Andrew J Carroll

List of Publications by Year
in descending order

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		94433	32842
162	10,584	37	100
papers	citations	h-index	g-index

163	163	163	11105
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical and molecular relevance of genetic variants in the non-coding transcriptome of patients with cytogenetically normal acute myeloid leukemia. Haematologica, 2022, 107, 1034-1044.	3.5	4
2	Outcomes in adolescent and young adult patients (16 to 30 years) compared to younger patients treated for high-risk B-lymphoblastic leukemia: report from Childrenâ€™s Oncology Group Study AALL0232. Leukemia, 2022, 36, 648-655.	7.2	14
3	Molecular, clinical, and prognostic implications of PTPN11 mutations in acute myeloid leukemia. Blood Advances, 2022, 6, 1371-1380.	5.2	16
4	Sexâ€based disparities in outcome in pediatric acute lymphoblastic leukemia: a Childrenâ€™s Oncology Group report. Cancer, 2022, 128, 1863-1870.	4.1	12
5	Outstanding outcomes in infants with <i>KMT2A</i>-germline acute lymphoblastic leukemia treated with chemotherapy alone: results of the Childrenâ€™s Oncology Group AALL0631 trial. Haematologica, 2022, 107, 1205-1208.	3.5	11
6	Poor Survival and Differential Impact of Genetic Features of Black Patients with Acute Myeloid Leukemia. Cancer Discovery, 2021, 11, 626-637.	9.4	41
7	FLT3 inhibitor lestaurtinib plus chemotherapy for newly diagnosed KMT2A-rearranged infant acute lymphoblastic leukemia: Childrenâ€™s Oncology Group trial AALL0631. Leukemia, 2021, 35, 1279-1290.	7.2	46
8	Prognostic impact of minimal residual disease at the end of consolidation in NCI standardâ€risk Bâ€lymphoblastic leukemia: A report from the Children's Oncology Group. Pediatric Blood and Cancer, 2021, 68, e28929.	1.5	9
9	Gene expression signature predicts relapse in adult patients with cytogenetically normal acute myeloid leukemia. Blood Advances, 2021, 5, 1474-1482.	5.2	20
10	Establishment and genomic characterization of a sporadic malignant peripheral nerve sheath tumor cell line. Scientific Reports, 2021, 11, 5690.	3.3	9
11	Excellent Outcomes With Reduced Frequency of Vincristine and Dexamethasone Pulses in Standard-Risk B-Lymphoblastic Leukemia: Results From Children's Oncology Group AALL0932. Journal of Clinical Oncology, 2021, 39, 1437-1447.	1.6	56
12	Favorable Trisomies and ETV6-RUNX1 Predict Cure in Low-Risk B-Cell Acute Lymphoblastic Leukemia: Results From Children's Oncology Group Trial AALL0331. Journal of Clinical Oncology, 2021, 39, 1540-1552.	1.6	19
13	Phase 3 randomized trial of chemotherapy with or without oblimersen in older AML patients: CALGB 10201 (Alliance). Blood Advances, 2021, 5, 2775-2787.	5.2	15
14	VpreB Surrogate Light Chain Expression in B-Lineage ALL: A Report from the Children's Oncology Group. Blood Advances, 2021, , .	5.2	1
15	HUGO Gene Nomenclature Committee (HGNC) recommendations for the designation of gene fusions. Leukemia, 2021, 35, 3040-3043.	7.2	42
16	Epigenetic Phenocopying Expands Molecular Risk Assessment in Acute Myeloid Leukemia (Alliance). Blood, 2021, 138, 803-803.	1.4	0
17	High Early Death Rates, Treatment Resistance and Short Survival of Black Adolescent and Young Adults (AYAs) with Acute Myeloid Leukemia (AML) (Alliance). Blood, 2021, 138, 221-221.	1.4	2
18	Multi-Dimensional Analysis of Adult Acute Myeloid Leukemia (AML) Landscape Cross-Continents Reveals Age Associated Trends in Mutations and Outcomes. Blood, 2021, 138, 685-685.	1.4	0

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19	Comparative Outcomes and Molecular Response Predictors of IDH1/2-Mutated Adult Acute Myeloid Leukemia (AML) Patients (Pts) after Frontline Treatment with Intensive Induction Chemotherapy (IC), Targeted Inhibitors, or Hypomethylating Agents (HMA) (Alliance). Blood, 2021, 138, 226-226.	1.4	0
20	White Blood Cell Count (WBC) Levels Are Associated with Molecular Profiles and Are Independent Outcome Predictors in Acute Myeloid Leukemia (AML) Patients (Pts) (Alliance). Blood, 2021, 138, 3369-3369.	1.4	0
21	Mutations associated with a 17-gene leukemia stem cell score and the scoreâ€™s prognostic relevance in the context of the European LeukemiaNet classification of acute myeloid leukemia. Haematologica, 2020, 105, 721-729.	3.5	21
22	Mixedâ€phenotype acute leukemia: A cohort and consensus research strategy from the Childrenâ€™s Oncology Group Acute Leukemia of Ambiguous Lineage Task Force. Cancer, 2020, 126, 593-601.	4.1	32
23	Randomized assessment of delayed intensification and two methods for parenteral methotrexate delivery in childhood B-ALL: Childrenâ€™s Oncology Group Studies P9904 and P9905. Leukemia, 2020, 34, 1006-1016.	7.2	8
24	Outcome in Children With Standard-Risk B-Cell Acute Lymphoblastic Leukemia: Results of Childrenâ€™s Oncology Group Trial AALL0331. Journal of Clinical Oncology, 2020, 38, 602-612.	1.6	107
25	Mutational landscape and clinical outcome of patients with de novo acute myeloid leukemia and rearrangements involving 11q23/ KMT2A. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 26340-26346.	7.1	59
26	Childrenâ€™s Oncology Group AALL0434: A Phase III Randomized Clinical Trial Testing Nelarabine in Newly Diagnosed T-Cell Acute Lymphoblastic Leukemia. Journal of Clinical Oncology, 2020, 38, 3282-3293.	1.6	136
27	Impact of Intrathecal Triple Therapy Versus Intrathecal Methotrexate on Disease-Free Survival for High-Risk B-Lymphoblastic Leukemia: Childrenâ€™s Oncology Group Study AALL1131. Journal of Clinical Oncology, 2020, 38, 2628-2638.	1.6	41
28	Clinical and functional significance of circular RNAs in cytogenetically normal AML. Blood Advances, 2020, 4, 239-251.	5.2	29
29	Outcomes of Patients with CRLF2-Overexpressing Acute Lymphoblastic Leukemia without Down Syndrome: A Report from the Children's Oncology Group. Blood, 2020, 136, 45-46.	1.4	6
30	Clinical and Prognostic Implications of PTPN11 Mutations in Acute Myeloid Leukemia (Alliance). Blood, 2020, 136, 20-21.	1.4	2
31	Cytogenetic Subgroups Drive Risk Stratification and Response to Chemotherapy and Blinatumomab in Children and Young Adults with Relapsed B-ALL: A Children's Oncology Group Study. Blood, 2020, 136, 16-17.	1.4	1
32	Poor Treatment Outcomes of Young (<60 Years) African American Patients (Pts) Diagnosed with Acute Myeloid Leukemia (AML) (Alliance). Blood, 2020, 136, 5-7.	1.4	4
33	Outcomes of Patients with Down Syndrome and CRLF2-Overexpressing Acute Lymphoblastic Leukemia (ALL): A Report from the Children's Oncology Group (COG). Blood, 2020, 136, 44-45.	1.4	1
34	Meta-Analysis of Genome-Wide Association Studies of Acute Myeloid Leukemia (AML) Patients Identifies Variants Associated with Risk of 11q23/KMT2A-Translocated and Core-Binding Factor (CBF) AML and Suggests a Role for Transcription Elongation in Leukemogenesis. Blood, 2020, 136, 29-30.	1.4	0
35	Enhanced Risk Stratification of 21,178 Children, Adolescents, and Young Adults with Acute Lymphoblastic Leukemia (ALL) Incorporating White Blood Count (WBC), Age, and Minimal Residual Disease (MRD) at Day 8 and 29 As Continuous Variables: A Children's Oncology Group (COG) Report. Blood, 2020, 136, 39-40.	1.4	2
36	Differential Impact of Prognostically Significant Gene Mutations in Acute Myeloid Leukemia (AML) Patients (Pts) Older Than 70 Years (y) Treated with Cytarabine-Based Induction Therapy. Blood, 2020, 136, 40-41.	1.4	0

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37	Genetic Characterization and Prognostic Relevance of Acquired Uniparental Disomies in Cytogenetically Normal Acute Myeloid Leukemia. Clinical Cancer Research, 2019, 25, 6524-6531.	7.0	12
38	Masked hypodiploidy: Hypodiploid acute lymphoblastic leukemia (ALL) mimicking hyperdiploid ALL in children: A report from the Children's Oncology Group. Cancer Genetics, 2019, 238, 62-68.	0.4	32
39	Inherited genetic susceptibility to acute lymphoblastic leukemia in Down syndrome. Blood, 2019, 134, 1227-1237.	1.4	37
40	Response to interferon treatment in essential thrombocythemia with inv(3)(q21q26). Annals of Hematology, 2019, 98, 2845-2846.	1.8	0
41	Prognostic and Biologic Relevance of Clinically Applicable Long Noncoding RNA Profiling in Older Patients with Cytogenetically Normal Acute Myeloid Leukemia. Molecular Cancer Therapeutics, 2019, 18, 1451-1459.	4.1	7
42	Impact of corticosteroid pretreatment in pediatric patients with newly diagnosed B-lymphoblastic leukemia: a report from the Childrenâ€™s Oncology Group. Haematologica, 2019, 104, e517-e520.	3.5	11
43	Relapse after Prolonged Remission in Philadelphia-Like Acute Lymphoblastic Leukemia. Case Reports in Hematology, 2019, 2019, 1-3.	0.4	1
44	Replacing cyclophosphamide/cytarabine/mercaptopurine with cyclophosphamide/etoposide during consolidation/delayed intensification does not improve outcome for pediatric B-cell acute lymphoblastic leukemia: a report from the COG. Haematologica, 2019, 104, 986-992.	3.5	25
45	Targeting EIF4E signaling with ribavirin in infant acute lymphoblastic leukemia. Oncogene, 2019, 38, 2241-2262.	5.9	29
46	Genome-wide association study identifies an acute myeloid leukemia susceptibility locus near BICRA. Leukemia, 2019, 33, 771-775.	7.2	15
47	Excellent Outcomes with Reduced Frequency of Vincristine and Dexamethasone Pulses in Children with National Cancer Institute (NCI) Standard-Risk B Acute Lymphoblastic Leukemia (SR B-ALL): A Report from Children's Oncology Group (COG) Study AALL0932. Blood, 2019, 134, 824-824.	1.4	6
48	The 2017 European Leukemianet Genetic Risk Classification Performs Poorly in Older Patients with Acute Myeloid Leukemia (AML) and Should be Refined to Identify Patients Requiring Additional or Alternative Treatment. Blood, 2019, 134, 2681-2681.	1.4	1
49	FLT3 Inhibitor Correlative Laboratory Assays Impact Outcomes in KMT2A-Rearranged Infant Acute Lymphoblastic Leukemia (ALL) Patients Treated with Lestaurtinib: AALL0631, a Children's Oncology Group Study. Blood, 2019, 134, 1293-1293.	1.4	4
50	Next-Generation RNA Sequencing-Based Analysis Identifies a Novel Set of Prognostic Micrornas (miRs) in Cytogenetically Normal Acute Myeloid Leukemia (CN-AML). Blood, 2019, 134, 2694-2694.	1.4	0
51	Distinct Gene Expression Profiles and Mutations Associate with Outcome in Younger Adults with De Novo Cytogenetically Normal Acute Myeloid Leukemia (CN-AML) (Alliance). Blood, 2019, 134, 1247-1247.	1.4	1
52	Personalized Oncology in Acute Myeloid Leukemia (AML): Validation of the Prognostic Value of the Knowledge Bank Algorithm in Patients (Pts) Treated on Cancer and Leukemia Group B (CALGB)/Alliance Protocols. Blood, 2019, 134, 182-182.	1.4	0
53	Outcome in Adolescent and Young Adult (AYA) Patients Compared to Younger Patients Treated for High-Risk B-Lymphoblastic Leukemia (HR B-ALL): Report from the Children's Oncology Group Study AALL0232. Blood, 2019, 134, 286-286.	1.4	0
54	The Genomic Landscape of Childhood Acute Lymphoblastic Leukemia. Blood, 2019, 134, 649-649.	1.4	5

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55	Mutation patterns identify adult patients with de novo acute myeloid leukemia aged 60 years or older who respond favorably to standard chemotherapy: an analysis of Alliance studies. Leukemia, 2018, 32, 1338-1348.	7.2	80
56	Toxicity associated with intensive postinduction therapy incorporating clofarabine in the very highâ€risk stratum of patients with newly diagnosed highâ€risk Bâ€lymphoblastic leukemia: A report from the Children's Oncology Group study AALL1131. Cancer, 2018, 124, 1150-1159.	4.1	46
57	Ten-year outcome of patients with acute myeloid leukemia not treated with allogeneic transplantation in first complete remission. Blood Advances, 2018, 2, 1645-1650.	5.2	85
58	Genomic and outcome analyses of Ph-like ALL in NCI standard-risk patients: a report from the Childrenâ€™s Oncology Group. Blood, 2018, 132, 815-824.	1.4	97
59	NF1 mutations are recurrent in adult acute myeloid leukemia and confer poor outcome. Leukemia, 2018, 32, 2536-2545.	7.2	33
60	Triple Intrathecal Therapy (Methotrexate/Hydrocortisone/Cytarabine) Does Not Improve Disease-Free Survival Versus Intrathecal Methotrexate Alone in Children with High Risk B-Lymphoblastic Leukemia: Results of Children's Oncology Group Study AALL1131. Blood, 2018, 132, 35-35.	1.4	7
61	Additional Gene Mutations Refine the 2017 European Leukemianet (ELN) Classification of Adult Patients (Pts) with De Novo Acute Myeloid Leukemia (AML) Aged <60 Years: An Analysis of Alliance for Clinical Trials in Oncology (Alliance) Studies. Blood, 2018, 132, 2740-2740.	1.4	1
62	Mutations in Genes Associated with Familial Predisposition to Myeloid Neoplasms: Their Frequency and Associations with Pretreatment Characteristics in Adult Patients (Pts) with Presumably Sporadic De Novo Acute Myeloid Leukemia (AML). Blood, 2018, 132, 1478-1478.	1.4	13
63	Uniparental Disomies (UPD) of Chromosome 13q Is Associated with Shorter Disease-Free Survival in Adult Patients (Pts) with De Novo Cytogenetically Normal Acute Myeloid Leukemia (CN-AML). Blood, 2018, 132, 2777-2777.	1.4	0
64	Prognostic and Biologic Significance of Long Non-Coding RNA (lncRNA) Profiling in Cytogenetically Abnormal Acute Myeloid Leukemia (CA-AML). Blood, 2018, 132, 2767-2767.	1.4	0
65	Genome-Wide Association Study (GWAS) Identifies a Significant Acute Myeloid Leukemia (AML) Susceptibility Locus Near BICRA. Blood, 2018, 132, 85-85.	1.4	0
66	Targetable kinase gene fusions in high-risk B-ALL: a study from the Childrenâ€™s Oncology Group. Blood, 2017, 129, 3352-3361.	1.4	236
67	Prognostic and biologic significance of long non-coding RNA profiling in younger adults with cytogenetically normal acute myeloid leukemia. Haematologica, 2017, 102, 1391-1400.	3.5	28
68	The genomic landscape of pediatric and young adult T-lineage acute lymphoblastic leukemia. Nature Genetics, 2017, 49, 1211-1218.	21.4	693
69	Klinefelter syndrome and 47,<scp>XYY</scp> syndrome in children with B cell acute lymphoblastic leukaemia. British Journal of Haematology, 2017, 179, 843-846.	2.5	4
70	Clinical relevance of small copy-number variants in chromosomal microarray clinical testing. Genetics in Medicine, 2017, 19, 377-385.	2.4	24
71	Outcome of Children with Standardâ€Risk Tâ€Lineage Acute Lymphoblastic Leukemiaâ€”Comparison among Different Treatment Strategies. Pediatric Blood and Cancer, 2016, 63, 255-261.	1.5	17
72	Overview of Clinical Cytogenetics. Current Protocols in Human Genetics, 2016, 89, 8.1.1-8.1.13.	3.5	7

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73	Family history of hematologic malignancies and risk of multiple myeloma: differences by race and clinical features. Cancer Causes and Control, 2016, 27, 81-91.	1.8	35
74	Outcomes of Children, Adolescents, and Young Adults with Acute Lymphoblastic Leukemia Based on Blast Genotype at Diagnosis: A Report from the Children's Oncology Group. Blood, 2016, 128, 451-451.	1.4	4
75	Minimal Residual Disease Assessment of Remission after Induction Therapy Is Superior to Morphologic Assessment for Risk Stratification in Childhood Acute Lymphoblastic Leukemia: A Report from the Children's Oncology Group (COG). Blood, 2016, 128, 758-758.	1.4	1
76	Integrated Genomic Analysis of Down Syndrome Acute Lymphoblastic Leukemia Reveals Recurrent Cancer Gene Alterations and Evidence of Frequent Subclonal Driver Events. Blood, 2016, 128, 4083-4083.	1.4	0
77	New Insights into Deregulated Gene Expression Pathways in MLL- and AF10-Rearranged T-Lineage Acute Lymphoblastic Leukemia. Blood, 2016, 128, 2906-2906.	1.4	0
78	Improved Diagnosis of Intrachromosomal Amplification of Chromosome 21 (iAMP21) By Copy Number Profiling. Blood, 2016, 128, 1733-1733.	1.4	0
79	The Mutational Patterns Associated with Cytogenetic Subsets of De Novo Acute Myeloid Leukemia (AML): A Study of 1603 Adult Patients (Pts). Blood, 2016, 128, 287-287.	1.4	0
80	CCND1 and CCND2 Mutations Are Frequent in Adults with Core-Binding Factor Acute Myeloid Leukemia (CBF-AML) with t(8;21)(q22;q22). Blood, 2016, 128, 2740-2740.	1.4	0
81	Whole Exome Sequencing of Pediatric Acute Lymphoblastic Leukemia Patients Identify Mutations in 11 Pathways: A Report from the Children's Oncology Group. Blood, 2016, 128, 455-455.	1.4	1
82	A genome-wide association study of susceptibility to acute lymphoblastic leukemia in adolescents and young adults. Blood, 2015, 125, 680-686.	1.4	110
83	Prognostic significance of minimal residual disease in high risk B-ALL: a report from Childrenâ€™s Oncology Group study AALL0232. Blood, 2015, 126, 964-971.	1.4	287
84	Integration of cytogenomic data for furthering the characterization of pediatric B-cell acute lymphoblastic leukemia: a multi-institution, multi-platform microarray study. Cancer Genetics, 2015, 208, 1-18.	0.4	30
85	Mixed Lineage Leukemia Rearrangements (MLL-R) Are Determinants of High Risk Disease in Homeobox A (HOXA)-deregulated T-Lineage Acute Lymphoblastic Leukemia: A Children's Oncology Group Study. Blood, 2015, 126, 694-694.	1.4	2
86	Capizzi-Style Methotrexate with Pegasparagase (C-MTX) Is Superior to High-Dose Methotrexate (HDMTX) in T-Lineage Acute Lymphoblastic Leukemia (T-ALL): Results from Children's Oncology Group (COG) AALL0434. Blood, 2015, 126, 794-794.	1.4	12
87	Genetic and Response-Based Risk Classification Identifies a Subgroup of NCI High Risk Childhood B-Lymphoblastic Leukemia (HR B-ALL) with Outstanding Outcomes: A Report from the Children's Oncology Group (COG). Blood, 2015, 126, 807-807.	1.4	5
88	Expression and prognostic impact of lncRNAs in acute myeloid leukemia. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 18679-18684.	7.1	214
89	The recurrent distal 22q11.2 microdeletions are often de novo and do not represent a single clinical entity: a proposed categorization system. Genetics in Medicine, 2014, 16, 92-100.	2.4	49
90	Constitutional and somatic rearrangement of chromosome 21 in acute lymphoblastic leukaemia. Nature, 2014, 508, 98-102.	27.8	261

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91	Epigenetics Meets Genetics in Acute Myeloid Leukemia: Clinical Impact of a Novel Seven-Gene Score. Journal of Clinical Oncology, 2014, 32, 548-556.	1.6	134
92	Prognostic gene mutations and distinct gene- and microRNA-expression signatures in acute myeloid leukemia with a sole trisomy 8. Leukemia, 2014, 28, 1754-1758.	7.2	24
93	Targetable Kinase-Activating Lesions in Ph-like Acute Lymphoblastic Leukemia. New England Journal of Medicine, 2014, 371, 1005-1015.	27.0	1,161
94	Clinical Role of microRNAs in Cytogenetically Normal Acute Myeloid Leukemia: miR-155 Upregulation Independently Identifies High-Risk Patients. Journal of Clinical Oncology, 2013, 31, 2086-2093.	1.6	165
95	Genomic Characterization and Experimental Modeling Of BCR-ABL1-Like Acute Lymphoblastic Leukemia. Blood, 2013, 122, 232-232.	1.4	8
96	Genome-Wide DNA Methylation Analysis Reveals Biological and Clinical Insights In Relapsed Childhood Acute Lymphoblastic Leukemia: A Report From The COG ALL Target Project. Blood, 2013, 122, 3736-3736.	1.4	1
97	Integrated Genomic and Mutational Profiling Of Adolescent and Young Adult ALL Identifies a High Frequency Of BCR-ABL1-Like ALL with Very Poor Outcome. Blood, 2013, 122, 825-825.	1.4	8
98	Development and Validation Of a Highly Sensitive and Specific Gene Expression Classifier To Prospectively Screen and Identify B-Precursor Acute Lymphoblastic Leukemia (ALL) Patients With a Philadelphia Chromosome-Like (â€œPh-likeâ€•or â€œBCR-ABL1-Likeâ€) Signature For Therapeutic Targeting and Clinical Intervention. Blood, 2013, 122, 826-826.	1.4	65
99	Excellent Event Free (EFS) and Overall Survival (OS) For Children With Standard Risk Acute Lymphoblastic Leukemia (SR ALL) Despite The Absence Of a Significant Impact On Outcome With The Addition Of An Intensified Consolidation: Results Of Childrenâ€™s Oncology Group (COG) AALL0331. Blood, 2013, 122, 837-837.	1.4	13
100	Genomic- and Transcriptomic Profiling Of Acute Lymphoblastic Leukemia With Dicentric Chromosomes. Blood, 2013, 122, 234-234.	1.4	1
101	Differential Clinical Impact Of Gene Mutations and Their Combinations In Primary Cytogenetically Normal Acute Myeloid Leukemia (CN-AML). Blood, 2013, 122, 2540-2540.	1.4	0
102	PI3K/AKT/mTOR Signaling Is a Significant Druggable Pathway In Infant Acute Lymphoblastic Leukemia (ALL). Blood, 2013, 122, 1669-1669.	1.4	7
103	Cytogenetics and Outcome Of Infants With Acute Lymphoblastic Leukemia and Absence Of MLL Rearrangements. Blood, 2013, 122, 1349-1349.	1.4	0
104	Leukemic Blasts With The PNH Phenotype: Correlation With Cytogenetics In ALL. Blood, 2013, 122, 2628-2628.	1.4	0
105	RUNX1 Mutations Are Associated With Poor Outcome in Younger and Older Patients With Cytogenetically Normal Acute Myeloid Leukemia and With Distinct Gene and MicroRNA Expression Signatures. Journal of Clinical Oncology, 2012, 30, 3109-3118.	1.6	242
106	Continuous Dose Dasatinib Is Safe and Feasible in Combination with Intensive Chemotherapy in Pediatric Philadelphia Chromosome Positive Acute Lymphoblastic Leukemia (Ph+ ALL): Children's Oncology Group (COG) Trial AALL0622. Blood, 2012, 120, 137-137.	1.4	7
107	The Clinical Role of Micrornas (miRs) in Cytogenetically Normal (CN) Acute Myeloid Leukemia (AML): miR-155 Upregulation Independently Identifies High-Risk Patients (Pts). Blood, 2012, 120, 1387-1387.	1.4	1
108	Intrachromosomal Amplification of Chromosome 21(iAMP21): Cytogenetic Characterisation and Outcome in Childhood B-Cell Precursor Acute Lymphoblastic Leukaemia (BCP-ALL). A Study On Behalf of the Ponte Di Legno International Childhood ALL Workshop. Blood, 2012, 120, 293-293.	1.4	0

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109	Expression Profiling for MEIS1 and HOXA9/10 Identifies an Increased Incidence of MLL Rearrangements in T-ALL: A Children's Oncology Group Study.. Blood, 2012, 120, 2505-2505.	1.4	0
110	ASXL1 mutations identify a high-risk subgroup of older patients with primary cytogenetically normal AML within the ELN Favorable genetic category. Blood, 2011, 118, 6920-6929.	1.4	246
111	Improved Post-Induction Chemotherapy Does Not Abrogate Prognostic Significance of Minimal Residual Disease (MRD) for Children and Young Adults with High Risk Acute Lymphoblastic Leukemia (ALL). A Report From Children's Oncology Group (COG) Study AALL0232. Blood, 2011, 118, 1440-1440.	1.4	3
112	Predicting Clinical Dose-Exposure and Exposure-Response Relationships of Pan-Antiapoptotic BCL-2 Family Inhibitor Obatoclax in MLL Rearranged Infant Leukemias From Preclinical Disease Models and Adult Experience. Blood, 2011, 118, 2580-2580.	1.4	2
113	Prognostic Utility of the European LeukemiaNet (ELN) Genetic-Risk Classification in Adults with De Novo Acute Myeloid Leukemia (AML): A Study of 1,550 Patients (Pts). Blood, 2011, 118, 414-414.	1.4	2
114	A BCR-ABL1-Like Gene Expression Profile Confers a Poor Prognosis In Patients with High-Risk Acute Lymphoblastic Leukemia (HR-ALL): A Report From Children's Oncology Group (COG) AALL0232. Blood, 2011, 118, 743-743.	1.4	3
115	Poor Outcome of RUNX1-Mutated (RUNX1-mut) Patients (Pts) with Primary, Cytogenetically Normal Acute Myeloid Leukemia (CN-AML) and Associated Gene- and MicroRNA (miR) Expression Signatures,. Blood, 2011, 118, 3454-3454.	1.4	0
116	Cytogenetic, Molecular and Clinical Features Associated with Rare CBFB-MYH11 Fusion Transcripts in Patients (Pts) with Acute Myeloid Leukemia (AML) and inv(16)/t(16;16). Blood, 2011, 118, 2514-2514.	1.4	0
117	MLL Rearrangement and Age At Diagnosis Are Strongly Associated with High Level Surface FLT3 Expression and Ex Vivo Sensitivity to FLT3 Inhibition: A Prospective Analysis of 54 Consecutive Infants with ALL Enrolled in Children's Oncology Group (COG) Trial AALL0631. Blood, 2011, 118, 568-568.	1.4	0
118	iAMP21 Is Associated with Inferior Outcomes in Children with Acute Lymphoblastic Leukemia (ALL) on Contemporary Children's Oncology Group (COG) Studies. Blood, 2011, 118, 739-739.	1.4	2
119	ASXL1 Mutations Identify a High-Risk Subgroup of Older Patients with Primary Cytogenetically Normal Acute Myeloid Leukemia within the European LeukemiaNet â€˜Favorable' Genetic Category. Blood, 2011, 118, 417-417.	1.4	0
120	FLT3 internal tandem duplication associates with adverse outcome and gene- and microRNA-expression signatures in patients 60 years of age or older with primary cytogenetically normal acute myeloid leukemia: a Cancer and Leukemia Group B study. Blood, 2010, 116, 3622-3626.	1.4	201
121	Prognostic Significance of Expression of a Single MicroRNA, miR-181a, in Cytogenetically Normal Acute Myeloid Leukemia: A Cancer and Leukemia Group B Study. Journal of Clinical Oncology, 2010, 28, 5257-5264.	1.6	176
122	IDH1 and IDH2 Gene Mutations Identify Novel Molecular Subsets Within De Novo Cytogenetically Normal Acute Myeloid Leukemia: A Cancer and Leukemia Group B Study. Journal of Clinical Oncology, 2010, 28, 2348-2355.	1.6	699
123	Infant Acute Lymphoblastic Leukemias Are Pan-Sensitive to Obatoclax Across molecular/Cytogenetic Subtypes, Especially MLL-ENL, and gene Expression Profiles Determine Obatoclax IC50: A Report on the Children's Oncology Group (COG) P9407 Trial. Blood, 2010, 116, 2757-2757.	1.4	1
124	Genome-Wide Analysis of Genetic Alterations In Hypodiploid Acute Lymphoblastic Leukemia Identifies a High Frequency of Mutations Targeting the IKAROS Gene Family and Ras Signaling. Blood, 2010, 116, 411-411.	1.4	3
125	Sole Trisomy 8 In Patients (pts) with De Novo Acute Myeloid Leukemia (AML) Is Associated with Age-Independent Poor Outcome That Is Modified by Molecular Markers and with Unique Gene- and Microrna (miR)-Signatures: a Cancer and Leukemia Group B (CALGB) Study. Blood, 2010, 116, 577-577.	1.4	2
126	Clonal Markers In Relapsed Acute Promyelocytic Leukemia (APL): Clinicopathological Associations and Relation to All-Trans Retinoic Acid (ATRA) Treatment on Intergroup Phase III Trial C9710.. Blood, 2010, 116, 1038-1038.	1.4	0

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127	Targeted Microarray Analyses Augment the Clinical Cytogenetic Diagnosis of Acute Lymphoblastic Leukemia (ALL): Submicroscopic Genetic Events Improve Diagnosis, Contribute to Risk Stratification, and Provide Genetic Markers for Minimal Residual Disease (MRD) Testing. Blood, 2010, 116, 2690-2690.	1.4	0
128	Mutations In the Tet Oncogene Family Member 2 (TET2) Gene Refine the New European LeukemiaNet Risk Classification of Primary, Cytogenetically Normal Acute Myeloid Leukemia (CN-AML) In Adults: A Cancer and Leukemia Group B (CALGB) Study. Blood, 2010, 116, 98-98.	1.4	0
129	Gene Expression Profiling Reveals Genes Predictive of Outcome In Infant Acute Lymphoblastic Leukemia (ALL) and Distinctive Age-Related Gene Expression Profiles (< 90 Days vs. > 90 Days): A Children's Oncology Group Study. Blood, 2010, 116, 412-412.	1.4	0
130	Prognostic Importance of MN1 Transcript Levels, and Biologic Insights From MN1-Associated Gene and MicroRNA Expression Signatures in Cytogenetically Normal Acute Myeloid Leukemia: A Cancer and Leukemia Group B Study. Journal of Clinical Oncology, 2009, 27, 3198-3204.	1.6	149
131	Adverse Prognostic Impact of FLT3 Internal Tandem Duplication (ITD) Is Age-Associated in Older [â‰¥60 Years (Y)] De Novo cytogenetically Normal Acute Myeloid Leukemia (CN-AML) Patients (Pts): a Cancer and Leukemia Group B (CALGB) Study.. Blood, 2009, 114, 1579-1579.	1.4	2
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