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List of Publications by Year
in descending order

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45 papers	1,142 citations	361413 20 h-index	395702 33 g-index
46 all docs	46 docs citations	46 times ranked	2007 citing authors

#	Article	IF	Citations
1	Clinicopathological evaluation of the programmed cell death 1 (PD1)/programmed cell deathâ€ligand 1 (PDâ€L1) axis in postâ€transplant lymphoproliferative disorders: association with Epsteinâ€“Barr virus, PDâ€L1 copy number alterations, and outcome. Histopathology, 2019, 75, 799-812.	2.9	29
2	Refining the Breakpoints of Three New Translocations Identified in Myelodysplastic Syndromes. Acta Haematologica, 2016, 135, 94-100.	1.4	2
3	NOTCH1, TP53, and MAP2K1 Mutations in Splenic Diffuse Red Pulp Small B-cell Lymphoma Are Associated With Progressive Disease. American Journal of Surgical Pathology, 2016, 40, 192-201.	3.7	40
4	Detection of chromothripsisâ€like patterns with a custom array platform for chronic lymphocytic leukemia. Genes Chromosomes and Cancer, 2015, 54, 668-680.	2.8	23
5	Paraspinal extramedullary hematopoiesis in hereditary spherocytosis with a concurrent follicular lymphoma: case report and review of the literature. Diagnostic Pathology, 2015, 10, 158.	2.0	2
6	Exuberant complex metaplastic carcinoma of the breast with SOX2 expression: Cove ring the full spectrum of ductal neoplasia of the breast. Revista Espanola De Patologia, 2015, 48, 170-175.	0.2	0
7	Interstitial 13q14 deletions detected in the karyotype and translocations with concomitant deletion at 13q14 in chronic lymphocytic leukemia: Different genetic mechanisms but equivalent poorer clinical outcome. Genes Chromosomes and Cancer, 2014, 53, 788-797.	2.8	15
8	Genomic complexity and IGHV mutational status are key predictors of outcome of chronic lymphocytic leukemia patients with TP53 disruption. Haematologica, 2014, 99, e231-e234.	3.5	33
9	Reciprocal translocations in myelodysplastic syndromes and chronic myelomonocytic leukemias: Review of 5,654 patients with an evaluable karyotype. Genes Chromosomes and Cancer, 2013, 52, 753-763.	2.8	15
10	Clonal evolution in chronic lymphocytic leukemia: Analysis of correlations with IGHV mutational status, NOTCH1 mutations and clinical significance. Genes Chromosomes and Cancer, 2013, 52, 920-927.	2.8	15
11	Refining the Diagnosis and Prognostic Categorization of Acute Myeloid Leukemia Patients with an Integrated Use of Cytogenetic and Molecular Studies. Acta Haematologica, 2013, 129, 65-71.	1.4	3
12	Biallelic losses of 13q do not confer a poorer outcome in chronic lymphocytic leukaemia: analysis of 627 patients with isolated 13q deletion. British Journal of Haematology, 2013, 163, 47-54.	2.5	13
13	Different distribution of NOTCH1 mutations in chronic lymphocytic leukemia with isolated trisomy 12 or associated with other chromosomal alterations. Genes Chromosomes and Cancer, 2012, 51, 881-889.	2.8	57
14	NOTCH1 mutations in chronic lymphocytic leukemia with trisomy 12. Genes Chromosomes and Cancer, 2012, 51, 1064-1065.	2.8	0
15	12p13 rearrangements: 6â€‰Mb deletion responsible for ID/MCA and reciprocal duplication without clinical responsibility. American Journal of Medical Genetics, Part A, 2012, 158A, 1071-1076.	1.2	10
16	A new genetic abnormality leading to TP53 gene deletion in chronic lymphocytic leukaemia. British Journal of Haematology, 2012, 156, 612-618.	2.5	7
17	Response to lenalidomide in patients with myelodysplastic syndrome with deletion 5q: clinical and cytogenetic analysis of a single centre series. Annals of Hematology, 2010, 89, 1069-1070.	1.8	0
18	Incidence and prognostic impact of secondary cytogenetic aberrations in a series of 145 patients with mantle cell lymphoma. Genes Chromosomes and Cancer, 2010, 49, 439-451.	2.8	68

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19	Do we need to do fluorescence in situ hybridization analysis in myelodysplastic syndromes as often as we do?. Leukemia Research, 2010, 34, 1437-1441.	0.8	27
20	Multiple recurrent chromosomal breakpoints in mantle cell lymphoma revealed by a combination of molecular cytogenetic techniques. Genes Chromosomes and Cancer, 2008, 47, 1086-1097.	2.8	28
21	New chromosomal alterations in a series of 23 splenic marginal zone lymphoma patients revealed by Spectral Karyotyping (SKY). Leukemia Research, 2008, 32, 727-736.	0.8	20
22	A Retrospective and Theoretical Evaluation of Rapid Methods for Detecting Chromosome Abnormalities and Their Implications on Genetic Counseling Based on a Series of 3868 CVS Diagnoses. Fetal Diagnosis and Therapy, 2008, 23, 126-131.	1.4	4
23	Transient donor cell-derived myelodysplastic syndrome with monosomy 7 after unrelated cord blood transplantation. European Journal of Haematology, 2006, 77, 259-263.	2.2	34
24	Studies of complex Ph translocations in cases with chronic myelogenous leukemia and one with acute lymphoblastic leukemia. Cancer Genetics and Cytogenetics, 2006, 166, 89-93.	1.0	12
25	Fluorescence in situ hybridization studies using BAC clones of the EVI1 locus in hematological malignancies with 3q rearrangements. Cancer Genetics and Cytogenetics, 2006, 170, 115-120.	1.0	12
26	Clinical implications of ZAP-70 expressionin chronic lymphocytic leukemia. Cytometry Part B - Clinical Cytometry, 2006, 70B, 214-217.	1.5	26
27	Gene Expression Profiling of Acute Myeloid Leukemia with Translocation t(8;16)(p11;p13) and MYST3-CREBBP Rearrangement Reveals a Distinctive Signature with a Specific Pattern of HOX Gene Expression. Cancer Research, 2006, 66, 6947-6954.	0.9	127
28	Comparative genomic hybridisation identifies two variants of smoldering multiple myeloma. British Journal of Haematology, 2005, 130, 729-732.	2.5	40
29	Recombination in a male carrier of two reciprocal translocations involving chromosomes 14, 14â€², 15, and 21 leading to balanced and unbalanced rearrangements in offspring. , 2005, 134A, 309-314.		9
30	46,XY,18q+/46,XY,18qâˆ’ mosaicism in a fragile X prenatal diagnosis. Prenatal Diagnosis, 2005, 25, 448-450.	2.3	4
31	Gene Expression Signature of Acute Myeloid Leukemia (AML) with T(8;16)(P11;P13) and MYST3-CREBBP Rearrangement: A Microarray Study Validated by Multiple Real-Time PCR.. Blood, 2005, 106, 3009-3009.	1.4	0
32	A Novel Elastin Gene Mutation Resulting in an Autosomal Dominant Form of Cutis Laxa. Archives of Dermatology, 2004, 140, 1135-9.	1.4	73
33	Fluorescence in situ hybridization analysis of matched primary tumour and lymph-node metastasis of D1 (pT2-3pN1M0) prostate cancer. BJU International, 2004, 94, 407-411.	2.5	7
34	Type IMOZ/CBP (MYST3/CREBBP)is the most common chimeric transcript in acute myeloid leukemia with t(8;16)(p11;p13) translocation. Genes Chromosomes and Cancer, 2004, 40, 140-145.	2.8	72
35	Chimeric BCR/ABL gene detected by fluorescence in situ hybridization in three new cases of Philadelphia chromosome-negative chronic myelocytic leukemia. Cancer Genetics and Cytogenetics, 2003, 141, 114-119.	1.0	16
36	High levels of chromosomal imbalances in typical and small-cell variants of T-cell prolymphocytic leukemia. Cancer Genetics and Cytogenetics, 2003, 147, 36-43.	1.0	30

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37	Fetoplacental discrepancy involving structural abnormalities of chromosome 8 detected by prenatal diagnosis. Prenatal Diagnosis, 2003, 23, 319-322.	2.3	24
38	Genetic Imbalances in Progressed B-Cell Chronic Lymphocytic Leukemia and Transformed Large-Cell Lymphoma (Richter's Syndrome). American Journal of Pathology, 2002, 161, 957-968.	3.8	86
39	Large de novo deletion in chromosome 12 affecting the PAH, IGF1, ASCL1, and TRA1 genes. Journal of Molecular Medicine, 2001, 78, 721-724.	3.9	6
40	High-grade prostate intraepithelial neoplasia shares cytogenetic alterations with invasive prostate cancer. Prostate, 2001, 47, 29-35.	2.3	25
41	Highâ€grade prostate intraepithelial neoplasia shares cytogenetic alterations with invasive prostate cancer. Prostate, 2001, 47, 29-35.	2.3	1
42	Paternal isodisomy 13 in a normal newborn infant after trisomy rescue evidenced by prenatal diagnosis. , 2000, 90, 291-293.		23
43	Prevalence of Y chromosome microdeletions in oligospermic and azoospermic candidates for intracytoplasmic sperm injection. Fertility and Sterility, 1998, 70, 506-510.	1.0	75
44	Brachycephaly is ineffective for detection of Down syndrome in early midtrimester fetuses. Early Human Development, 1997, 47, 57-61.	1.8	9
45	Prenatal diagnosis of fragile x syndrome: (cgg)n expansion and methylation of chorionic villus samples. Prenatal Diagnosis, 1995, 15, 801-807.	2.3	20