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List of Publications by Year
in descending order

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136 papers	4,385 citations	109264 35 h-index	128225 60 g-index
137 all docs	137 docs citations	137 times ranked	5297 citing authors

#	Article	IF	Citations
1	Chromosome banding analysis and genomic microarrays are both useful but not equivalent methods for genomic complexity risk stratification in chronic lymphocytic leukemia patients. Haematologica, 2022, 107, 593-603.	1.7	18
2	Balanced and unbalanced translocations in a multicentric series of 2843 patients with chronic lymphocytic leukemia. Genes Chromosomes and Cancer, 2022, 61, 37-43.	1.5	10
3	Molecular and cytogenetic characterization of myelodysplastic syndromes in cell-free DNA. Blood Advances, 2022, 6, 3178-3188.	2.5	6
4	Cytogenetics in Chronic Lymphocytic Leukemia: ERIC Perspectives and Recommendations. HemaSphere, 2022, 6, e707.	1.2	17
5	Validation and functional characterization of GWAS-identified variants for chronic lymphocytic leukemia: a CRuCIAL study. Blood Cancer Journal, 2022, 12, 79.	2.8	1
6	Outcomes and molecular profile of oligomonocytic CMML support its consideration as the first stage in the CMML continuum. Blood Advances, 2022, 6, 3921-3931.	2.5	7
7	Optical Genome Mapping: A Promising New Tool to Assess Genomic Complexity in Chronic Lymphocytic Leukemia (CLL). Cancers, 2022, 14, 3376.	1.7	18
8	Higher-order connections between stereotyped subsets: implications for improved patient classification in CLL. Blood, 2021, 137, 1365-1376.	0.6	72
9	Reduced expansion of CD94/NKG2C⁺ NK cells in chronic lymphocytic leukemia and CLLâ€like monoclonal Bâ€cell lymphocytosis is not related to increased human cytomegalovirus seronegativity or NKG2C deletions. International Journal of Laboratory Hematology, 2021, 43, 1032-1040.	0.7	6
10	Chronic lymphocytic leukemiaâ€“like monoclonal B-cell lymphocytosis exhibits an increased inflammatory signature that is reduced in early-stage chronic lymphocytic leukemia. Experimental Hematology, 2021, 95, 68-80.	0.2	6
11	Lack of expression of LMO2 clone SP51 identifies MYC rearrangements in aggressive large B-cell lymphomas. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2021, , 1.	1.4	1
12	COVID-19 severity and mortality in patients with CLL: an update of the international ERIC and Campus CLL study. Leukemia, 2021, 35, 3444-3454.	3.3	57
13	Cryptic insertions of the immunoglobulin light chain enhancer region near CCND1 in t(11;14)-negative mantle cell lymphoma. Haematologica, 2020, 105, e408-e411.	1.7	13
14	Oligomonocytic and overt chronic myelomonocytic leukemia show similar clinical, genomic, and immunophenotypic features. Blood Advances, 2020, 4, 5285-5296.	2.5	27
15	Genomic and epigenomic insights into the origin, pathogenesis, and clinical behavior of mantle cell lymphoma subtypes. Blood, 2020, 136, 1419-1432.	0.6	131
16	Prognosis Assessment of Early-Stage Chronic Lymphocytic Leukemia: Are We Ready to Predict Clinical Evolution Without a Crystal Ball?. Clinical Lymphoma, Myeloma and Leukemia, 2020, 20, 548-555.e4.	0.2	10
17	Genomic arrays identify high-risk chronic lymphocytic leukemia with genomic complexity: a multi-center study. Haematologica, 2020, 106, 87-97.	1.7	43
18	Conventional and molecular cytogenetic studies to characterize 2 complex variant Philadelphia translocations in patients with chronic myeloid leukemia. Oncology Letters, 2019, 17, 5705-5710.	0.8	5

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19	Pemphigusâ€like hypereosinophilic syndrome with <scp>FIP</scp>1L1â€“<scp>PDGFRA</scp> fusion gene: A challenging and uncommon clinical presentation. Journal of Dermatology, 2019, 46, 531-534.	0.6	5
20	CCND2 and CCND3 hijack immunoglobulin light-chain enhancers in cyclin D1â^' mantle cell lymphoma. Blood, 2019, 133, 940-951.	0.6	77
21	Cytogenetic complexity in chronic lymphocytic leukemia: definitions, associations, and clinical impact. Blood, 2019, 133, 1205-1216.	0.6	164
22	Diseaseâ€biased and shared characteristics of the immunoglobulin gene repertoires in marginal zone B cell lymphoproliferations. Journal of Pathology, 2019, 247, 416-421.	2.1	25
23	Restricted T cell receptor repertoire in CLL-like monoclonal B cell lymphocytosis and early stage CLL. OncoImmunology, 2018, 7, e1432328.	2.1	20
24	Characterizing patients with multiple chromosomal aberrations detected by FISH in chronic lymphocytic leukemia. Leukemia and Lymphoma, 2018, 59, 633-642.	0.6	8
25	<scp>CD</scp>274 (<scp>PDL</scp>1) and <scp>JAK</scp>2 genomic amplifications in pulmonary squamousâ€cell and adenocarcinoma patients. Histopathology, 2018, 72, 259-269.	1.6	27
26	An Integrated Data Resource for Genomic AnalysisÂof Cutaneous T-Cell Lymphoma. Journal of Investigative Dermatology, 2018, 138, 2681-2683.	0.3	38
27	A gene signature that distinguishes conventional and leukemic nonnodal mantle cell lymphoma helps predict outcome. Blood, 2018, 132, 413-422.	0.6	89
28	LMO2-negative Expression Predicts the Presence of MYC Translocations in Aggressive B-Cell Lymphomas. American Journal of Surgical Pathology, 2017, 41, 877-886.	2.1	19
29	Chronic lymphocytic leukemia with isochromosome 17q: An aggressive subgroup associated with TP53 mutations and complex karyotypes. Cancer Letters, 2017, 409, 42-48.	3.2	6
30	Clinical and biological significance of isolated Y chromosome loss in myelodysplastic syndromes and chronic myelomonocytic leukemia. A report from the Spanish MDS Group. Leukemia Research, 2017, 63, 85-89.	0.4	9
31	Patients with chronic lymphocytic leukemia and complex karyotype show an adverse outcome even in absence of TP53/ATM FISH deletions. Oncotarget, 2017, 8, 54297-54303.	0.8	44
32	Imatinib Treatment of Lymphomatoid Papulosis Associated with Myeloproliferative Hypereosinophilic Syndrome Presenting the FIP1L1-PDGFRA Fusion Gene. Acta Dermato-Venereologica, 2017, 97, 855-857.	0.6	4
33	Karyotypic complexity rather than chromosome 8 abnormalities aggravates the outcome of chronic lymphocytic leukemia patients with TP53 aberrations. Oncotarget, 2016, 7, 80916-80924.	0.8	29
34	A high proportion of cells carrying trisomy 12 is associated with a worse outcome in patients with chronic lymphocytic leukemia. Hematological Oncology, 2016, 34, 84-92.	0.8	26
35	Additional trisomies amongst patients with chronic lymphocytic leukemia carrying trisomy 12: the accompanying chromosome makes a difference. Haematologica, 2016, 101, e299-e302.	1.7	35
36	Identification of Gene Mutations and Fusion Genes in Patients with SÃ©zary Syndrome. Journal of Investigative Dermatology, 2016, 136, 1490-1499.	0.3	77

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37	Refining the Breakpoints of Three New Translocations Identified in Myelodysplastic Syndromes. Acta Haematologica, 2016, 135, 94-100.	0.7	2
38	MiR-204 silencing in intraepithelial to invasive cutaneous squamous cell carcinoma progression. Molecular Cancer, 2016, 15, 53.	7.9	48
39	Guidelines for cytogenetic investigations in tumours. European Journal of Human Genetics, 2016, 24, 6-13.	1.4	28
40	ROS1 copy number alterations are frequent in non-small cell lung cancer. Oncotarget, 2016, 7, 8019-8028.	0.8	24
41	A Low Frequency of Losses in 11q Chromosome Is Associated with Better Outcome and Lower Rate of Genomic Mutations in Patients with Chronic Lymphocytic Leukemia. PLoS ONE, 2015, 10, e0143073.	1.1	24
42	Notch1 Pathway Activation Results from the Epigenetic Abrogation of Notch-Related MicroRNAs in Mycosis Fungoides. Journal of Investigative Dermatology, 2015, 135, 3144-3152.	0.3	31
43	MicroRNA Expression Profiling and DNA Methylation Signature for Deregulated MicroRNA in Cutaneous T-Cell Lymphoma. Journal of Investigative Dermatology, 2015, 135, 1128-1137.	0.3	87
44	Trisomy 8, a Cytogenetic Abnormality in Myelodysplastic Syndromes, Is Constitutional or Not?. PLoS ONE, 2015, 10, e0129375.	1.1	19
45	MET expression and copy number heterogeneity in nonsquamous non-small cell lung cancer (nsNSCLC). Oncotarget, 2015, 6, 16215-16226.	0.8	54
46	Genetic Abnormalities in Chronic Lymphocytic Leukemia: Where We Are and Where We Go. BioMed Research International, 2014, 2014, 1-13.	0.9	106
47	Distinction between Asymptomatic Monoclonal B-cell Lymphocytosis with Cyclin D1 Overexpression and Mantle Cell Lymphoma: From Molecular Profiling to Flow Cytometry. Clinical Cancer Research, 2014, 20, 1007-1019.	3.2	44
48	Assessment of ALK Status by FISH on 1000 Spanish Non-Small Cell Lung Cancer Patients. Journal of Thoracic Oncology, 2014, 9, 1816-1820.	0.5	23
49	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.	1.5	15
50	Application of FISH 7q in MDS patients without monosomy 7 or 7q deletion by conventional G-banding cytogenetics: Does â˜7/7qâ˜ detection by FISH have prognostic value?. Leukemia Research, 2013, 37, 416-421.	0.4	16
51	Mycosis Fungoides and SÃ©zary Syndrome. Methods in Molecular Biology, 2013, 973, 175-188.	0.4	10
52	Genomic arrays in chronic lymphocytic leukemia routine clinical practice: are we ready to substitute conventional cytogenetics and fluorescencein situhybridization techniques?. Leukemia and Lymphoma, 2013, 54, 986-995.	0.6	18
53	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.	1.2	13
54	Primary Bone Marrow Lymphoma. American Journal of Surgical Pathology, 2012, 36, 296-304.	2.1	59

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55	In situ mantle cell lymphoma: clinical implications of an incidental finding with indolent clinical behavior. Haematologica, 2012, 97, 270-278.	1.7	146
56	Are ER+PR+ and ER+PRâ˜' breast tumors genetically different? A CGH array study. Cancer Genetics, 2012, 205, 138-146.	0.2	11
57	MYC Copy Number Gains are Associated with Poor Outcome in Penile Squamous Cell Carcinoma. Journal of Urology, 2012, 188, 1965-1971.	0.2	24
58	Chronic lymphocytic leukaemia with 17p deletion: a retrospective analysis of prognostic factors and therapy results. British Journal of Haematology, 2012, 157, 67-74.	1.2	39
59	Absence of TCR loci chromosomal translocations in cutaneous T-cell lymphomas. Cancer Genetics, 2011, 204, 405-409.	0.2	9
60	Identification of t(17;22)(q22;q13) (COL1A1/PDGFB) in dermatofibrosarcoma protuberans by fluorescence in situ hybridization in paraffin-embedded tissue microarrays. Human Pathology, 2011, 42, 176-184.	1.1	43
61	Primary Cutaneous CD30+ Anaplastic Large-Cell Lymphomas Show a Heterogeneous Genomic Profile: An Oligonucleotide ArrayCGH Approach. Journal of Investigative Dermatology, 2011, 131, 269-271.	0.3	14
62	Deletion of TET2 gene in an acute myeloid leukemia case with a t(4;15)(q24;q26) characterized by glass needle based chromosome microdissection and oligonucleotide array. Leukemia Research, 2011, 35, e161-e163.	0.4	2
63	Differential expression of JAK2 and Src kinase genes in response to hydroxyurea treatment in polycythemia vera and essential thrombocythemia. Annals of Hematology, 2011, 90, 939-946.	0.8	7
64	Molecular diagnosis of dermatofibrosarcoma protuberans: A comparison between reverse transcriptaseâ€polymerase chain reaction and fluorescence in situ hybridization methodologies. Genes Chromosomes and Cancer, 2011, 50, 510-517.	1.5	69
65	A multicolor fluorescence in situ hybridization assay: A monitoring tool in the surveillance of patients with a history of nonâ€“muscleâ€invasive urothelial cell carcinoma. Cancer Cytopathology, 2011, 119, 395-403.	1.4	24
66	Cryptic IGH/BCL2 rearrangements with variant FISH patterns in follicular lymphoma. Leukemia Research, 2011, 35, 256-259.	0.4	11
67	Absence of mutations of the histone methyltransferase gene EZH2 in splenic b-cell marginal zone lymphoma. Leukemia Research, 2011, 35, e23-e24.	0.4	5
68	Increased MLL gene rearrangements in amniocytes from fetuses of mothers who smoke. Leukemia Research, 2011, 35, 1066-1069.	0.4	5
69	Prospective study of clinical and biological prognostic factors at diagnosis in patients with early stage B-cell chronic lymphocytic leukemia. Leukemia and Lymphoma, 2011, 52, 429-435.	0.6	6
70	FOXP1 molecular cytogenetics and protein expression analyses in primary cutaneous large B cell lymphoma, leg-type. Histology and Histopathology, 2011, 26, 213-21.	0.5	16
71	Cytogenetic aberrations and their prognostic value in a series of 330 splenic marginal zone B-cell lymphomas: a multicenter study of the Splenic B-Cell Lymphoma Group. Blood, 2010, 116, 1479-1488.	0.6	174
72	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.	1.5	68

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73	CKS1B amplification is a frequent event in cutaneous squamous cell carcinoma with aggressive clinical behaviour. Genes Chromosomes and Cancer, 2010, 49, 1054-1061.	1.5	10
74	Does monosomy 5 really exist in myelodysplastic syndromes and acute myeloid leukemia?. Leukemia Research, 2010, 34, 1242-1245.	0.4	23
75	Epidermal growth factor receptor gene numerical aberrations are frequent events in actinic keratoses and invasive cutaneous squamous cell carcinomas. Experimental Dermatology, 2010, 19, 151-153.	1.4	77
76	Oligonucleotide Array-CGH Identifies Genomic Subgroups and Prognostic Markers for Tumor Stage Mycosis Fungoides. Journal of Investigative Dermatology, 2010, 130, 1126-1135.	0.3	71
77	â€œEruptive postoperative squamous cell carcinomasâ€•or â€œHypertrophic lichen planusâ€“like reactions combined with infundibulocystic hyperplasiaâ€?. Journal of the American Academy of Dermatology, 2010, 63, 910-911.	0.6	2
78	Genomic and Gene Expression Profiling Defines Indolent Forms of Mantle Cell Lymphoma. Cancer Research, 2010, 70, 1408-1418.	0.4	429
79	Cytogenetic characterization of NCI-H69 and NCI-H69AR small cell lung cancer cell lines by spectral karyotyping. Cancer Genetics and Cytogenetics, 2009, 191, 97-101.	1.0	7
80	Fine-Mapping Chromosomal Loss at 9p21: Correlation with Prognosis in Primary Cutaneous Diffuse Large B-Cell Lymphoma, Leg Type. Journal of Investigative Dermatology, 2009, 129, 1149-1155.	0.3	84
81	Cutaneous Desmoid Tumor. Dermatologic Surgery, 2009, 35, 1582-1587.	0.4	5
82	3q26 (hTERC) gain studied by fluorescence in situ hybridization as a persistence-progression indicator in low-grade squamous intraepithelial lesion cases. Human Pathology, 2009, 40, 1474-1478.	1.1	29
83	FISH and immunohistochemical status of the hepatocyte growth factor receptor (c-Met) in 184 invasive breast tumors. Breast Cancer Research, 2009, 11, 402.	2.2	22
84	HPV Determination in the Control After LEEP Due to CIN II-III: Prospective Study and Predictive Model. International Journal of Gynecological Pathology, 2009, 28, 120-126.	0.9	11
85	Multiple recurrent chromosomal breakpoints in mantle cell lymphoma revealed by a combination of molecular cytogenetic techniques. Genes Chromosomes and Cancer, 2008, 47, 1086-1097.	1.5	28
86	FISH is better than BIOMED-2 PCR to detect IgH/BCL2 translocation in follicular lymphoma at diagnosis using paraffin-embedded tissue sections. Leukemia Research, 2008, 32, 737-742.	0.4	37
87	New chromosomal alterations in a series of 23 splenic marginal zone lymphoma patients revealed by Spectral Karyotyping (SKY). Leukemia Research, 2008, 32, 727-736.	0.4	20
88	Blast cells with nuclear extrusions in the form of micronuclei are associated with MYC amplification in acute myeloid leukemia. Cancer Genetics and Cytogenetics, 2008, 185, 32-36.	1.0	11
89	Fluorescence in situ hybridization improves the detection of 5q31 deletion in myelodysplastic syndromes without cytogenetic evidence of 5q-. Haematologica, 2008, 93, 1001-1008.	1.7	36
90	Monosomy 7 with severe myelodysplasia developing during imatinib treatment of Philadelphia-positive chronic myeloid leukemia: Two cases with a different outcome. American Journal of Hematology, 2007, 82, 849-851.	2.0	33

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91	Gain of multiple copies of the CBFB gene: a new genetic aberration in a case of granulocytic sarcoma. Cancer Genetics and Cytogenetics, 2007, 179, 62-65.	1.0	6
92	Aberrant nuclear BCL10 expression and lack of t(11;18)(q21;q21) in primary cutaneous marginal zone B-cell lymphoma. Human Pathology, 2006, 37, 867-873.	1.1	36
93	Gastrointestinal Involvement in Mantle Cell Lymphoma: A Prospective Clinic, Endoscopic, and Pathologic Study. American Journal of Surgical Pathology, 2006, 30, 1274-1280.	2.1	121
94	Study of chromosomal abnormalities in 11 cases of cervical dysplasia using comparative genomic hybridization on cotton-lint cervical samples. Cancer Genetics and Cytogenetics, 2006, 164, 61-65.	1.0	2
95	Detection of abnormalities of PRV-1, TPO, and c-MPL genes detected by fluorescence in situ hybridization in essential thrombocythemia. Cancer Genetics and Cytogenetics, 2006, 167, 39-42.	1.0	6
96	Methotrexate resistancein vitro is achieved by a dynamic selectionprocess of tumor cell variants emerging during treatment. International Journal of Cancer, 2006, 119, 1607-1615.	2.3	9
97	Comparative genomic hybridization analysis of cutaneous large B-cell lymphomas. Experimental Dermatology, 2005, 14, 883-890.	1.4	17
98	Insertion (8;11) in a renal oncocytoma with multifocal transformation to chromophobe renal cell carcinoma. Cancer Genetics and Cytogenetics, 2005, 163, 160-163.	1.0	11
99	Clonal proliferation of cyclin D1â€“positive mantle lymphocytes in an asymptomatic patient: an early-stage event in the development or an indolent form of a mantle cell lymphoma?. Human Pathology, 2005, 36, 1232-1237.	1.1	41
100	Identification of Male Cardiomyocytes of Extracardiac Origin in the Hearts of Women with Male Progeny: Male Fetal Cell Microchimerism of the Heart. Journal of Heart and Lung Transplantation, 2005, 24, 2179-2183.	0.3	78
101	Correlation between histologic findings and cytogenetic abnormalities in bladder carcinoma: A FISH study. Urology, 2005, 65, 913-918.	0.5	14
102	Polysomy of chromosome 17 in breast cancer tumors showing an overexpression of ERBB2: a study of 175 cases using fluorescence in situhybridization and immunohistochemistry. Breast Cancer Research, 2005, 7, R267-73.	2.2	76
103	Identification of novel cytogenetic markers with prognostic significance in a series of 968 patients with primary myelodysplastic syndromes. Haematologica, 2005, 90, 1168-78.	1.7	163
104	Activation of mitochondrial apoptotic pathway in mantle cell lymphoma: high sensitivity to mitoxantrone in cases with functional DNA-damage response genes. Oncogene, 2004, 23, 8941-8949.	2.6	23
105	Cytogenetic and fluorescence in situ hybridization studies in 60 patients with multiple myeloma and plasma cell leukemia. Cancer Genetics and Cytogenetics, 2004, 148, 71-76.	1.0	18
106	Is fluorescence in situ hybridization a useful method in diagnosis of polycythemia vera patients?. Cancer Genetics and Cytogenetics, 2004, 151, 139-145.	1.0	14
107	Endogenous erythroid and megakaryocytic circulating progenitors, HUMARA clonality assay, and PRV-1 expression are useful tools for diagnosis of polycythemia vera and essential thrombocythemia. Blood, 2004, 103, 2427-2428.	0.6	18
108	Genetic characterization of SÃ©zary's syndrome by conventional cytogenetics and cross-species color banding fluorescent in situhybridization. Haematologica, 2004, 89, 165-73.	1.7	17

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109	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
110	Additional i(1)(q10) in a primitive neuroectodermal tumor type Merkel cell carcinoma as a primary cytogenetic change. Cancer Genetics and Cytogenetics, 2003, 142, 165-167.	1.0	5
111	Cytogenetic findings in familial B-cell chronic lymphocytic leukemia. Cancer Genetics and Cytogenetics, 2003, 143, 172-173.	1.0	5
112	RxFISH karyotype andMYCamplification in the HT-29 colon adenocarcinoma cell line. Genes Chromosomes and Cancer, 2003, 36, 319-320.	1.5	6
113	Comparative Analysis of TCR-Î³ Gene Rearrangements by Genescan and Polyacrylamide Gel-electrophoresis in Cutaneous T-cell Lymphoma. Acta Dermato-Venereologica, 2003, 84, 6-11.	0.6	11
114	Analysis of T-Cell Receptor Î³ Gene Rearrangements by PCR-Genescan and PCR-Polyacrylamide Gel Electrophoresis in Early-Stage Mycosis fungoides/Large-Plaque Parapsoriasis. Dermatology, 2003, 207, 418-419.	0.9	8
115	Incidence of trisomy 8 and 9, deletion of D13S319 and D20S108 loci and BCR/ABL translocation in non-treated essential thrombocythemia patients: an analysis of bone marrow cells using interphase fluorescence in situ hybridization. Haematologica, 2003, 88, 110-1.	1.7	10
116	Nodal marginal zone lymphoma in AIDS patients. Aids, 2002, 16, 2232-2234.	1.0	2
117	Genetic characterization of the paraimmunoblastic variant of small lymphocytic lymphoma/chronic lymphocytic leukemia: A case report and review of the literature. Human Pathology, 2002, 33, 1145-1148.	1.1	5
118	Contribution of cytogenetics and in situ hybridization to the study of monoclonal gammopathies of undetermined significance. Cancer Genetics and Cytogenetics, 2002, 132, 25-29.	1.0	10
119	Translocation (5;17)(q13;q21) in a case with precursor T-lymphoblastic lymphoma/leukemia. Cancer Genetics and Cytogenetics, 2002, 132, 81-82.	1.0	1
120	Pentasomy 21 with two isochromosomes 21 in a case of acute myeloid leukemia without maturation. Cancer Genetics and Cytogenetics, 2002, 132, 71-73.	1.0	7
121	A new case of acute nonlymphocytic leukemia (Frenchâ€“Americanâ€“British subtype M1) with double minutes and c-MYC amplification. Cancer Genetics and Cytogenetics, 2002, 132, 161-164.	1.0	6
122	Monosomy 15 in chronic myelomonocytic leukemia. Cancer Genetics and Cytogenetics, 2002, 134, 165-167.	1.0	2
123	Clinical Utility of a Multiprobe FISH Assay in Voided Urine Specimens for the Detection of Bladder Cancer and its Recurrences, Compared with Urinary Cytology. European Urology, 2002, 42, 547-552.	0.9	74
124	Report of 46,XX/46,XY/47,XXY/48,XXYY mosaicism in an adult phenotypic male. American Journal of Medical Genetics Part A, 2002, 111, 215-217.	2.4	8
125	New t(11;12)(q12;q11) characterized by RxFISH in a patient with T-cell large granular lymphocyte leukemia. Cancer Genetics and Cytogenetics, 2001, 125, 70-73.	1.0	3
126	Isochromosome +i(3)(q10) in a new case of persistent polyclonal B-cell lymphocytosis (PPBL). European Journal of Haematology, 2000, 64, 344-346.	1.1	0

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127	Cytogenetic and Fluorescence In Situ Hybridization Studies in Four Cases of Plasma Cell Leukemia. Cancer Genetics and Cytogenetics, 2000, 121, 163-166.	1.0	1
128	Dicentric (17;18) in a Case of Atypical B-Cell Chronic Lymphocytic Leukemia. Cancer Genetics and Cytogenetics, 2000, 121, 194-197.	1.0	6
129	A New Case of Turner Syndrome Associated with Multiple Myeloma. Cancer Genetics and Cytogenetics, 2000, 117, 80-81.	1.0	4
130	Translocation t(6;14)(p12;q32): a novel cytogenetic abnormality in splenic lymphoma with villous lymphocytes. British Journal of Haematology, 2000, 110, 241-243.	1.2	8
131	Cytogenetic Findings in Five Patients with Hairy Cell Leukemia. Cancer Genetics and Cytogenetics, 1999, 110, 41-43.	1.0	16
132	Translocation (11;14)(q13;q32) and Preferential Involvement of Chromosomes 1, 2, 9, 13, and 17 in Mantle Cell Lymphoma. Cancer Genetics and Cytogenetics, 1999, 111, 92-98.	1.0	31
133	Two New Cases of Near-Tetraploidy in Adult Acute Myeloid Leukemia. Cancer Genetics and Cytogenetics, 1998, 102, 131-134.	1.0	11
134	Cytogenetic Abnormalities in Three Patients with B-Cell Prolymphocytic Leukemia. Cancer Genetics and Cytogenetics, 1998, 103, 43-45.	1.0	12
135	Coexistence of tetrasomy 8 and trisomy 8 in a case with myeloid metaplasia with myelofibrosis. Cancer Genetics and Cytogenetics, 1997, 94, 147-150.	1.0	9
136	Frequent involvement of chromosomes 1, 3, 7 and 8 in splenic marginal zone Bâ€cell lymphoma. British Journal of Haematology, 1997, 98, 446-449.	1.2	56