

Shinsaku Imashuku

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2296164/publications.pdf>

Version: 2024-02-01

161
papers

11,677
citations

93792

39
h-index

34195

103
g-index

163
all docs

163
docs citations

163
times ranked

7539
citing authors

#	ARTICLE	IF	CITATIONS
1	Bone lesions of Langerhans cell histiocytosis triggered by trauma in children. <i>Pediatrics International</i> , 2022, 64, .	0.2	1
2	Multiple polypoid lesions with erosion of the gastric mucosa in adult Tâ€cell lymphoma/leukemia superimposed on cytomegalovirus infection. <i>EJHaem</i> , 2022, 3, 1054-1055.	0.4	1
3	Longâ€term complications in uniformly treated paediatric Langerhans histiocytosis patients disclosed by 12 years of followâ€up of the JLSGâ€96/02 studies. <i>British Journal of Haematology</i> , 2021, 192, 615-620.	1.2	11
4	Pentasony 21 in an older adult case of AML with myelodysplastic changes. <i>International Journal of Laboratory Hematology</i> , 2021, 43, e122-e123.	0.7	0
5	Congenital Hypofibrinogenemia in a Neonate with a Novel Mutation in the FGB Gene. <i>Pediatric Reports</i> , 2021, 13, 113-117.	0.5	0
6	Protein C Gene Mutation in an Older Adult Patient with Clostridium perfringens Septicemia-Related Visceral Vein Thrombosis. <i>TH Open</i> , 2021, 05, e171-e173.	0.7	0
7	Virusâ€triggered secondary hemophagocytic lymphohistiocytosis. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2021, 110, 2729-2736.	0.7	30
8	COVID-19 vaccination: effective utilization of low dead space (LDS) syringes. <i>International Journal of Infectious Diseases</i> , 2021, 113, 90-92.	1.5	5
9	Aortic Mural Thrombus Associated with Congenital Protein C Deficiency in an Elderly Patient. <i>Journal of Atherosclerosis and Thrombosis</i> , 2020, 27, 100-103.	0.9	5
10	CD4â~/CD8+ adult Tâ€cell leukemia/lymphoma with unusual morphology presenting as ascites and pleural effusion. <i>International Journal of Laboratory Hematology</i> , 2020, 42, e105-e106.	0.7	1
11	Nationwide retrospective review of hematopoietic stem cell transplantation in children with refractory Langerhans cell histiocytosis. <i>International Journal of Hematology</i> , 2020, 111, 137-148.	0.7	9
12	Rivaroxaban-Related Traumatic Large Subcutaneous Hematoma in the Calf Requiring Surgical Repair in an Elderly Patient. <i>TH Open</i> , 2020, 04, e104-e106.	0.7	1
13	Natural history of epithelioid hemangioendothelioma that progressed over 20 years. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28261.	0.8	0
14	Management of Chronic Disseminated Intravascular Coagulation Associated with Aortic Aneurysm/Dissection. <i>Case Reports in Hematology</i> , 2019, 2019, 1-6.	0.3	6
15	Central diabetes insipidus in pediatric patients with Langerhans cell histiocytosis: Results from the JLSGâ€96/02 studies. <i>Pediatric Blood and Cancer</i> , 2019, 66, e27454.	0.8	15
16	Intensification of induction therapy and prolongation of maintenance therapy did not improve the outcome of pediatric Langerhans cell histiocytosis with single-system multifocal bone lesions: results of the Japan Langerhans Cell Histiocytosis Study Group-02 Protocol Study. <i>International Journal of Hematology</i> , 2018, 108, 192-198.	0.7	23
17	Splenic mass in a case of CALR -mutated essential thrombocythemia. <i>Clinical Case Reports (discontinued)</i> , 2018, 6, 2291-2292.	0.2	3
18	Management of Acquired Hemophilia A in Elderly Patients. <i>Case Reports in Hematology</i> , 2018, 2018, 1-5.	0.3	3

#	ARTICLE	IF	CITATIONS
19	Bacteremia and meningitis caused by a novel clone of <i>Neisseria meningitidis</i> serogroup B. <i>Pediatrics International</i> , 2018, 60, 1093-1094.	0.2	2
20	Plasmapheresis for Spur Cell Anemia in a Patient with Alcoholic Liver Cirrhosis. <i>Case Reports in Hematology</i> , 2018, 2018, 1-4.	0.3	6
21	Importance of the Average Glucose Level and Estimated Glycated Hemoglobin in a Diabetic Patient with Hereditary Hemolytic Anemia and Liver Cirrhosis. <i>Internal Medicine</i> , 2018, 57, 537-543.	0.3	9
22	Topical Imiquimod for the Treatment of Relapsed Cutaneous Langerhans Cell Histiocytosis after Chemotherapy in an Elderly Patient. <i>Case Reports in Dermatological Medicine</i> , 2018, 2018, 1-5.	0.1	1
23	Merkel cell polyomavirus and Langerhans cell neoplasm. <i>Cell Communication and Signaling</i> , 2018, 16, 49.	2.7	10
24	Fatal Invasive Cryptococcal Infection in an HIV-Negative Elderly Patient with Decompensated Hepatic Cirrhosis. <i>Case Reports in Hepatology</i> , 2018, 2018, 1-6.	0.4	2
25	CNS-directed Prophylactic Approach to Langerhans Cell Histiocytosis. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, 321-322.	0.3	3
26	Waldenstrom's Macroglobulinemia: A Report of Two Cases, One with Severe Retinopathy and One with Renal Failure. <i>Case Reports in Hematology</i> , 2017, 2017, 1-6.	0.3	5
27	Inflammatory serum cytokines and chemokines increase associated with the disease extent in pediatric Langerhans cell histiocytosis. <i>Cytokine</i> , 2017, 97, 73-79.	1.4	37
28	A Fatal Case of Congenital Langerhans Cell Histiocytosis with Disseminated Cutaneous Lesions in a Premature Neonate. <i>Case Reports in Pediatrics</i> , 2016, 2016, 1-4.	0.2	6
29	Acute Monocytic Leukemia Masquerading Behçet's Disease-Like Illness at Onset in an Elderly Female. <i>Case Reports in Hematology</i> , 2016, 2016, 1-5.	0.3	2
30	Intensified and prolonged therapy comprising cytarabine, vincristine and prednisolone improves outcome in patients with multisystem Langerhans cell histiocytosis: results of the Japan Langerhans Cell Histiocytosis Study Group-02 Protocol Study. <i>International Journal of Hematology</i> , 2016, 104, 99-109.	0.7	68
31	Severe <i>Helicobacter pylori</i> gastritis-related thrombocytopenia and iron deficiency anemia in an adolescent female. <i>Annals of Hematology</i> , 2016, 95, 835-836.	0.8	4
32	PIEZO1 gene mutation in a Japanese family with hereditary high phosphatidylcholine hemolytic anemia and hemochromatosis-induced diabetes mellitus. <i>International Journal of Hematology</i> , 2016, 104, 125-129.	0.7	25
33	Complete Response of Adult-Onset CNS Langerhans Cell Histiocytosis Documented on 18F-FDG PET/CT. <i>Clinical Nuclear Medicine</i> , 2015, 40, 981-982.	0.7	8
34	Acute-phase ITIH4 levels distinguish multi-system from single-system Langerhans cell histiocytosis via plasma peptidomics. <i>Clinical Proteomics</i> , 2015, 12, 16.	1.1	8
35	Sporadic Epstein syndrome with macrothrombocytopenia, sensorineural hearing loss and renal failure. <i>Pediatrics International</i> , 2015, 57, 977-981.	0.2	5
36	Commentary: Intravenous Immunoglobulin (IVIG) Therapy for Patients with Langerhans Cell Histiocytosis (LCH)-Related Neurodegenerative Diseases of the CNS. <i>CNS and Neurological Disorders - Drug Targets</i> , 2015, 14, 688-690.	0.8	0

#	ARTICLE	IF	CITATIONS
37	Successful Treatment of Leukemic Mature B-Cell Lymphoid Neoplasm with Similar Features to Splenic Marginal Zone Lymphoma Possessing Aberrant Myeloid Markers. Case Reports in Hematology, 2015, 2015, 1-4.	0.3	0
38	Two Cases of Primary Cold Agglutinin Disease Associated with Megaloblastic Anemia. Case Reports in Hematology, 2015, 2015, 1-5.	0.3	3
39	Opportunistic Infections in Patients with HTLV-1 Infection. Case Reports in Hematology, 2015, 2015, 1-5.	0.3	12
40	Interleukin-1 loop model for pathogenesis of Langerhans cell histiocytosis. Cell Communication and Signaling, 2015, 13, 13.	2.7	30
41	Follow-up of pediatric patients treated by IVIG for Langerhans cell histiocytosis (LCH)-related neurodegenerative CNS disease. International Journal of Hematology, 2015, 101, 191-197.	0.7	40
42	Strategies for the Prevention of Central Nervous System Complications in Patients with Langerhans Cell Histiocytosis. Hematology/Oncology Clinics of North America, 2015, 29, 875-893.	0.9	15
43	Treatment of Epstein-Barr virus-related hemophagocytic lymphohistiocytosis: Study protocol of a prospective pilot study. World Journal of Hematology, 2015, 4, 69.	0.1	1
44	High serum osteopontin levels in pediatric patients with high risk Langerhans cell histiocytosis. Cytokine, 2014, 70, 194-197.	1.4	17
45	Expansion of Natural Killer Cells in Peripheral Blood in a Japanese Elderly with Human T-Cell Lymphotropic Virus Type 1-Related Skin Lesions. Case Reports in Dermatological Medicine, 2014, 2014, 1-4.	0.1	1
46	Merkel cell polyomavirus DNA sequences in peripheral blood and tissues from patients with Langerhans cell histiocytosis. Human Pathology, 2014, 45, 119-126.	1.1	24
47	IL-17A receptor expression differs between subclasses of Langerhans cell histiocytosis, which might settle the IL-17A controversy. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2013, 462, 219-228.	1.4	22
48	Therapeutic outcome of multifocal Langerhans cell histiocytosis in adults treated with the Special C regimen formulated by the Japan LCH Study Group. International Journal of Hematology, 2013, 97, 103-108.	0.7	29
49	Myeloid/NK cell acute leukemia with unique blast morphology: de novo or secondary leukemia?. International Journal of Hematology, 2013, 98, 509-511.	0.7	1
50	Are regimens containing rituximab effective in the initial treatment of Epstein-Barr virus-positive natural killer cell lymphoproliferative disease-associated hemophagocytic lymphohistiocytosis?. International Journal of Hematology, 2013, 98, 375-377.	0.7	8
51	Rituximab for managing acquired hemophilia A in a case of chronic neutrophilic leukemia with the JAK2 kinase V617F mutation. Journal of Blood Medicine, 2012, 3, 157.	0.7	15
52	Immune Dysregulation Diseases. , 2012, , 233-277.		0
53	Persistent thrombocytosis in elderly patients with rare hyposplenias that mimic essential thrombocythemia. International Journal of Hematology, 2012, 95, 702-705.	0.7	8
54	Chemoimmunotherapy for hemophagocytic lymphohistiocytosis: long-term results of the HLH-94 treatment protocol. Blood, 2011, 118, 4577-4584.	0.6	493

#	ARTICLE	IF	CITATIONS
55	Effectiveness of a combination of cyclosporine A, suplatast tosilate and prednisolone on periodic oscillating hypereosinophilia. <i>International Medical Case Reports Journal</i> , 2011, 4, 79.	0.3	2
56	Management of Langerhans Cell Histiocytosis (LCH)-Induced Central Diabetes Insipidus and Its Associated Endocrinological/Neurological Sequelae. , 2011, , .		0
57	Treatment of Epstein-Barr Virus-related Hemophagocytic Lymphohistiocytosis (EBV-HLH); Update 2010. <i>Journal of Pediatric Hematology/Oncology</i> , 2011, 33, 35-39.	0.3	91
58	Tyrosine phosphatase SHP-1 is expressed higher in multisystem than in single-system Langerhans cell histiocytosis by immunohistochemistry. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2011, 459, 227-234.	1.4	3
59	Treatment of patients with hypothalamic-pituitary lesions as adult-onset Langerhans cell histiocytosis. <i>International Journal of Hematology</i> , 2011, 94, 556-560.	0.7	23
60	Analysis of 43 cases of Langerhans cell histiocytosis (LCH)-induced central diabetes insipidus registered in the JLSG-96 and JLSG-02 studies in Japan. <i>International Journal of Hematology</i> , 2011, 94, 545-551.	0.7	23
61	Hematopoietic stem cell transplantation for familial hemophagocytic lymphohistiocytosis and Epstein-Barr virus-associated hemophagocytic lymphohistiocytosis in Japan. <i>Pediatric Blood and Cancer</i> , 2010, 54, 299-306.	0.8	104
62	VCR/AraC chemotherapy and ND-CNS-LCH. <i>Pediatric Blood and Cancer</i> , 2010, 55, 215-216.	0.8	8
63	Comprehensive analyses and characterization of haemophagocytic lymphohistiocytosis in Vietnamese children. <i>British Journal of Haematology</i> , 2010, 148, 301-310.	1.2	33
64	M-protein-positive chronic active Epstein-Barr virus infection: features mimicking HIV-1 infection. <i>International Journal of Hematology</i> , 2009, 90, 235-238.	0.7	1
65	Langerhans cell histiocytosis with multifocal bone lesions: comparative clinical features between single and multi-systems. <i>International Journal of Hematology</i> , 2009, 90, 506-512.	0.7	38
66	Infection of T lymphocytes in non-Asian patients with Epstein-Barr virus-associated hemophagocytic lymphohistiocytosis. <i>Pediatric Blood and Cancer</i> , 2009, 53, 1359-1359.	0.8	1
67	High Dose Immunoglobulin (IVIg) May Reduce the Incidence of Langerhans Cell Histiocytosis (LCH)-Associated Central Nervous System Involvement. <i>CNS and Neurological Disorders - Drug Targets</i> , 2009, 8, 380-386.	0.8	16
68	Treatment of neurodegenerative CNS disease in Langerhans cell histiocytosis with a combination of intravenous immunoglobulin and chemotherapy. <i>Pediatric Blood and Cancer</i> , 2008, 50, 308-311.	0.8	46
69	Impact of reactivation on the sequelae of multi-system Langerhans cell histiocytosis patients. <i>Pediatric Blood and Cancer</i> , 2008, 50, 931-932.	0.8	13
70	Hyperferritinemia in hemophagocytic lymphohistiocytosis and related diseases. <i>Pediatric Blood and Cancer</i> , 2008, 51, 442-442.	0.8	6
71	Is cytokine-induced sensorineural hearing loss reversible?. <i>Arthritis and Rheumatism</i> , 2008, 58, 3970-3970.	6.7	1
72	Frequency and spectrum of central nervous system involvement in 193 children with haemophagocytic lymphohistiocytosis. <i>British Journal of Haematology</i> , 2008, 140, 327-335.	1.2	217

#	ARTICLE	IF	CITATIONS
73	FATAL SIBLING CASES OF FAMILIAL HEMOPHAGOCYTIC LYMPHOHISTIOCYTOSIS (FHL) WITH MUNC13 ⁴ MUTATIONS: Case Reports. <i>Pediatric Hematology and Oncology</i> , 2008, 25, 171-180.	0.3	13
74	Neurodegenerative central nervous system disease as late sequelae of Langerhans cell histiocytosis. Report from the Japan LCH Study Group. <i>Haematologica</i> , 2008, 93, 615-618.	1.7	31
75	SYSTEMIC TYPE EPSTEIN-BARR VIRUS-RELATED LYMPHOPROLIFERATIVE DISEASES IN CHILDREN AND YOUNG ADULTS: Challenges for Pediatric Hemato-Oncologists and Infectious Disease Specialists. <i>Pediatric Hematology and Oncology</i> , 2007, 24, 563-568.	0.3	26
76	Association of transforming growth factor- β 1 gene polymorphism in the development of Epstein-Barr virus-related hematologic diseases. <i>Haematologica</i> , 2007, 92, 1470-1474.	1.7	26
77	Late-onset cases of familial hemophagocytic lymphohistiocytosis with missense perforin gene mutations. <i>American Journal of Hematology</i> , 2007, 82, 427-432.	2.0	72
78	Sensorineural hearing loss in a case of familial hemophagocytic lymphohistiocytosis. <i>Pediatric Blood and Cancer</i> , 2007, 49, 856-858.	0.8	7
79	HLH-2004: Diagnostic and therapeutic guidelines for hemophagocytic lymphohistiocytosis. <i>Pediatric Blood and Cancer</i> , 2007, 48, 124-131.	0.8	4,018
80	Etoposide-related secondary acute myeloid leukemia (t-AML) in hemophagocytic lymphohistiocytosis. <i>Pediatric Blood and Cancer</i> , 2007, 48, 121-123.	0.8	15
81	Nationwide Survey of Hemophagocytic Lymphohistiocytosis in Japan. <i>International Journal of Hematology</i> , 2007, 86, 58-65.	0.7	360
82	Isolated Central Nervous System Hemophagocytic Lymphohistiocytosis: Case Report. <i>Neurosurgery</i> , 2006, 58, E590.	0.6	2
83	Correlation between phenotypic heterogeneity and gene mutational characteristics in familial hemophagocytic lymphohistiocytosis (FHL). <i>Pediatric Blood and Cancer</i> , 2006, 46, 482-488.	0.8	76
84	High serum values of soluble CD154, IL-2 receptor, RANKL and osteoprotegerin in Langerhans cell histiocytosis. <i>Pediatric Blood and Cancer</i> , 2006, 47, 194-199.	0.8	44
85	Improved outcome in the treatment of pediatric multifocal Langerhans cell histiocytosis. <i>Cancer</i> , 2006, 107, 613-619.	2.0	124
86	Genetic subtypes of familial hemophagocytic lymphohistiocytosis: correlations with clinical features and cytotoxic T lymphocyte/natural killer cell functions. <i>Blood</i> , 2005, 105, 3442-3448.	0.6	126
87	Review of hemophagocytic lymphohistiocytosis (HLH) in children with focus on Japanese experiences. <i>Critical Reviews in Oncology/Hematology</i> , 2005, 53, 209-223.	2.0	63
88	Response to Imatinib Mesylate in a Patient with Idiopathic Hypereosinophilic Syndrome Associated with Cyclic Eosinophil Oscillations. <i>International Journal of Hematology</i> , 2005, 81, 310-314.	0.7	7
89	Proposed guidelines for diagnosing chronic active Epstein-Barr virus infection. <i>American Journal of Hematology</i> , 2005, 80, 64-69.	2.0	246
90	Occurrence of haemophagocytic lymphohistiocytosis at less than 1 year of age: analysis of 96 patients. <i>European Journal of Pediatrics</i> , 2005, 164, 315-319.	1.3	61

#	ARTICLE	IF	CITATIONS
91	Clinical aspects and therapy of hemophagocytic lymphohistiocytosis. , 2005, , 353-379.		10
92	Healing hemophagocytosis. <i>Clinical Immunology</i> , 2005, 117, 121-124.	1.4	6
93	Pineal dysfunction in Langerhans cell histiocytosis?. <i>Pediatric Blood and Cancer</i> , 2004, 43, 95-95.	0.8	0
94	Secondary Acute Promyelocytic Leukemia Following Chemotherapy for Non-Hodgkin's Lymphoma in a Child. <i>Journal of Pediatric Hematology/Oncology</i> , 2004, 26, 427-430.	0.3	11
95	Cerebellar Ataxia in Pediatric Patients With Langerhans Cell Histiocytosis. <i>Journal of Pediatric Hematology/Oncology</i> , 2004, 26, 735-739.	0.3	19
96	Longitudinal follow-up of patients with Epstein-Barr virus-associated hemophagocytic lymphohistiocytosis. <i>Haematologica</i> , 2004, 89, 183-8.	1.7	67
97	Pineal dysfunction (low melatonin production) as a cause of sudden death in a long-term survivor of langerhans cell histiocytosis?. <i>Medical and Pediatric Oncology</i> , 2003, 41, 151-153.	1.0	5
98	Treatment of Epstein-Barr virus-associated hemophagocytic lymphohistiocytosis (EBV-HLH) in young adults: A report from the HLH study center. <i>Medical and Pediatric Oncology</i> , 2003, 41, 103-109.	1.0	133
99	Chronic myeloid leukemia in a patient with chronic idiopathic thrombocytopenic purpura: Rapid response to imatinib mesylate (STI571). <i>Medical and Pediatric Oncology</i> , 2003, 41, 159-160.	1.0	2
100	Engraftment and dissemination of T lymphocytes from primary haemophagocytic lymphohistiocytosis in scid mice. <i>British Journal of Haematology</i> , 2003, 121, 349-358.	1.2	13
101	Characteristic perforin gene mutations of haemophagocytic lymphohistiocytosis patients in Japan. <i>British Journal of Haematology</i> , 2003, 121, 503-510.	1.2	87
102	Fatal Hemophagocytic Lymphohistiocytosis with Clonal and Granular T Cell Proliferation in an Infant. <i>Acta Haematologica</i> , 2003, 110, 217-219.	0.7	4
103	Prognostic Factors for Chronic Active Epstein-Barr Virus Infection. <i>Journal of Infectious Diseases</i> , 2003, 187, 527-533.	1.9	207
104	Quantitative Analysis of Cell-free Epstein-Barr Virus Genome Copy Number in Patients with EBV-associated Hemophagocytic Lymphohistiocytosis. <i>Leukemia and Lymphoma</i> , 2002, 43, 173-179.	0.6	60
105	Treatment of hemophagocytic lymphohistiocytosis with HLH-94 immunochemotherapy and bone marrow transplantation. <i>Blood</i> , 2002, 100, 2367-2373.	0.6	737
106	Intravenous Immunoglobulin for Hemophagocytic Lymphohistiocytosis?. <i>Journal of Clinical Oncology</i> , 2002, 20, 599-601.	0.8	50
107	Low natural killer activity and central nervous system disease as a high-risk prognostic indicator in young patients with hemophagocytic lymphohistiocytosis. <i>Cancer</i> , 2002, 94, 3023-3031.	2.0	50
108	Risk of Etoposide-Related Acute Myeloid Leukemia in the Treatment of Epstein-Barr Virus-Associated Hemophagocytic Lymphohistiocytosis. <i>International Journal of Hematology</i> , 2002, 75, 174-177.	0.7	37

#	ARTICLE	IF	CITATIONS
109	Clinical features and treatment strategies of Epstein-Barr virus-associated hemophagocytic lymphohistiocytosis. <i>Critical Reviews in Oncology/Hematology</i> , 2002, 44, 259-272.	2.0	245
110	Recent developments in the management of haemophagocytic lymphohistiocytosis. <i>Expert Opinion on Pharmacotherapy</i> , 2001, 2, 1437-1448.	0.9	41
111	Hemophagocytic lymphohistiocytosis due to germline mutations in SH2D1A, the X-linked lymphoproliferative disease gene. <i>Blood</i> , 2001, 97, 1131-1133.	0.6	148
112	Requirement for Etoposide in the Treatment of Epstein-Barr Virus-Associated Hemophagocytic Lymphohistiocytosis. <i>Journal of Clinical Oncology</i> , 2001, 19, 2665-2673.	0.8	232
113	Effect of chemotherapy and stem cell transplantation on T-lymphocyte clones in familial haemophagocytic lymphohistiocytosis. <i>British Journal of Haematology</i> , 2001, 113, 822-831.	1.2	5
114	Letter to the editor: Eyelid leukemia as a relapse sign of B-cell type acute lymphoblastic leukemia. <i>Medical and Pediatric Oncology</i> , 2001, 36, 505-506.	1.0	21
115	Secondary acute myelocytic leukemia after successful chemotherapy with etoposide for Epstein-Barr virus-associated hemophagocytic lymphohistiocytosis. <i>Medical and Pediatric Oncology</i> , 2001, 37, 153-154.	1.0	19
116	Hemophagocytosis by leukemic blasts in 7 acute myeloid leukemia cases with t(16;21)(p11;q22). , 2000, 88, 1970-1975.		39
117	Sudden death of a 21-year-old female with Williams syndrome showing rare complications. <i>Pediatrics International</i> , 2000, 42, 322-324.	0.2	17
118	Splenectomy in haemophagocytic lymphohistiocytosis: report of histopathological changes with CD19+ B-cell depletion and therapeutic results. <i>British Journal of Haematology</i> , 2000, 108, 505-510.	1.2	15
119	Management of Severe Neutropenia With Cyclosporin During Initial Treatment of Epstein-Barr Virus-Related Hemophagocytic Lymphohistiocytosis. <i>Leukemia and Lymphoma</i> , 2000, 36, 339-346.	0.6	34
120	Molecular Analysis of Latent Membrane Protein 1 in Patients with Epstein-Barr Virus-Associated Hemophagocytic Lymphohistiocytosis in Japan. <i>Leukemia and Lymphoma</i> , 2000, 38, 373-380.	0.6	17
121	Outcome of Clonal Hemophagocytic Lymphohistiocytosis: Analysis of 32 Cases. <i>Leukemia and Lymphoma</i> , 2000, 37, 577-584.	0.6	51
122	Treatment Strategies for Epstein-Barr Virus-Associated Hemophagocytic Lymphohistiocytosis (EBV-HLH). <i>Leukemia and Lymphoma</i> , 2000, 39, 37-49.	0.6	46
123	Hemophagocytosis by leukemic blasts in 7 acute myeloid leukemia cases with t(16;21)(p11;q22). , 2000, 88, 1970.		6
124	Mixed-lineage leukemia with t(10;11)(p13;q21): An analysis of AF10-CALM and CALM-AF10 fusion mRNAs and clinical features. , 1999, 25, 33-39.		46
125	Hemophagocytic syndrome in five patients with Epstein-Barr virus negative B-cell lymphoma. <i>Cancer</i> , 1999, 85, 2298-2299.	2.0	12
126	Clinical and epidemiologic studies of familial hemophagocytic lymphohistiocytosis in Japan. , 1998, 30, 276-283.		48

#	ARTICLE	IF	CITATIONS
127	Biomarker and morphological characteristics of Epstein-Barr virus-related hemophagocytic lymphohistiocytosis. , 1998, 31, 131-137.		39
128	FAMILIAL HEMOPHAGOCYTIC LYMPHOHISTIOCYTOSIS. Hematology/Oncology Clinics of North America, 1998, 12, 417-433.	0.9	241
129	INFECTION- AND MALIGNANCY-ASSOCIATED HEMOPHAGOCYTIC SYNDROMES. Hematology/Oncology Clinics of North America, 1998, 12, 435-444.	0.9	376
130	Heterogeneity of Immune Markers in Hemophagocytic Lymphohistiocytosis. Journal of Pediatric Hematology/Oncology, 1998, 20, 207-214.	0.3	26
131	Hemophagocytic lymphohistiocytosis in infancy and childhood. Journal of Pediatrics, 1997, 130, 352-357.	0.9	123
132	Haemophagocytic lymphohistiocytosis in association with granular lymphocyte proliferative disorders in early childhood: characteristic bone marrow morphology. British Journal of Haematology, 1997, 96, 708-714.	1.2	21
133	Philadelphia chromosome-positive acute lymphoblastic leukemia after therapy for langerhans cell histiocytosis. , 1997, 54, 88-88.		10
134	HLH-94: A treatment protocol for hemophagocytic lymphohistiocytosis. , 1997, 28, 342-347.		417
135	Effects of thrombopoietin (<i>câ€mpl</i> ligand) on growth of blast cells from patients with transient abnormal myelopoiesis and acute myeloblastic leukemia. European Journal of Haematology, 1997, 59, 38-46.	1.1	18
136	Hyperâ€interleukin (IL)â€naemia in haemophagocytic lymphohistiocytosis. British Journal of Haematology, 1996, 93, 803-807.	1.2	40
137	Myelodysplasia and acute myeloid leukaemia in cases of aplastic anaemia and congenital neutropenia following Câ€CSF administration. British Journal of Haematology, 1995, 89, 188-190.	1.2	55
138	Serum and urine beta---2-microglobulin in hemophagocytic syndrome. Cancer, 1995, 75, 1700-1705.	2.0	36
139	Haemophagocytic lymphohistiocytosis, interferon-gamma-naemia and Epstein-Barr virus involvement. British Journal of Haematology, 1994, 88, 656-658.	1.2	86
140	Effect of Recombinant Human Granulocyte-Colony Stimulation Factor (rhG-CSF) on Immune System in Pediatric Patients with Aplastic Anemia. Pediatric Hematology and Oncology, 1994, 11, 319-323.	0.3	2
141	INCREASE OF GAMMA/DELTA T CELLS IN A CASE OF TYPHOID FEVER. British Journal of Haematology, 1993, 83, 177-178.	1.2	3
142	Hypercytokinemia in Hemophagocytic Syndrome. Journal of Pediatric Hematology/Oncology, 1993, 15, 92-98.	0.3	216
143	Unique expression of integrin fibronectin receptors in human neuroblastoma cell lines. International Journal of Cancer, 1992, 51, 620-626.	2.3	16
144	CYTOKINE LEVELS IN AGGRESSIVE NATURAL KILLER CELL LEUKAEMIA AND MALIGNANT HISTIOCYTOSIS. British Journal of Haematology, 1991, 79, 132-133.	1.2	12

#	ARTICLE	IF	CITATIONS
145	CYTOKINES IN HAEMOPHAGOCYTIC SYNDROME. British Journal of Haematology, 1991, 77, 438-440.	1.2	48
146	Hemophagocytic syndrome associated with aggressive natural killer cell leukemia. American Journal of Hematology, 1991, 38, 321-323.	2.0	66
147	Serum Levels of Interferon-gamma, Cytotoxic Factor and Soluble Interleukin-2 Receptor in Childhood Hemophagocytic Syndromes. Leukemia and Lymphoma, 1991, 3, 287-292.	0.6	71
148	Target cell of leukemic transformation in acute megakaryoblastic leukemia. American Journal of Hematology, 1990, 34, 252-258.	2.0	18
149	Impaired natural killer activity and expression of interleukin-2 receptor antigen in familial erythrophagocytic lymphohistiocytosis. Cancer, 1990, 65, 1937-1941.	2.0	35
150	Allogeneic Bone Marrow Transplantation for Familial Erythrophagocytic Lymphohistiocytosis, with High Dose VP16-Containing Conditioning Regimen. Leukemia and Lymphoma, 1990, 1, 361-364.	0.6	9
151	Procoagulant Activity of Human Neuroblastoma Cell Lines, in Relation to Cell Growth, Differentiation and Cytogenetic Abnormalities. Japanese Journal of Cancer Research, 1989, 80, 438-443.	1.7	10
152	High serum ferritin level as a marker of malignant histiocytosis and virus-associated hemophagocytic syndrome. Cancer, 1988, 61, 2071-2076.	2.0	109
153	Diagnosis of neuroblastoma metastasis in bone marrow with a panel of monoclonal antibodies. Medical and Pediatric Oncology, 1988, 16, 190-196.	1.0	14
154	Malignant Histiocytosis in Childhood Clinical Features and Therapeutic Results by Combination Chemotherapy. Journal of Pediatric Hematology/Oncology, 1986, 8, 300-307.	0.3	18
155	Enzyme Defect in a Case of Tyrosinemia Type I, Acute Form. Pediatric Research, 1984, 18, 463-466.	1.1	9
156	Studies on tyrosine hydroxylase in neuroblastoma, in relation to urinary levels of catecholamine metabolites. Cancer, 1975, 36, 450-457.	2.0	15
157	Ultrastructural Observations of Functional Neural Tumor. Pediatrics International, 1975, 17, 46-46.	0.2	0
158	Studies on the Levels of Urinary VMA and HVA in Ten Cases of Neuroblastoma. Pediatrics International, 1972, 14, 19-19.	0.2	0
159	Cyclosporine Treatment in a Case of Acquired Hemophilia A Diagnosed After Tooth Extraction. Medical Science Case Reports, 0, 4, 24-27.	0.0	1
160	¹⁸ F-fluorodeoxyglucose-positron emission tomography/computed tomography delineates involved sites in the cervical spine in Langerhans cell histiocytosis. EJHaem, 0, , .	0.4	0
161	In situ diffuse large B-cell lymphoma in haemorrhoidectomy tissue. EJHaem, 0, , .	0.4	0