

Naoki Kakazu

List of Publications by Year in descending order

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Version: 2024-02-01

27
papers

751
citations

687363

13
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552781

26
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27
all docs

27
docs citations

27
times ranked

1193
citing authors

#	ARTICLE	IF	CITATIONS
1	Acute lymphoblastic leukemia (ALL) with t(8;14)(q11.2;q32) in an elderly patient. <i>Leukemia Research</i> , 2010, 34, e82-e84.	0.8	3
2	Chromosomal Manipulation by Site-Specific Recombinases and Fluorescent Protein-Based Vectors. <i>PLoS ONE</i> , 2010, 5, e9846.	2.5	11
3	A novel PAX3 rearrangement in embryonal rhabdomyosarcoma. <i>Cancer Genetics and Cytogenetics</i> , 2009, 189, 98-104.	1.0	12
4	Late appearance of a Philadelphia chromosome in a patient with therapy-related acute myeloid leukemia and high expression of EVI1. <i>Cancer Genetics and Cytogenetics</i> , 2008, 180, 115-120.	1.0	9
5	Establishment and characterization of the new splenic marginal zone lymphoma-derived cell line UCH1 carrying a complex rearrangement involving t(8;14) and chromosome 3. <i>Leukemia and Lymphoma</i> , 2007, 48, 767-773.	1.3	2
6	Establishment of a human herpes virus-8-negative malignant effusion lymphoma cell line (STR-428) carrying concurrent translocations of BCL2 and c-MYC genes. <i>Leukemia Research</i> , 2007, 31, 1285-1292.	0.8	13
7	Clinical Features of Hypereosinophilic Syndrome: FIP1L1-PDGFR α Fusion Gene-Positive Disease is a Distinct Clinical Entity with Myeloproliferative Features and a Poor Response to Corticosteroid. <i>International Journal of Hematology</i> , 2007, 85, 5-10.	1.6	6
8	Rad54 is dispensable for the ALT pathway. <i>Genes To Cells</i> , 2006, 11, 1305-1315.	1.2	7
9	Monoallelic yet combinatorial expression of variable exons of the protocadherin- β gene cluster in single neurons. <i>Nature Genetics</i> , 2005, 37, 171-176.	21.4	246
10	Chromosome 7 abnormalities in acute megakaryoblastic leukemia associated with Down syndrome. <i>Cancer Genetics and Cytogenetics</i> , 2005, 158, 184-187.	1.0	13
11	Developmental Pluripotency of the Nuclei of Neurons in the Cerebral Cortex of Juvenile Mice. <i>Journal of Neuroscience</i> , 2005, 25, 8368-8374.	3.6	12
12	Secondary Acute Monocytic Leukemia with a Translocation t(8;16)(p11;p13): Case Report and Review of the Literature. <i>Leukemia and Lymphoma</i> , 2004, 45, 621-625.	1.3	18
13	Rearrangement of the MOZ gene in pediatric therapy-related myelodysplastic syndrome with a novel chromosomal translocation t(2;8)(p23;p11). <i>Genes Chromosomes and Cancer</i> , 2003, 36, 413-419.	2.8	42
14	A new chromosome banding technique, spectral color banding (SCAN), for full characterization of chromosomal abnormalities. <i>Genes Chromosomes and Cancer</i> , 2003, 37, 412-416.	2.8	18
15	Marked Thrombocytosis Following Relapse of Acute Myeloblastic Leukemia Associated with Development of Translocation (2;14) (p13;q32). <i>Leukemia and Lymphoma</i> , 2002, 43, 2063-2065.	1.3	0
16	Establishment of a Primary Effusion Lymphoma Cell Line (RM-P1) and In Vivo Growth System Using SCID Mice. <i>International Journal of Hematology</i> , 2002, 76, 165-172.	1.6	13
17	Acute Myeloid Leukemia (FAB-M2) with a Masked Type of t(8;21) Translocation Revealed by Spectral Karyotyping. <i>International Journal of Hematology</i> , 2002, 76, 338-343.	1.6	5
18	Development of spectral colour banding in cytogenetic analysis. <i>Lancet, The</i> , 2001, 357, 529-530.	13.7	21

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19	Agranular CD4+CD56+ blastic natural killer leukemia/lymphoma. <i>Annals of Hematology</i> , 2001, 80, 228-231.	1.8	27
20	Involvement of the NUP98 Gene in a Chromosomal Translocation t(11;20)(p15;q11.2) in a Patient With Acute Monocytic Leukemia (FAB-M5b). <i>International Journal of Hematology</i> , 2001, 74, 53-57.	1.6	4
21	A Variant Form of Myelodysplastic Syndrome With Ph ⁺ Minor-BCR/ABL Transcript. <i>International Journal of Hematology</i> , 2001, 74, 58-63.	1.6	6
22	Identification and characterization of SEB, a novel protein that binds to the acute undifferentiated leukemia-associated protein SET. <i>FEBS Journal</i> , 2001, 268, 1340-1351.	0.2	81
23	Characterization of complex chromosomal abnormalities in B-cell lymphoma by a combined spectral karyotyping (SKY) analysis and fluorescence in situ hybridization (FISH) using a 14q telomere probe. <i>American Journal of Hematology</i> , 2000, 65, 291-297.	4.1	10
24	PEBP2 [±] A/CBFA1 mutations in Japanese cleidocranial dysplasia patients. <i>Gene</i> , 2000, 244, 21-28.	2.2	51
25	Combined spectral karyotyping and DAPI banding analysis of chromosome abnormalities in myelodysplastic syndrome. <i>Genes Chromosomes and Cancer</i> , 1999, 26, 336-345.	2.8	93
26	A complete Not I restriction map covering the entire long arm of human chromosome 11. <i>Genes To Cells</i> , 1997, 2, 345-357.	1.2	12
27	Precise localization of the human gene encoding cell adhesion kinase $\hat{2}$ (CAK $\hat{2}$ /PYK2) to chromosome 8 at p21.1 by fluorescence in situ hybridization. <i>Human Genetics</i> , 1996, 98, 508-510.	3.8	16