The FANCA gene in Japanese Fanconi anemia: Reports of eight novel mutations and analysis of sequence variability. Human Mutation, 1999, 13, 237.	2.5	4
13	Minimal residual disease with TELâ€AML1 fusion transcript in childhood acute lymphoblastic leukaemia with t(12;21). British Journal of Haematology, 1997, 97, 607-611.	2.5	32
14	Congenital leukaemia with a mixed phenotype of megakaryoblasts and erythroblasts: a case report and characterization of the blasts. British Journal of Haematology, 1997, 96, 740-742.	2.5	12
15	Leukemia in GH Deficient Children. Clinical Pediatric Endocrinology, 1994, 3, 53-60.	0.8	2
16	Deletion ofWT1andWIT1Genes and Loss of Heterozygosity on Chromosome 11p in Wilms Tumors in Japan. Japanese Journal of Cancer Research, 1993, 84, 616-624.	1.7	10
17	Hypofibrinogenemia in a girl with Langerhans cell histiocytosis during etoposide and prednisolone therapy. Pediatrics International, 1993, 35, 148-150.	0.5	5
18	Detection of Novel Germ-line p53 Mutations in Diverse-Cancer-Prone Families Identifiled by Selecting Patients With Childhood Adrencortical Carcinoma. Journal of the National Cancer Institute, 1992, 84, 703-707.	6.3	90

#	Article	IF	Citations
19	Familial Aggregation of Cancer from Proband Cases with Childhood Adrenal Cortical Carcinoma1. Japanese Journal of Cancer Research, 1991, 82, 893-900.	1.7	13
20	Infant leukemia in Japan: Clinical and biological analysis of 48 cases. Medical and Pediatric Oncology, 1991, 19, 28-32.	1.0	37
21	Risk Factors for Leukemia Occurrence among Growth Hormone Users. Japanese Journal of Cancer Research, 1989, 80, 822-825.	1.7	34
22	Megakaryocyte proliferative disorder in neonates with Down's syndrome.. Keio Journal of Medicine, 1987, 36, 57-61.	1.1	3
23	Detection of Cytomegalovirusâ€Specific RNA in Frozen Tissue Sections by Biotinâ€Labeled Hybridization Probe. Pediatrics International, 1986, 28, 586-590.	0.5	0
24	SELECTIVE MALABSORPTION OF VITAMIN B12: REPORT OF A CASE AND STUDIES ON VITAMIN B12 ABSORPTION. Keio Journal of Medicine, 1973, 22, 45-57.	1.1	0