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List of Publications by Year in descending order

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24
papers

494
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840776

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490
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Detection of Novel Germ-line p53 Mutations in Diverse-Cancer-Prone Families Identified by Selecting Patients With Childhood Adrenocortical Carcinoma. <i>Journal of the National Cancer Institute</i> , 1992, 84, 703-707. | 6.3 | 90 |
| 2 | 11p15 translocations involving theNUP98 gene in childhood therapy-related acute myeloid leukemia/myelodysplastic syndrome. <i>Genes Chromosomes and Cancer</i> , 1999, 26, 215-220. | 2.8 | 69 |
| 3 | Hemophagocytosis by leukemic blasts in 7 acute myeloid leukemia cases with t(16;21)(p11;q22)., 2000, 88, 1970-1975. | | 39 |
| 4 | Infant leukemia in Japan: Clinical and biological analysis of 48 cases. <i>Medical and Pediatric Oncology</i> , 1991, 19, 28-32. | 1.0 | 37 |
| 5 | Risk Factors for Leukemia Occurrence among Growth Hormone Users. <i>Japanese Journal of Cancer Research</i> , 1989, 80, 822-825. | 1.7 | 34 |
| 6 | Minimal residual disease with TELâ€AML1 fusion transcript in childhood acute lymphoblastic leukaemia with t(12;21). <i>British Journal of Haematology</i> , 1997, 97, 607-611. | 2.5 | 32 |
| 7 | TheFANCA gene in Japanese Fanconi anemia: Reports of eight novel mutations and analysis of sequence variability. <i>Human Mutation</i> , 1999, 13, 237-244. | 2.5 | 30 |
| 8 | 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. <i>Japanese Journal of Cancer Research</i> , 2001, 92, 854-862. | 1.7 | 30 |
| 9 | Germline p53 Mutation in a Case of Li-Fraumeni Syndrome Presenting Gastric Cancer. <i>Japanese Journal of Clinical Oncology</i> , 1999, 29, 513-516. | 1.3 | 22 |
| 10 | Mutations/deletions of theWT1 gene, loss of heterozygosity on chromosome arms 11p and 11q, chromosome ploidy and histology in Wilms' tumors in Japan. <i>International Journal of Cancer</i> , 2001, 94, 396-400. | 5.1 | 22 |
| 11 | Correlation of chromosome abnormalities with presence or absence ofWT1 deletions/mutations in Wilms tumor. , 1999, 25, 26-32. | | 15 |
| 12 | Familial Aggregation of Cancer from Proband Cases with Childhood Adrenal Cortical Carcinoma1. <i>Japanese Journal of Cancer Research</i> , 1991, 82, 893-900. | 1.7 | 13 |
| 13 | Congenital leukaemia with a mixed phenotype of megakaryoblasts and erythroblasts: a case report and characterization of the blasts. <i>British Journal of Haematology</i> , 1997, 96, 740-742. | 2.5 | 12 |
| 14 | Deletion ofWT1andWIT1Genes and Loss of Heterozygosity on Chromosome 11p in Wilms Tumors in Japan. <i>Japanese Journal of Cancer Research</i> , 1993, 84, 616-624. | 1.7 | 10 |
| 15 | Abnormalities of the p53 gene in juvenile myelomonocytic leukaemia. <i>British Journal of Haematology</i> , 1999, 106, 980-986. | 2.5 | 9 |
| 16 | Characteristics of Liâ€Fraumeni syndrome in Japan: A review study by the special committee of JSHT. <i>Cancer Science</i> , 2021, 112, 2821-2834. | 3.9 | 6 |
| 17 | Hemophagocytosis by leukemic blasts in 7 acute myeloid leukemia cases with t(16;21)(p11;q22). <i>Cancer</i> , 2000, 88, 1970-1975. | 4.1 | 6 |
| 18 | Hypofibrinogenemia in a girl with Langerhans cell histiocytosis during etoposide and prednisolone therapy. <i>Pediatrics International</i> , 1993, 35, 148-150. | 0.5 | 5 |

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 19 | 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 |
| 20 | 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 |
| 21 | Megakaryocyte proliferative disorder in neonates with Down's syndrome.. Keio Journal of Medicine, 1987, 36, 57-61. | 1.1 | 3 |
| 22 | Leukemia in GH Deficient Children. Clinical Pediatric Endocrinology, 1994, 3, 53-60. | 0.8 | 2 |
| 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 |