

Blanca Espinet

List of Publications by Year in descending order

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136
papers

4,385
citations

109321

35
h-index

128289

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. <i>Haematologica</i> , 2022, 107, 593-603.	3.5	18
2	Balanced and unbalanced translocations in a multicentric series of 2843 patients with chronic lymphocytic leukemia. <i>Genes Chromosomes and Cancer</i> , 2022, 61, 37-43.	2.8	10
3	Molecular and cytogenetic characterization of myelodysplastic syndromes in cell-free DNA. <i>Blood Advances</i> , 2022, 6, 3178-3188.	5.2	6
4	Cytogenetics in Chronic Lymphocytic Leukemia: ERIC Perspectives and Recommendations. <i>HemaSphere</i> , 2022, 6, e707.	2.7	17
5	Validation and functional characterization of GWAS-identified variants for chronic lymphocytic leukemia: a CRuCIAL study. <i>Blood Cancer Journal</i> , 2022, 12, 79.	6.2	1
6	Outcomes and molecular profile of oligomonocytic CMML support its consideration as the first stage in the CMML continuum. <i>Blood Advances</i> , 2022, 6, 3921-3931.	5.2	7
7	Optical Genome Mapping: A Promising New Tool to Assess Genomic Complexity in Chronic Lymphocytic Leukemia (CLL). <i>Cancers</i> , 2022, 14, 3376.	3.7	18
8	Higher-order connections between stereotyped subsets: implications for improved patient classification in CLL. <i>Blood</i> , 2021, 137, 1365-1376.	1.4	72
9	Reduced expansion of CD94/NKG2C ⁺ NK cells in chronic lymphocytic leukemia and CLL-like monoclonal B-cell lymphocytosis is not related to increased human cytomegalovirus seronegativity or <i>NKG2C</i> deletions. <i>International Journal of Laboratory Hematology</i> , 2021, 43, 1032-1040.	1.3	6
10	Chronic lymphocytic leukemia-like monoclonal B-cell lymphocytosis exhibits an increased inflammatory signature that is reduced in early-stage chronic lymphocytic leukemia. <i>Experimental Hematology</i> , 2021, 95, 68-80.	0.4	6
11	Lack of expression of LMO2 clone SP51 identifies MYC rearrangements in aggressive large B-cell lymphomas. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2021, , 1.	2.8	1
12	COVID-19 severity and mortality in patients with CLL: an update of the international ERIC and Campus CLL study. <i>Leukemia</i> , 2021, 35, 3444-3454.	7.2	57
13	Cryptic insertions of the immunoglobulin light chain enhancer region near <i>CCND1</i> in t(11;14)-negative mantle cell lymphoma. <i>Haematologica</i> , 2020, 105, e408-e411.	3.5	13
14	Oligomonocytic and overt chronic myelomonocytic leukemia show similar clinical, genomic, and immunophenotypic features. <i>Blood Advances</i> , 2020, 4, 5285-5296.	5.2	27
15	Genomic and epigenomic insights into the origin, pathogenesis, and clinical behavior of mantle cell lymphoma subtypes. <i>Blood</i> , 2020, 136, 1419-1432.	1.4	131
16	Prognosis Assessment of Early-Stage Chronic Lymphocytic Leukemia: Are We Ready to Predict Clinical Evolution Without a Crystal Ball?. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2020, 20, 548-555.e4.	0.4	10
17	Genomic arrays identify high-risk chronic lymphocytic leukemia with genomic complexity: a multi-center study. <i>Haematologica</i> , 2020, 106, 87-97.	3.5	43
18	Conventional and molecular cytogenetic studies to characterize 2 complex variant Philadelphia translocations in patients with chronic myeloid leukemia. <i>Oncology Letters</i> , 2019, 17, 5705-5710.	1.8	5

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19	Pemphigus-like hypereosinophilic syndrome with <i>FIP1L1-PDGFR</i> fusion gene: A challenging and uncommon clinical presentation. <i>Journal of Dermatology</i> , 2019, 46, 531-534.	1.2	5
20	CCND2 and CCND3 hijack immunoglobulin light-chain enhancers in cyclin D1 ^{hi} mantle cell lymphoma. <i>Blood</i> , 2019, 133, 940-951.	1.4	77
21	Cytogenetic complexity in chronic lymphocytic leukemia: definitions, associations, and clinical impact. <i>Blood</i> , 2019, 133, 1205-1216.	1.4	164
22	Disease-biased and shared characteristics of the immunoglobulin gene repertoires in marginal zone B cell lymphoproliferations. <i>Journal of Pathology</i> , 2019, 247, 416-421.	4.5	25
23	Restricted T cell receptor repertoire in CLL-like monoclonal B cell lymphocytosis and early stage CLL. <i>OncImmunology</i> , 2018, 7, e1432328.	4.6	20
24	Characterizing patients with multiple chromosomal aberrations detected by FISH in chronic lymphocytic leukemia. <i>Leukemia and Lymphoma</i> , 2018, 59, 633-642.	1.3	8
25	<i>CD274</i> (<i>PDL1</i>) and <i>JAK2</i> genomic amplifications in pulmonary squamous-cell and adenocarcinoma patients. <i>Histopathology</i> , 2018, 72, 259-269.	2.9	27
26	An Integrated Data Resource for Genomic Analysis of Cutaneous T-Cell Lymphoma. <i>Journal of Investigative Dermatology</i> , 2018, 138, 2681-2683.	0.7	38
27	A gene signature that distinguishes conventional and leukemic nonnodal mantle cell lymphoma helps predict outcome. <i>Blood</i> , 2018, 132, 413-422.	1.4	89
28	LMO2-negative Expression Predicts the Presence of MYC Translocations in Aggressive B-Cell Lymphomas. <i>American Journal of Surgical Pathology</i> , 2017, 41, 877-886.	3.7	19
29	Chronic lymphocytic leukemia with isochromosome 17q: An aggressive subgroup associated with TP53 mutations and complex karyotypes. <i>Cancer Letters</i> , 2017, 409, 42-48.	7.2	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. <i>Leukemia Research</i> , 2017, 63, 85-89.	0.8	9
31	Patients with chronic lymphocytic leukemia and complex karyotype show an adverse outcome even in absence of <i>TP53/ATM</i> FISH deletions. <i>Oncotarget</i> , 2017, 8, 54297-54303.	1.8	44
32	Imatinib Treatment of Lymphomatoid Papulosis Associated with Myeloproliferative Hypereosinophilic Syndrome Presenting the <i>FIP1L1-PDGFR</i> Fusion Gene. <i>Acta Dermato-Venereologica</i> , 2017, 97, 855-857.	1.3	4
33	Karyotypic complexity rather than chromosome 8 abnormalities aggravates the outcome of chronic lymphocytic leukemia patients with <i>TP53</i> aberrations. <i>Oncotarget</i> , 2016, 7, 80916-80924.	1.8	29
34	A high proportion of cells carrying trisomy 12 is associated with a worse outcome in patients with chronic lymphocytic leukemia. <i>Hematological Oncology</i> , 2016, 34, 84-92.	1.7	26
35	Additional trisomies amongst patients with chronic lymphocytic leukemia carrying trisomy 12: the accompanying chromosome makes a difference. <i>Haematologica</i> , 2016, 101, e299-e302.	3.5	35
36	Identification of Gene Mutations and Fusion Genes in Patients with Sezary Syndrome. <i>Journal of Investigative Dermatology</i> , 2016, 136, 1490-1499.	0.7	77

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37	Refining the Breakpoints of Three New Translocations Identified in Myelodysplastic Syndromes. <i>Acta Haematologica</i> , 2016, 135, 94-100.	1.4	2
38	MiR-204 silencing in intraepithelial to invasive cutaneous squamous cell carcinoma progression. <i>Molecular Cancer</i> , 2016, 15, 53.	19.2	48
39	Guidelines for cytogenetic investigations in tumours. <i>European Journal of Human Genetics</i> , 2016, 24, 6-13.	2.8	28
40	<i>ROS1</i> copy number alterations are frequent in non-small cell lung cancer. <i>Oncotarget</i> , 2016, 7, 8019-8028.	1.8	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. <i>PLoS ONE</i> , 2015, 10, e0143073.	2.5	24
42	Notch1 Pathway Activation Results from the Epigenetic Abrogation of Notch-Related MicroRNAs in Mycosis Fungoides. <i>Journal of Investigative Dermatology</i> , 2015, 135, 3144-3152.	0.7	31
43	MicroRNA Expression Profiling and DNA Methylation Signature for Deregulated MicroRNA in Cutaneous T-Cell Lymphoma. <i>Journal of Investigative Dermatology</i> , 2015, 135, 1128-1137.	0.7	87
44	Trisomy 8, a Cytogenetic Abnormality in Myelodysplastic Syndromes, Is Constitutional or Not?. <i>PLoS ONE</i> , 2015, 10, e0129375.	2.5	19
45	MET expression and copy number heterogeneity in nonsquamous non-small cell lung cancer (nsNSCLC). <i>Oncotarget</i> , 2015, 6, 16215-16226.	1.8	54
46	Genetic Abnormalities in Chronic Lymphocytic Leukemia: Where We Are and Where We Go. <i>BioMed Research International</i> , 2014, 2014, 1-13.	1.9	106
47	Distinction between Asymptomatic Monoclonal B-cell Lymphocytosis with Cyclin D1 Overexpression and Mantle Cell Lymphoma: From Molecular Profiling to Flow Cytometry. <i>Clinical Cancer Research</i> , 2014, 20, 1007-1019.	7.0	44
48	Assessment of ALK Status by FISH on 1000 Spanish Non-Small Cell Lung Cancer Patients. <i>Journal of Thoracic Oncology</i> , 2014, 9, 1816-1820.	1.1	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. <i>Genes Chromosomes and Cancer</i> , 2014, 53, 788-797.	2.8	15
50	Application of FISH 7q in MDS patients without monosomy 7 or 7q deletion by conventional G-banding cytogenetics: Does $\Delta(7)(q)$ detection by FISH have prognostic value?. <i>Leukemia Research</i> , 2013, 37, 416-421.	0.8	16
51	Mycosis Fungoides and Sézary Syndrome. <i>Methods in Molecular Biology</i> , 2013, 973, 175-188.	0.9	10
52	Genomic arrays in chronic lymphocytic leukemia routine clinical practice: are we ready to substitute conventional cytogenetics and fluorescence in situ hybridization techniques?. <i>Leukemia and Lymphoma</i> , 2013, 54, 986-995.	1.3	18
53	Biallelic losses of 13q do not confer a poorer outcome in chronic lymphocytic leukaemia: analysis of 627 patients with isolated 13q deletion. <i>British Journal of Haematology</i> , 2013, 163, 47-54.	2.5	13
54	Primary Bone Marrow Lymphoma. <i>American Journal of Surgical Pathology</i> , 2012, 36, 296-304.	3.7	59

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55	In situ mantle cell lymphoma: clinical implications of an incidental finding with indolent clinical behavior. <i>Haematologica</i> , 2012, 97, 270-278.	3.5	146
56	Are ER+PR+ and ER+PR+ breast tumors genetically different? A CGH array study. <i>Cancer Genetics</i> , 2012, 205, 138-146.	0.4	11
57	<i>MYC</i> Copy Number Gains are Associated with Poor Outcome in Penile Squamous Cell Carcinoma. <i>Journal of Urology</i> , 2012, 188, 1965-1971.	0.4	24
58	Chronic lymphocytic leukaemia with 17p deletion: a retrospective analysis of prognostic factors and therapy results. <i>British Journal of Haematology</i> , 2012, 157, 67-74.	2.5	39
59	Absence of TCR loci chromosomal translocations in cutaneous T-cell lymphomas. <i>Cancer Genetics</i> , 2011, 204, 405-409.	0.4	9
60	Identification of t(17;22)(q22;q13) (COL1A1/PDGFB) in dermatofibrosarcoma protuberans by fluorescence in situ hybridization in paraffin-embedded tissue microarrays. <i>Human Pathology</i> , 2011, 42, 176-184.	2.0	43
61	Primary Cutaneous CD30+ Anaplastic Large-Cell Lymphomas Show a Heterogeneous Genomic Profile: An Oligonucleotide ArrayCGH Approach. <i>Journal of Investigative Dermatology</i> , 2011, 131, 269-271.	0.7	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. <i>Leukemia Research</i> , 2011, 35, e161-e163.	0.8	2
63	Differential expression of JAK2 and Src kinase genes in response to hydroxyurea treatment in polycythemia vera and essential thrombocythemia. <i>Annals of Hematology</i> , 2011, 90, 939-946.	1.8	7
64	Molecular diagnosis of dermatofibrosarcoma protuberans: A comparison between reverse transcriptase-polymerase chain reaction and fluorescence in situ hybridization methodologies. <i>Genes Chromosomes and Cancer</i> , 2011, 50, 510-517.	2.8	69
65	A multicolor fluorescence in situ hybridization assay: A monitoring tool in the surveillance of patients with a history of non-muscle-invasive urothelial cell carcinoma. <i>Cancer Cytopathology</i> , 2011, 119, 395-403.	2.4	24
66	Cryptic IGH/BCL2 rearrangements with variant FISH patterns in follicular lymphoma. <i>Leukemia Research</i> , 2011, 35, 256-259.	0.8	11
67	Absence of mutations of the histone methyltransferase gene EZH2 in splenic b-cell marginal zone lymphoma. <i>Leukemia Research</i> , 2011, 35, e23-e24.	0.8	5
68	Increased MLL gene rearrangements in amniocytes from fetuses of mothers who smoke. <i>Leukemia Research</i> , 2011, 35, 1066-1069.	0.8	5
69	Prospective study of clinical and biological prognostic factors at diagnosis in patients with early stage B-cell chronic lymphocytic leukemia. <i>Leukemia and Lymphoma</i> , 2011, 52, 429-435.	1.3	6
70	FOXP1 molecular cytogenetics and protein expression analyses in primary cutaneous large B cell lymphoma, leg-type. <i>Histology and Histopathology</i> , 2011, 26, 213-21.	0.7	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. <i>Blood</i> , 2010, 116, 1479-1488.	1.4	174
72	Incidence and prognostic impact of secondary cytogenetic aberrations in a series of 145 patients with mantle cell lymphoma. <i>Genes Chromosomes and Cancer</i> , 2010, 49, 439-451.	2.8	68

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73	<i>CKS1B</i> amplification is a frequent event in cutaneous squamous cell carcinoma with aggressive clinical behaviour. <i>Genes Chromosomes and Cancer</i> , 2010, 49, 1054-1061.	2.8	10
74	Does monosomy 5 really exist in myelodysplastic syndromes and acute myeloid leukemia?. <i>Leukemia Research</i> , 2010, 34, 1242-1245.	0.8	23
75	Epidermal growth factor receptor gene numerical aberrations are frequent events in actinic keratoses and invasive cutaneous squamous cell carcinomas. <i>Experimental Dermatology</i> , 2010, 19, 151-153.	2.9	77
76	Oligonucleotide Array-CGH Identifies Genomic Subgroups and Prognostic Markers for Tumor Stage Mycosis Fungoides. <i>Journal of Investigative Dermatology</i> , 2010, 130, 1126-1135.	0.7	71
77	“Eruptive postoperative squamous cell carcinomas” or “Hypertrophic lichen planus” like reactions combined with infundibulocystic hyperplasia?. <i>Journal of the American Academy of Dermatology</i> , 2010, 63, 910-911.	1.2	2
78	Genomic and Gene Expression Profiling Defines Indolent Forms of Mantle Cell Lymphoma. <i>Cancer Research</i> , 2010, 70, 1408-1418.	0.9	429
79	Cytogenetic characterization of NCI-H69 and NCI-H69AR small cell lung cancer cell lines by spectral karyotyping. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Journal of Investigative Dermatology</i> , 2009, 129, 1149-1155.	0.7	84
81	Cutaneous Desmoid Tumor. <i>Dermatologic Surgery</i> , 2009, 35, 1582-1587.	0.8	5
82	3q26 (hTERC) gain studied by fluorescence in situ hybridization as a persistence-progression indicator in low-grade squamous intraepithelial lesion cases. <i>Human Pathology</i> , 2009, 40, 1474-1478.	2.0	29
83	FISH and immunohistochemical status of the hepatocyte growth factor receptor (c-Met) in 184 invasive breast tumors. <i>Breast Cancer Research</i> , 2009, 11, 402.	5.0	22
84	HPV Determination in the Control After LEEP Due to CIN II-III: Prospective Study and Predictive Model. <i>International Journal of Gynecological Pathology</i> , 2009, 28, 120-126.	1.4	11
85	Multiple recurrent chromosomal breakpoints in mantle cell lymphoma revealed by a combination of molecular cytogenetic techniques. <i>Genes Chromosomes and Cancer</i> , 2008, 47, 1086-1097.	2.8	28
86	FISH is better than BIOMED-2 PCR to detect IgH/BCL2 translocation in follicular lymphoma at diagnosis using paraffin-embedded tissue sections. <i>Leukemia Research</i> , 2008, 32, 737-742.	0.8	37
87	New chromosomal alterations in a series of 23 splenic marginal zone lymphoma patients revealed by Spectral Karyotyping (SKY). <i>Leukemia Research</i> , 2008, 32, 727-736.	0.8	20
88	Blast cells with nuclear extrusions in the form of micronuclei are associated with MYC amplification in acute myeloid leukemia. <i>Cancer Genetics and Cytogenetics</i> , 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-. <i>Haematologica</i> , 2008, 93, 1001-1008.	3.5	36
90	Monosomy 7 with severe myelodysplasia developing during imatinib treatment of Philadelphia-positive chronic myeloid leukemia: Two cases with a different outcome. <i>American Journal of Hematology</i> , 2007, 82, 849-851.	4.1	33

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91	Gain of multiple copies of the CBFβ gene: a new genetic aberration in a case of granulocytic sarcoma. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Human Pathology</i> , 2006, 37, 867-873.	2.0	36
93	Gastrointestinal Involvement in Mantle Cell Lymphoma: A Prospective Clinic, Endoscopic, and Pathologic Study. <i>American Journal of Surgical Pathology</i> , 2006, 30, 1274-1280.	3.7	121
94	Study of chromosomal abnormalities in 11 cases of cervical dysplasia using comparative genomic hybridization on cotton-lint cervical samples. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Cancer Genetics and Cytogenetics</i> , 2006, 167, 39-42.	1.0	6
96	Methotrexate resistance in vitro is achieved by a dynamic selection process of tumor cell variants emerging during treatment. <i>International Journal of Cancer</i> , 2006, 119, 1607-1615.	5.1	9
97	Comparative genomic hybridization analysis of cutaneous large B-cell lymphomas. <i>Experimental Dermatology</i> , 2005, 14, 883-890.	2.9	17
98	Insertion (8;11) in a renal oncocytoma with multifocal transformation to chromophobe renal cell carcinoma. <i>Cancer Genetics and Cytogenetics</i> , 2005, 163, 160-163.	1.0	11
99	Clonal proliferation of cyclin D1 ⁺ positive mantle lymphocytes in an asymptomatic patient: an early-stage event in the development or an indolent form of a mantle cell lymphoma?. <i>Human Pathology</i> , 2005, 36, 1232-1237.	2.0	41
100	Identification of Male Cardiomyocytes of Extracardiac Origin in the Hearts of Women with Male Progeny: Male Fetal Cell Microchimerism of the Heart. <i>Journal of Heart and Lung Transplantation</i> , 2005, 24, 2179-2183.	0.6	78
101	Correlation between histologic findings and cytogenetic abnormalities in bladder carcinoma: A FISH study. <i>Urology</i> , 2005, 65, 913-918.	1.0	14
102	Polysomy of chromosome 17 in breast cancer tumors showing an overexpression of ERBB2: a study of 175 cases using fluorescence in situ hybridization and immunohistochemistry. <i>Breast Cancer Research</i> , 2005, 7, R267-73.	5.0	76
103	Identification of novel cytogenetic markers with prognostic significance in a series of 968 patients with primary myelodysplastic syndromes. <i>Haematologica</i> , 2005, 90, 1168-78.	3.5	163
104	Activation of mitochondrial apoptotic pathway in mantle cell lymphoma: high sensitivity to mitoxantrone in cases with functional DNA-damage response genes. <i>Oncogene</i> , 2004, 23, 8941-8949.	5.9	23
105	Cytogenetic and fluorescence in situ hybridization studies in 60 patients with multiple myeloma and plasma cell leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2004, 148, 71-76.	1.0	18
106	Is fluorescence in situ hybridization a useful method in diagnosis of polycythemia vera patients?. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Blood</i> , 2004, 103, 2427-2428.	1.4	18
108	Genetic characterization of SÅžary's syndrome by conventional cytogenetics and cross-species color banding fluorescent in situ hybridization. <i>Haematologica</i> , 2004, 89, 165-73.	3.5	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. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Cancer Genetics and Cytogenetics</i> , 2003, 142, 165-167.	1.0	5
111	Cytogenetic findings in familial B-cell chronic lymphocytic leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2003, 143, 172-173.	1.0	5
112	RxFISH karyotype andMYC amplification in the HT-29 colon adenocarcinoma cell line. <i>Genes Chromosomes and Cancer</i> , 2003, 36, 319-320.	2.8	6
113	Comparative Analysis of TCR- β Gene Rearrangements by Genescan and Polyacrylamide Gel-electrophoresis in Cutaneous T-cell Lymphoma. <i>Acta Dermato-Venereologica</i> , 2003, 84, 6-11.	1.3	11
114	Analysis of T-Cell Receptor β Gene Rearrangements by PCR-Genescan and PCR-Polyacrylamide Gel Electrophoresis in Early-Stage Mycosis fungoides/Large-Plaque Parapsoriasis. <i>Dermatology</i> , 2003, 207, 418-419.	2.1	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. <i>Haematologica</i> , 2003, 88, 110-1.	3.5	10
116	Nodal marginal zone lymphoma in AIDS patients. <i>Aids</i> , 2002, 16, 2232-2234.	2.2	2
117	Genetic characterization of the paraimmunoblastic variant of small lymphocytic lymphoma/chronic lymphocytic leukemia: A case report and review of the literature. <i>Human Pathology</i> , 2002, 33, 1145-1148.	2.0	5
118	Contribution of cytogenetics and in situ hybridization to the study of monoclonal gammopathies of undetermined significance. <i>Cancer Genetics and Cytogenetics</i> , 2002, 132, 25-29.	1.0	10
119	Translocation (5;17)(q13;q21) in a case with precursor T-lymphoblastic lymphoma/leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2002, 132, 81-82.	1.0	1
120	Pentasomy 21 with two isochromosomes 21 in a case of acute myeloid leukemia without maturation. <i>Cancer Genetics and Cytogenetics</i> , 2002, 132, 71-73.	1.0	7
121	A new case of acute nonlymphocytic leukemia (French-American-British subtype M1) with double minutes and c-MYC amplification. <i>Cancer Genetics and Cytogenetics</i> , 2002, 132, 161-164.	1.0	6
122	Monosomy 15 in chronic myelomonocytic leukemia. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>European Urology</i> , 2002, 42, 547-552.	1.9	74
124	Report of 46,XX/46,XY/47,XXY/48,XXYY mosaicism in an adult phenotypic male. <i>American Journal of Medical Genetics Part A</i> , 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. <i>Cancer Genetics and Cytogenetics</i> , 2001, 125, 70-73.	1.0	3
126	Isochromosome +i(3)(q10) in a new case of persistent polyclonal B-cell lymphocytosis (PPBL). <i>European Journal of Haematology</i> , 2000, 64, 344-346.	2.2	0

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127	Cytogenetic and Fluorescence In Situ Hybridization Studies in Four Cases of Plasma Cell Leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2000, 121, 163-166.	1.0	1
128	Dicentric (17;18) in a Case of Atypical B-Cell Chronic Lymphocytic Leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2000, 121, 194-197.	1.0	6
129	A New Case of Turner Syndrome Associated with Multiple Myeloma. <i>Cancer Genetics and Cytogenetics</i> , 2000, 117, 80-81.	1.0	4
130	Translocation t(6;14)(p12;q32): a novel cytogenetic abnormality in splenic lymphoma with villous lymphocytes. <i>British Journal of Haematology</i> , 2000, 110, 241-243.	2.5	8
131	Cytogenetic Findings in Five Patients with Hairy Cell Leukemia. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Cancer Genetics and Cytogenetics</i> , 1999, 111, 92-98.	1.0	31
133	Two New Cases of Near-Tetraploidy in Adult Acute Myeloid Leukemia. <i>Cancer Genetics and Cytogenetics</i> , 1998, 102, 131-134.	1.0	11
134	Cytogenetic Abnormalities in Three Patients with B-Cell Prolymphocytic Leukemia. <i>Cancer Genetics and Cytogenetics</i> , 1998, 103, 43-45.	1.0	12
135	Coexistence of tetrasomy 8 and trisomy 8 in a case with myeloid metaplasia with myelofibrosis. <i>Cancer Genetics and Cytogenetics</i> , 1997, 94, 147-150.	1.0	9
136	Frequent involvement of chromosomes 1, 3, 7 and 8 in splenic marginal zone B-cell lymphoma. <i>British Journal of Haematology</i> , 1997, 98, 446-449.	2.5	56