MauriziÃ² AricÃ²

List of Publications by Year in descending order

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349 papers 26,281 citations

9786 73 h-index 153 g-index

351 all docs

351 docs citations

times ranked

351

17469 citing authors

#	Article	IF	Citations
1	HLHâ€2004: Diagnostic and therapeutic guidelines for hemophagocytic lymphohistiocytosis. Pediatric Blood and Cancer, 2007, 48, 124-131.	1.5	4,018
2	Contemporary classification of histiocytic disorders. Medical and Pediatric Oncology, 1997, 29, 157-166.	1.0	740
3	Treatment of hemophagocytic lymphohistiocytosis with HLH-94 immunochemotherapy and bone marrow transplantation. Blood, 2002, 100, 2367-2373.	1.4	737
4	Molecular response to treatment redefines all prognostic factors in children and adolescents with B-cell precursor acute lymphoblastic leukemia: results in 3184 patients of the AIEOP-BFM ALL 2000 study. Blood, 2010, 115, 3206-3214.	1.4	685
5	Outcome of Treatment in Children with Philadelphia Chromosome–Positive Acute Lymphoblastic Leukemia. New England Journal of Medicine, 2000, 342, 998-1006.	27.0	539
6	Chemoimmunotherapy for hemophagocytic lymphohistiocytosis: long-term results of the HLH-94 treatment protocol. Blood, 2011, 118, 4577-4584.	1.4	493
7	Langerhans cell histiocytosis in adultsReport from the International Registry of the Histiocyte Society. European Journal of Cancer, 2003, 39, 2341-2348.	2.8	450
8	A randomized trial of treatment for multisystem Langerhans' cell histiocytosis. Journal of Pediatrics, 2001, 138, 728-734.	1.8	427
9	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis: A European League Against Rheumatism/American College of Rheumatology/Paediatric Rheumatology International Trials Organisation Collaborative Initiative. Arthritis and Rheumatology. 2016. 68. 566-576.	5.6	427
10	Confirmed efficacy of etoposide and dexamethasone in HLH treatment: long-term results of the cooperative HLH-2004 study. Blood, 2017, 130, 2728-2738.	1.4	418
11	HLH-94: A treatment protocol for hemophagocytic lymphohistiocytosis. Medical and Pediatric Oncology, 1997, 28, 342-347.	1.0	417
12	Pulmonary Langerhans'-Cell Histiocytosis. New England Journal of Medicine, 2000, 342, 1969-1978.	27.0	411
13	Central Diabetes Insipidus in Children and Young Adults. New England Journal of Medicine, 2000, 343, 998-1007.	27.0	402
14	Late MRD response determines relapse risk overall and in subsets of childhood T-cell ALL: results of the AIEOP-BFM-ALL 2000 study. Blood, 2011, 118, 2077-2084.	1.4	370
15	NK-dependent DC maturation is mediated by TNFÎ \pm and IFNÎ 3 released upon engagement of the NKp30 triggering receptor. Blood, 2005, 106, 566-571.	1.4	365
16	Therapy prolongation improves outcome in multisystem Langerhans cell histiocytosis. Blood, 2013, 121, 5006-5014.	1.4	343
17	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Annals of the Rheumatic Diseases, 2016, 75, 481-489.	0.9	338
18	Hemophagocytic lymphohistiocytosis. Report of 122 children from the International Registry. FHL Study Group of the Histiocyte Society. Leukemia, 1996, 10, 197-203.	7.2	338

#	Article	IF	CITATIONS
19	Chronic myelomonocytic leukemia in childhood: a retrospective analysis of 110 cases. European Working Group on Myelodysplastic Syndromes in Childhood (EWOG-MDS). Blood, 1997, 89, 3534-43.	1.4	320
20	Improved outcome in multisystem Langerhans cell histiocytosis is associated with therapy intensification. Blood, 2008, 111, 2556-2562.	1.4	287
21	Imatinib after induction for treatment of children and adolescents with Philadelphia-chromosome-positive acute lymphoblastic leukaemia (EsPhALL): a randomised, open-label, intergroup study. Lancet Oncology, The, 2012, 13, 936-945.	10.7	282
22	Management of adult patients with Langerhans cell histiocytosis: recommendations from an expert panel on behalf of Euro-Histio-Net. Orphanet Journal of Rare Diseases, 2013, 8, 72.	2.7	281
23	Risk factors for diabetes insipidus in langerhans cell histiocytosis. Pediatric Blood and Cancer, 2006, 46, 228-233.	1.5	271
24	Genetic evidence for lineage-related and differentiation stage-related contribution of somatic PTPN11 mutations to leukemogenesis in childhood acute leukemia. Blood, 2004, 104, 307-313.	1.4	265
25	A prospective evaluation of degranulation assays in the rapid diagnosis of familial hemophagocytic syndromes. Blood, 2012, 119, 2754-2763.	1.4	263
26	Effectiveness of Immunoglobulin Replacement Therapy on Clinical Outcome in Patients with Primary Antibody Deficiencies: Results from a Multicenter Prospective Cohort Study. Journal of Clinical Immunology, 2011, 31, 315-322.	3.8	252
27	Risk of Relapse of Childhood Acute Lymphoblastic Leukemia Is Predicted By Flow Cytometric Measurement of Residual Disease on Day 15 Bone Marrow. Journal of Clinical Oncology, 2009, 27, 5168-5174.	1.6	247
28	FAMILIAL HEMOPHAGOCYTIC LYMPHOHISTIOCYTOSIS. Hematology/Oncology Clinics of North America, 1998, 12, 417-433.	2.2	241
29	Frequency and spectrum of central nervous system involvement in 193 children with haemophagocytic lymphohistiocytosis. British Journal of Haematology, 2008, 140, 327-335.	2.5	217
30	Dexamethasone vs prednisone in induction treatment of pediatric ALL: results of the randomized trial AIEOP-BFM ALL 2000. Blood, 2016, 127, 2101-2112.	1.4	208
31	Clinical Outcome of Children With Newly Diagnosed Philadelphia Chromosome–Positive Acute Lymphoblastic Leukemia Treated Between 1995 and 2005. Journal of Clinical Oncology, 2010, 28, 4755-4761.	1.6	203
32	Clinical relevance of BCL-2 overexpression in childhood acute lymphoblastic leukemia. Blood, 1996, 87, 1140-1146.	1.4	201
33	GIMEMA-AIEOPAIDA protocol for the treatment of newly diagnosed acute promyelocytic leukemia (APL) in children. Blood, 2005, 106, 447-453.	1.4	196
34	Pathogenesis of haemophagocytic lymphohistiocytosis. British Journal of Haematology, 2001, 114, 761-769.	2.5	189
35	Acute lymphoblastic leukemia in children with Down syndrome: a retrospective analysis from the Ponte di Legno study group. Blood, 2014, 123, 70-77.	1.4	189
36	RAS mutations and clonality analysis in children with juvenile myelomonocytic leukemia (JMML). Leukemia, 1999, 13, 32-37.	7.2	186

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37	A proportion of patients with lymphoma may harbor mutations of the perforin gene. Blood, 2005, 105, 4424-4428.	1.4	182
38	Long-Term Results of a Randomized Trial on Extended Use of High Dose l-Asparaginase for Standard Risk Childhood Acute Lymphoblastic Leukemia. Journal of Clinical Oncology, 2005, 23, 7161-7167.	1.6	180
39	Langerhans cell histiocytosis reveals a new IL-17A–dependent pathway of dendritic cell fusion. Nature Medicine, 2008, 14, 81-87.	30.7	180
40	SAP controls the cytolytic activity of CD8+ T cells against EBV-infected cells. Blood, 2005, 105, 4383-4389.	1.4	167
41	THE RELATION OF LANGERHANS CELL HISTIOCYTOSIS TO ACUTE LEUKEMIA, LYMPHOMAS, AND OTHER SOLID TUMORS. Hematology/Oncology Clinics of North America, 1998, 12, 369-378.	2.2	165
42	Analysis of natural killer–cell function in familial hemophagocytic lymphohistiocytosis (FHL): defective CD107a surface expression heralds Munc13-4 defect and discriminates between genetic subtypes of the disease. Blood, 2006, 108, 2316-2323.	1.4	161
43	CLINICAL ASPECTS OF LANGERHANS CELL HISTIOCYTOSIS. Hematology/Oncology Clinics of North America, 1998, 12, 247-258.	2.2	157
44	Philadelphia chromosome-positive (Ph+) childhood acute lymphoblastic leukemia: good initial steroid response allows early prediction of a favorable treatment outcome. Blood, 1998, 92, 2730-41.	1.4	156
45	Adult onset and atypical presentation of hemophagocytic lymphohistiocytosis in siblings carryingPRF1 mutations. Blood, 2002, 100, 2266-2266.	1.4	155
46	Hemophagocytic lymphohistiocytosis due to germline mutations inSH2D1A, the X-linked lymphoproliferative disease gene. Blood, 2001, 97, 1131-1133.	1.4	148
47	Long-term results of the Italian Association of Pediatric Hematology and Oncology (AIEOP) Studies 82, 87, 88, 91 and 95 for childhood acute lymphoblastic leukemia. Leukemia, 2010, 24, 255-264.	7.2	148
48	Myelodysplastic syndrome, juvenile myelomonocytic leukemia, and acute myeloid leukemia associated with complete or partial monosomy 7. Leukemia, 1999, 13, 376-385.	7.2	142
49	Prospective comparative study of bone marrow transplantation and postremission chemotherapy for childhood acute myelogenous leukemia. The Associazione Italiana Ematologia ed Oncologia Pediatrica Cooperative Group Journal of Clinical Oncology, 1993, 11, 1046-1054.	1.6	139
50	Genetic predisposition to hemophagocytic lymphohistiocytosis: Report on 500 patients from the Italian registry. Journal of Allergy and Clinical Immunology, 2016, 137, 188-196.e4.	2.9	139
51	Prolonged survival of B-lineage acute lymphoblastic leukemia cells is accompanied by overexpression of bcl-2 protein. Blood, 1993, 81, 1025-1031.	1.4	127
52	Genotype phenotype study of familial haemophagocytic lymphohistiocytosis due to perforin mutations. Journal of Medical Genetics, 2007, 45, 15-21.	3.2	118
53	Cladribine and cytarabine in refractory multisystem Langerhans cell histiocytosis: results of an international phase 2 study. Blood, 2015, 126, 1415-1423.	1.4	117
54	Familial clustering of Langerhans cell histiocytosis. British Journal of Haematology, 1999, 107, 883-888.	2.5	116

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55	LCH-I: A randomized trial of etoposide vs. vinblastine in disseminated langerhans cell histiocytosis. Medical and Pediatric Oncology, 1994, 23, 107-110.	1.0	111
56	Osteonecrosis: An emerging complication of intensive chemotherapy for childhood acute lymphoblastic leukemia. Haematologica, 2003, 88, 747-53.	3.5	97
57	Correlation between magnetic resonance imaging of posterior pituitary and neurohypophyseal function in children with diabetes insipidus. Journal of Clinical Endocrinology and Metabolism, 1992, 74, 795-800.	3.6	95
58	Multidrug resistant Pseudomonas aeruginosa infection in children undergoing chemotherapy and hematopoietic stem cell transplantation. Haematologica, 2010, 95, 1612-1615.	3.5	93
59	Familial Hemophagocytic Lymphohistiocytosis: When Rare Diseases Shed Light on Immune System Functioning. Frontiers in Immunology, 2014, 5, 167.	4.8	93
60	Long-term results of the Italian Association of Pediatric Hematology and Oncology (AIEOP) Acute Lymphoblastic Leukemia Studies, 1982–1995. Leukemia, 2000, 14, 2196-2204.	7.2	92
61	Improved outcome in high-risk childhood acute lymphoblastic leukemia defined by prednisone-poor response treated with double Berlin-Frankfurt-Muenster protocol II. Blood, 2002, 100, 420-426.	1.4	92
62	Extended intrathecal methotrexate may replace cranial irradiation for prevention of CNS relapse in children with intermediate-risk acute lymphoblastic leukemia treated with Berlin-Frankfurt-Münster-based intensive chemotherapy. The Associazione Italiana di Ematologia ed Oncologia Pediatrica Journal of Clinical Oncology, 1995, 13, 2497-2502.	1.6	91
63	Acute leukemia in association with langerhans cell histiocytosis. Medical and Pediatric Oncology, 1994, 23, 81-85.	1.0	90
64	Good steroid response in vivo predicts a favorable outcome in children with T-cell acute lymphoblastic leukemia. Cancer, 1995, 75, 1684-1693.	4.1	90
65	L-asparagine depletion and L-asparaginase activity in children with acute lymphoblastic leukemia receiving i.m. or i.v. Erwinia C. or E. coli L-asparaginase as first exposure. Annals of Oncology, 2000, 11, 189-193.	1.2	90
66	Reactivations in Multisystem Langerhans Cell Histiocytosis: Data of the International LCH Registry. Journal of Pediatrics, 2008, 153, 700-705.e2.	1.8	88
67	Human herpesvirus type 8 DNA sequences in biological samples of HIV-positive and negative individuals in Sicily. Aids, 1997, 11, 607-612.	2.2	87
68	Slp1 and Slp2â€a Localize to the Plasma Membrane of CTL and Contribute to Secretion from the Immunological Synapse. Traffic, 2008, 9, 446-457.	2.7	87
69	Secondary cytogenetic aberrations in childhood Philadelphia chromosome positive acute lymphoblastic leukemia are nonrandom and may be associated with outcome. Leukemia, 2004, 18, 693-702.	7.2	81
70	A single amino acid change, A91V, leads to conformational changes that can impair processing to the active form of perforin. Blood, 2005, 106, 932-937.	1.4	80
71	Somatic <i>PTPN11</i> mutations in childhood acute myeloid leukaemia. British Journal of Haematology, 2005, 129, 333-339.	2.5	78
72	Familial Hemophagocytic Lymphohistiocytosis May Present during Adulthood: Clinical and Genetic Features of a Small Series. PLoS ONE, 2012, 7, e44649.	2.5	77

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73	Hepatitis C virus infection in children treated for acute lymphoblastic leukemia. Blood, 1994, 84, 2919-2922.	1.4	76
74	Role of cranial radiotherapy for childhood T-cell acute lymphoblastic leukemia with high WBC count and good response to prednisone. Associazione Italiana Ematologia Oncologia Pediatrica and the Berlin-Frankfurt-Mýnster groups Journal of Clinical Oncology, 1997, 15, 2786-2791.	1.6	76
75	Genotype-phenotype study of familial haemophagocytic lymphohistiocytosis type 3. Journal of Medical Genetics, 2011, 48, 343-352.	3.2	76
76	Natural cytotoxicity impairment in familial haemophagocytic lymphohistiocytosis Archives of Disease in Childhood, 1988, 63, 292-296.	1.9	75
77	Langerhans cell histiocytosis in adults: more questions than answers?. European Journal of Cancer, 2004, 40, 1467-1473.	2.8	73
78	Novel Munc13-4 mutations in children and young adult patients with haemophagocytic lymphohistiocytosis. Journal of Medical Genetics, 2006, 43, 953-960.	3.2	71
79	Marriage and parenthood among childhood cancer survivors: a report from the Italian AIEOP Off-Therapy Registry. Haematologica, 2011, 96, 744-751.	3.5	71
80	Molecular basis of familial hemophagocytic lymphohistiocytosis. Haematologica, 2010, 95, 538-541.	3 . 5	70
81	Syntaxin binding mechanism and disease-causing mutations in Munc18-2. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, E4482-91.	7.1	70
82	Six novel mutations in the PRF1 gene in children with haemophagocytic lymphohistiocytosis. Journal of Medical Genetics, 2001, 38, 643-646.	3.2	69
83	Long-Term Results of the AIEOP-ALL-95 Trial for Childhood Acute Lymphoblastic Leukemia: Insight on the Prognostic Value of DNA Index in the Framework of Berlin-Frankfurt-Muenster–Based Chemotherapy. Journal of Clinical Oncology, 2008, 26, 283-289.	1.6	69
84	Risk factors for early death in children with haemophagocytic lymphohistiocytosis. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, 313-318.	1.5	69
85	Childhood high-risk acute lymphoblastic leukemia in first remission: results after chemotherapy or transplant from the AIEOP ALL 2000 study. Blood, 2014, 123, 1470-1478.	1.4	69
86	Predictive value of minimal residual disease in Philadelphia-chromosome-positive acute lymphoblastic leukemia treated with imatinib in the European intergroup study of post-induction treatment of Philadelphia-chromosome-positive acute lymphoblastic leukemia, based on immunoglobulin/T-cell receptor and BCR/ABL1 methodologies. Haematologica, 2018, 103, 107-115.	3 . 5	68
87	Germline mutations of the perforin gene are a frequent occurrence in childhood anaplastic large cell lymphoma. Cancer, 2007, 109, 2566-2571.	4.1	64
88	Treatment of pediatric hodgkin disease tailored to stage, mediastinal mass, and age an italian (aieop) multicenter study on 215 patients. Cancer, 1993, 72, 2049-2057.	4.1	63
89	Evolving pituitary hormone deficiency is associated with pituitary vasculopathy: dynamic MR study in children with hypopituitarism, diabetes insipidus, and Langerhans cell histiocytosis Radiology, 1994, 193, 493-499.	7.3	62
90	Hematopoietic stem cell transplantation for hemophagocytic lymphohistiocytosis: a retrospective analysis of data from the Italian Association of Pediatric Hematology Oncology (AIEOP). Haematologica, 2008, 93, 1694-1701.	3. 5	62

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91	Familial hemophagocytic lymphohistiocytosis: how late can the onset be?. Haematologica, 2001, 86, 499-503.	3.5	62
92	Adolescents with cancer in Italy: Entry into the national cooperative paediatric oncology group AIEOP trials. European Journal of Cancer, 2009, 45, 328-334.	2.8	61
93	Centriole polarisation to the immunological synapse directs secretion from cytolytic cells of both the innate and adaptive immune systems. BMC Biology, 2011, 9, 45.	3.8	60
94	Lack of clinically significant cardiac dysfunction during intermediate dobutamine doses in long-term childhood cancer survivors exposed to anthracyclines. American Heart Journal, 2000, 140, 315-323.	2.7	59
95	Dynamic Endocrine Testing and Magnetic Resonance Imaging in the Long Term Follow-Up of Childhood Langerhans Cell Histiocytosis. Journal of Clinical Endocrinology and Metabolism, 1998, 83, 3089-3094.	3.6	58
96	Haemophagocytic lymphohistiocytosis: proposal of a diagnostic algorithm based on perforin expression. British Journal of Haematology, 2002, 119, 180-188.	2.5	58
97	Prophylactic co-trimoxazole versus norfloxacin in neutropenic children — perspective randomized study. Infection, 1989, 17, 65-69.	4.7	57
98	Expert consensus on dynamics of laboratory tests for diagnosis of macrophage activation syndrome complicating systemic juvenile idiopathic arthritis. RMD Open, 2016, 2, e000161.	3.8	57
99	Effect of Protracted High-Dose I-Asparaginase Given as a Second Exposure in a Berlin-Frankfurt-Münster–Based Treatment: Results of the Randomized 9102 Intermediate-Risk Childhood Acute Lymphoblastic Leukemia Study—A Report From the Associazione Italiana Ematologia Oncologia Pediatrica, lournal of Clinical Oncology, 2001, 19, 1297-1303.	1.6	54
100	Cytoreduction and prognosis in childhood acute lymphoblastic leukemia Journal of Clinical Oncology, 1996, 14, 2403-2406.	1.6	53
101	Juvenile Myelomonocytic Leukemia. Blood, 1997, 90, 479-488.	1.4	52
102	Correlation between magnetic resonance imaging of posterior pituitary and neurohypophyseal function in children with diabetes insipidus. Journal of Clinical Endocrinology and Metabolism, 1992, 74, 795-800.	3.6	52
103	Human Immunodeficiency Virus–Related Cancer in Children: Incidence and Treatment Outcome—Report of the Italian Register. Journal of Clinical Oncology, 2000, 18, 3854-3861.	1.6	51
104	Acute lymphoblastic leukemia and Down syndrome. Cancer, 2008, 113, 515-521.	4.1	51
105	Malignancies in children with human immunodeficiency virus type 1 infection. Cancer, 1991, 68, 2473-2477.	4.1	50
106	Development and Initial Validation of the Macrophage Activation Syndrome/Primary Hemophagocytic Lymphohistiocytosis Score, a Diagnostic Tool that Differentiates Primary Hemophagocytic Lymphohistiocytosis from Macrophage Activation Syndrome. Journal of Pediatrics, 2017, 189, 72-78.e3.	1.8	50
107	Single-Day Trimethoprim/Sulfamethoxazole Prophylaxis for Pneumocystis Pneumonia in Children with Cancer. Journal of Pediatrics, 2014, 164, 389-392.e1.	1.8	49
108	cAMP response element binding protein (CREB) overexpression CREB has been described as critical for leukemia progression. Haematologica, 2007, 92, 1435-1437.	3. 5	48

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109	STXBP2 mutations in children with familial haemophagocytic lymphohistiocytosis type 5. Journal of Medical Genetics, 2010, 47, 595-600.	3.2	48
110	Minimal Disseminated Disease in High-Risk Burkitt's Lymphoma Identifies Patients With Different Prognosis. Journal of Clinical Oncology, 2011, 29, 1779-1784.	1.6	48
111	Plasma Cell-Free DNA in Paediatric Lymphomas. Journal of Cancer, 2013, 4, 323-329.	2.5	48
112	A single amino acid change A91V in perforin: a novel, frequent predisposing factor to childhood acute lymphoblastic leukemia?. Haematologica, 2005, 90, 697-8.	3.5	48
113	GHRH Plus Arginine in the Diagnosis of Acquired GH Deficiency of Childhood-Onset. Journal of Clinical Endocrinology and Metabolism, 2002, 87, 2740-2744.	3.6	47
114	Hepatitis C virus (HCV) core serotypes in chronic HCV infection. Journal of Clinical Microbiology, 1994, 32, 2523-2527.	3.9	47
115	Langerhans cell histiocytosis in children: from the bench to bedside for an updated therapy. British Journal of Haematology, 2016, 173, 663-670.	2.5	46
116	Evolution of childhood central diabetes insipidus into panhypopituitarism with a large hypothalamic mass: is 'lymphocytic infundibuloneurohypophysitis' in children a different entity?. European Journal of Endocrinology, 1998, 139, 635-640.	3.7	44
117	Juvenile myelomonocytic leukemia. Blood, 1997, 90, 479-88.	1.4	43
118	Blood spotlight on Langerhans cell histiocytosis. Blood, 2014, 124, 867-872.	1.4	41
119	MR of the hypothalamic-pituitary axis in Langerhans cell histiocytosis. American Journal of Neuroradiology, 1992, 13, 1365-71.	2.4	41
120	Autologous bone marrow transplantation for treatment of isolated central nervous system relapse of childhood acute lymphoblastic leukemia. Bone Marrow Transplantation, 1998, 21, 9-14.	2.4	40
121	A prospective, randomized study of empirical antifungal therapy for the treatment of chemotherapyâ€induced febrile neutropenia in children. British Journal of Haematology, 2012, 158, 249-255.	2.5	40
122	Familial hemophagocytic lymphohistiocytosis: a model for understanding the human machinery of cellular cytotoxicity. Cellular and Molecular Life Sciences, 2012, 69, 29-40.	5.4	40
123	Patients with Griscelli syndrome and normal pigmentation identify RAB27A mutations that selectively disrupt MUNC13-4 binding. Journal of Allergy and Clinical Immunology, 2015, 135, 1310-1318.e1.	2.9	40
124	Outcomes of Children with Hemophagocytic Lymphohistiocytosis Given Allogeneic Hematopoietic Stem Cell Transplantation in Italy. Biology of Blood and Marrow Transplantation, 2018, 24, 1223-1231.	2.0	39
125	Morbidity of pandemic H1N1 influenza in children with cancer. Pediatric Blood and Cancer, 2010, 55, 226-228.	1.5	38
126	Cytogenetic abnormalities in PHA-stimulated lymphocytes from patients with Langerhans cell histiocytosis. British Journal of Haematology, 2000, 111, 258-262.	2.5	38

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127	Allogeneic bone marrow transplantation versus chemotherapy in high-risk childhood acute lymphoblastic leukaemia in first remission. British Journal of Haematology, 1997, 96, 387-394.	2.5	37
128	Detection of prognostic factors in children and adolescents with Burkitt and Diffuse Large B ell Lymphoma treated with the <scp>AIEOP LNH</scp> â€97 protocol. British Journal of Haematology, 2016, 175, 467-475.	2.5	37
129	Intensive BFM chemotherapy for childhood ALL: interim analysis of the AIEOP-ALL 91 study. Associazione Italiana Ematologia Oncologia Pediatrica. Haematologica, 1998, 83, 791-9.	3.5	37
130	Successful treatment of Griscelli syndrome with unrelated donor allogeneic hematopoietic stem cell transplantation. Bone Marrow Transplantation, 2002, 29, 995-998.	2.4	36
131	Incidence of colonization and bloodstream infection with carbapenem-resistant <i>Enterobacteriaceae</i> in children receiving antineoplastic chemotherapy in Italy. Infectious Diseases, 2016, 48, 152-155.	2.8	36
132	Treatment of isolated testicular relapse in childhood acute lymphoblastic leukemia: an Italian multicenter study. Associazione Italiana Ematologia ed Oncologia Pediatrica Journal of Clinical Oncology, 1990, 8, 672-677.	1.6	34
133	Prospective molecular monitoring of BCR/ABL transcript in children with Ph+ acute lymphoblastic leukaemia unravels differences in treatment response. British Journal of Haematology, 2002, 119, 445-453.	2.5	34
134	Mutations affecting mRNA splicing are the most common molecular defect in patients with familial hemophagocytic lymphohistiocytosis type 3. Haematologica, 2008, 93, 1086-1090.	3.5	34
135	The Quality of Life of Children and Adolescents with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2009, 29, 501-507.	3.8	34
136	Stem cell transplantation for children with hemophagocytic lymphohistiocytosis: results from the HLH-2004 study. Blood Advances, 2020, 4, 3754-3766.	5.2	34
137	Long-term pulmonary sequelae after treatment of childhood Hodgkin's disease. Annals of Oncology, 1997, 8, S19-S24.	1.2	33
138	Risk of Seizures in Children Receiving Busulphan-Containing Regimens for Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2014, 20, 282-285.	2.0	33
139	Severe acute encephalopathy following inadvertent intrathecal doxorubicin administration. Medical and Pediatric Oncology, 1990, 18, 261-263.	1.0	32
140	Neuroradiologic findings and followâ€up with magnetic resonance imaging of the genetic forms of haemophagocytic lymphohistiocytosis with CNS involvement. Pediatric Blood and Cancer, 2012, 58, 810-814.	1.5	32
141	No evidence of SARSâ€CoVâ€2 infection by polymerase chain reaction or serology in children with pseudoâ€chilblain. British Journal of Dermatology, 2020, 183, 784-785.	1.5	32
142	Insulin-Like Growth Factor I (IGF-I) and IGF-Binding Protein 3 Response to Growth Hormone Is Impaired in HIV-Infected Children. AIDS Research and Human Retroviruses, 2002, 18, 331-339.	1.1	31
143	Specific polymorphisms of cytokine genes are associated with different risks to develop single-system or multi-system childhood Langerhans cell histiocytosis. British Journal of Haematology, 2006, 132, 784-787.	2.5	31
144	Cyclosporine therapy for refractory langerhans cell histiocytosis. Medical and Pediatric Oncology, 1995, 25, 12-16.	1.0	30

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145	Langerhans cell histiocytosis in two generations: A new family and review of the literature. Medical and Pediatric Oncology, 2001, 36, 314-316.	1.0	30
146	Upper Age Limits for Accessing Pediatric Oncology Centers in Italy: A Barrier Preventing Adolescents with Cancer from Entering National Cooperative AIEOP Trials. Pediatric Hematology and Oncology, 2012, 29, 55-61.	0.8	30
147	Dexamethasone in Induction Can Eliminate One Third of All Relapses in Childhood Acute Lymphoblastic Leukemia (ALL): Results of An International Randomized Trial in 3655 Patients (Trial AIEOP-BFM ALL) Tj ETQq1 1	0.718 4 314	rg Bō /Overlo
148	Langerhans' cell histiocytosis: is there a role for genetics?. Haematologica, 2001, 86, 1009-14.	3.5	30
149	Langerhans cell histiocytosis and acute leukemia: Unusual association in two cases. Medical and Pediatric Oncology, 1993, 21, 271-273.	1.0	28
150	Echocardiographic evaluation of patients cured of childhood cancer: A single center study of 117 subjects who received anthracyclines. Medical and Pediatric Oncology, 2001, 36, 593-600.	1.0	28
151	Primary cutaneous plasmacytosis in a child. Is this a new entity?. Journal of the European Academy of Dermatology and Venereology, 2002, 16, 164-167.	2.4	28
152	Postinduction Minimal Residual Disease Monitoring by Polymerase Chain Reaction in Children With Acute Lymphoblastic Leukemia. Journal of Clinical Oncology, 2014, 32, 3553-3558.	1.6	28
153	Primary Sacral Bone Tumours in Children (Report of 16 cases with a short literature review). Journal of Medical Imaging and Radiation Oncology, 1990, 34, 142-149.	0.6	27
154	A91V perforin variation in healthy subjects and FHLH patients. International Journal of Immunogenetics, 2006, 33, 123-125.	1.8	27
155	Munc18â€2 is required for Syntaxin 11 Localization on the Plasma Membrane inÂCytotoxic Tâ€Lymphocytes. Traffic, 2015, 16, 1330-1341.	2.7	27
156	Dup(3)(p2â†'pter) in two families, including one infant with cyclopia. American Journal of Medical Genetics Part A, 1985, 20, 341-348.	2.4	26
157	2019-nCoV: Polite with Children!. Mental Illness, 2020, 12, 8495.	0.8	26
158	Detection of PICALM-MLLT10 (CALM-AF10) and outcome in children with T-lineage acute lymphoblastic leukemia. Leukemia, 2013, 27, 2419-2421.	7.2	25
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