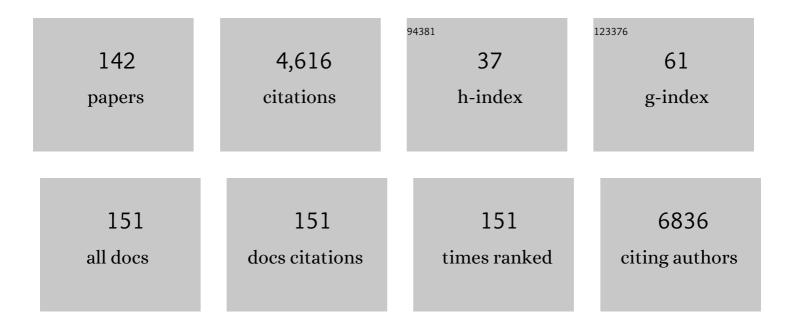
Markus Metzler

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Age†and sexâ€specific pediatric reference intervals for neutrophilâ€toâ€lymphocyte ratio, lymphocyteâ€toâ€monocyte ratio, and plateletâ€toâ€lymphocyte ratio. International Journal of Laboratory Hematology, 2022, 44, 296-301.	0.7	20
2	The Cytogenetic Landscape of Pediatric Chronic Myeloid Leukemia Diagnosed in Chronic Phase. Cancers, 2022, 14, 1712.	1.7	3
3	High-Dose Treosulfan and Melphalan as Consolidation Therapy Versus Standard Therapy for High-Risk (Metastatic) Ewing Sarcoma. Journal of Clinical Oncology, 2022, 40, 2307-2320.	0.8	24
4	Rare pediatric tumors in Germany–Ânot as rare as expected: a study based on data from the Bavarian Cancer Registry and the German Childhood Cancer Registry. European Journal of Pediatrics, 2022, , 1.	1.3	0
5	Genetic testing and surveillance in infantile myofibromatosis: a report from the SIOPE Host Genome Working Group. Familial Cancer, 2021, 20, 327-336.	0.9	13
6	Distribution and Cytokine Profile of Peripheral B Cell Subsets Is Perturbed in Pediatric IBD and Partially Restored During a Successful IFX Therapy. Inflammatory Bowel Diseases, 2021, 27, 224-235.	0.9	4
7	Pediatric Mesothelioma With ALK Fusions. American Journal of Surgical Pathology, 2021, 45, 653-661.	2.1	22
8	Definition, Epidemiology, Pathophysiology, and Essential Criteria for Diagnosis of Pediatric Chronic Myeloid Leukemia. Cancers, 2021, 13, 798.	1.7	18
9	High-resolution pediatric reference intervals for 15 biochemical analytes described using fractional polynomials. Clinical Chemistry and Laboratory Medicine, 2021, 59, 1267-1278.	1.4	15
10	Paediatric chronic myeloid leukaemia presenting in <i>de novo</i> or secondary blast phase ―a comparison of clinical and genetic characteristics. British Journal of Haematology, 2021, 193, 613-618.	1.2	6
11	Haematological characteristics and spontaneous haematological recovery in Pearson syndrome. British Journal of Haematology, 2021, 193, 1283-1287.	1.2	8
12	Ewing Sarcoma—Diagnosis, Treatment, Clinical Challenges and Future Perspectives. Journal of Clinical Medicine, 2021, 10, 1685.	1.0	101
13	Initial presenting manifestations in 16,486 patients with inborn errors of immunity include infections and noninfectious manifestations. Journal of Allergy and Clinical Immunology, 2021, 148, 1332-1341.e5.	1.5	75
14	Multimodal analysis of cell-free DNA whole-genome sequencing for pediatric cancers with low mutational burden. Nature Communications, 2021, 12, 3230.	5.8	95
15	Course of renal allograft function after diagnosis and treatment of postâ€ŧransplant lymphoproliferative disorders in pediatric kidney transplant recipients. Pediatric Transplantation, 2021, 25, e14042.	0.5	4
16	Quantification of Translocation-Specific ctDNA Provides an Integrating Parameter for Early Assessment of Treatment Response and Risk Stratification in Ewing Sarcoma. Clinical Cancer Research, 2021, 27, 5922-5930.	3.2	14
17	The age of the bone marrow microenvironment influences B-cell acute lymphoblastic leukemia progression via CXCR5-CXCL13. Blood, 2021, 138, 1870-1884.	0.6	20
18	Influence of Turkish origin on hematology reference intervals in the German population. Scientific Reports, 2021, 11, 21074.	1.6	4

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19	Liquid Biopsies in Ewing Sarcoma. Methods in Molecular Biology, 2021, 2226, 39-45.	0.4	1
20	Side Effects and Sequelae of Treatment for Chronic Myeloid Leukemia in Childhood and Adolescence. , 2021, , 189-205.		0
21	Data mining of pediatric reference intervals. Journal of Laboratory Medicine, 2021, 45, 311-317.	1.1	8
22	Genotyping circulating tumor DNA of pediatric Hodgkin lymphoma. Leukemia, 2020, 34, 151-166.	3.3	53
23	Comprehensive assessments and related interventions to enhance the long-term outcomes of child, adolescent and young adult cancer survivors – presentation of the CARE for CAYA-Program study protocol and associated literature review. BMC Cancer, 2020, 20, 16.	1.1	25
24	Sickle cell disease in Germany: Results from a national registry. Pediatric Blood and Cancer, 2020, 67, e28130.	0.8	20
25	Assessment of treatment responses in children and adolescents with Ewing sarcoma with metabolic tumor parameters derived from 18F-FDG-PET/CT and circulating tumor DNA. European Journal of Nuclear Medicine and Molecular Imaging, 2020, 47, 1564-1575.	3.3	14
26	Latent class distributional regression for the estimation of non-linear reference limits from contaminated data sources. BMC Bioinformatics, 2020, 21, 524.	1.2	8
27	Low-frequency variation near common germline susceptibility loci are associated with risk of Ewing sarcoma. PLoS ONE, 2020, 15, e0237792.	1.1	6
28	Exploitable metabolic dependencies in MLL-ENL–induced leukemia. Blood Advances, 2020, 4, 3626-3638.	2.5	2
29	Malignant teratoid tumor of the thyroid gland: an aggressive primitive multiphenotypic malignancy showing organotypical elements and frequent DICER1 alterations—is the term "thyroblastoma―more appropriate?. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 477, 787-798.	1.4	45
30	Bone marrow niche-derived extracellular matrix-degrading enzymes influence the progression of B-cell acute lymphoblastic leukemia. Leukemia, 2020, 34, 1540-1552.	3.3	46
31	Reference Interval Estimation from Mixed Distributions using Truncation Points and the Kolmogorov-Smirnov Distance (kosmic). Scientific Reports, 2020, 10, 1704.	1.6	42
32	Social inequalities in the participation and activity of children and adolescents with leukemia, brain tumors, and sarcomas (SUPATEEN): a protocol for a multicenter longitudinal prospective observational study. BMC Pediatrics, 2020, 20, 48.	0.7	3
33	Vaccination With Live Attenuated Vaccines in Four Children With Chronic Myeloid Leukemia While on Imatinib Treatment. Frontiers in Immunology, 2020, 11, 628.	2.2	7
34	Model-Based Simulation of Maintenance Therapy of Childhood Acute Lymphoblastic Leukemia. Frontiers in Physiology, 2020, 11, 217.	1.3	10
35	Blood counts in adult and elderly individuals: defining the norms over eight decades of life. British Journal of Haematology, 2020, 189, 777-789.	1.2	22
36	Horn of plenty: Value of the international registry for pediatric chronic myeloid leukemia. World Journal of Clinical Oncology, 2020, 11, 308-319.	0.9	10

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37	Large amplicon droplet digital PCR for DNAâ€based monitoring of pediatric chronic myeloid leukaemia. Journal of Cellular and Molecular Medicine, 2019, 23, 4955-4961.	1.6	17
38	The German National Registry of Primary Immunodeficiencies (2012–2017). Frontiers in Immunology, 2019, 10, 1272.	2.2	71
39	The preleukemic TCF3-PBX1 gene fusion can be generated in utero and is present in â‰^0.6% of healthy newborns. Blood, 2019, 134, 1355-1358.	0.6	28
40	Data mining of reference intervals for coagulation screening tests in adult patients. Clinica Chimica Acta, 2019, 499, 108-114.	0.5	15
41	<i>TRIM28</i> haploinsufficiency predisposes to Wilms tumor. International Journal of Cancer, 2019, 145, 941-951.	2.3	45
42	Next-generation reference intervals for pediatric hematology. Clinical Chemistry and Laboratory Medicine, 2019, 57, 1595-1607.	1.4	42
43	High sensitivity and clonal stability of the genomic fusion as single marker for response monitoring in <i>ETV6â€RUNX1</i> â€positive acute lymphoblastic leukemia. Pediatric Blood and Cancer, 2019, 66, e27780.	0.8	10
44	Brainstem biopsy in pediatric diffuse intrinsic pontine glioma in the era of precision medicine: the INFORM study experience. European Journal of Cancer, 2019, 114, 27-35.	1.3	51
45	Computed Tomography-Guided Wire-Marking for Thoracoscopic Resection of Small Lung Nodules in Children. Journal of Laparoendoscopic and Advanced Surgical Techniques - Part A, 2019, 29, 688-693.	0.5	2
46	Molecular Composition of Genomic <i>TMPRSS2-ERG</i> Rearrangements in Prostate Cancer. Disease Markers, 2019, 2019, 1-8.	0.6	8
47	A mathematical model of white blood cell dynamics during maintenance therapy of childhood acute lymphoblastic leukemia. Mathematical Medicine and Biology, 2019, 36, 471-488.	0.8	9
48	Indirect determination of hematology reference intervals in adult patients on Beckman Coulter UniCell DxH 800 and Abbott CELL-DYN Sapphire devices. Clinical Chemistry and Laboratory Medicine, 2019, 57, 730-739.	1.4	18
49	High Platelet Counts, Thrombosis, Bleeding Signs, and Acquired Von Willebrand Syndrome at Diagnosis of Pediatric Chronic Myeloid Leukemia. Blood, 2019, 134, 4152-4152.	0.6	3
50	Abstract 2506: Exploring the complex etiology of oncogenic fusions in childhood cancer. , 2019, , .		0
51	Vaccination with Live Attenuated Virus Vaccines in Four Pediatric Patients with CML While on Imatinib Treatment. Blood, 2019, 134, 5903-5903.	0.6	0
52	The Influence of the Age of the Bone Marrow Microenvironment on Leukaemia Progression. Blood, 2019, 134, 2748-2748.	0.6	0
53	A Comparison of GFR Estimation Formulae in Pediatric Oncology. Klinische Padiatrie, 2018, 230, 142-150.	0.2	2
54	Pharmacology and pharmacokinetics of imatinib in pediatric patients. Expert Review of Clinical Pharmacology, 2018, 11, 219-231.	1.3	43

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55	Generic formulations of imatinib for treatment of Philadelphia chromosome–positive leukemia in pediatric patients. Pediatric Blood and Cancer, 2018, 65, e27431.	0.8	11
56	Rearrangement bursts generate canonical gene fusions in bone and soft tissue tumors. Science, 2018, 361, .	6.0	121
57	MECOM-associated syndrome: a heterogeneous inherited bone marrow failure syndrome with amegakaryocytic thrombocytopenia. Blood Advances, 2018, 2, 586-596.	2.5	75
58	Low incidence of symptomatic osteonecrosis after allogeneic <scp>HSCT</scp> in children with highâ€risk or relapsed <scp>ALL</scp> – results of the <scp>ALL</scp> â€ <scp>SCT</scp> 2003 trial. British Journal of Haematology, 2018, 183, 104-109.	1.2	12
59	Genome-wide association study identifies multiple new loci associated with Ewing sarcoma susceptibility. Nature Communications, 2018, 9, 3184.	5.8	50
60	Multidisciplinary Late Effects Clinics for Childhood Cancer Survivors in Germany - a Two-Center Study. Oncology Research and Treatment, 2018, 41, 430-436.	0.8	22
61	Front-line imatinib treatment in children and adolescents with chronic myeloid leukemia: results from a phase III trial. Leukemia, 2018, 32, 1657-1669.	3.3	86
62	Characterization and diagnostic application of genomic NPM-ALK fusion sequences in anaplastic large-cell lymphoma. Oncotarget, 2018, 9, 26543-26555.	0.8	14
63	Adrenal crisis in a 14-year-old boy 12 years after hematopoietic stem cell transplantation. Endocrinology, Diabetes and Metabolism Case Reports, 2018, 2018, .	0.2	0
64	Abstract 2970: Multiple new susceptibility loci identified in genome-wide association study of Ewing sarcoma. , 2018, , .		0
65	Abstract A13: Genome-wide association study identifies multiple new loci associated with Ewing sarcoma susceptibility. , 2018, , .		0
66	Childhood cancer predisposition syndromes—A concise review and recommendations by the Cancer Predisposition Working Group of the Society for Pediatric Oncology and Hematology. American Journal of Medical Genetics, Part A, 2017, 173, 1017-1037.	0.7	200
67	Systematic comparison of donor chimerism in peripheral blood and bone marrow after hematopoietic stem cell transplantation. Blood Cancer Journal, 2017, 7, e566-e566.	2.8	14
68	Recurrent Somatic PDGFRB Mutations in Sporadic Infantile/Solitary Adult Myofibromas But Not in Angioleiomyomas and Myopericytomas. American Journal of Surgical Pathology, 2017, 41, 195-203.	2.1	76
69	Long-term positive and negative psychosocial outcomes in young childhood cancer survivors, type 1 diabetics and their healthy peers. International Journal of Adolescent Medicine and Health, 2017, 29, .	0.6	4
70	Pediatric reference intervals for alkaline phosphatase. Clinical Chemistry and Laboratory Medicine, 2017, 55, 102-110.	1.4	78
71	The second European interdisciplinary Ewing sarcoma research summit - A joint effort to deconstructing the multiple layers of a complex disease. Oncotarget, 2016, 7, 8613-8624.	0.8	55
72	Pediatric Colorectal Carcinoma is Associated With Excellent Outcome in the Context of Cancer Predisposition Syndromes. Pediatric Blood and Cancer, 2016, 63, 611-617.	0.8	22

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73	Paediatric and adult soft tissue sarcomas with <i>NTRK1</i> gene fusions: a subset of spindle cell sarcomas unified by a prominent myopericytic/haemangiopericytic pattern. Journal of Pathology, 2016, 238, 700-710.	2.1	108
74	Genotype, Clinical Course, and Therapeutic Decision Making in 76 Infants with SevereÂGeneralized Junctional EpidermolysisÂBullosa. Journal of Investigative Dermatology, 2016, 136, 2150-2157.	0.3	58
75	Pediatric chronic myeloid leukemia is a unique disease that requires a different approach. Blood, 2016, 127, 392-399.	0.6	141
76	Hyperactive mTOR pathway promotes lymphoproliferation and abnormal differentiation in autoimmune lymphoproliferative syndrome. Blood, 2016, 128, 227-238.	0.6	77
77	The ENCCA-WP7/EuroSarc/EEC/PROVABES/EURAMOS 3rd European Bone Sarcoma Networking Meeting/Joint Workshop of EU Bone Sarcoma Translational Research Networks; Vienna, Austria, September 24–25, 2015. Workshop Report. Clinical Sarcoma Research, 2016, 6, 3.	2.3	14
78	Genomic <i>EWSR1</i> Fusion Sequence as Highly Sensitive and Dynamic Plasma Tumor Marker in Ewing Sarcoma. Clinical Cancer Research, 2016, 22, 4356-4365.	3.2	68
79	Adamantinomatous and papillary craniopharyngiomas are characterized by distinct epigenomic as well as mutational and transcriptomic profiles. Acta Neuropathologica Communications, 2016, 4, 20.	2.4	136
80	Therapy with lowâ€dose azacitidine for <scp>MDS</scp> in children and young adults: a retrospective analysis of the <scp>EWOG</scp> â€ <scp>MDS</scp> study group. British Journal of Haematology, 2016, 172, 930-936.	1.2	31
81	Nivolumab As Salvage Therapy in Pediatric Patients with Relapsed and Refractory Lymphomas. Blood, 2016, 128, 5414-5414.	0.6	5
82	Age- and Sex-Specific Dynamics in 22 Hematologic and Biochemical Analytes from Birth to Adolescence. Clinical Chemistry, 2015, 61, 964-973.	1.5	132
83	Sclerosing epithelioid fibrosarcoma of the kidney: clinicopathologic and molecular study of a rare neoplasm at a novel location. Annals of Diagnostic Pathology, 2015, 19, 221-225.	0.6	19
84	Response monitoring of infant acute myeloid leukemia treatment by quantification of the tumor specific <i>MLL–FNBP1</i> fusion gene. Leukemia and Lymphoma, 2015, 56, 793-796.	0.6	4
85	From initiation to eradication: the lifespan of an MLL-rearranged therapy-related paediatric AML. Bone Marrow Transplantation, 2015, 50, 1382-1384.	1.3	3
86	Insights into the Infiltrative Behavior of Adamantinomatous Craniopharyngioma in a New Xenotransplant Mouse Model. Brain Pathology, 2015, 25, 1-10.	2.1	42
87	Feasibility of VECOPA, a dose-intensive chemotherapy regimen for children and adolescents with intermediate and advanced stage Hodgkin lymphoma: results of the GPOH-HD-2002/VECOPA pilot trial. Leukemia and Lymphoma, 2015, 56, 1308-1314.	0.6	4
88	Hyperactive mTOR Pathway Promotes Lymphoproliferation and Abnormal Differentiation in Human Autoimmune Lymphoproliferative Syndrome. Blood, 2015, 126, 1020-1020.	0.6	1
89	Spleen Size at Diagnosis and Genomic BCR-ABL1 Breakpoint Distribution Differ Age-Dependently in Pediatric Patients with Chronic Myeloid Leukemia (CML). Blood, 2015, 126, 4827-4827.	0.6	0
90	Frequent and sex-biased deletion of SLX4IP by illegitimate V(D)J-mediated recombination in childhood acute lymphoblastic leukemia. Human Molecular Genetics, 2014, 23, 590-601.	1.4	13

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91	Clustering of genomic breakpoints at the <i>MLL</i> locus in therapyâ€related acute leukemia with t(4;11)(q21;q23). Genes Chromosomes and Cancer, 2014, 53, 248-254.	1.5	5
92	<scp>DNA</scp> copy number alterations mark disease progression in paediatric chronic myeloid leukaemia. British Journal of Haematology, 2014, 166, 250-253.	1.2	9
93	Malignant Epithelioid Peripheral Nerve Sheath Tumor With Prominent Reticular/Microcystic Pattern in a Child. Applied Immunohistochemistry and Molecular Morphology, 2014, 22, 627-633.	0.6	2
94	Sustained complete molecular remission after imatinib discontinuation in children with chronic myeloid leukemia. Pediatric Blood and Cancer, 2014, 61, 2080-2082.	0.8	11
95	Decisions Taken in Children and Adolescents with Chronic Myeloid Leukemia (CML) at Failure of Imatinib Treatment. Blood, 2014, 124, 1798-1798.	0.6	0
96	Indirect determination of pediatric blood count reference intervals. Clinical Chemistry and Laboratory Medicine, 2013, 51, 863-872.	1.4	65
97	Identification of the genomic BCR-ABL1 fusion sequence from blood specimen stored on filter paper. Leukemia Research, 2013, 37, 117-119.	0.4	2
98	Novel MLL2 Mutation in Kabuki Syndrome With Hypogammaglobulinemia and Severe Chronic Thrombopenia. Journal of Pediatric Hematology/Oncology, 2013, 35, e314-e316.	0.3	4
99	Copy Number Variations and IKZF1 Mutations In Pediatric CML. Blood, 2013, 122, 1473-1473.	0.6	2
100	Clinical Impact Of Post-Transplant Chimerism Monitoring In CD33/34 Bone Marrow Subpopulations and Whole Blood In Pediatric AML: Prospective Comparison Of Highly Sensitive Real Time Sequence Polymorphism PCR Versus Gold-Standard Conventional STR-PCR. Blood, 2013, 122, 411-411.	0.6	3
101	Genomic EWS-FLI1 Fusion Sequences in Ewing Sarcoma Resemble Breakpoint Characteristics of Immature Lymphoid Malignancies. PLoS ONE, 2013, 8, e56408.	1.1	30
102	B-Cell Precursor Acute Lymphoblastic Leukemia (BCP-ALL) Specific Copy Number Alterations Are Unique For Progressive Pediatric Chronic Myeloid Leukemia (CML): A Large Cohort Study. Blood, 2013, 122, 2715-2715.	0.6	0
103	The First European Interdisciplinary Ewing Sarcoma Research Summit. Frontiers in Oncology, 2012, 2, 54.	1.3	32
104	Targeted Therapeutics in Treatment of Children and Young Adults with Solid Tumors: an Expert Survey and Review of the Literature. Klinische Padiatrie, 2012, 224, 124-131.	0.2	15
105	Malignant Peritoneal Mesothelioma in a 16-Year-Old Girl: Presentation of a Rare Disease. Klinische Padiatrie, 2012, 224, 170-173.	0.2	9
106	Genomic <i>BCR</i> â€ <i>ABL1</i> breakpoints in pediatric chronic myeloid leukemia. Genes Chromosomes and Cancer, 2012, 51, 1045-1053.	1.5	50
107	ETV6/RUNX1-positive relapses evolve from an ancestral clone and frequently acquire deletions of genes implicated in glucocorticoid signaling. Blood, 2011, 117, 2658-2667.	0.6	83
108	The turnover of synovial T cells is higher than in T cells in the peripheral blood in persistent oligoarticular juvenile idiopathic arthritis. Rheumatology International, 2010, 30, 1529-1532.	1.5	1

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109	Bimodal distribution of genomic MLL breakpoints in infant acute lymphoblastic leukemia treatment. Leukemia, 2010, 24, 903-907.	3.3	16
110	Recurrent Fever Episodes with Arthralgia or Hyperesthesia – Have You Ruled Out Parvovirus B19?. Klinische Padiatrie, 2010, 222, 397-398.	0.2	2
111	HLA-DQA1 gene expression profiling in oligoarticular JIA. Autoimmunity, 2009, 42, 389-391.	1.2	9
112	Differences in DNA Methylation Patterns and Expression of the CCRK Gene in Human and Nonhuman Primate Cortices. Molecular Biology and Evolution, 2009, 26, 1379-1389.	3.5	47
113	Prognostic significance of minimal residual disease in infants with acute lymphoblastic leukemia treated within the Interfant-99 protocol. Leukemia, 2009, 23, 1073-1079.	3.3	137
114	Cluster analysis of genomic ETV6–RUNX1 (TEL–AML1) fusion sites in childhood acute lymphoblastic leukemia. Leukemia Research, 2009, 33, 1082-1088.	0.4	18
115	Inv(11)(q21q23) fuses MLL to the Notch co-activator mastermind-like 2 in secondary T-cell acute lymphoblastic leukemia. Leukemia, 2008, 22, 1807-1811.	3.3	23
116	Modeling Chromosomal Translocations Using Conditional Alleles to Recapitulate Initiating Events in Human Leukemias. Journal of the National Cancer Institute Monographs, 2008, 2008, 58-63.	0.9	26
117	Screening for leukemia- and clone-specific markers at birth in children with T-cell precursor ALL suggests a predominantly postnatal origin. Blood, 2007, 110, 3036-3038.	0.6	26
118	Temporary blast reduction after immunoglobulin administration for congenital cytomegalovirus infection masking infant leukemia with cryptic MLL rearrangement. Leukemia Research, 2007, 31, 553-557.	0.4	0
119	A conditional model of MLL-AF4 B-cell tumourigenesis using invertor technology. Oncogene, 2006, 25, 3093-3103.	2.6	95
120	No Evidence for Angiotensin Type 2 Receptor Gene Polymorphism in Intron 1 in Patients with Coarctation of the Aorta and Ullrich–Turner Syndrome. Pediatric Cardiology, 2006, 27, 636-639.	0.6	4
121	Low prevalence of Gs α mutations in śomatotroph adenomas of children and adolescents. Cancer Genetics and Cytogenetics, 2006, 166, 146-151.	1.0	12
122	Inv(11)(q21q23) Fuses MLL to the NOTCH Co-Activator Mastermind-Like 2 in Secondary T Cell Acute Lymphoblastic Leukemia Blood, 2006, 108, 4284-4284.	0.6	2
123	Minimal residual disease analysis in children with t(12;21)-positive acute lymphoblastic leukemia: comparison of Ig/TCR rearrangements and the genomic fusion gene. Haematologica, 2006, 91, 683-6.	1.7	13
124	Chromosomal Translocation Engineering to Recapitulate Primary Events of Human Cancer. Cold Spring Harbor Symposia on Quantitative Biology, 2005, 70, 275-282.	2.0	20
125	The invertor knock-in conditional chromosomal translocation mimic. Nature Methods, 2005, 2, 27-30.	9.0	36
126	Use ofMLL/GRAF fusion mRNA for measurement of minimal residual disease during chemotherapy in an infant with acute monoblastic leukemia (AML-M5). Genes Chromosomes and Cancer, 2005, 43, 424-426.	1.5	10

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127	The Ews-ERG Fusion Protein Can Initiate Neoplasia from Lineage-Committed Haematopoietic Cells. PLoS Biology, 2005, 3, e242.	2.6	39
128	Asymmetric multiplex-polymerase chain reaction - a high throughput method for detection and sequencing genomic fusion sites in t(4;11). British Journal of Haematology, 2004, 124, 47-54.	1.2	6
129	High expression of precursor microRNA-155/BIC RNA in children with Burkitt lymphoma. Genes Chromosomes and Cancer, 2004, 39, 167-169.	1.5	530
130	Protracted postnatal natural histories in childhood leukemia. Genes Chromosomes and Cancer, 2004, 39, 335-340.	1.5	57
131	Emergence of translocation t(9;11)-positive leukemia during treatment of childhood acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2004, 41, 291-296.	1.5	11
132	Analysis of t(9;11) chromosomal breakpoint sequences in childhood acute leukemia: Almost identicalMLL breakpoints in therapy-related AML after treatment without etoposides. Genes Chromosomes and Cancer, 2003, 36, 393-401.	1.5	70
133	Occurrence of anMLL/LAF4 fusion gene caused by the insertion ins(11;2)(q23;q11.2q11.2) in an infant with acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2003, 37, 106-109.	1.5	17
134	A New Missense Mutation of the Vasopressin-Neurophysin II Gene in a Family with Neurohypophyseal Diabetes insipidus. Hormone Research in Paediatrics, 2003, 60, 143-147.	0.8	8
135	Hematologic Features and Clinical Course of an Infant With Pearson Syndrome Caused by a Novel Deletion of Mitochondrial DNA. Journal of Pediatric Hematology/Oncology, 2003, 25, 948-951.	0.3	20
136	Late relapses evolve from slow-responding subclones in t(12;21)-positive acute lymphoblastic leukemia: evidence for the persistence of a preleukemic clone. Blood, 2003, 101, 3635-3640.	0.6	84
137	An unusual manifestation of Wegener's granulomatosis in a 4-year-old girl. Pediatric Neurology, 2002, 27, 71-74.	1.0	13
138	Adrenomedullin, calcitonin gene-related peptide and their receptors: evidence for a decreased placental mRNA content in preeclampsia and HELLP syndrome. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2002, 101, 47-53.	0.5	38
139	Identification of Leptin in Human Saliva. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 5234-5239.	1.8	27
140	Quantitative DNA Fragment Analysis for Detecting Low Amounts of Hepatitis B Virus Deletion Mutants in Highly Viremic Carriers. Hepatology, 2000, 32, 1096-1105.	3.6	11
141	Nuclear transport of oligonucleotides in HepG2-cells mediated by protamine sulfate and negatively charged liposomes. Pharmaceutical Research, 2000, 17, 1206-1211.	1.7	23
142	A new fingerprint method for sequence analysis of chromosomal translocations at the genomic DNA level. Leukemia, 1998, 12, 758-763.	3.3	7