Jan-Inge Henter

List of Publications by Year
in descending order

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142 papers	19,959 citations	30070 54 h-index	11939 134 g-index
145 all docs	145 docs citations	145 times ranked	17272 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	Robust T Cell Immunity in Convalescent Individuals with Asymptomatic or Mild COVID-19. Cell, 2020, 183, 158-168.e14.	28.9	1,561
3	Perforin Gene Defects in Familial Hemophagocytic Lymphohistiocytosis. Science, 1999, 286, 1957-1959.	12.6	1,074
4	Revised classification of histiocytoses and neoplasms of the macrophage-dendritic cell lineages. Blood, 2016, 127, 2672-2681.	1.4	1,040
5	Contemporary classification of histiocytic disorders. Medical and Pediatric Oncology, 1997, 29, 157-166.	1.0	740
6	Treatment of hemophagocytic lymphohistiocytosis with HLH-94 immunochemotherapy and bone marrow transplantation. Blood, 2002, 100, 2367-2373.	1.4	737
7	Recommendations for the management of hemophagocytic lymphohistiocytosis in adults. Blood, 2019, 133, 2465-2477.	1.4	587
8	Linkage of familial hemophagocytic lymphohistiocytosis (FHL) type-4 to chromosome 6q24 and identification of mutations in syntaxin 11. Human Molecular Genetics, 2005, 14, 827-834.	2.9	502
9	Chemoimmunotherapy for hemophagocytic lymphohistiocytosis: long-term results of the HLH-94 treatment protocol. Blood, 2011, 118, 4577-4584.	1.4	493
10	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
11	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
12	HLH-94: A treatment protocol for hemophagocytic lymphohistiocytosis. Medical and Pediatric Oncology, 1997, 28, 342-347.	1.0	417
13	INFECTION- AND MALIGNANCY-ASSOCIATED HEMOPHAGOCYTIC SYNDROMES. Hematology/Oncology Clinics of North America, 1998, 12, 435-444.	2.2	376
14	Incidence in Sweden and Clinical Features of Familial Hemophagocytic Lymphohistiocytosis. Acta Paediatrica, International Journal of Paediatrics, 1991, 80, 428-435.	1.5	362
15	Therapy prolongation improves outcome in multisystem Langerhans cell histiocytosis. Blood, 2013, 121, 5006-5014.	1.4	343
16	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Annals of the Rheumatic Diseases, 2016, 75, 481-489.	0.9	338
17	Defective cytotoxic lymphocyte degranulation in syntaxin-11â€“deficient familial hemophagocytic lymphohistiocytosis 4 (FHL4) patients. Blood, 2007, 110, 1906-1915.	1.4	272
18	A prospective evaluation of degranulation assays in the rapid diagnosis of familial hemophagocytic syndromes. Blood, 2012, 119, 2754-2763.	1.4	263

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19	Spectrum of Perforin Gene Mutations in Familial Hemophagocytic Lymphohistiocytosis. American Journal of Human Genetics, 2001, 68, 590-597.	6.2	246
20	FAMILIAL HEMOPHAGOCYTIC LYMPHOHISTIOCYTOSIS. Hematology/Oncology Clinics of North America, 1998, 12, 417-433.	2.2	241
21	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
22	Haematopoietic stem cell transplantation in haemophagocytic lymphohistiocytosis. British Journal of Haematology, 2005, 129, 622-630.	2.5	206
23	Central Nervous System Disease in Langerhans Cell Histiocytosis. Journal of Pediatrics, 2010, 156, 873-881.e1.	1.8	193
24	Neuropathologic findings and neurologic symptoms in twenty-three children with hemophagocytic lymphohistiocytosis. Journal of Pediatrics, 1997, 130, 358-365.	1.8	189
25	Langerhans cell histiocytosis reveals a new IL-17Aâ€“dependent pathway of dendritic cell fusion. Nature Medicine, 2008, 14, 81-87.	30.7	180
26	Incidence of Langerhans cell histiocytosis in children: A populationâ€based study. Pediatric Blood and Cancer, 2008, 51, 76-81.	1.5	179
27	Vascular endothelial growth factor prolongs survival in a transgenic mouse model of ALS. Annals of Neurology, 2004, 56, 564-567.	5.3	145
28	Consensus recommendations for the diagnosis and management of hemophagocytic lymphohistiocytosis associated with malignancies. Haematologica, 2015, 100, 997-1004.	3.5	135
29	Familial hemophagocytic lymphohistiocytosis and viral infections. Acta Paediatrica, International Journal of Paediatrics, 1993, 82, 369-372.	1.5	123
30	Comparison of primary human cytotoxic T-cell and natural killer cell responses reveal similar molecular requirements for lytic granule exocytosis but differences in cytokine production. Blood, 2013, 121, 1345-1356.	1.4	122
31	Familial hemophagocytic lymphohistiocytosis type 3 (FHL3) caused by deep intronic mutation and inversion in UNC13D. Blood, 2011, 118, 5783-5793.	1.4	115
32	Recommendations for the Use of Etoposide-Based Therapy and Bone Marrow Transplantation for the Treatment of HLH: Consensus Statements by the HLH Steering Committee of the Histiocyte Society. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 1508-1517.	3.8	112
33	Spectrum of clinical presentations in familial hemophagocytic lymphohistiocytosis type 5 patients with mutations in STXBP2. Blood, 2010, 116, 2635-2643.	1.4	108
34	Major alterations in the mononuclear phagocyte landscape associated with COVID-19 severity. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	104
35	Targeting SAMHD1 with the Vpx protein to improve cytarabine therapy for hematological malignancies. Nature Medicine, 2017, 23, 256-263.	30.7	102
36	Different NK cellâ€“activating receptors preferentially recruit Rab27a or Munc13-4 to perforin-containing granules for cytotoxicity. Blood, 2009, 114, 4117-4127.	1.4	90

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37	How to Treat Involvement of the Central Nervous System in Hemophagocytic Lymphohistiocytosis?. Current Treatment Options in Neurology, 2017, 19, 3.	1.8	90
38	Tissue-infiltrating neutrophils represent the main source of IL-23 in the colon of patients with IBD. Gut, 2016, 65, 1632-1641.	12.1	87
39	Cytotoxic therapy for severe avian influenza A (H5N1) infection. Lancet, The, 2006, 367, 870-873.	13.7	85
40	The minimum required level of donor chimerism in hereditary hemophagocytic lymphohistiocytosis. Blood, 2016, 127, 3281-3290.	1.4	83
41	Clinical presentation of Griscelli syndrome type 2 and spectrum of RAB27A mutations. Pediatric Blood and Cancer, 2010, 54, 563-572.	1.5	82
42	Successful Treatment of Langerhans'-Cell Histiocytosis with Etanercept. New England Journal of Medicine, 2001, 345, 1577-1578.	27.0	78
43	Characterization of PRF1, STX11 and UNC13D genotypeâ€phenotype correlations in familial hemophagocytic lymphohistiocytosis. British Journal of Haematology, 2008, 143, 75-83.	2.5	78
44	Genotype-phenotype study of familial haemophagocytic lymphohistiocytosis type 3. Journal of Medical Genetics, 2011, 48, 343-352.	3.2	76
45	Biology and treatment of familial hemophagocytic lymphohistiocytosis: Importance of perforin in lymphocyte-mediated cytotoxicity and triggering of apoptosis. Medical and Pediatric Oncology, 2002, 38, 305-309.	1.0	72
46	Risk factors for early death in children with haemophagocytic lymphohistiocytosis. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, 313-318.	1.5	69
47	The need for worldwide policy and action plans for rare diseases. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, 805-807.	1.5	69
48	Hemophagocytic lymphohistiocytosis in 2 patients with underlying IFN-Î³ receptor deficiency. Journal of Allergy and Clinical Immunology, 2015, 135, 1638-1641.e5.	2.9	69
49	Incidence and clinical presentation of primary hemophagocytic lymphohistiocytosis in Sweden. Pediatric Blood and Cancer, 2015, 62, 346-352.	1.5	63
50	Fatal agranulocytosis after deferiprone therapy in a child with Diamond-Blackfan anemia. Blood, 2007, 109, 5157-5159.	1.4	61
51	VEGF reduces astrogliosis and preserves neuromuscular junctions in ALS transgenic mice. Biochemical and Biophysical Research Communications, 2007, 363, 989-993.	2.1	59
52	Frequency and development of CNS involvement in Chinese children with hemophagocytic lymphohistiocytosis. Pediatric Blood and Cancer, 2010, 54, 408-415.	1.5	58
53	Incidence and pattern of radiological central nervous system Langerhans cell histiocytosis in children: A population based study. Pediatric Blood and Cancer, 2011, 56, 250-257.	1.5	58
54	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

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55	SAMHD1 protects cancer cells from various nucleoside-based antimetabolites. Cell Cycle, 2017, 16, 1029-1038.	2.6	56
56	Familial Hemophagocytic Lymphohistiocytosis: Clinical Review Based on the Findings in Seven Children. Acta Paediatrica, International Journal of Paediatrics, 1991, 80, 269-277.	1.5	55
57	High-dimensional profiling reveals phenotypic heterogeneity and disease-specific alterations of granulocytes in COVID-19. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	52
58	Children with cancer share their views: tell the truth but leave room for hope. Acta Paediatrica, International Journal of Paediatrics, 2016, 105, 1094-1099.	1.5	51
59	Pulmonary abnormalities at long-term follow-up of patients with Langerhans cell histiocytosis. Medical and Pediatric Oncology, 2001, 36, 459-468.	1.0	50
60	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
61	High Systemic Levels of the Cytokine-Inducing HMGB1 Isoform Secreted in Severe Macrophage Activation Syndrome. Molecular Medicine, 2014, 20, 538-547.	4.4	45
62	Induction of apoptosis and caspase activation in cells obtained from familial haemophagocytic lymphohistiocytosis patients. British Journal of Haematology, 1999, 106, 406-415.	2.5	44
63	Elevated circulating levels of interleukin-1 receptor antagonist but not IL-1 agonists in hemophagocytic lymphohistiocytosis. , 1996, 27, 21-25.		43
64	Long-term follow-up of Langerhans cell histiocytosis: 39 yearsâ€™ experience at a single centre. Acta Paediatrica, International Journal of Paediatrics, 2005, 94, 1073-1084.	1.5	43
65	Longâ€term followâ€up of Langerhans cell histiocytosis: 39 yearsâ€™ experience at a single centre. Acta Paediatrica, International Journal of Paediatrics, 2005, 94, 1073-1084.	1.5	43
66	With me or against me: Tumor suppressor and drug resistance activities of SAMHD1. Experimental Hematology, 2017, 52, 32-39.	0.4	43
67	SUCCESSFUL INDUCTION WITH CHEMOTHERAPY INCLUDING TENIPOSIDE IN FAMILIAL ERYTHROPHAGOCYTIC LYMPHOHISTIOCYTOSIS. Lancet, The, 1986, 328, 1402.	13.7	42
68	Persistent B19 parvovirus infection in pediatric malignancies. , 1998, 31, 66-72.		39
69	Novel deep intronic and missense <scp>UNC</scp>13<scp>D</scp> mutations in familial haemophagocytic lymphohistiocytosis type 3. British Journal of Haematology, 2013, 162, 415-418.	2.5	39
70	Pathophysiology and spectrum of diseases caused by defects in lymphocyte cytotoxicity. Experimental Cell Research, 2014, 325, 10-17.	2.6	38
71	Spectrum of Atypical Clinical Presentations in Patients with Biallelic PRF1 Missense Mutations. Pediatric Blood and Cancer, 2015, 62, 2094-2100.	1.5	38
72	Cytotoxic therapy for severe swine flu A/H1N1. Lancet, The, 2010, 376, 2116.	13.7	37

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73	Targeted high-throughput sequencing for genetic diagnostics of hemophagocytic lymphohistiocytosis. Genome Medicine, 2015, 7, 130.	8.2	37
74	Brain 18-FDG PET scan in central nervous system langerhans cell histiocytosis. Journal of Pediatrics, 2002, 141, 435-440.	1.8	36
75	Development of classical Hodgkin's lymphoma in an adult with biallelic STXBP2 mutations. Haematologica, 2013, 98, 760-764.	3.5	35
76	Ribonucleotide reductase inhibitors suppress <scp>SAMHD</scp> 1 araâ€•<scp>CTP</scp> ase activity enhancing cytarabine efficacy. EMBO Molecular Medicine, 2020, 12, e10419.	6.9	35
77	Stem cell transplantation for children with hemophagocytic lymphohistiocytosis: results from the HLH-2004 study. Blood Advances, 2020, 4, 3754-3766.	5.2	34
78	Subtyping of natural killer cell cytotoxicity deficiencies in haemophagocytic lymphohistocytosis provides therapeutic guidance. British Journal of Haematology, 2005, 129, 658-666.	2.5	33
79	Treatment of the Xâ€linked lymphoproliferative, Griscelli and ChÃ©diakâ€“Higashi syndromes by HLH directed therapy. Pediatric Blood and Cancer, 2009, 52, 268-272.	1.5	33
80	Langerhans-cell histiocytosis: neoplasia or unbridled inflammation?. Trends in Immunology, 2003, 24, 409-410.	6.8	32
81	Cancer risk in relatives of patients with a primary disorder of lymphocyte cytotoxicity: a retrospective cohort study. Lancet Haematology,the, 2015, 2, e536-e542.	4.6	32
82	Dengue Infection Complicated by Hemophagocytic Lymphohistiocytosis: Experiences From 180 Patients With Severe Dengue. Clinical Infectious Diseases, 2020, 70, 2247-2255.	5.8	32
83	Kalaâ€azar in a oneâ€yearâ€old Swedish child. Diagnostic difficulties because of active hemophagocytosis. Acta Paediatrica, International Journal of Paediatrics, 1993, 82, 794-796.	1.5	31
84	Successful Hematopoietic Stem Cell Transplantation in a Patient with LPS-Responsive Beige-Like Anchor (LRBA) Gene Mutation. Journal of Clinical Immunology, 2016, 36, 480-489.	3.8	30
85	Tumor Necrosis Factor, Interleukin 11, and Leukemia Inhibitory Factor Produced by Langerhans Cells in Langerhans Cell Histiocytosis. Journal of Pediatric Hematology/Oncology, 2004, 26, 706-711.	0.6	29
86	Consensus-Based Guidelines for the Recognition, Diagnosis, and Management of Hemophagocytic Lymphohistiocytosis in Critically Ill Children and Adults. Critical Care Medicine, 2022, 50, 860-872.	0.9	29
87	Familial Hemophagocytic Lymphohistiocytosis: Too Little Cell Death Can Seriously Damage Your Health. Leukemia and Lymphoma, 2001, 42, 13-20.	1.3	28
88	Low-level expression of SAMHD1 in acute myeloid leukemia (AML) blasts correlates with improved outcome upon consolidation chemotherapy with high-dose cytarabine-based regimens. Blood Cancer Journal, 2018, 8, 98.	6.2	28
89	Successful extracorporeal membrane oxygenation in four children with malignant disease and severePneumocystis carinii pneumonia. , 1999, 32, 25-31.		27
90	Differences in Granule Morphology yet Equally Impaired Exocytosis among Cytotoxic T Cells and NK Cells from Chediakâ€“Higashi Syndrome Patients. Frontiers in Immunology, 2017, 8, 426.	4.8	26

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91	Efficacy of Moderately Dosed Etoposide in Macrophage Activation Syndromeâ€“Hemophagocytic Lymphohistiocytosis. Journal of Rheumatology, 2021, 48, 1596-1602.	2.0	26
92	Detection of IL-17A-producing peripheral blood monocytes in Langerhans cell histiocytosis patients. Clinical Immunology, 2014, 153, 112-122.	3.2	24
93	HLHâ€94: A treatment protocol for hemophagocytic lymphohistiocytosis. Medical and Pediatric Oncology, 1997, 28, 342-347.	1.0	24
94	Clinical presentation of hemophagocytic lymphohistiocytosis in adults is less typical than in children. Clinics, 2016, 71, 205-209.	1.5	23
95	Immunogenetic Heterogeneity in Single-System and Multisystem Langerhans Cell Histiocytosis. Pediatric Research, 2003, 54, 30-36.	2.3	22
96	Biomarkers in the cerebrospinal fluid and neurodegeneration in Langerhans cell histiocytosis. Pediatric Blood and Cancer, 2009, 53, 1264-1270.	1.5	22
97	A RAB27A 5â€² untranslated region structural variant associated with late-onset hemophagocytic lymphohistiocytosis and normal pigmentation. Journal of Allergy and Clinical Immunology, 2018, 142, 317-321.e8.	2.9	22
98	An N-Terminal Missense Mutation in STX11 Causative of FHL4 Abrogates Syntaxin-11 Binding to Munc18-2. Frontiers in Immunology, 2014, 4, 515.	4.8	20
99	Combined newborn screening for familial hemophagocytic lymphohistiocytosis and severe T- and B-cell immunodeficiencies. Journal of Allergy and Clinical Immunology, 2014, 134, 226-228.e7.	2.9	20
100	Treatment of Familial Hemophagocytic Lymphohistiocytosis with Third-Party Mesenchymal Stromal Cells. Stem Cells and Development, 2012, 21, 3147-3151.	2.1	19
101	Neuropsychological sequelae in patients with neurodegenerative Langerhans cell histiocytosis. Pediatric Blood and Cancer, 2008, 51, 669-674.	1.5	16
102	Detection of Langerhans cell histiocytosis lesions with somatostatin analogue scintigraphy?a preliminary report. Medical and Pediatric Oncology, 2000, 35, 462-467.	1.0	15
103	Langerhans cell histiocytosis in children born 1982â€“2005 after in vitro fertilization. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, 1151-1155.	1.5	15
104	Severe bacteriaâ€associated hemophagocytic lymphohistiocytosis in an extremely premature infant. Acta Paediatrica, International Journal of Paediatrics, 2007, 96, 1703-1706.	1.5	14
105	Foxp3+ Tregs from Langerhans cell histiocytosis lesions co-express CD56 and have a definitively regulatory capacity. Clinical Immunology, 2020, 215, 108418.	3.2	14
106	Clinical and Genetic Studies of Familial Hemophagocytic Lymphohistiocytosis in Oman: Need for Early Treatment. Pediatric Hematology and Oncology, 2003, 20, 603-609.	0.8	13
107	Elevated ferritin and soluble CD25 in critically ill patients are associated with parameters of (hyper) inflammation and lymphocyte cytotoxicity. Minerva Anestesiologica, 2019, 85, 1289-1298.	1.0	13
108	Elevated Serum Levels of the Decoy Receptor Osteoprotegerin in Children with Langerhans Cell Histiocytosis. Pediatric Research, 2006, 59, 281-286.	2.3	12

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109	Pronounced hyperferritinemia: Expanding the field of hemophagocytic lymphohistiocytosis. Pediatric Blood and Cancer, 2008, 50, 1127-1129.	1.5	11
110	Adsorptive depletion of blood monocytes reduces the levels of circulating interleukin-17A in Langerhans cell histiocytosis. Blood, 2016, 128, 1302-1305.	1.4	11
111	Emapalumab in Primary Hemophagocytic Lymphohistiocytosis. New England Journal of Medicine, 2020, 383, 596-599.	27.0	11
112	Pulmonary function testing and pulmonary Langerhans cell histiocytosis. Pediatric Blood and Cancer, 2007, 49, 323-328.	1.5	10
113	High levels of plasma interleukin-17A are associated with severe neurological sequelae in Langerhans cell histiocytosis. Cytokine, 2020, 126, 154877.	3.2	10
114	Lipoprotein alterations in children with bacterial meningitis. Acta Paediatrica, International Journal of Paediatrics, 1993, 82, 694-698.	1.5	9
115	HAEMOPHAGOCYTICLYMPHOHISTIOCYTOSIS: AN INHERITED PRIMARY FORM AND A REACTIVE SECONDARY FORM. British Journal of Haematology, 1995, 91, 774-775.	2.5	9
116	Killer cell immunoglobulinâ€like receptor gene polymorphisms predispose susceptibility to Epsteinâ€Barr virus associated hemophagocytic lymphohistiocytosis in Chinese children. Microbiology and Immunology, 2012, 56, 378-384.	1.4	9
117	Response to mitogenâ€activated protein kinase inhibition of neurodegeneration in Langerhans cell histiocytosis monitored by cerebrospinal fluid neurofilament light as a biomarker: a pilot study. British Journal of Haematology, 2022, 196, 248-254.	2.5	9
118	Haploinsufficiency of UNC13D increases the risk of lymphoma. Cancer, 2019, 125, 1848-1854.	4.1	8
119	Patients with both Langerhans cell histiocytosis and Crohnâ€™s disease highlight a common role of interleukinâ€23. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 1315-1321.	1.5	8
120	Additive Prognostic Impact of Gastrointestinal Involvement in Severe Multisystem Langerhans Cell Histiocytosis. Journal of Pediatrics, 2021, 237, 65-70.e3.	1.8	8
121	Severe congenital neutropeniaâ€associated JAGN1 mutations unleash a calpainâ€dependent cell death programme in myeloid cells. British Journal of Haematology, 2021, 192, 200-211.	2.5	7
122	Is neutralization of IFNâ€Î³ sufficient to control inflammation in HLH?. Pediatric Blood and Cancer, 2021, 68, e28886.	1.5	7
123	Therapeutic administration of etoposide coincides with reduced systemic HMGB1 levels in macrophage activation syndrome. Molecular Medicine, 2021, 27, 48.	4.4	7
124	Screening for neurodegeneration in Langerhans cell histiocytosis with neurofilament light in plasma. British Journal of Haematology, 2022, , .	2.5	7
125	Reply to: "Interleukin-17A is not expressed by CD207+ cells in Langerhans cell histiocytosis lesions". Nature Medicine, 2009, 15, 484-485.	30.7	6
126	Polyclonal T-Cells Express CD1a in Langerhans Cell Histiocytosis (LCH) Lesions. PLoS ONE, 2014, 9, e109586.	2.5	6

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127	Sequence analysis of the SRGN, AP3B1, ARF6, and SH2D1A genes in familial hemophagocytic lymphohistiocytosis. Pediatric Blood and Cancer, 2008, 50, 1067-1069.	1.5	4
128	Molecular Genetics Diversity of Primary Hemophagocytic Lymphohistiocytosis among Polish Pediatric Patients. Archivum Immunologiae Et Therapiae Experimentalis, 2021, 69, 31.	2.3	4
129	Diagnostic challenges for a novel SH2D1A mutation associated with Xâ€linked lymphoproliferative disease. Pediatric Blood and Cancer, 2020, 67, e28184.	1.5	4
130	Simple Evaluation of Clinical Situation and Subtypes of Pediatric Hemophagocytic Lymphohistiocytosis by Cytokine Patterns. Frontiers in Immunology, 2022, 13, 850443.	4.8	3
131	Persistent B19 parvovirus infection in pediatric malignancies. Medical and Pediatric Oncology, 1998, 31, 66-72.	1.0	2
132	Plasma Signaling Factors in Patients With Langerhans Cell Histiocytosis (LCH) Correlate With Relative Frequencies of LCH Cells and T Cells Within Lesions. Frontiers in Pediatrics, 0, 10, .	1.9	2
133	Elevated circulating levels of interleukinâ€1 receptor antagonist but not ILâ€1 agonists in hemophagocytic lymphohistiocytosis. Medical and Pediatric Oncology, 1996, 27, 21-25.	1.0	1
134	Clinical and laboratory signs of haemophagocytic lymphohistiocytosis associated with pandemic influenza A (H1N1) infection in patients needing extracorporeal membrane oxygenation. European Journal of Anaesthesiology, 2021, 38, 692-701.	1.7	1
135	Neutralizing Anti-IL-17A Antibody Demonstrates Preclinical Activity Enhanced by Vinblastine in Langerhans Cell Histiocytosis. Frontiers in Oncology, 2021, 11, 780191.	2.8	1
136	Clinical and Genetic Studies of Familial Hemophagocytic Lymphohistiocytosis in Oman: Need for Early Treatment. Pediatric Hematology and Oncology, 2003, 20, 603-609.	0.8	1
137	Treatment of Newly Diagnosed HLH and Refractory Disease. , 2018, , 247-263.		0
138	Comparison of three different ELISAs for the detection of recombinant, native and plasma IL-17A. MethodsX, 2020, 7, 100997.	1.6	0
139	Identification of a MEF Gene Mutation in a Familial Hemophagocytic Lymphohistiocytosis Patient That Decreases MEF Transcriptional Activity.. Blood, 2005, 106, 3013-3013.	1.4	0
140	Etoposide Therapy of Cytokine Storm Syndromes. , 2019, , 521-547.		0
141	Expression of concern to: High systematic levels of the cytokine-inducing HMGB1 isoform secreted in severe macrophage activation syndrome. Molecular Medicine, 2020, 26, 17.	4.4	0
142	Seasonality of birth month in patients diagnosed with Langerhans cell histiocytosis (LCH). Pediatric Research, 2022, , .	2.3	0