MauriziÃ² AricÃ²

List of Publications by Year in descending order

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Version: 2024-02-01

349 papers 26,281 citations

73 h-index

9784

153 g-index

351 all docs

 $\begin{array}{c} 351 \\ \text{docs citations} \end{array}$

times ranked

351

17469 citing authors

#	Article	IF	Citations
1	Limited Additive Diagnostic Impact of Isolated Gastrointestinal Involvement for the Triage of Children with Suspected COVID-19. Children, 2022, 9, 41.	1.5	O
2	STOP Pain Projectâ€"Opioid Response in Pediatric Cancer Patients and Gene Polymorphisms of Cytokine Pathways. Pharmaceutics, 2022, 14, 619.	4.5	2
3	Diagnosis of HLH: two siblings, two distinct genetic causes. Clinical and Experimental Immunology, 2022, 207, 205-207.	2.6	1
4	Hypereosinophilia, Abdominal Pain, and Diarrhea: Anisakiasis, a Difficult Diagnosis in a Child. Pediatric Infectious Disease Journal, 2022, 41, e343-e344.	2.0	0
5	Acute, Severe Hepatitis of Unknown Origin: Should We Really Be Afraid of Another Obscure Enemy of Our Children?. Pediatric Reports, 2022, 14, 217-219.	1.3	7
6	SARS-CoV-2 Vaccination for Children—An Open Issue. Pediatric Reports, 2021, 13, 95-97.	1.3	3
7	Effectiveness of Preventive Measures in Keeping Low Prevalence of SARS-CoV-2 Infection in Health Care Workers in a Referral Children's Hospital in Southern Italy. Pediatric Reports, 2021, 13, 118-124.	1.3	4
8	Clinical features, diagnosis and therapy of familial haemophagocytic lymphohistiocytosis. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 2723-2728.	1.5	4
9	Imaging Appropriateness in Pediatric Radiology during COVID-19 Pandemic: A Retrospective Comparison with No COVID-19 Period. Children, 2021, 8, 463.	1.5	10
10	COVID-19 and Inherited Metabolic Disorders: One-Year Experience of a Referral Center. Children, 2021, 8, 781.	1.5	9
11	Additive Prognostic Impact of Gastrointestinal Involvement in Severe Multisystem Langerhans Cell Histiocytosis. Journal of Pediatrics, 2021, 237, 65-70.e3.	1.8	8
12	Neutralizing Anti-IL-17A Antibody Demonstrates Preclinical Activity Enhanced by Vinblastine in Langerhans Cell Histiocytosis. Frontiers in Oncology, 2021, 11, 780191.	2.8	1
13	High levels of plasma interleukin-17A are associated with severe neurological sequelae in Langerhans cell histiocytosis. Cytokine, 2020, 126, 154877.	3.2	10
14	Comparison of three different ELISAs for the detection of recombinant, native and plasma IL-17A. MethodsX, 2020, 7, 100997.	1.6	0
15	Covid-19 pandemic and pediatric healthcare policy in Italy: time for a change. Mental Illness, 2020, 12, 8823.	0.8	4
16	Introducing Pediatric Reports: Only Another Pediatric Journal?. Pediatric Reports, 2020, 12, 124-126.	1.3	0
17	Response to â€No evidence of SARSâ€CoVâ€2 infection by polymerase chain reaction or serology in children with pseudoâ€chilblain'. Reply from the authors. British Journal of Dermatology, 2020, 183, 1156-1157.	1.5	2
18	Stem cell transplantation for children with hemophagocytic lymphohistiocytosis: results from the HLH-2004 study. Blood Advances, 2020, 4, 3754-3766.	5.2	34

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19	SARS-CoV-2 Infection in Children in Southern Italy: A Descriptive Case Series. International Journal of Environmental Research and Public Health, 2020, 17, 6080.	2.6	12
20	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
21	2019-nCoV: Polite with Children!. Mental Illness, 2020, 12, 8495.	0.8	26
22	Childhood Histiocytoses., 2020,, 162-168.		0
23	Opioid response in paediatric cancer patients and the Val158Met polymorphism of the human catechol-O-methyltransferase (COMT) gene: an Italian study on 87 cancer children and a systematic review. BMC Cancer, 2019, 19, 113.	2.6	9
24	2B4 dysfunction in XLP1 NK cells: More than inability to control EBV infection. Clinical Immunology, 2019, 204, 31-36.	3.2	11
25	Long-term results of the AIEOP MH'96 childhood Hodgkin's lymphoma trial and focus on significance of response to chemotherapy and its implication in low risk patients to avoid radiotherapy. Leukemia and Lymphoma, 2018, 59, 2612-2621.	1.3	4
26	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
27	Primary mediastinal large Bâ€cell lymphoma: Outcome of a series of pediatric patients treated with highâ€dose methotrexate and cytarabine plus antiâ€CD20. Pediatric Blood and Cancer, 2018, 65, e26855.	1.5	10
28	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
29	A systematic review of the risk factors for clinical response to opioids for all-age patients with cancer-related pain and presentation of the paediatric STOP pain study. BMC Cancer, 2018, 18, 568.	2.6	10
30	Inhibitory 2B4 contributes to NK cell education and immunological derangements in XLP1 patients. European Journal of Immunology, 2017, 47, 1051-1061.	2.9	15
31	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
32	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
33	Biosimilars in the management of neutropenia: focus on filgrastim. Biologics: Targets and Therapy, 2016, 10, 17.	3.2	8
34	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
35	Road Traffic Pollution and Childhood Leukemia: A Nationwide Case-control Study in Italy. Archives of Medical Research, 2016, 47, 694-705.	3.3	10
36	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

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37	Timely follow-up of a GATA2 deficiency patient allows successful treatment. Journal of Allergy and Clinical Immunology, 2016, 138, 1480-1483.e4.	2.9	7
38	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
39	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
40	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
41	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Annals of the Rheumatic Diseases, 2016, 75, 481-489.	0.9	338
42	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
43	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
44	Reduced Intensity Delayed Intensification in Standard-Risk Patients Defined By Minimal Residual Disease in Childhood Acute Lymphoblastic Leukemia: Results of an International Randomized Trial in 1164 Patients (Trial AIEOP-BFM ALL 2000). Blood, 2016, 128, 4-4.	1.4	6
45	<i>CRLF2</i> over-expression is a poor prognostic marker in children with high risk T-cell acute lymphoblastic leukemia. Oncotarget, 2016, 7, 59260-59272.	1.8	24
46	Munc18â€2 is required for Syntaxin 11 Localization on the Plasma Membrane inÂCytotoxic T‣ymphocytes. Traffic, 2015, 16, 1330-1341.	2.7	27
47	Cladribine and cytarabine in refractory multisystem Langerhans cell histiocytosis: results of an international phase 2 study. Blood, 2015, 126, 1415-1423.	1.4	117
48	Non-Hodgkin lymphoma in children with an associated inherited condition: A retrospective analysis of the Associazione Italiana Ematologia Oncologia Pediatrica (AIEOP). Pediatric Blood and Cancer, 2015, 62, 1782-1789.	1.5	15
49	Lack of bone lesions at diagnosis is associated with inferior outcome in multisystem langerhans cell histiocytosis of childhood. British Journal of Haematology, 2015, 169, 241-248.	2.5	25
50	Longâ€ŧerm results of the AIEOP LNHâ€97 protocol for childhood lymphoblastic lymphoma. Pediatric Blood and Cancer, 2015, 62, 1388-1394.	1.5	25
51	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
52	Human monocyte-derived dendritic cells turn into foamy dendritic cells with IL-17A. Journal of Lipid Research, 2015, 56, 1110-1122.	4.2	21
53	Early Diagnosis and Monitoring of Neurodegenerative Langerhans Cell Histiocytosis. PLoS ONE, 2015, 10, e0131635.	2.5	13
54	Hematopoietic Stem Cell Transplantation for Hemophagocytic Lymphohistiocitosis: A National Retrospective Analysis of Data from the Italian Association of Pediatric Hematology Oncology (AIEOP). Blood, 2015, 126, 621-621.	1.4	0

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55	Blood spotlight on Langerhans cell histiocytosis. Blood, 2014, 124, 867-872.	1.4	41
56	Mutations of familial hemophagocytic lymphohistiocytosis (FHL) related genes and abnormalities of cytotoxicity function tests in patients with macrophage activation syndrome (MAS) occurring in systemic juvenile idiopathic arthritis (sJIA). Pediatric Rheumatology, 2014, 12, .	2.1	21
57	Highâ€dose chemotherapy with autologous stem cell rescue for treatment of retinoblastoma: Report of five cases. Pediatric Transplantation, 2014, 18, 631-636.	1.0	4
58	Mediastinal Burkitt lymphoma in childhood. Pediatric Blood and Cancer, 2014, 61, 2127-2128.	1.5	4
59	Development of new classification criteria for macrophage activation syndrome complicating systemic juvenile idiopathic arthritis. Pediatric Rheumatology, 2014, 12, .	2.1	6
60	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
61	Familial Hemophagocytic Lymphohistiocytosis: When Rare Diseases Shed Light on Immune System Functioning. Frontiers in Immunology, 2014, 5, 167.	4.8	93
62	Single-Day Trimethoprim/Sulfamethoxazole Prophylaxis for Pneumocystis Pneumonia in Children with Cancer. Journal of Pediatrics, 2014, 164, 389-392.e1.	1.8	49
63	<scp>XLP</scp> 1 inhibitory effect by 2 <scp>B</scp> 4 does not affect <scp>DNAM</scp> â€1 and <scp>NKG</scp> 2 <scp>D</scp> activating pathways in <scp>NK</scp> cells. European Journal of Immunology, 2014, 44, 1526-1534.	2.9	20
64	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
65	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
66	Diagnosing XLP1 in patients with hemophagocytic lymphohistiocytosis. Journal of Allergy and Clinical Immunology, 2014, 134, 1381-1387.e7.	2.9	14
67	Cytophagic histiocytic panniculitis, hemophagocytic lymphohistiocytosis and undetermined autoimmune disorder: reconciling the puzzle. Italian Journal of Pediatrics, 2014, 40, 17.	2.6	12
68	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
69	Detection of IL-17A-producing peripheral blood monocytes in Langerhans cell histiocytosis patients. Clinical Immunology, 2014, 153, 112-122.	3.2	24
70	Familial Hemophagocytic Lymphohistiocytosis Type 3 Diagnosed at School Age. Journal of Pediatric Hematology/Oncology, 2014, 36, e128-e130.	0.6	6
71	Monoallelic Mutations of the Perforin Gene may Represent a Predisposing Factor to Childhood Anaplastic Large Cell Lymphoma. Journal of Pediatric Hematology/Oncology, 2014, 36, e359-e365.	0.6	22
72	Outcome of Early T-Cell Precursor Acute Lymphoblastic Leukemia in AIEOP Patients Treated with the AIEOP-BFM ALL 2000 Study. Blood, 2014, 124, 3780-3780.	1.4	1

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73	Comparison of MRD Monitoring By IG/TR Rearrangement and BCR/ABL1 Transcript in Ph+ Pediatric Acute Lymphoblastic Leukemia Patients Treated with Imatinib in the Esphall Study. Blood, 2014, 124, 3782-3782.	1.4	O
74	PReS-FINAL-2186: Monoallelic mutations of familial hlh-related genes associated to macrophage activation syndrome. Pediatric Rheumatology, 2013, 11 , .	2.1	1
75	Detection of PICALM-MLLT10 (CALM-AF10) and outcome in children with T-lineage acute lymphoblastic leukemia. Leukemia, 2013, 27, 2419-2421.	7.2	25
76	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
77	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
78	Mutation of <i>FAS</i> , <i>XIAP</i> , and <i>UNC13D</i> Genes in a Patient With a Complex Lymphoproliferative Phenotype. Pediatrics, 2013, 132, e1052-e1058.	2.1	16
79	Therapy prolongation improves outcome in multisystem Langerhans cell histiocytosis. Blood, 2013, 121, 5006-5014.	1.4	343
80	Adenovirus pneumonia during induction therapy for childhood acute lymphoblastic leukemia. Pediatric Blood and Cancer, 2013, 60, 1390-1391.	1.5	4
81	Expression of \hat{l}^2 -adrenergic receptors in pediatric malignant brain tumors. Oncology Letters, 2013, 5, 221-225.	1.8	14
82	Variations of the UNC13D Gene in Patients with Autoimmune Lymphoproliferative Syndrome. PLoS ONE, 2013, 8, e68045.	2.5	20
83	Plasma Cell-Free DNA in Paediatric Lymphomas. Journal of Cancer, 2013, 4, 323-329.	2.5	48
84	Targeting BCL2 Family in Human Myeloid Dendritic Cells: A Challenge to Cure Diseases with Chronic Inflammations Associated with Bone Loss. Clinical and Developmental Immunology, 2013, 2013, 1-11.	3.3	7
85	Chemoresistance of Human Monocyte-Derived Dendritic Cells Is Regulated by IL-17A. PLoS ONE, 2013, 8, e56865.	2.5	22
86	Minimal Residual Disease Monitoring In Acute Lymphoblastic Leukemia Patients Using Immune Receptor Sequencing. Blood, 2013, 122, 1381-1381.	1.4	0
87	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
88	Deregulation of Ion Channel and Transporter Encoding Genes in Pediatric Gliomas. Frontiers in Oncology, 2012, 2, 53.	2.8	7
89	Pediatric Primary Anaplastic Ganglioglioma: A Case Report and Review of the Literature. Pediatric Neurosurgery, 2012, 48, 35-41.	0.7	13
90	Acute Encephalopathy in the Immune-compromised Child. Journal of Pediatric Hematology/Oncology, 2012, 34, 383-386.	0.6	17

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91	Simultaneous Diagnosis of Acute Lymphoblastic Leukemia and Peripheral Neuroblastic Tumor in a Child. Journal of Pediatric Hematology/Oncology, 2012, 34, 72-75.	0.6	1
92	Successful treatment with a low-dose cisplatin–etoposide regimen for patients with diencephalic syndrome. Journal of Neuro-Oncology, 2012, 109, 375-383.	2.9	10
93	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
94	Glioneuronal Tumor with Neuropil-Like Islands: Clinical, Morphologic, Immunohistochemical, and Molecular Features of Three Pediatric Cases. Pediatric and Developmental Pathology, 2012, 15, 352-360.	1.0	7
95	A prospective evaluation of degranulation assays in the rapid diagnosis of familial hemophagocytic syndromes. Blood, 2012, 119, 2754-2763.	1.4	263
96	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
97	Cord blood hematopoietic stem cell transplantation in an adolescent with haemophilia. Haemophilia, 2012, 18, e48-9.	2.1	6
98	Risk factors for early death in children with haemophagocytic lymphohistiocytosis. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, 313-318.	1.5	69
99	Reflectance mode confocal microscopy and digital image analysis in naevus of Hori and pathogenetic evaluation. British Journal of Dermatology, 2012, 167, 692-694.	1.5	2
100	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
101	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
102	Familial Hemophagocytic Lymphohistiocytosis May Present during Adulthood: Clinical and Genetic Features of a Small Series. PLoS ONE, 2012, 7, e44649.	2.5	77
103	Role of MRD in Ph+ Pediatric Acute Lymphoblastic Leukemia Patients Treated with and without Tirosine Kinase Inhibitor in the Esphall Study. Blood, 2012, 120, 1467-1467.	1.4	0
104	Chemoimmunotherapy for hemophagocytic lymphohistiocytosis: long-term results of the HLH-94 treatment protocol. Blood, 2011, 118, 4577-4584.	1.4	493
105	Genotype-phenotype study of familial haemophagocytic lymphohistiocytosis type 3. Journal of Medical Genetics, 2011, 48, 343-352.	3.2	76
106	Multidrug resistant Pseudomonas aeruginosa bloodstream infection: comparison between children and adult patients with cancer (reply). Haematologica, 2011, 96, e4-e4.	3.5	1
107	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
108	Primary Cerebral Myxofibrosarcoma. Journal of Pediatric Hematology/Oncology, 2011, 33, e279-e283.	0.6	9

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109	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
110	Detection of doxorubicin hydrochloride accumulation in the rat brain after morphine treatment by mass spectrometry. Cancer Chemotherapy and Pharmacology, 2011, 67, 1333-1340.	2.3	20
111	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
112	Foreign children with cancer in Italy. Italian Journal of Pediatrics, 2011, 37, 44.	2.6	8
113	Longâ€ŧerm results of AIEOPâ€8805 protocol for acute B ell lymphoblastic leukemia of childhood. Pediatric Blood and Cancer, 2011, 56, 544-550.	1.5	3
114	Marriage and parenthood among childhood cancer survivors: a report from the Italian AIEOP Off-Therapy Registry. Haematologica, 2011, 96, 744-751.	3.5	71
115	Metastatic Spread Outside the Central Nervous System of Anaplastic Medulloblastoma Associated with a Spinal-Peritoneal Shunt Immediately after Radiotherapy. Pediatric Neurosurgery, 2011, 47, 235-237.	0.7	2
116	Preventing Transmission of Infectious Agents in the Pediatric in-Patients Hematology-Oncology Setting: What Is the Role for Nonpharmacological Prophylaxis?. Mental Illness, 2011, 3, e9.	0.8	9
117	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
118	Partial Rescue of Biochemical Parameters After Hematopoietic Stem Cell Transplantation in a Patient with Prolidase Deficiency Due to Two Novel PEPD Mutations. JIMD Reports, 2011, 3, 71-77.	1.5	11
119	Acute Lymphoblastic Leukemia in Children with Down Syndrome: A Report From the Ponte Di Legno Study Group,. Blood, 2011, 118, 3579-3579.	1.4	8
120	Efficacy and Safety of Imatinib on Top of BFM-Like Chemotherapy in Pediatric Patients with Ph+/BCR-ABL+ Acute Lymphoblastic Leukemia (Ph+ALL). the EsPhALL Study. Blood, 2011, 118, 873-873.	1.4	3
121	Familial hemophagocytic lymphohistiocytosis: clinical and neuroradiological findings and review of the literature. Child's Nervous System, 2010, 26, 121-127.	1.1	20
122	Morbidity of pandemic H1N1 influenza in children with cancer. Pediatric Blood and Cancer, 2010, 55, 226-228.	1.5	38
123	The prevalence of the obesity in patients with moderate to severe psoriasis in Sicily populations. Journal of the European Academy of Dermatology and Venereology, 2010, 24, 92-93.	2.4	14
124	correspondence: A novel assay for investigation of suspected familial haemophagocytic lymphohistiocytosis. British Journal of Haematology, 2010, 150, 727-730.	2.5	20
125	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
126	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

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127	STXBP2 mutations in children with familial haemophagocytic lymphohistiocytosis type 5. Journal of Medical Genetics, 2010, 47, 595-600.	3.2	48
128	Multidrug resistant Pseudomonas aeruginosa infection in children undergoing chemotherapy and hematopoietic stem cell transplantation. Haematologica, 2010, 95, 1612-1615.	3.5	93
129	Molecular basis of familial hemophagocytic lymphohistiocytosis. Haematologica, 2010, 95, 538-541.	3.5	70
130	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
131	Key Role of Staff Competencies for Patient and Donor Safety in a Bone Marrow Transplantation Unit: Design and Implementation of an Accredited Training and Self-Assessment Program. Transplantation Proceedings, 2010, 42, 2254-2256.	0.6	1
132	Juvenile Myelomonocytic Leukemia. Paediatric Drugs, 2010, 12, 11-21.	3.1	23
133	Respiratory depression and somnolence in children receiving dimethylsulfoxide and morphine during hematopoietic stem cells transplantation. Haematologica, 2009, 94, 152-153.	3.5	12
134	HYPEREOSINOPHILIC SYNDROME IN CHILDHOOD: Clinical and Molecular Features of Two Cases. Pediatric Hematology and Oncology, 2009, 26, 129-135.	0.8	6
135	A Rare Presentation of Mycosis Fungoides Mimicking Psoriasis Vulgaris. Case Reports in Dermatology, 2009, 1, 60-65.	0.8	12
136	Promoter methylation and expression analysis of MGMT in advanced pediatric brain tumors. Oncology Reports, 2009, 22, 773-9.	2.6	13
137	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
138	The Quality of Life of Children and Adolescents with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2009, 29, 501-507.	3.8	34
139	Monotherapy with thalidomide for treatment of spinal cord hemangioblastomas in a patient with von Hippel–Lindau disease. Pediatric Blood and Cancer, 2009, 53, 464-467.	1.5	22
140	Altered mRNA expression of <i>PAX5</i> is a common event in acute lymphoblastic leukaemia. British Journal of Haematology, 2009, 146, 686-689.	2.5	10
141	Reply to: "Interleukin-17A is not expressed by CD207+ cells in Langerhans cell histiocytosis lesions". Nature Medicine, 2009, 15, 484-485.	30.7	6
142	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
143	The use of B-type natriuretic peptide in paediatric patients: a review of literature. Journal of Cardiovascular Medicine, 2009, 10, 298-302.	1.5	15
144	Acute lymphoblastic leukemia and Down syndrome. Cancer, 2008, 113, 515-521.	4.1	51

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145	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
146	The role of BMT in childhood histiocytoses. Bone Marrow Transplantation, 2008, 41, S8-S13.	2.4	19
147	Continuous intravenous infusion of lorazepam as seizure prophylaxis in children treated with high-dose busulfan. Bone Marrow Transplantation, 2008, 42, 135-136.	2.4	10
148	Langerhans cell histiocytosis reveals a new IL-17A–dependent pathway of dendritic cell fusion. Nature Medicine, 2008, 14, 81-87.	30.7	180
149	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
150	Reactivations in Multisystem Langerhans Cell Histiocytosis: Data of the International LCH Registry. Journal of Pediatrics, 2008, 153, 700-705.e2.	1.8	88
151	Patterns of domestic migrations and access to childhood cancer care centres in Italy: A report from the hospital based registry of the Italian Association of Pediatric Hematology and Oncology (AIEOP). European Journal of Cancer, 2008, 44, 2101-2105.	2.8	13
152	Treatment of pediatric acute lymphoblastic leukemia. Haematologica, 2008, 93, 1124-1128.	3.5	20
153	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
154	SYNOVIAL SARCOMA OF THE NECK IN A CHILD: A Multidisciplinary Approach. Pediatric Hematology and Oncology, 2008, 25, 431-437.	0.8	5
155	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
156	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
157	Improved outcome in multisystem Langerhans cell histiocytosis is associated with therapy intensification. Blood, 2008, 111, 2556-2562.	1.4	287
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