List of Publications by Year in descending order

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Δείδα ζηιμαρά

#	Article	IF	CITATIONS
1	Acute kidney injury in Japanese type 2 diabetes patients receiving sodium–glucose cotransporter 2 inhibitors: A nationwide cohort study. Journal of Diabetes Investigation, 2022, 13, 42-46.	1.1	5
2	Dupilumab-related type 1 diabetes in a patient with atopic dermatitis: a case report. Diabetology International, 2022, 13, 300-303.	0.7	4
3	Japanese Type 1 Diabetes Database Study (TIDE-J): rationale and study design. Diabetology International, 2022, 13, 288-294.	0.7	4
4	Current clinical state of type 1 diabetes in Saitama prefecture. Diabetology International, 2022, 13, 436-446.	0.7	2
5	Genome-wide DNA methylation analysis in pediatric acute myeloid leukemia. Blood Advances, 2022, 6, 3207-3219.	2.5	7
6	Predisposition to prolonged neutropenia after chemotherapy for paediatric acute myeloid leukaemia is associated with better prognosis in the Japanese Paediatric Leukaemia/Lymphoma Study Group AMLâ€05 study. British Journal of Haematology, 2021, 193, 176-180.	1.2	3
7	The outcomes of relapsed acute myeloid leukemia in children: Results from the Japanese Pediatric Leukemia/Lymphoma Study Group AMLâ€05R study. Pediatric Blood and Cancer, 2021, 68, e28736.	0.8	11
8	Bone marrow transplantation from a human leukocyte antigen-mismatched unrelated donor in a case with C1q deficiency associated with refractory systemic lupus erythematosus. International Journal of Hematology, 2021, 113, 302-307.	0.7	4
9	Onâ€label use of sodium–glucose cotransporterÂ2 inhibitors might increase the risk of diabetic ketoacidosis in patients with typeÂ1 diabetes. Journal of Diabetes Investigation, 2021, 12, 1586-1593.	1.1	8
10	Post-induction MRD by FCM and GATA1-PCR are significant prognostic factors for myeloid leukemia of Down syndrome. Leukemia, 2021, 35, 2508-2516.	3.3	5
11	Clinical significance of RAS pathway alterations in pediatric acute myeloid leukemia. Haematologica, 2021, , .	1.7	10
12	Clinical features resembling subcutaneous insulin resistance observed in a patient with type 2 diabetes and severe COVID-19-associated pneumonia: a case report. Diabetology International, 2021, 12, 474-479.	0.7	6
13	Glycemic control status, diabetes management patterns, and clinical characteristics of adults with type 1 diabetes in Japan: Study of Adults' Glycemia in T1DM subanalysis. Diabetology International, 2021, 12, 460-473.	0.7	4
14	Profile of down syndrome–associated malignancies: Epidemiology, clinical features and therapeutic aspects. Pediatric Hematology Oncology Journal, 2021, 6, 63-72.	0.1	3
15	Investigation of the molecular causes underlying physical abnormalities in Diamondâ€Blackfan anemia patients with RPL5 haploinsufficiency. Pathology International, 2021, , .	0.6	1
16	Effects of Ppar <sup>ĵ</sup> 31 deletion on late-stage murine embryogenesis and cells that undergo endocycle. Developmental Biology, 2021, 478, 222-235.	0.9	2
17	Anagliptin Monotherapy for Six Months in Patients With Type 2 Diabetes Mellitus and Hyper-Low-Density Lipoprotein Cholesterolemia Reduces Plasma Levels of Fasting Low-Density Lipoprotein Cholesterol and Lathosterol: A Single-Arm Intervention Trial. Journal of Clinical Medicine Research. 2021. 13. 502-509.	0.6	2
18	Clonal Evolution Pattern and Prognostic Significance of Clonal Architecture in KMT2A-Rearranged Acute Myeloid Leukemia. Blood, 2021, 138, 2358-2358.	0.6	0

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19	Clinical Significance of Insulin Peptide–specific Interferon-γ–related Immune Responses in Ketosis-prone Type 2 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2021, , .	1.8	2
20	Reduced oxygenation but not fibrosis defined by functional magnetic resonance imaging predicts the long-term progression of chronic kidney disease. Nephrology Dialysis Transplantation, 2020, 35, 964-970.	0.4	40
21	Bodyweight threshold for sudden onset of ketosis might exist in ketosisâ€prone typeÂ2 diabetes patients. Journal of Diabetes Investigation, 2020, 11, 499-501.	1.1	3
22	Hematopoietic Stem Cell Transplantation in Solid Organ Recipients with Emphasis on Transplant Complications: A Nationwide Retrospective Survey on Behalf of the Japan Society for Hematopoietic Stem Cell Transplantation Transplant Complications Working Group. Biology of Blood and Marrow Transplantation, 2020, 26, 66-75.	2.0	4
23	The effect of graftâ€versusâ€host disease on outcomes after allogeneic stem cell transplantation for refractory lymphoblastic lymphoma in children and young adults. Pediatric Blood and Cancer, 2020, 67, e28129.	0.8	5
24	Nationwide retrospective review of hematopoietic stem cell transplantation in children with refractory Langerhans cell histiocytosis. International Journal of Hematology, 2020, 111, 137-148.	0.7	9
25	Pediatric growing teratoma syndrome of the ovary. Medicine (United States), 2020, 99, e22297.	0.4	8
26	Ezetimibe impairs transcellular lipid trafficking and induces large lipid droplet formation in intestinal absorptive epithelial cells. Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids, 2020, 1865, 158808.	1.2	6
27	Real-world risk of hypoglycemia-related hospitalization in Japanese patients with type 2 diabetes using SGLT2 inhibitors: a nationwide cohort study. BMJ Open Diabetes Research and Care, 2020, 8, e001856.	1.2	18
28	Attempts to optimize postinduction treatment in childhood acute myeloid leukemia without coreâ€binding factors: A report from the Japanese Pediatric Leukemia/Lymphoma Study Group (JPLSG). Pediatric Blood and Cancer, 2020, 67, e28692.	0.8	8
29	Possible involvement of autoimmunity in fulminant type 1 diabetes. Diabetology International, 2020, 11, 329-335.	0.7	8
30	Pluripotent stem cell model of Shwachman–Diamond syndrome reveals apoptotic predisposition of hemoangiogenic progenitors. Scientific Reports, 2020, 10, 14859.	1.6	4
31	Clinical features of children with polycythemia vera, essential thrombocythemia, and primary myelofibrosis in Japan: A retrospective nationwide survey. EJHaem, 2020, 1, 86-93.	0.4	3
32	A case of alveolar rhabdomyosarcoma showing concurrent responsive bone marrow lesions and refractory pancreatic lesions to pazopanib monotherapy. Pediatric Blood and Cancer, 2020, 67, e28323.	0.8	1
33	Severe acute intestinal graft versus host disease requiring surgical resection. EJHaem, 2020, 1, 328-329.	0.4	0
34	Simultaneous development of Graves' disease and typeÂ1 diabetes during antiâ€programmed cell deathâ€1 therapy: A case report. Journal of Diabetes Investigation, 2020, 11, 1006-1009.	1.1	17
35	A Sudden Onset of Severe Thrombocytopenia While Using Evolocumab. Case Reports in Hematology, 2020, 1-4.	0.3	1
36	Panel‑based next‑generation sequencing facilitates the characterization of childhood acute myeloid leukemia in clinical settings. Biomedical Reports, 2020, 13, 1-1.	0.9	5

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37	RARE-32. PEDIATRIC METASTATIC SKULL BASE CHORDOMA WITH TP53 MUTATION – A CASE REPORT AND REVIEW OF THE LITERATURE. Neuro-Oncology, 2020, 22, iii449-iii449.	0.6	2
38	Etoposide, Cytarabine and Mitoxantrone- or Fludarabine, Cytarabine and Granulocyte Colony-Stimulating Factor-Based Intensive Reinduction Chemotherapy Is Recommended for Children with Relapsed Acute Myeloid Leukemia: The Results from the Japanese Pediatric Leukemia/Lymphoma Study Group (JPLSG) AML-05R Study. Blood, 2020, 136, 6-6.	0.6	0
39	Delayed Methotrexate Elimination after Administration of a Medium Dose of Methotrexate in a Patient with Genetic Variants Associated with Methotrexate Clearance. Acta Medica Okayama, 2020, 74, 545-550.	0.1	1
40	Lupus anticoagulant-hypoprothrombinemia syndrome and immunoglobulin-A vasculitis: a report of Japanese sibling cases and review of the literature. Rheumatology International, 2019, 39, 1811-1819.	1.5	9
41	Unique Inflammatory Changes in Exocrine and Endocrine Pancreas in Enterovirus-Induced Fulminant Type 1 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 4282-4294.	1.8	14
42	Significance of peripheral mononuclear cells producing interferon-γ in response to insulin B:9–23-related peptides in subtypes of type 1 diabetes. Clinical Immunology, 2019, 208, 108260.	1.4	2
43	Hematological malignancies and molecular targeting therapy. European Journal of Pharmacology, 2019, 862, 172641.	1.7	44
44	Discontinuation of l-asparaginase and poor response to prednisolone are associated with poor outcome of ETV6-RUNX1-positive pediatric B-cell precursor acute lymphoblastic leukemia. International Journal of Hematology, 2019, 109, 477-482.	0.7	16
45	Detailed Time Course of Decline in Serum C-Peptide Levels in Anti–Programmed Cell Death-1 Therapy–Induced Fulminant Type 1 Diabetes. Diabetes Care, 2019, 42, e40-e41.	4.3	12
46	High-grade glioneuronal tumor with an ARHGEF2–NTRK1 fusion gene. Brain Tumor Pathology, 2019, 36, 121-128.	1.1	18
47	Selective laser trabeculoplasty for steroid glaucoma in a child with leukemia. Pediatrics International, 2019, 61, 208-210.	0.2	2
48	Clinical characteristics of anti-glutamic acid decarboxylase antibody-positive fulminant type 1 diabetes. Endocrine Journal, 2019, 66, 329-336.	0.7	3
49	Clinical and biological features of paediatric acute myeloid leukaemia ( AML ) with primary induction failure in the Japanese Paediatric Leukaemia/Lymphoma Study Group AML â€05 study. British Journal of Haematology, 2019, 185, 284-288.	1.2	12
50	Transcriptome analysis offers a comprehensive illustration of the genetic background of pediatric acute myeloid leukemia. Blood Advances, 2019, 3, 3157-3169.	2.5	51
51	Characteristics and clinical course of type 1 diabetes mellitus related to anti-programmed cell death-1 therapy. Diabetology International, 2019, 10, 58-66.	0.7	65
52	Panel-based next-generation sequencing identifies prognostic and actionable genes in childhood acute lymphoblastic leukemia and is suitable for clinical sequencing. Annals of Hematology, 2019, 98, 657-668.	0.8	7
53	Actual condition survey regarding mismatch of measurements between radioimmunoassay and enzymeâ€linked immunosorbent assay tests for antiâ€glutamic acid decarboxylase antibody in realâ€world clinical practice. Journal of Diabetes Investigation, 2019, 10, 685-689.	1.1	4
54	Post-Induction Minimal Residual Disease Measured By Flow Cytometry and Deep Sequencing of Mutant GATA1 Are Both Significant Prognostic Factors for Children with Myeloid Leukemia and Down Syndrome: A Nationwide Prospective Study of the Japanese Pediatric Leukemia/Lymphoma Study Group. Blood. 2019, 134, 3848-3848.	0.6	1

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55	Clinical Features of Children with Polycythemia Vera, Essential Thrombocythemia, and Primary Myelofibrosis in Japan: Retrospective Nationwide Survey. Blood, 2019, 134, 2958-2958.	0.6	1
56	Remission of Congenital Multi-system Type Langerhans Cell Histiocytosis with Chemotherapy. Acta Medica Okayama, 2019, 73, 61-65.	0.1	3
57	Significant Features of DNA Methylation at Bivalent Promotor and Repressed Polycomb Regions in Pediatric AML-the Jccg Study, JPLSG AML-05 Blood, 2019, 134, 2739-2739.	0.6	Ο
58	Influence of Cyclophosphamide on L-Asparaginase-Induced Allergy in Animal Model. Blood, 2019, 134, 5119-5119.	0.6	0
59	Clinical Features of Pediatric Acute Myeloid Leukemia with TP53 and CDKN2A/2B copy Number Alterations. Blood, 2019, 134, 2727-2727.	0.6	Ο
60	Recurrent Gene Mutations in Pediatric Patients with AML By Targeted Sequencing ―the Jccg Study, JPLSG AML-05―. Blood, 2019, 134, 2697-2697.	0.6	0
61	Exchange Transfusion and Cytarabine for Transient Abnormal Myelopoiesis in Hydrops Fetalis. Acta Medica Okayama, 2019, 73, 181-188.	0.1	2
62	Pharmacological inhibition of JAK3 enhances the antitumor activity of imatinib in human chronic myeloid leukemia. European Journal of Pharmacology, 2018, 825, 28-33.	1.7	7
63	Simultaneous detection of <i><scp>ABL</scp>1</i> mutation and <i><scp>IKZF</scp>1</i> deletion in Philadelphia chromosomeâ€positive acute lymphoblastic leukemia using a customized target enrichment system panel. International Journal of Laboratory Hematology, 2018, 40, 427-436.	0.7	6
64	Risk-stratified therapy for children with FLT3-ITD-positive acute myeloid leukemia: results from the JPLSG AML-05 study. International Journal of Hematology, 2018, 107, 586-595.	0.7	20
65	Monitoring of fusion gene transcripts to predict relapse in pediatric acute myeloid leukemia. Pediatrics International, 2018, 60, 41-46.	0.2	13
66	Prognostic value of genetic mutations in adolescent and young adults with acute myeloid leukemia. International Journal of Hematology, 2018, 107, 201-210.	0.7	15
67	Insulin degludec overdose may lead to long-lasting hypoglycaemia through its markedly prolonged half-life. Diabetic Medicine, 2018, 35, 277-280.	1.2	5
68	Multiplex fusion gene testing in pediatric acute myeloid leukemia. Pediatrics International, 2018, 60, 47-51.	0.2	12
69	Luminal plant sterol promotes brush border membrane-to-lumen cholesterol efflux in the small intestine. Journal of Clinical Biochemistry and Nutrition, 2018, 63, 102-105.	0.6	9
70	Wernicke's encephalopathy in a child with autism during chemotherapy for Tâ€cell acute leukemia. Pediatrics International, 2018, 60, 757-758.	0.2	3
71	Copy number abnormality of acute lymphoblastic leukemia cell lines based on their genetic subtypes. International Journal of Hematology, 2018, 108, 312-318.	0.7	10
72	Enhanced AKT Phosphorylation of Circulating B Cells in Patients With Activated PI3Kδ Syndrome. Frontiers in Immunology, 2018, 9, 568.	2.2	15

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73	Hematopoietic Stem Cell Transplantation in Children with Refractory Langerhans Cell Histiocytosis. Blood, 2018, 132, 4657-4657.	0.6	1
74	Comprehensive Analysis of 343 Genes Using Targeted Sequencing Panel By Next-Generation Sequencer in 77 Pediatric AML Patients with Normal and Complex Karyotypes: Jccg Study, JPLSG AML-05. Blood, 2018, 132, 1530-1530.	0.6	0
75	Long-term Remission of Hepatitis-associated Aplastic Anemia Possibly due to Immunosuppressive Therapy after Liver Transplantation. Acta Medica Okayama, 2018, 72, 515-518.	0.1	0
76	Germline IKAROS mutation associated with primary immunodeficiency that progressed to T-cell acute lymphoblastic leukemia. Leukemia, 2017, 31, 1221-1223.	3.3	56
77	Prognostic impact of specific molecular profiles in pediatric acute megakaryoblastic leukemia in nonâ€Down syndrome. Genes Chromosomes and Cancer, 2017, 56, 394-404.	1.5	51
78	Congenital Glioblastoma with Distinct Clinical and Molecular Characteristics: Case Reports and a Literature Review. World Neurosurgery, 2017, 101, 817.e5-817.e14.	0.7	20
79	Outcome of relapsed core binding factor acute myeloid leukemia in children: A result from the Japanese Pediatric Leukemia/Lymphoma Study Group (JPLSG) AMLâ€05R study. Pediatric Blood and Cancer, 2017, 64, e26491.	0.8	5
80	Desmoidâ€ŧype fibromatosis in a boy with Down syndrome. Pediatrics International, 2017, 59, 624-626.	0.2	1
81	Everolimus for Treatment of Pseudomyogenic Hemangioendothelioma. Journal of Pediatric Hematology/Oncology, 2017, 39, e328-e331.	0.3	35
82	Gene expression analysis of hypersensitivity to mosquito bite, chronic active EBV infection and NK/T-lymphoma/leukemia. Leukemia and Lymphoma, 2017, 58, 2683-2694.	0.6	6
83	Pediatric intestinal Behçet disease complicated by myeloid malignancies. International Journal of Hematology, 2017, 105, 377-382.	0.7	7
84	<i><scp>MSH</scp>2</i> deletion with <i><scp>CREBBP</scp></i> and <i><scp>KRAS</scp></i> mutations in pediatric highâ€hyperdiploid acute lymphoblastic leukemia. Pediatrics International, 2017, 59, 1103-1105.	0.2	1
85	Childhood cancer survivors: Anxieties felt after treatment and the need for continued support. Pediatrics International, 2017, 59, 1140-1150.	0.2	5
86	Possible Long-Term Efficacy of Sitagliptin, a Dipeptidyl Peptidase-4 Inhibitor, for Slowly Progressive Type 1 Diabetes (SPIDDM) in the Stage of Non-Insulin-Dependency: An Open-Label Randomized Controlled Pilot Trial (SPAN-S). Diabetes Therapy, 2017, 8, 1123-1134.	1.2	36
87	Fludarabine, cytarabine, granulocyte colonyâ€stimulating factor and idarubicin for relapsed childhood acute myeloid leukemia. Pediatrics International, 2017, 59, 1046-1052.	0.2	7
88	Clinical features of cases of seroconversion of anti-glutamic acid decarboxylase antibody during the clinical course of type 2 diabetes: a nationwide survey in Japan. Diabetology International, 2017, 8, 306-315.	0.7	1
89	Sorafenib treatment for papillary thyroid carcinoma with diffuse lung metastases in a child with autism spectrum disorder: a case report. BMC Cancer, 2017, 17, 775.	1.1	6
90	Positive Minimal Residual Disease of FLT3-ITD before Hematopoietic Stem Cell Transplantation Resulted in a Poor Prognosis of an Acute Myeloid Leukemia. Acta Medica Okayama, 2017, 71, 79-83.	0.1	3

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91	A Case of Refractory Langerhans Cell Histiocytosis Complicated with Hemophagocytic Lymphohistiocytosis Rescued by Cord Blood Transplantation with Reduced-intensity Conditioning. Acta Medica Okayama, 2017, 71, 249-254.	0.1	3
92	Relapsed infant <i>MLL</i> â€rearranged acute lymphoblastic leukemia with additional genetic alterations. Pediatric Blood and Cancer, 2016, 63, 2059-2060.	0.8	5
93	High eventâ€free survival rate with minimumâ€doseâ€anthracycline treatment in childhood acute promyelocytic leukaemia: a nationwide prospective study by the Japanese Paediatric Leukaemia/Lymphoma Study Group. British Journal of Haematology, 2016, 174, 437-443.	1.2	16
94	Verification of risk scores to predict i.v. immunoglobulin resistance in incomplete Kawasaki disease. Pediatrics International, 2016, 58, 146-151.	0.2	14
95	High <i><scp>PRDM</scp>16</i> expression identifies a prognostic subgroup of pediatric acute myeloid leukaemia correlated to <i><scp>FLT</scp>3</i> â€ <scp>ITD</scp> , <i><scp>KMT</scp>2A</i> â€ <scp>PTD</scp> , and <i><scp>NUP</scp>98â€<scp>NSD</scp>1</i> the results of the Japanese Paediatric Leukaemia/Lymphoma	1.2	41
96	Wholeâ€exome sequencing reveals the spectrum of gene mutations and the clonal evolution patterns in paediatric acute myeloid leukaemia. British Journal of Haematology, 2016, 175, 476-489.	1.2	60
97	Persistent clonal chromosomal abnormalities in a chronic myeloid leukemia patient. Pediatrics International, 2016, 58, 53-56.	0.2	1
98	JAK2, MPL, and CALR mutations in children with essential thrombocythemia. International Journal of Hematology, 2016, 104, 266-267.	0.7	10
99	Relapsed childhood acute myeloid leukemia patient with inversion of chromosome 16 harboring a low <i>FLT3</i> internal tandem duplication allelic burden and <i>KIT</i> mutations. Pediatrics International, 2016, 58, 905-908.	0.2	1
100	Preserved High Probability of Overall Survival with Significant Reduction of Chemotherapy for Myeloid Leukemia in Down Syndrome: A Nationwide Prospective Study in Japan. Pediatric Blood and Cancer, 2016, 63, 248-254.	0.8	33
101	Adults with germline CBL mutation complicated with juvenile myelomonocytic leukemia at infancy. Journal of Human Genetics, 2016, 61, 523-526.	1.1	12
102	L-Asparaginase-Induced Allergy in Mice: Effects of Concomitant Drugs and Anti-IgE Antibody. Blood, 2016, 128, 1632-1632.	0.6	1
103	Clinical and Biological Features of Pediatric Acute Myeloid Leukemia with Primary Induction Failure in the Japanese Pediatric Leukemia/Lymphoma Study Group (JPLSG) AML-05 Study. Blood, 2016, 128, 1610-1610.	0.6	1
104	Retrospective Evaluation of Correlations Between Genetic Backgrounds and Stem Cell Transplantation for De Novo Pediatric Acute Myeloid Leukemia: A Study from the Japan Pediatric Leukemia/Lymphoma Study Group (JPLSG) AML-05 Clinical Trial. Blood, 2016, 128, 2904-2904.	0.6	0
105	A Long-term Survivor after Congenital Acute Myeloid Leukemia with t(8 ; 16)(p11 ; p13). Acta Medica Okayama, 2016, 70, 31-5.	0.1	3
106	Two Relapsed Stage III Childhood Anaplastic Large Cell Lymphoma Patients with NPM-ALK Fusion in Bone Marrow from Initial Diagnosis. Acta Medica Okayama, 2016, 70, 503-506.	0.1	0
107	Suspected early onset of congenital Langerhans cell histiocytosis involving ectopic cervical thymus and mediastinal thymus, simultaneously. Pediatric Blood and Cancer, 2015, 62, 1491-1492.	0.8	1
108	Transient myeloproliferative disorder with partial trisomy 21. Pediatric Blood and Cancer, 2015, 62, 2021-2024.	0.8	5

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109	Japanese family with congenital factor VII deficiency. Pediatrics International, 2015, 57, 1023-1024.	0.2	0
110	Outcome of adolescent patients with acute myeloid leukemia treated with pediatric protocols. International Journal of Hematology, 2015, 102, 318-326.	0.7	17
111	Adverse prognostic impact of KIT mutations in childhood CBF-AML: the results of the Japanese Pediatric Leukemia/Lymphoma Study Group AML-05 trial. Leukemia, 2015, 29, 2438-2441.	3.3	28
112	The Outcome of Relapsed Childhood Core Binding Factor Acute Myeloid Leukemia: A Report from the JPLSG AML-05R Study. Blood, 2015, 126, 2516-2516.	0.6	0
113	Distinct Clinical and Cytogenetic Characteristics and Poor Prognosis in Children with Acute Erythroid Leukemia: A Report from the JPLSG AML-05 Study. Blood, 2015, 126, 4945-4945.	0.6	0
114	Analysis of Copy Number Abnormalities of 86 Acute Lymphoblastic Leukemia Cell Lines Based on the Genetic Subtypes. Blood, 2015, 126, 1424-1424.	0.6	6
115	Outcome of children with relapsed acute myeloid leukemia following initial therapy under the AML99 protocol. International Journal of Hematology, 2014, 100, 171-179.	0.7	31
116	RUNX1 mutation associated with clonal evolution in relapsed pediatric acute myeloid leukemia with t(16;21)(p11;q22). International Journal of Hematology, 2014, 99, 169-174.	0.7	9
117	ABL kinase mutation and relapse in 4 pediatric Philadelphia chromosome-positive acute lymphoblastic leukemia cases. International Journal of Hematology, 2014, 99, 609-615.	0.7	7
118	EVI1 overexpression is a poor prognostic factor in pediatric patients with mixed lineage leukemia-AF9 rearranged acute myeloid leukemia. Haematologica, 2014, 99, e225-e227.	1.7	35
119	Long-Term Parvovirus B19 Infections With Genetic Drift After Cord Blood Transplantation Complicated by Persistent CD4+ Lymphocytopenia. Journal of Pediatric Hematology/Oncology, 2014, 36, e65-e68.	0.3	5
120	Detection of RBM15-MKL1 fusion was useful for diagnosis and monitoring of minimal residual disease in infant acute megakaryoblastic leukemia. Acta Medica Okayama, 2014, 68, 119-23.	0.1	4
121	Outcome of Adolescent and Young Adults with Acute Myeloid Leukemia Treated with Pediatric Protocols: A Report from the 3 Japanese Cooperative Studies. Blood, 2014, 124, 374-374.	0.6	20
122	<i>IKZF1</i> and <i>CRLF2</i> gene alterations correlate with poor prognosis in Japanese <i>BCRâ€ABL1</i> â€negative highâ€risk Bâ€cell precursor acute lymphoblastic leukemia. Pediatric Blood and Cancer, 2013, 60, 1587-1592.	0.8	61
123	WT1 mutation in pediatric patients with acute myeloid leukemia: a report from the Japanese Childhood AML Cooperative Study Group. International Journal of Hematology, 2013, 98, 437-445.	0.7	16
124	<i>NUP98</i> â€ <i>NSD1</i> gene fusion and its related gene expression signature are strongly associated with a poor prognosis in pediatric acute myeloid leukemia. Genes Chromosomes and Cancer, 2013, 52, 683-693.	1.5	76
125	Appropriate dose reduction in induction therapy is essential for the treatment of infants with acute myeloid leukemia: a report from the Japanese Pediatric Leukemia/Lymphoma Study Group. International Journal of Hematology, 2013, 98, 578-588.	0.7	47
126	Cytomegalovirus Retinitis During Maintenance Therapy for T-Cell Acute Lymphoblastic Leukemia. Journal of Pediatric Hematology/Oncology, 2013, 35, 162-163.	0.3	21

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127	Excess treatment reduction including anthracyclines results in higher incidence of relapse in core binding factor acute myeloid leukemia in children. Leukemia, 2013, 27, 2413-2416.	3.3	52
128	Correlation of CYP2C19 Phenotype With Voriconazole Plasma Concentration in Children. Journal of Pediatric Hematology/Oncology, 2013, 35, e219-e223.	0.3	46
129	Whole Exome Sequencing Reveals Clonal Evolution Pattern and Driver Mutations Of Relapsed Pediatric AML. Blood, 2013, 122, 1410-1410.	0.6	1
130	Poor Prognosis With Different Induction Rate Was Observed In Children With Acute Myeloid Leukemia and FLT3-ITD According To The ITD/WT Allelic Ratio: A Result From The Japanese Pediatric Leukemia/Lymphoma Study Group. Blood, 2013, 122, 3891-3891.	0.6	0
131	Autoimmune-like hepatitis following unrelated BMT successfully treated with rituximab. Bone Marrow Transplantation, 2012, 47, 600-602.	1.3	9
132	Somatic mosaicism for oncogenic NRAS mutations in juvenile myelomonocytic leukemia. Blood, 2012, 120, 1485-1488.	0.6	27
133	Clinical characteristics and outcome of refractory/relapsed myeloid leukemia in children with Down syndrome. Blood, 2012, 120, 1810-1815.	0.6	46
134	Outcome in 146 patients with paediatric acute myeloid leukaemia treated according to the <scp>AML</scp> 99 protocol in the period 2003–06 from the <scp>J</scp> apan Association of Childhood Leukaemia Study. British Journal of Haematology, 2012, 159, 204-210.	1.2	24
135	High WT1 mRNA expression after induction chemotherapy and FLT3-ITD have prognostic impact in pediatric acute myeloid leukemia: a study of the Japanese Childhood AML Cooperative Study Group. International Journal of Hematology, 2012, 96, 469-476.	0.7	21
136	Incidence, clinical features, and risk factors of idiopathic pneumonia syndrome following hematopoietic stem cell transplantation in children. Pediatric Blood and Cancer, 2012, 58, 780-784.	0.8	31
137	Congenital pancreatoblastoma associated with βâ€catenin mutation. Pediatric Blood and Cancer, 2012, 58, 827-827.	0.8	7
138	Mutations profile of polycythemia vera and essential thrombocythemia among Japanese children. Pediatric Blood and Cancer, 2012, 59, 530-535.	0.8	18
139	RAS mutations are frequent in FAB type M4 and M5 of acute myeloid leukemia, and related to late relapse: a study of the Japanese Childhood AML Cooperative Study Group. International Journal of Hematology, 2012, 95, 509-515.	0.7	33
140	Childhood acute myeloid leukemia with bone marrow eosinophilia caused by t(16;21)(q24;q22). International Journal of Hematology, 2012, 95, 577-580.	0.7	9
141	Excellent outcome of allogeneic bone marrow transplantation for Fanconi anemia using fludarabine-based reduced-intensity conditioning regimen. International Journal of Hematology, 2012, 95, 675-679.	0.7	26
142	<i>DNMT3A</i> mutations are rare in childhood acute myeloid leukaemia, myelodysplastic syndromes and juvenile myelomonocytic leukaemia. British Journal of Haematology, 2012, 156, 413-414.	1.2	19
143	Molecular lesions in childhood and adult acute megakaryoblastic leukaemia. British Journal of Haematology, 2012, 156, 316-325.	1.2	18
144	De novo childhood myelodysplastic/myeloproliferative disease with unique molecular characteristics. British Journal of Haematology, 2012, 158, 129-137.	1.2	9

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145	Mutation in the <i><scp>THPO</scp></i> gene is not associated with aplastic anaemia in <scp>J</scp> apanese children. British Journal of Haematology, 2012, 158, 553-555.	1.2	3
146	Excess Reduction of Anthracyclines Results in Inferior Event-Free Survival in Core Binding Factor Acute Myeloid Leukemia in Children; A Report From the Japanese Pediatric Leukemia/Lymphoma Study Group (JPLSG). Blood, 2012, 120, 409-409.	0.6	1
147	Appropriate Dose Modification in Induction Therapy Is Essential for the Treatment of Infants with Acute Myeloid Leukemia; A Report From the Japanese Pediatric Leukemia/Lymphoma Study Group (JPLSG) Blood, 2012, 120, 2615-2615.	0.6	0
148	Effects of a Histone Deacetylase Inhibitor On Myeloid Leukemia Cell Line with Mixed-Lineage Leukemia Partial Tandem Duplication. Blood, 2012, 120, 4331-4331.	0.6	0
149	Attempts to Optimize Post-Induction Treatment in Childhood Acute Myeloid Leukemia without Core Binding Factors: A Report From the Japanese Pediatric Leukemia/Lymphoma Study Group (JPLSG). Blood, 2012, 120, 3545-3545.	0.6	1
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