

# Jose Cervera

## List of Publications by Year in descending order

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138  
papers

5,141  
citations

136885

32  
h-index

102432

66  
g-index

142  
all docs

142  
docs citations

142  
times ranked

5612  
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular International Prognostic Scoring System for Myelodysplastic Syndromes. , 2022, 1, .		259
2	Genomic characterization of patients with polycythemia vera developing resistance to hydroxyurea. <i>Leukemia</i> , 2021, 35, 623-627.	3.3	12
3	Impact of combinations of single-nucleotide polymorphisms of anthracycline transporter genes upon the efficacy and toxicity of induction chemotherapy in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2021, 62, 659-668.	0.6	10
4	Healthcare resource utilization in adult patients with relapsed/refractory FLT3 mutated acute myeloid leukemia: A retrospective chart review from Spain. <i>European Journal of Haematology</i> , 2021, 106, 724-733.	1.1	1
5	A Single-Run Next-Generation Sequencing (NGS) Assay for the Simultaneous Detection of Both Gene Mutations and Large Chromosomal Abnormalities in Patients with Myelodysplastic Syndromes (MDS) and Related Myeloid Neoplasms. <i>Cancers</i> , 2021, 13, 1947.	1.7	5
6	Aberrant Alternative Splicing in U2af1/Tet2 Double Mutant Mice Contributes to Major Hematological Phenotypes. <i>International Journal of Molecular Sciences</i> , 2021, 22, 6963.	1.8	5
7	Adverse prognostic impact of complex karyotype (≥3 cytogenetic alterations) in adult T-cell acute lymphoblastic leukemia (T-ALL). <i>Leukemia Research</i> , 2021, 109, 106612.	0.4	11
8	Genome-wide association study identifies susceptibility loci for acute myeloid leukemia. <i>Nature Communications</i> , 2021, 12, 6233.	5.8	17
9	Spanish Guidelines for the use of targeted deep sequencing in myelodysplastic syndromes and chronic myelomonocytic leukaemia. <i>British Journal of Haematology</i> , 2020, 188, 605-622.	1.2	25
10	Implications of TP53 allelic state for genome stability, clinical presentation and outcomes in myelodysplastic syndromes. <i>Nature Medicine</i> , 2020, 26, 1549-1556.	15.2	372
11	Helpful Criteria When Implementing NGS Panels in Childhood Lymphoblastic Leukemia. <i>Journal of Personalized Medicine</i> , 2020, 10, 244.	1.1	1
12	Acute Promyelocytic Leukemia: A Constellation of Molecular Events around a Single PML-RARA Fusion Gene. <i>Cancers</i> , 2020, 12, 624.	1.7	77
13	Factor XIII deficiency in two Spanish families with a novel variant in gene F13A1 detected by next-generation sequencing; symptoms and clinical management. <i>Journal of Thrombosis and Thrombolysis</i> , 2020, 50, 686-688.	1.0	4
14	Impact of somatic mutations in myelodysplastic patients with isolated partial or total loss of chromosome 7. <i>Leukemia</i> , 2020, 34, 2441-2450.	3.3	14
15	Analysis of SNP Array Abnormalities in Patients with DE NOVO Acute Myeloid Leukemia with Normal Karyotype. <i>Scientific Reports</i> , 2020, 10, 5904.	1.6	8
16	Chronic graft-versus-host disease could ameliorate the impact of adverse somatic mutations in patients with myelodysplastic syndromes and hematopoietic stem cell transplantation. <i>Annals of Hematology</i> , 2019, 98, 2151-2162.	0.8	2
17	Clinical and molecular characterization by next generation sequencing of Spanish patients affected by congenital deficiencies of fibrinogen. <i>Thrombosis Research</i> , 2019, 180, 115-117.	0.8	8
18	Incidence and outcome of invasive fungal disease after front-line intensive chemotherapy in patients with acute myeloid leukemia: impact of antifungal prophylaxis. <i>Annals of Hematology</i> , 2019, 98, 2081-2088.	0.8	16

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19	RNA Sequencing Analysis for the Identification of a PCM1/PDGFRB Fusion Gene Responsive to Imatinib. <i>Acta Haematologica</i> , 2019, 142, 92-97.	0.7	6
20	CRISPR to fix bad blood: a new tool in basic and clinical hematology. <i>Haematologica</i> , 2019, 104, 881-893.	1.7	26
21	Clinical Utility of a Next-Generation Sequencing Panel for Acute Myeloid Leukemia Diagnostics. <i>Journal of Molecular Diagnostics</i> , 2019, 21, 228-240.	1.2	24
22	Clinical significance of complex karyotype at diagnosis in pediatric and adult patients with de novo acute promyelocytic leukemia treated with ATRA and chemotherapy. <i>Leukemia and Lymphoma</i> , 2019, 60, 1146-1155.	0.6	12
23	TP53 State Dictates Genome Stability, Clinical Presentation and Outcomes in Myelodysplastic Syndromes. <i>Blood</i> , 2019, 134, 675-675.	0.6	17
24	Germinal Predisposition in Myelodysplastic Syndromes in Young Adults without a Preexisting Disorder or Organ Dysfunction: Identification of Deleterious Variants in Microsatellite Instability Genes. <i>Blood</i> , 2019, 134, 4226-4226.	0.6	0
25	Clonal genetic evolution at relapse of favorable-risk acute myeloid leukemia with NPM1 mutation is associated with phenotypic changes and worse outcomes. <i>Haematologica</i> , 2018, 103, e400-e403.	1.7	9
26	Guía práctica de tratamiento urgente de la microangiopatía trombótica. <i>Medicina Clínica</i> , 2018, 151, 123.e1-123.e9.	0.3	7
27	Clonal architecture in patients with myelodysplastic syndromes and double or minor complex abnormalities: Detailed analysis of clonal composition, involved abnormalities, and prognostic significance. <i>Genes Chromosomes and Cancer</i> , 2018, 57, 547-556.	1.5	3
28	The modular network structure of the mutational landscape of Acute Myeloid Leukemia. <i>PLoS ONE</i> , 2018, 13, e0202926.	1.1	5
29	Frequency and prognostic significance of t(v;11q23)/KMT2A rearrangements in adult patients with acute lymphoblastic leukemia treated with risk-adapted protocols. <i>Leukemia and Lymphoma</i> , 2017, 58, 145-152.	0.6	7
30	Influence of cytarabine metabolic pathway polymorphisms in acute myeloid leukemia induction treatment. <i>Leukemia and Lymphoma</i> , 2017, 58, 2880-2894.	0.6	12
31	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.	0.4	9
32	Negative impact on clinical outcome of the mutational co-occurrence of SF3B1 and DNMT3A in refractory anemia with ring sideroblasts (RARS). <i>Leukemia and Lymphoma</i> , 2017, 58, 1686-1693.	0.6	14
33	Impact of ABC single nucleotide polymorphisms upon the efficacy and toxicity of induction chemotherapy in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2017, 58, 1197-1206.	0.6	33
34	Prognostic impact of gene mutations in myelodysplastic syndromes with ring sideroblasts. <i>Blood Cancer Journal</i> , 2017, 7, 630.	2.8	11
35	Mutations in the DNA methylation pathway and number of driver mutations predict response to azacitidine in myelodysplastic syndromes. <i>Oncotarget</i> , 2017, 8, 106948-106961.	0.8	38
36	Karyotypic complexity rather than chromosome 8 abnormalities aggravates the outcome of chronic lymphocytic leukemia patients with TP53 aberrations. <i>Oncotarget</i> , 2016, 7, 80916-80924.	0.8	29

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37	Impact of <sc>SNP</sc> array karyotyping on the diagnosis and the outcome of chronic myelomonocytic leukemia with low risk cytogenetic features or no metaphases. American Journal of Hematology, 2016, 91, 185-192.	2.0	18
38	Analysis of the Possible Persistent Genotoxic Damage in Workers Linked to the Ardystil Syndrome. Genetic Testing and Molecular Biomarkers, 2016, 20, 94-97.	0.3	1
39	Integrating clinical features and genetic lesions in the risk assessment of patients with chronic myelomonocytic leukemia. Blood, 2016, 128, 1408-1417.	0.6	249
40	Prognostic impact of chromosomal translocations in myelodysplastic syndromes and chronic myelomonocytic leukemia patients. A study by the spanish group of myelodysplastic syndromes. Genes Chromosomes and Cancer, 2016, 55, 322-327.	1.5	7
41	Study of the S427C polymorphism and of MYBL2 variants in patients with acute myeloid leukemia. Leukemia and Lymphoma, 2016, 57, 429-435.	0.6	10
42	The Mutational Landscape of Acute Promyelocytic Leukemia Reveals an Interacting Network of Co-Occurrences and Recurrent Mutations. PLoS ONE, 2016, 11, e0148346.	1.1	23
43	Comparison of the Molecular Spectrum of Lenalidomide-Treated Myelodysplastic Syndrome with and without Del(5q). Blood, 2016, 128, 3172-3172.	0.6	0
44	Clonal Hematopoiesis Landscape in Patients with De Novo Acute Myeloid Leukemia By Deep Sequencing. Blood, 2016, 128, 598-598.	0.6	0
45	Influence of Cytarabine Metabolic Pathway Polymorphisms in Acute Myeloid Leukemia Induction Treatment. Blood, 2016, 128, 5130-5130.	0.6	0
46	CIP2A high expression is a poor prognostic factor in normal karyotype acute myeloid leukemia. Haematologica, 2015, 100, e183-e185.	1.7	20
47	Correlation of myelodysplastic syndromes with i(17)(q10) and <i><sc>TP</sc>53</i> and <i><sc>SETBP</sc>1</i> mutations. British Journal of Haematology, 2015, 171, 137-141.	1.2	11
48	Somatic Genetic and Epigenetic Architecture of Myelodysplastic Syndromes Arising from GATA2 Deficiency. Blood, 2015, 126, 299-299.	0.6	10
49	Impact of Transporter Genes Polymorphisms in Standard Induction of Acute Myeloid Leukemia. Blood, 2015, 126, 4842-4842.	0.6	1
50	Influence of Single Nucleotide Polymorphisms in Anthracycline Metabolism Pathway in Standard Induction of Acute Myeloid Leukemia. Blood, 2015, 126, 4845-4845.	0.6	1
51	Validation of a Next-Generation Sequencing Panel for AML Routine Diagnosis. Blood, 2015, 126, 1403-1403.	0.6	0
52	Simplified in-House Deep Sequencing Method of Immunoglobulin Genes for Minimal Residual Disease Quantification and Risk Stratification in Multiple Myeloma. Blood, 2015, 126, 2972-2972.	0.6	0
53	Single-Nucleotide Polymorphism Array-Based Karyotyping of Acute Promyelocytic Leukemia. PLoS ONE, 2014, 9, e100245.	1.1	7
54	Prognostic significance of complex karyotype and monosomal karyotype in adult patients with acute lymphoblastic leukemia treated with risk-adapted protocols. Cancer, 2014, 120, 3958-3964.	2.0	24

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55	In vitro all-trans retinoic acid sensitivity of acute myeloid leukemia blasts with NUP98/RARG fusion gene. <i>Annals of Hematology</i> , 2014, 93, 1931-1933.	0.8	21
56	Sister chromatid exchange, (SCE), High-Frequency Cells (HFCs) and SCE distribution patterns in peripheral blood lymphocytes of Spanish adult smokers compared to non-smokers. <i>Food and Chemical Toxicology</i> , 2014, 66, 107-112.	1.8	9
57	Prognostic Impact of Anthracycline Metabolism Gene Polymorphisms in Newly Diagnosed Acute Myeloid Leukemia Adults. <i>Blood</i> , 2014, 124, 2237-2237.	0.6	1
58	Utility of SNP Arrays in Chronic Myelomonocytic Leukemia with Low Risk Cytogenetic Features or No Metaphases. <i>Blood</i> , 2014, 124, 4659-4659.	0.6	0
59	Somatic Mutations of ASXL1, RUNX1 and SETBP1 Improve Prognostic Stratification of Patients with Chronic Myelomonocytic Leukemia. <i>Blood</i> , 2014, 124, 1915-1915.	0.6	6
60	Genomic Microarray Alterations Add Prognostic Power to the IPSS-R in MDS with Normal Karyotype. <i>Blood</i> , 2014, 124, 3262-3262.	0.6	0
61	Prognostic Impact of Cytarabine Pathway Gene Polymorphisms in Acute Myeloid Leukemia Adults Undergoing Induction Chemotherapy. <i>Blood</i> , 2014, 124, 5223-5223.	0.6	0
62	Novel Real-Time Polymerase Chain Reaction Assay for Simultaneous Detection of Recurrent Fusion Genes in Acute Myeloid Leukemia. <i>Journal of Molecular Diagnostics</i> , 2013, 15, 678-686.	1.2	8
63	Characterization and prognostic implication of 17 chromosome abnormalities in myelodysplastic syndrome. <i>Leukemia Research</i> , 2013, 37, 769-776.	0.4	11
64	Reciprocal translocations in myelodysplastic syndromes and chronic myelomonocytic leukemias: Review of 5,654 patients with an evaluable karyotype. <i>Genes Chromosomes and Cancer</i> , 2013, 52, 753-763.	1.5	15
65	Frequency of dicentrics and contamination levels in Ukrainian children and adolescents from areas near Chernobyl 20 years after the nuclear plant accident. <i>International Journal of Radiation Biology</i> , 2013, 89, 944-949.	1.0	2
66	Radioprotective activity and cytogenetic effect of resveratrol in human lymphocytes: An in vitro evaluation. <i>Food and Chemical Toxicology</i> , 2013, 51, 391-395.	1.8	32
67	Application of FISH 7q in MDS patients without monosomy 7 or 7q deletion by conventional G-banding cytogenetics: Does $\sim 7/7q$ detection by FISH have prognostic value?. <i>Leukemia Research</i> , 2013, 37, 416-421.	0.4	16
68	Adverse prognostic value of MYBL2 overexpression and association with microRNA-30 family in acute myeloid leukemia patients. <i>Leukemia Research</i> , 2013, 37, 1690-1696.	0.4	36
69	WT1 isoform expression pattern in acute myeloid leukemia. <i>Leukemia Research</i> , 2013, 37, 1744-1749.	0.4	11
70	Response to lenalidomide in myelodysplastic syndromes with del(5q): influence of cytogenetics and mutations. <i>British Journal of Haematology</i> , 2013, 162, 74-86.	1.2	73
71	Screening for IDH mutations in chronic myelomonocytic leukemia. <i>Leukemia and Lymphoma</i> , 2013, 54, 406-407.	0.6	4
72	Assessing an Improved Protocol for Plasma microRNA Extraction. <i>PLoS ONE</i> , 2013, 8, e82753.	1.1	81

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73	<i>BRAF</i> V600E mutation in adult acute lymphoblastic leukemia. <i>Leukemia and Lymphoma</i> , 2013, 54, 1105-1106.	0.6	4
74	Single nucleotide polymorphism array karyotyping: A diagnostic and prognostic tool in myelodysplastic syndromes with unsuccessful conventional cytogenetic testing. <i>Genes Chromosomes and Cancer</i> , 2013, 52, 1167-1177.	1.5	44
75	Development and validation of a prognostic scoring system for patients with chronic myelomonocytic leukemia. <i>Blood</i> , 2013, 121, 3005-3015.	0.6	251
76	Prognostic value of trisomy 8 as a single anomaly and the influence of additional cytogenetic aberrations in primary myelodysplastic syndromes. <i>British Journal of Haematology</i> , 2012, 159, 311-321.	1.2	25
77	Rapid Screening of ASXL1, IDH1, IDH2, and c-CBL Mutations in de Novo Acute Myeloid Leukemia by High-Resolution Melting. <i>Journal of Molecular Diagnostics</i> , 2012, 14, 594-601.	1.2	12
78	New Comprehensive Cytogenetic Scoring System for Primary Myelodysplastic Syndromes (MDS) and Oligoblastic Acute Myeloid Leukemia After MDS Derived From an International Database Merge. <i>Journal of Clinical Oncology</i> , 2012, 30, 820-829.	0.8	584
79	In vitro cytogenetic and genotoxic effects of curcumin on human peripheral blood lymphocytes. <i>Food and Chemical Toxicology</i> , 2012, 50, 3229-3233.	1.8	19
80	Analysis of SNP rs16754 of WT1 gene in a series of de novo acute myeloid leukemia patients. <i>Annals of Hematology</i> , 2012, 91, 1845-1853.	0.8	8
81	Functional polymorphisms in SOCS1 and PTPN22 genes correlate with the response to imatinib treatment in newly diagnosed chronic-phase chronic myeloid leukemia. <i>Leukemia Research</i> , 2012, 36, 174-181.	0.4	17
82	Absence of mutations in the activation loop and juxtamembrane domains of VEGFR-1 and VEGFR-2 gene in chronic myelomonocytic leukemia (CMML). <i>Leukemia Research</i> , 2012, 36, e50-e51.	0.4	0
83	Better prognosis for patients with del(7q) than for patients with monosomy 7 in myelodysplastic syndrome. <i>Cancer</i> , 2012, 118, 127-133.	2.0	43
84	Fragment length analysis screening for detection of CEBPA mutations in intermediate-risk karyotype acute myeloid leukemia. <i>Annals of Hematology</i> , 2012, 91, 1-7.	0.8	13
85	Prognostic value of cytogenetics in adult patients with Philadelphia-negative acute lymphoblastic leukemia. <i>Annals of Hematology</i> , 2012, 91, 19-25.	0.8	5
86	Association of MDM2 Gene Polymorphisms SNP285 and 309 with Myelodysplastic Syndromes (MDS) Susceptibility and Outcome. <i>Blood</i> , 2012, 120, 2823-2823.	0.6	0
87	Cytogenetic risk stratification in chronic myelomonocytic leukemia. <i>Haematologica</i> , 2011, 96, 375-383.	1.7	226
88	A novel NUP98/RARG gene fusion in acute myeloid leukemia resembling acute promyelocytic leukemia. <i>Blood</i> , 2011, 117, 242-245.	0.6	65
89	Down-regulation of EVI1 is associated with epigenetic alterations and good prognosis in patients with acute myeloid leukemia. <i>Haematologica</i> , 2011, 96, 1448-1456.	1.7	45
90	CYP2C8 gene polymorphism and bisphosphonate-related osteonecrosis of the jaw in patients with multiple myeloma. <i>Haematologica</i> , 2011, 96, 1557-1559.	1.7	31

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91	Correlation Between Preoperative Magnetic Resonance Imaging and Surgical Margins with Modified Mohs for Dermatofibrosarcoma Protuberans. <i>Dermatologic Surgery</i> , 2011, 37, 1638-1645.	0.4	53
92	Assessment inÂvitro of radioprotective efficacy of curcumin and resveratrol. <i>Radiation Measurements</i> , 2011, 46, 962-966.	0.7	29
93	Aberrant methylation of tumor suppressor genes in patients with refractory anemia with ring sideroblasts. <i>Leukemia Research</i> , 2011, 35, 479-483.	0.4	9
94	Clinical impact of the clone size in MDS cases with monosomy 7 or 7q deletion, trisomy 8, 20q deletion and loss of Y chromosome. <i>Leukemia Research</i> , 2011, 35, 834-836.	0.4	11
95	Gene Microdeletions in Adult and Pediatric Acute Lymphoblastic Leukemia,. <i>Blood</i> , 2011, 118, 3539-3539.	0.6	2
96	Quantitative Expression Analysis of WT1 Main Isoforms in AML. <i>Blood</i> , 2011, 118, 1469-1469.	0.6	0
97	Analysis of ASXL1, IDH1, IDH2, c-CBL, and WT1 Mutations in De Novo Acute Myeloid Leukaemia. <i>Blood</i> , 2011, 118, 1450-1450.	0.6	0
98	Additional chromosome abnormalities in patients with acute promyelocytic leukemia treated with all-trans retinoic acid and chemotherapy. <i>Haematologica</i> , 2010, 95, 424-431.	1.7	84
99	Epigenetic regulation of the non- canonical Wnt pathway in acute myeloid leukemia. <i>Cancer Science</i> , 2010, 101, 425-432.	1.7	43
100	New Chemistry Cartridge for the GEM Premier 4000 Critical Care Analyzer, Including Electrochemical Sensors for Creatinine, Urea, and Measured Bicarbonate. <i>Point of Care</i> , 2010, 9, 1-7.	0.5	1
101	A new reliable fluorescence in situ hybridization method for identifying multiple specific cytogenetic abnormalities in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2010, 51, 680-685.	0.6	3
102	DNA Methylation Profiles and Their Relationship with Cytogenetic Status in Adult Acute Myeloid Leukemia. <i>PLoS ONE</i> , 2010, 5, e12197.	1.1	73
103	Loss of the Y Chromosome in MDS - Age-Related Phenomenon or Clonal Abnormality?. <i>Blood</i> , 2010, 116, 4008-4008.	0.6	0
104	Impact of hematopoietic chimerism at day +14 on engraftment after unrelated donor umbilical cord blood transplantation for hematologic malignancies. <i>Haematologica</i> , 2009, 94, 827-832.	1.7	29
105	Rapid Detection of KIT Mutations in Core-Binding Factor Acute Myeloid Leukemia Using High-Resolution Melting Analysis. <i>Journal of Molecular Diagnostics</i> , 2009, 11, 458-463.	1.2	24
106	Minimal residual disease detection in acute myeloid leukemia by mutant nucleophosmin (NPM1): Comparison with WT1 gene expression. <i>Clinica Chimica Acta</i> , 2008, 395, 120-123.	0.5	32
107	Measurement of Creatinine in Whole Blood Samples Supplemented to Achieve Increased Creatinine Concentrations. <i>Clinical Chemistry</i> , 2008, 54, 451-452.	1.5	6
108	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.	1.7	36

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109	Molecular analysis of t(15;17) genomic breakpoints in secondary acute promyelocytic leukemia arising after treatment of multiple sclerosis. <i>Blood</i> , 2008, 112, 3383-3390.	0.6	74
110	Lack of RPS14 promoter aberrant methylation supports the haploinsufficiency model for the 5q-syndrome. <i>Blood</i> , 2008, 112, 918-918.	0.6	78
111	A comprehensive strategy for the subtyping of patients with Fanconi anaemia: conclusions from the Spanish Fanconi Anemia Research Network. <i>Journal of Medical Genetics</i> , 2007, 44, 241-249.	1.5	47
112	The potential effect of gender in combination with common genetic polymorphisms of drug-metabolizing enzymes on the risk of developing acute leukemia. <i>Haematologica</i> , 2007, 92, 308-314.	1.7	76
113	1: Analysis of Risk Factors in Adults Transplanted with UCB for Treatment of Hematologic Malignancy. <i>Biology of Blood and Marrow Transplantation</i> , 2007, 13, 1393.	2.0	0
114	The GST deletions and NQO1*2 polymorphism confers interindividual variability of response to treatment in patients with acute myeloid leukemia. <i>Leukemia Research</i> , 2007, 31, 947-953.	0.4	40
115	Could ATRA/Idarubicin treatment of acute promyelocytic leukemia induce the appearance of new clonal cytogenetic abnormalities in patients in complete remission?. <i>Leukemia Research</i> , 2007, 31, 1315-1317.	0.4	2
116	Profile of polymorphisms of drug-metabolising enzymes and the risk of therapy-related leukaemia. <i>British Journal of Haematology</i> , 2007, 136, 590-596.	1.2	75
117	MLL amplification in acute myeloid leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2007, 174, 127-131.	1.0	17
118	Incidence and Characteristics of Acute Promyelocytic Leukemia in Adult Patients. A Single Centre Registry Study.. <i>Blood</i> , 2007, 110, 4301-4301.	0.6	0
119	Influence of genetic polymorphisms on the risk of developing leukemia and on disease progression. <i>Leukemia Research</i> , 2006, 30, 1471-1491.	0.4	94
120	A t(17;20)(q21;q12) masking a variant t(15;17)(q22;q21) in a patient with acute promyelocytic leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2006, 168, 73-76.	1.0	9
121	High-level amplification of the RUNX1 gene in two cases of childhood acute lymphoblastic leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2006, 170, 171-174.	1.0	9
122	EVII is consistently expressed as principal transcript in common and rare recurrent 3q26 rearrangements. <i>Genes Chromosomes and Cancer</i> , 2006, 45, 349-356.	1.5	51
123	Long-Term Outcome and Prognostic Factors after Single-Unit Umbilical Cord-Blood Transplantation (UCBT) for Adults with Hematologic Malignancies.. <i>Blood</i> , 2006, 108, 3129-3129.	0.6	0
124	The JAK2 V617F Mutation Identifies Specific Subgroups of Myelodysplastic Syndrome (MDS).. <i>Blood</i> , 2006, 108, 2610-2610.	0.6	0
125	SHP1 expression in bone marrow biopsies of myelodysplastic syndrome patients: a new prognostic factor. <i>British Journal of Haematology</i> , 2005, 129, 791-794.	1.2	22
126	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.	1.7	163



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127	Immunofluorescent analysis with the anti-PML monoclonal antibody PG-M3 for rapid and accurate genetic diagnosis of acute promyelocytic leukemia. <i>Annals of Hematology</i> , 2004, 83, 687-690.	0.8	21
128	Locked Nucleic Acid-Enhanced Detection of 1100delC*CHEK2 Germ-Line Mutation in Spanish Patients with Hematologic Malignancies. <i>Clinical Chemistry</i> , 2004, 50, 2201-2204.	1.5	7
129	Additional Chromosome Abnormalities Have No Prognostic Value in Acute Promyelocytic Leukemia Patients Treated with Simultaneous ATRA and Anthracycline-Based Chemotherapy: An Update of the APL96 and APL99 Pethema Protocols.. <i>Blood</i> , 2004, 104, 2019-2019.	0.6	5
130	Prognostic implications of Wilms' tumor gene (WT1) expression in patients with de novo acute myeloid leukemia. <i>Haematologica</i> , 2004, 89, 926-33.	1.7	97
131	Incidence and prognostic value of FLT3 internal tandem duplication and D835 mutations in acute myeloid leukemia. <i>Haematologica</i> , 2003, 88, 19-24.	1.7	135
132	The t(4;22)(q12;q11) in atypical chronic myeloid leukaemia fuses BCR to PDGFRA. <i>Human Molecular Genetics</i> , 2002, 11, 1391-1397.	1.4	139
133	Identification of two atypical PML-RAR± transcripts in two patients with acute promyelocytic leukemia. <i>Leukemia Research</i> , 2002, 26, 439-442.	0.4	18
134	FLAG-IDA regimen (fludarabine, cytarabine, idarubicin and G-CSF) in the treatment of patients with high-risk myeloid malignancies. <i>Leukemia Research</i> , 2002, 26, 725-730.	0.4	41
135	Standardized, unrelated donor cord blood transplantation in adults with hematologic malignancies. <i>Blood</i> , 2001, 98, 2332-2338.	0.6	220
136	Recent improvements in outcome for elderly patients with de novo acute myeloblastic leukemia. <i>Leukemia Research</i> , 2001, 25, 685-692.	0.4	12
137	Graft-versus-tumour effect in non-small-cell lung cancer after allogeneic peripheral blood stem cell transplantation. SHORT REPORT. <i>British Journal of Haematology</i> , 2000, 111, 708-710.	1.2	29
138	Liver disease as primary manifestation of multiple myeloma in a young man. <i>Leukemia Research</i> , 1999, 23, 403-405.	0.4	14