Christine J Harrison

List of Publications by Year
in descending order

195 papers	21,011 citations	11908 72 h-index	11946 139 g-index
203 all docs	203 docs citations	203 times ranked	14467 citing authors

#	Article	IF	Citations
1	Prognostic significance of chromosomal abnormalities at relapse in children with relapsed acute myeloid leukemia: A retrospective cohort study of the Relapsed AML 2001/01 Study. Pediatric Blood and Cancer, 2022, 69, e29341.	0.8	5
2	Prognostic impact of chromosomal abnormalities and copy number alterations in adult B-cell precursor acute lymphoblastic leukaemia: a UKALL14 study. Leukemia, 2022, 36, 625-636.	3.3	25
3	Genetic characterisation of childhood Bâ€otherâ€acute lymphoblastic leukaemia in UK patients by fluorescence inÂsitu hybridisation and Multiplex Ligationâ€dependent Probe Amplification. British Journal of Haematology, 2022, 196, 753-763.	1.2	12
4	The 5th edition of the World Health Organization Classification of Haematolymphoid Tumours: Lymphoid Neoplasms. Leukemia, 2022, 36, 1720-1748.	3.3	1,023
5	Time to Cure for Childhood and Young Adult Acute Lymphoblastic Leukemia Is Independent of Early Risk Factors: Long-Term Follow-Up of the UKALL2003 Trial. Journal of Clinical Oncology, 2022, 40, 4228-4239.	0.8	8
6	Characterization of unusual <scp>iAMP21</scp> Bâ€lymphoblastic leukemia (<scp>iAMP21â€ALL</scp>) from the Mayo Clinic and Children's Oncology Group. Genes Chromosomes and Cancer, 2022, 61, 710-719.	1.5	14
7	SSBP2-CSF1R is a recurrent fusion in B-lineage acute lymphoblastic leukemia with diverse genetic presentation and variable outcome. Blood, 2021, 137, 1835-1838.	0.6	6
8	<scp>MYB</scp> rearrangements and overâ€expression in Tâ€cell acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2021, 60, 482-488.	1.5	12
9	14q32 rearrangements deregulating BCL11B mark a distinct subgroup of T and myeloid immature acute leukemia. Blood, 2021, 138, 773-784.	0.6	19
10	Single nucleotide polymorphism arrayâ€based signature of low hypodiploidy in acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2021, 60, 604-615.	1.5	12
11	Cytogenetics of Pediatric Acute Myeloid Leukemia: A Review of the Current Knowledge. Genes, 2021, 12, 924.	1.0	39
12	Defining low-risk high hyperdiploidy in patients with paediatric acute lymphoblastic leukaemia: a retrospective analysis of data from the UKALL97/99 and UKALL2003 clinical trials. Lancet Haematology,the, 2021, 8, e828-e839.	2.2	25
13	PCR amplicons identify widespread copy number variation in human centromeric arrays and instability in cancer. Cell Genomics, 2021, 1, 100064.	3.0	14
14	Concordance of copy number abnormality detection using SNP arrays and Multiplex Ligation-dependent Probe Amplification (MLPA) in acute lymphoblastic leukaemia. Scientific Reports, 2020, 10, 45.	1.6	7
15	Adjuvant tyrosine kinase inhibitor therapy improves outcome for children and adolescents with acute lymphoblastic leukaemia who have an ABLâ€class fusion. British Journal of Haematology, 2020, 191, 844-851.	1.2	31
16	MLPA and DNA index improve the molecular diagnosis of childhood B-cell acute lymphoblastic leukemia. Scientific Reports, 2020, 10, 11501.	1.6	13
17	Design of a Comprehensive Fluorescence in Situ Hybridization Assay for Genetic Classification of T-Cell Acute Lymphoblastic Leukemia. Journal of Molecular Diagnostics, 2020, 22, 629-639.	1.2	18
18	A validated novel continuous prognostic index to deliver stratified medicine in pediatric acute lymphoblastic leukemia. Blood, 2020, 135, 1438-1446.	0.6	25

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19	Genetic characterization and therapeutic targeting of MYCâ€rearranged T cell acute lymphoblastic leukaemia. British Journal of Haematology, 2019, 185, 169-174.	1.2	9
20	IKZF1 Deletions with COBL Breakpoints Are Not Driven by RAG-Mediated Recombination Events in Acute Lymphoblastic Leukemia. Translational Oncology, 2019, 12, 726-732.	1.7	7
21	SH2B3 inactivation through CN-LOH 12q is uniquely associated with B-cell precursor ALL with iAMP21 or other chromosome 21 gain. Leukemia, 2019, 33, 1881-1894.	3.3	26
22	Outcome of Children With Hypodiploid Acute Lymphoblastic Leukemia: A Retrospective Multinational Study. Journal of Clinical Oncology, 2019, 37, 770-779.	0.8	64
23	Validation of the United Kingdom copy-number alteration classifier in 3239 children with B-cell precursor ALL. Blood Advances, 2019, 3, 148-157.	2.5	48
24	Prognostic Impact of Chromosomal Abnormalities and Copy Number Alterations Among Adults with B-Cell Precursor Acute Lymphoblastic Leukaemia Treated on UKALL14. Blood, 2019, 134, 288-288.	0.6	6
25	Genetic and Genomic Characterisation of Older Adults with Acute Lymphoblastic Leukemia Treated on the UKALL14 and UKALL60+ Clinical Trials. Blood, 2019, 134, 2746-2746.	0.6	1
26	Dynamic clonal progression in xenografts of acute lymphoblastic leukemia with intrachromosomal amplification of chromosome 21. Haematologica, 2018, 103, 634-644.	1.7	13
27	Genome-wide association study identifies susceptibility loci for B-cell childhood acute lymphoblastic leukemia. Nature Communications, 2018, 9, 1340.	5.8	58
28	Mutant JAK3 signaling is increased by loss of wild-type JAK3 or by acquisition of secondary JAK3 mutations in T-ALL. Blood, 2018, 131, 421-425.	0.6	30
29	Genotype-Specific Minimal Residual Disease Interpretation Improves Stratification in Pediatric Acute Lymphoblastic Leukemia. Journal of Clinical Oncology, 2018, 36, 34-43.	0.8	147
30	Section 3: Ethics of initiation of long-term ventilation in children at home. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2018, 2, 16-22.	0.2	2
31	Advances in Bâ€cell Precursor Acute Lymphoblastic Leukemia Genomics. HemaSphere, 2018, 2, e53.	1.2	49
32	Use of Minimal Residual Disease Assessment to Redefine Induction Failure in Pediatric Acute Lymphoblastic Leukemia. Journal of Clinical Oncology, 2017, 35, 660-667.	0.8	76
33	Characterisation of the genomic landscape of CRLF2â€rearranged acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2017, 56, 363-372.	1.5	49
34	CREBBP knockdown enhances RAS/RAF/MEK/ERK signaling in Ras pathway mutated acute lymphoblastic leukemia but does not modulate chemotherapeutic response. Haematologica, 2017, 102, 736-745.	1.7	17
35	Digital Multiplex Ligation-Dependent Probe Amplification for Detection of Key Copy Number Alterations in T- and B-Cell Lymphoblastic Leukemia. Journal of Molecular Diagnostics, 2017, 19, 659-672.	1.2	30
36	Unlocking the potential of anti-CD33 therapy in adult and childhood acute myeloid leukemia. Experimental Hematology, 2017, 54, 40-50.	0.2	28

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37	Pediatric home mechanical ventilation: A Canadian Thoracic Society clinical practice guideline executive summary. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2017, 1, 7-36.	0.2	41
38	Intragenic amplification of PAX5: a novel subgroup in B-cell precursor acute lymphoblastic leukemia?. Blood Advances, 2017, 1, 1473-1477.	2.5	25
39	Cytogenetics and Molecular Genetics. , 2017, , 61-98.		2
40	Chronic myeloid leukemia: reminiscences and dreams. Haematologica, 2016, 101, 541-558.	1.7	92
41	Integration of genetic and clinical risk factors improves prognostication in relapsed childhood B-cell precursor acute lymphoblastic leukemia. Blood, 2016, 128, 911-922.	0.6	103
42	Deletions of the long arm of chromosome 5 define subgroups of T-cell acute lymphoblastic leukemia. Haematologica, 2016, 101, 951-958.	1.7	18
43	EBF1-PDGFRB fusion in pediatric B-cell precursor acute lymphoblastic leukemia (BCP-ALL): genetic profile and clinical implications. Blood, 2016, 127, 2214-2218.	0.6	108
44	Constitutional abnormalities of chromosome 21 predispose to iAMP21-acute lymphoblastic leukaemia. European Journal of Medical Genetics, 2016, 59, 162-165.	0.7	31
45	Quantitative proteomic analysis reveals maturation as a mechanism underlying glucocorticoid resistance in B lineage ALL and reâ€sensitization by JNK inhibition. British Journal of Haematology, 2015, 171, 595-605.	1.2	15
46	Blood Spotlight on iAMP21 acute lymphoblastic leukemia (ALL), a high-risk pediatric disease. Blood, 2015, 125, 1383-1386.	0.6	97
47	The 9p21.3 risk of childhood acute lymphoblastic leukaemia is explained by a rare high-impact variant in CDKN2A. Scientific Reports, 2015, 5, 15065.	1.6	24
48	Clinical Impact of Additional Cytogenetic Aberrations, cKIT and RAS Mutations, and Treatment Elements in Pediatric t(8;21)-AML: Results From an International Retrospective Study by the International Berlin-Frankfurt-MÃ¼nster Study Group. Journal of Clinical Oncology, 2015, 33, 4247-4258.	0.8	75
49	ZEB2 drives immature T-cell lymphoblastic leukaemia development via enhanced tumour-initiating potential and IL-7 receptor signalling. Nature Communications, 2015, 6, 5794.	5.8	75
50	Epigenetic landscape correlates with genetic subtype but does not predict outcome in childhood acute lymphoblastic leukemia. Epigenetics, 2015, 10, 717-726.	1.3	26
51	Targeted sequencing identifies associations between IL7R-JAK mutations and epigenetic modulators in T-cell acute lymphoblastic leukemia. Haematologica, 2015, 100, 1301-1310.	1.7	151
52	A novel integrated cytogenetic and genomic classification refines risk stratification in pediatric acute lymphoblastic leukemia. Blood, 2014, 124, 1434-1444.	0.6	178
53	IGH@ translocations co-exist with other primary rearrangements in B-cell precursor acute lymphoblastic leukemia. Haematologica, 2014, 99, 1334-1342.	1.7	20
54	t(6;9)(p22;q34)/DEK-NUP214-rearranged pediatric myeloid leukemia: an international study of 62 patients. Haematologica, 2014, 99, 865-872.	1.7	77

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55	Constitutional and somatic rearrangement of chromosome 21 in acute lymphoblastic leukaemia. Nature, 2014, 508, 98-102.	13.7	261
56	An international study of intrachromosomal amplification of chromosome 21 (iAMP21): cytogenetic characterization and outcome. Leukemia, 2014, 28, 1015-1021.	3.3	175
57	IGH@ Translocations Are Prevalent in Teenagers and Young Adults With Acute Lymphoblastic Leukemia and Are Associated With a Poor Outcome. Journal of Clinical Oncology, 2014, 32, 1453-1462.	0.8	87
58	IKZF1 status as a prognostic feature in BCR-ABL1â€"positive childhood ALL. Blood, 2014, 123, 1691-1698.	0.6	129
59	Ras pathway mutations are prevalent in relapsed childhood acute lymphoblastic leukemia and confer sensitivity to MEK inhibition. Blood, 2014, 124, 3420-3430.	0.6	209
60	Genetic profile of T-cell acute lymphoblastic leukemias with MYC translocations. Blood, 2014, 124, 3577-3582.	0.6	49
61	The B-13 hepatocyte progenitor cell resists pluripotency induction and differentiation to non-hepatocyte cells. Toxicology Research, 2013, 2, 308.	0.9	12
62	Abnormalities of the der(12)t(12;21) in ETV6â€RUNX1 acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2013, 52, 202-213.	1.5	22
63	Chromosomal translocations involving the IGH@ locus in B-cell precursor acute lymphoblastic leukemia: 29 new cases and a review of the literature. Cancer Genetics, 2013, 206, 162-173.	0.2	29
64	Targeting signaling pathways in acute lymphoblastic leukemia: new insights. Hematology American Society of Hematology Education Program, 2013, 2013, 118-125.	0.9	42
65	Risk-Directed Treatment Intensification Significantly Reduces the Risk of Relapse Among Children and Adolescents With Acute Lymphoblastic Leukemia and Intrachromosomal Amplification of Chromosome 21: A Comparison of the MRC ALL97/99 and UKALL2003 Trials. Journal of Clinical Oncology, 2013, 31, 3389-3396.	0.8	111
66	Genes commonly deleted in childhood B-cell precursor acute lymphoblastic leukemia: association with cytogenetics and clinical features. Haematologica, 2013, 98, 1081-1088.	1.7	139
67	Independent prognostic value of BCR-ABL1-like signature and IKZF1 deletion, but not high CRLF2 expression, in children with B-cell precursor ALL. Blood, 2013, 122, 2622-2629.	0.6	248
68	Pediatric acute myeloid leukemia with t(8;16)(p11;p13), a distinct clinical and biological entity: a collaborative study by the International-Berlin-Frankfurt-MÃ¼nster AML-study group. Blood, 2013, 122, 2704-2713.	0.6	86
69	The clinical characteristics, therapy and outcome of 85 adults with acute lymphoblastic leukemia and t(4;11)(q21;q23)/MLL-AFF1 prospectively treated in the UKALLXII/ECOG2993 trial. Haematologica, 2013, 98, 945-952.	1.7	54
70	Outcome in children with Down's syndrome and acute lymphoblastic leukemia: role of IKZF1 deletions and CRLF2 aberrations. Leukemia, 2012, 26, 2204-2211.	3.3	91
71	IGH@ Translocations, CRLF2 Deregulation, and Microdeletions in Adolescents and Adults With Acute Lymphoblastic Leukemia. Journal of Clinical Oncology, 2012, 30, 3100-3108.	0.8	120
72	Burkitt's lymphoma. Lancet, The, 2012, 379, 1234-1244.	6.3	486

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73	Treatment outcome of CRLF2-rearranged childhood acute lymphoblastic leukaemia: a comparative analysis of the AIEOP-BFM and UK NCRI-CCLG study groups. British Journal of Haematology, 2012, 158, 772-777.	1.2	39
74	Genomic Analysis Drives Tailored Therapy in Poor Risk Childhood Leukemia. Cancer Cell, 2012, 22, 139-140.	7.7	4
75	Episomal amplification of NUP214-ABL1 fusion gene in B-cell acute lymphoblastic leukemia. Blood, 2012, 120, 4441-4443.	0.6	21
76	Immunoglobulin Heavy Chain Locus (IGH@) Translocations in Childhood B-Cell Precursor Acute Lymphoblastic Leukemia (BCP-ALL): Incidence and Risk Stratification. Blood, 2012, 120, 1274-1274.	0.6	9
77	Acute Lymphoblastic Leukemia. Clinics in Laboratory Medicine, 2011, 31, 631-647.	0.7	31
78	Demographic, clinical, and outcome features of children with acute lymphoblastic leukemia and CRLF2 deregulation: results from the MRC ALL97 clinical trial. Blood, 2011, 117, 2129-2136.	0.6	133
79	Prognostic significance of additional cytogenetic aberrations in 733 de novo pediatric 11q23/MLL-rearranged AML patients: results of an international study. Blood, 2011, 117, 7102-7111.	0.6	58
80	Genomic characterization implicates iAMP21 as a likely primary genetic event in childhood B-cell precursor acute lymphoblastic leukemia. Blood, 2011, 117, 6848-6855.	0.6	108
81	Key pathways as therapeutic targets. Blood, 2011, 118, 2935-2936.	0.6	8
82	Results of a randomized trial in children with Acute Myeloid Leukaemia: Medical Research Council AML12 trial. British Journal of Haematology, 2011, 155, 366-376.	1.2	167
83	Analysis of a breakpoint cluster reveals insight into the mechanism of intrachromosomal amplification in a lymphoid malignancy. Human Molecular Genetics, 2011, 20, 2591-2602.	1.4	29
84	New genetics and diagnosis of childhood B-cell precursor acute lymphoblastic leukemia. Mental Illness, 2011, 3, 4.	0.8	2
85	Acute Lymphoblastic Leukaemia. Methods in Molecular Biology, 2011, 730, 99-117.	0.4	10
86	Cytogenetics of longâ€term survivors of ETV6â€RUNX1 fusion positive acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2010, 49, 253-259.	1.5	5
87	The t(14;20) is a poor prognostic factor in myeloma but is associated with long-term stable disease in monoclonal gammopathies of undetermined significance. Haematologica, 2010, 95, 1221-1225.	1.7	84
88	Down syndrome acute lymphoblastic leukemia, a highly heterogeneous disease in which aberrant expression of CRLF2 is associated with mutated JAK2: a report from the International BFM Study Group. Blood, 2010, 115, 1006-1017.	0.6	305
89	Acquisition of genome-wide copy number alterations in monozygotic twins with acute lymphoblastic leukemia. Blood, 2010, 115, 3553-3558.	0.6	87
90	Evaluation of multiplex ligationâ€dependent probe amplification as a method for the detection of copy number abnormalities in Bâ€cell precursor acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2010, 49, 1104-1113.	1.5	101

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91	Detection of prognostically relevant genetic abnormalities in childhood Bâ€cell precursor acute lymphoblastic leukaemia: recommendations from the Biology and Diagnosis Committee of the International Berlinâ€FrankfÃ¼rtâ€MÃ¼nster study group. British Journal of Haematology, 2010, 151, 132-142.	1.2	108
92	Long-term follow-up of the United Kingdom medical research council protocols for childhood acute lymphoblastic leukaemia, 1980â€“2001. Leukemia, 2010, 24, 406-418.	3.3	158
93	A new recurrent translocation t(11;14)(q24;q32) involving IGH@ and miR-125b-1 in B-cell progenitor acute lymphoblastic leukemia. Leukemia, 2010, 24, 1362-1364.	3.3	82
94	PHF6 mutations in T-cell acute lymphoblastic leukemia. Nature Genetics, 2010, 42, 338-342.	9.4	282
95	Variation in CDKN2A at 9p21.3 influences childhood acute lymphoblastic leukemia risk. Nature Genetics, 2010, 42, 492-494.	9.4	248
96	Cytogenetics of Childhood Acute Myeloid Leukemia: United Kingdom Medical Research Council Treatment Trials AML 10 and 12. Journal of Clinical Oncology, 2010, 28, 2674-2681.	0.8	256
97	Refinement of cytogenetic classification in acute myeloid leukemia: determination of prognostic significance of rare recurring chromosomal abnormalities among 5876 younger adult patients treated in the United Kingdom Medical Research Council trials. Blood, 2010, 116, 354-365.	0.6	1,661
98	Genetic aberrations in paediatric acute leukaemias and implications for management of patients. Lancet Oncology, The, 2010, 11, 880-889.	5.1	82
99	Prognostic effect of chromosomal abnormalities in childhood B-cell precursor acute lymphoblastic leukaemia: results from the UK Medical Research Council ALL97/99 randomised trial. Lancet Oncology, The, 2010, 11, 429-438.	5.1	338
100	Loss of 1p and rearrangement of MYC are associated with progression of smouldering myeloma to myeloma: sequential analysis of a single case. Haematologica, 2009, 94, 1024-1028.	1.7	47
101	Modeling the molecular consequences of unbalanced translocations in cancer: Lessons from acute lymphoblastic leukemia. Cell Cycle, 2009, 8, 2175-2184.	1.3	13
102	Heterogeneous breakpoints in patients with acute lymphoblastic leukemia and the dic(9;20)(p11Â13;q11) show recurrent involvement of genes at 20q11.21. Haematologica, 2009, 94, 1164-1169.	1.7	43
103	Frequent upregulation of MYC in plasma cell leukemia. Genes Chromosomes and Cancer, 2009, 48, 624-636.	1.5	65
104	Cytogenetics of paediatric and adolescent acute lymphoblastic leukaemia. British Journal of Haematology, 2009, 144, 147-156.	1.2	196
105	Methylation of tumour suppressor gene promoters in the presence and absence of transcriptional silencing in high hyperdiploid acute lymphoblastic leukaemia. British Journal of Haematology, 2009, 144, 838-847.	1.2	27
106	Specific JAK2 mutation (JAK2R683) and multiple gene deletions in Down syndrome acute lymphoblastic leukemia. Blood, 2009, 113, 646-648.	0.6	169
107	Timing of acquisition of deletion 13 in plasma cell dyscrasias is dependent on genetic context. Haematologica, 2009, 94, 1708-1713.	1.7	68
108	Novel prognostic subgroups in childhood 11q23/MLL-rearranged acute myeloid leukemia: results of an international retrospective study. Blood, 2009, 114, 2489-2496.	0.6	383

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109	A comprehensive analysis of the CDKN2A gene in childhood acute lymphoblastic leukemia reveals genomic deletion, copy number neutral loss of heterozygosity, and association with specific cytogenetic subgroups. Blood, 2009, 113, 100-107.	0.6	167
110	Deregulated expression of cytokine receptor gene, CRLF2, is involved in lymphoid transformation in B-cell precursor acute lymphoblastic leukemia. Blood, 2009, 114, 2688-2698.	0.6	445
111	Fluorescence In Situ Hybridization (FISH) as a Tool for the Detection of Significant Chromosomal Abnormalities in Childhood Leukaemia. Methods in Molecular Biology, 2009, 538, 29-55.	0.4	4
112	HLA-DPB1 supertype-associated protection from childhood leukaemia: relationship to leukaemia karyotype and implications for prevention. Cancer Immunology, Immunotherapy, 2008, 57, 53-61.	2.0	12
113	The complex genomic profile of ETV6â€RUNX1 positive acute lymphoblastic leukemia highlights a recurrent deletion of TBL1XR1. Genes Chromosomes and Cancer, 2008, 47, 1118-1125.	1.5	58
114	Cytogenetic and genomic characterization of cell line ARH77. Cancer Genetics and Cytogenetics, 2008, 181, 40-45.	1.0	0
115	Variable breakpoints target PAX5 in patients with dicentric chromosomes: A model for the basis of unbalanced translocations in cancer. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 17050-17054.	3.3	77
116	Mutation of Genes Affecting the RAS Pathway Is Common in Childhood Acute Lymphoblastic Leukemia. Cancer Research, 2008, 68, 6803-6809.	0.4	129
117	t(6;14)(p22;q32): a new recurrent IGH@ translocation involving ID4 in B-cell precursor acute lymphoblastic leukemia (BCP-ALL). Blood, 2008, 111, 387-391.	0.6	59
118	Cytogenetic features of acute lymphoblastic and myeloid leukemias in pediatric patients with Down syndrome: an iBFM-SG study. Blood, 2008, 111, 1575-1583.	0.6	149
119	Re: Faith held by Jehovah's Witnesses does not always forbid blood transfusions. Paediatrics and Child Health, 2008, 13, 341-341.	0.3	0
120	Re: Teenage decision-making in the context of the Jehovah's Witness faith (again). Paediatrics and Child Health, 2008, 13, 332-334.	0.3	1
121	Karyotype is an independent prognostic factor in adult acute lymphoblastic leukemia (ALL): analysis of cytogenetic data from patients treated on the Medical Research Council (MRC) UKALLXII/Eastern Cooperative Oncology Group (ECOG) 2993 trial. Blood, 2007, 109, 3189-3197.	0.6	655
122	Monosomy 7 and deletion 7q in children and adolescents with acute myeloid leukemia: an international retrospective study. Blood, 2007, 109, 4641-4647.	0.6	126
123	Outcome of treatment in children with hypodiploid acute lymphoblastic leukemia. Blood, 2007, 110, 1112-1115.	0.6	250
124	Prognosis of children with acute lymphoblastic leukemia (ALL) and intrachromosomal amplification of chromosome 21 (iAMP21). Blood, 2007, 109, 2327-2330.	0.6	200
125	Five members of the CEBP transcription factor family are targeted by recurrent IGH translocations in B-cell precursor acute lymphoblastic leukemia (BCP-ALL). Blood, 2007, 109, 3451-3461.	0.6	188
126	Advances in Molecular Cytogenetics to Study the Leukemia Genome. Laboratory Medicine, 2007, 38, 527-535.	0.8	1

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127	â€œOnly flesh with its soul â€“ its blood â€“ you must not eatâ€•(Genesis 9.3:4). Paediatrics and Child Health, 2007, 12, 867-868.	0.3	4
128	Primum non nocere is only the beginning. Paediatrics and Child Health, 2007, 12, 379-380.	0.3	4
129	Intrachromosomal amplification of chromosome 21 (iAMP21) may arise from a breakageâ€“fusionâ€“bridge cycle. Genes Chromosomes and Cancer, 2007, 46, 318-326.	1.5	64
130	Molecular cytogenetic characterization ofTCF3 (E2A)/19p13.3 rearrangements in B-cell precursor acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2007, 46, 478-486.	1.5	67
131	Genome complexity in acute lymphoblastic leukemia is revealed by array-based comparative genomic hybridization. Oncogene, 2007, 26, 4306-4318.	2.6	95
132	A multicenter evaluation of comprehensive analysis of MLL translocations and fusion gene partners in acute leukemia using the MLL FusionChip device. Cancer Genetics and Cytogenetics, 2007, 173, 17-22.	1.0	9
133	Overexpression of CEBPA resulting from the translocation t(14;19)(q32;q13) of human precursor B acute lymphoblastic leukemia. Blood, 2006, 108, 3560-3563.	0.6	67
134	Deletion of chromosome 13 detected by conventional cytogenetics is a critical prognostic factor in myeloma. Leukemia, 2006, 20, 1610-1617.	3.3	141
135	A diminutive chromosome 21 centromere in acute lymphoblastic leukemia. Cancer Genetics and Cytogenetics, 2006, 167, 78-81.	1.0	2
136	Complex genomic alterations and gene expression in acute lymphoblastic leukemia with intrachromosomal amplification of chromosome 21. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 8167-8172.	3.3	146
137	Del (9q) AML: clinical and cytological characteristics and prognostic implications. British Journal of Haematology, 2005, 129, 210-220.	1.2	44
138	Interphase molecular cytogenetic screening for chromosomal abnormalities of prognostic significance in childhood acute lymphoblastic leukaemia: a UK Cancer Cytogenetics Group Study. British Journal of Haematology, 2005, 129, 520-530.	1.2	137
139	Age has a profound effect on the incidence and significance of chromosome abnormalities in myeloma. Leukemia, 2005, 19, 1634-1642.	3.3	73
140	Treatment strategy and long-term results in paediatric patients treated in consecutive UK AML trials. Leukemia, 2005, 19, 2130-2138.	3.3	277
141	ETV6/RUNX1 fusion at diagnosis and relapse: Some prognostic indications. Genes Chromosomes and Cancer, 2005, 43, 54-71.	1.5	17
142	Detection of genomic aberrations in older patients with acute myeloid leukemia. Haematologica, 2005, 90, 147.	1.7	2
143	A Fluorescence in Situ Hybridization Map of 6q Deletions in Acute Lymphocytic Leukemia. Cancer Research, 2004, 64, 4089-4098.	0.4	49
144	Three distinct subgroups of hypodiploidy in acute lymphoblastic leukaemia. British Journal of Haematology, 2004, 125, 552-559.	1.2	184

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145	Complex chromosomal abnormalities in utero , 5â€ƒyears before leukaemia*. British Journal of Haematology, 2004, 126, 307-312.	1.2	18
146	Secondary cytogenetic aberrations in childhood Philadelphia chromosome positive acute lymphoblastic leukemia are nonrandom and may be associated with outcome. Leukemia, 2004, 18, 693-702.	3.3	81
147	Is trisomy 5 a distinct cytogenetic subgroup in acute lymphoblastic leukemia?. Cancer Genetics and Cytogenetics, 2004, 148, 159-162.	1.0	21
148	Successful treatment without cranial radiotherapy of children receiving intensified chemotherapy for acute lymphoblastic leukaemia: results of the risk-stratified randomized central nervous system treatment trial MRC UKALL XI (ISRC TN 16757172). British Journal of Haematology, 2004, 124, 33-46.	1.2	71
149	t(14;19)(q32;q13): A recurrent translocation in B-cell precursor acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2004, 39, 88-92.	1.5	25
150	MLL translocations with concurrent 3? deletions: Interpretation of FISH results. Genes Chromosomes and Cancer, 2004, 41, 266-271.	1.5	23
151	Comparative expressed sequence hybridization studies of high-hyperdiploid childhood acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2004, 41, 191-202.	1.5	28
152	Cytogenetics of multiple myeloma: interpretation of fluorescence in situ hybridization results. British Journal of Haematology, 2003, 120, 944-952.	1.2	32
153	Amplification of AML1 on a duplicated chromosome 21 in acute lymphoblastic leukemia: a study of 20 cases. Leukemia, 2003, 17, 547-553.	3.3	153
154	Clinical heterogeneity in childhood acute lymphoblastic leukemia with 11q23 rearrangements. Leukemia, 2003, 17, 700-706.	3.3	216
155	Prenatal origin of hyperdiploid acute lymphoblastic leukemia in identical twins. Leukemia, 2003, 17, 2202-2206.	3.3	84
156	Amplification of AML1 in acute lymphoblastic leukemia is associated with a poor outcome. Leukemia, 2003, 17, 2249-2250.	3.3	103
157	Outcome heterogeneity in childhood high-hyperdiploid acute lymphoblastic leukemia. Blood, 2003, 102, 2756-2762.	0.6	165
158	Clinical features, cytogenetics and outcome in acute lymphoblastic and myeloid leukaemia of infancy: report from the MRC Childhood Leukaemia working party. Leukemia, 2002, 16, 776-784.	3.3	94
159	Involvement of the MLL gene in T-lineage acute lymphoblastic leukemia. Blood, 2002, 100, 2273-2273.	0.6	16
160	Cytogenetics and molecular genetics of acute lymphoblastic leukemia. Reviews in Clinical and Experimental Hematology, 2002, 6, 91-113.	0.1	93
161	Translocations of 14q32 and deletions of 13q14 are common chromosomal abnormalities in systemic amyloidosis. British Journal of Haematology, 2002, 117, 427-435.	1.2	65
162	Treatment of infants with lymphoblastic leukaemia: results of the UK Infant Protocols 1987-1999. British Journal of Haematology, 2002, 117, 306-314.	1.2	62

#	Article	IF	Citations
163	Acute lymphoblastic leukaemia. Best Practice and Research in Clinical Haematology, 2001, 14, 593-607.	0.7	30
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