Hirokazu Kanegane

List of Publications by Year
in descending order

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167 papers	5,066 citations	126907 33 h-index	106344 65 g-index
178 all docs	178 docs citations	178 times ranked	5747 citing authors

#	Article	IF	Citations
1	Phenotype, penetrance, and treatment of 133 cytotoxic T-lymphocyte antigen 4â€“insufficient subjects. Journal of Allergy and Clinical Immunology, 2018, 142, 1932-1946.	2.9	344
2	Clinical similarities and differences of patients with X-linked lymphoproliferative syndrome type 1 (XLP-1/SAP deficiency) versus type 2 (XLP-2/XIAP deficiency). Blood, 2011, 117, 1522-1529.	1.4	320
3	X-linked lymphoproliferative disease due to SAP/SH2D1A deficiency: a multicenter study on the manifestations, management and outcome of the disease. Blood, 2011, 117, 53-62.	1.4	268
4	Long-term follow-up of IPEX syndrome patients after different therapeutic strategies: An international multicenter retrospective study. Journal of Allergy and Clinical Immunology, 2018, 141, 1036-1049.e5.	2.9	233
5	XIAP Restricts TNF- and RIP3-Dependent Cell Death and Inflammasome Activation. Cell Reports, 2014, 7, 1796-1808.	6.4	210
6	Genetic analysis of patients with defects in early Bâ€cell development. Immunological Reviews, 2005, 203, 216-234.	6.0	170
7	Dominant-negative IKZF1 mutations cause a T, B, and myeloid cell combined immunodeficiency. Journal of Clinical Investigation, 2018, 128, 3071-3087.	8.2	133
8	Allogeneic hematopoietic cell transplantation for XIAP deficiency: an international survey reveals poor outcomes. Blood, 2013, 121, 877-883.	1.4	132
9	Sustained elevation of serum interleukin-18 and its association with hemophagocytic lymphohistiocytosis in XIAP deficiency. Cytokine, 2014, 65, 74-78.	3.2	112
10	Identification of Severe Combined Immunodeficiency by T-Cell Receptor Excision Circles Quantification Using Neonatal Guthrie Cards. Journal of Pediatrics, 2009, 155, 829-833.	1.8	108
11	Characterization of Crohn disease in X-linked inhibitor of apoptosisâ€deficient male patients and female symptomatic carriers. Journal of Allergy and Clinical Immunology, 2014, 134, 1131-1141.e9.	2.9	101
12	The kinase Btk negatively regulates the production of reactive oxygen species and stimulation-induced apoptosis in human neutrophils. Nature Immunology, 2012, 13, 369-378.	14.5	100
13	Haploinsufficiency of A20 causes autoinflammatory and autoimmune disorders. Journal of Allergy and Clinical Immunology, 2018, 141, 1485-1488.e11.	2.9	100
14	Abnormal hematopoiesis and autoimmunity in human subjects with germline IKZF1 mutations. Journal of Allergy and Clinical Immunology, 2017, 140, 223-231.	2.9	99
15	Flow cytometry-based diagnosis of primary immunodeficiency diseases. Allergology International, 2018, 67, 43-54.	3.3	97
16	Haploinsufficiency of TNFAIP3 (A20) by germline mutation is involved in autoimmune lymphoproliferative syndrome. Journal of Allergy and Clinical Immunology, 2017, 139, 1914-1922.	2.9	91
17	Clinical and mutational characteristics of X-linked agammaglobulinemia and its carrier identified by flow cytometric assessment combined with genetic analysis. Journal of Allergy and Clinical Immunology, 2001, 108, 1012-1020.	2.9	87
18	Clinical and Genetic Characteristics of XIAP Deficiency in Japan. Journal of Clinical Immunology, 2012, 32, 411-420.	3.8	84

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19	Nationwide Survey of Patients with Primary Immunodeficiency Diseases in Japan. Journal of Clinical Immunology, 2011, 31, 968-976.	3.8	77
20	Identification of FOXP3-negative regulatory T-like (CD4+CD25+CD127low) cells in patients with immune dysregulation, polyendocrinopathy, enteropathy, X-linked syndrome. Clinical Immunology, 2011, 141, 111-120.	3.2	74
21	Neutrophils and Mononuclear Cells Express Vascular Endothelial Growth Factor in Acute Kawasaki Disease: Its Possible Role in Progression of Coronary Artery Lesions. Pediatric Research, 2001, 49, 74-80.	2.3	72
22	Human CD19 and CD40L deficiencies impair antibody selection and differentially affect somatic hypermutation. Journal of Allergy and Clinical Immunology, 2014, 134, 135-144.e7.	2.9	71
23	Gain-of-function IKBKB mutation causes human combined immune deficiency. Journal of Experimental Medicine, 2018, 215, 2715-2724.	8.5	69
24	Hematopoietic Stem Cell Transplantation for XIAP Deficiency in Japan. Journal of Clinical Immunology, 2017, 37, 85-91.	3.8	63
25	Biological aspects of Epsteinâ€"Barr virus (EBV)-infected lymphocytes in chronic active EBV infection and associated malignancies. Critical Reviews in Oncology/Hematology, 2002, 44, 239-249.	4.4	59
26	Female agammaglobulinemia due to the Bruton tyrosine kinase deficiency caused by extremely skewed X-chromosome inactivation. Blood, 2004, 103, 185-187.	1.4	59
27	Activation-dependent T cell expression of the X-linked lymphoproliferative disease gene product SLAM-associated protein and its assessment for patient detection. International Immunology, 2002, 14, 1215-1223.	4.0	54
28	Multicolor Flow Cytometry for the Diagnosis of Primary Immunodeficiency Diseases. Journal of Clinical Immunology, 2017, 37, 486-495.	3.8	42
29	Ionizing radiation induces apoptotic cell death in human TcR-Î³/Î´+ T and natural killer cells without detectable p53 protein. European Journal of Immunology, 1994, 24, 2914-2917.	2.9	40
30	Atypical X-linked Agammaglobulinemia Diagnosed in Three Adults.. Internal Medicine, 1999, 38, 722-725.	0.7	39
31	EBV-NK Cells Interactions and Lymphoproliferative Disorders. Leukemia and Lymphoma, 1998, 29, 491-498.	1.3	38
32	X-linked lymphoproliferative syndrome presenting with systemic lymphocytic vasculitis. American Journal of Hematology, 2005, 78, 130-133.	4.1	36
33	Targeted Sequencing and Immunological Analysis Reveal the Involvement of Primary Immunodeficiency Genes in Pediatric IBD: a Japanese Multicenter Study. Journal of Clinical Immunology, 2017, 37, 67-79.	3.8	36
34	Heterozygous OAS1 gain-of-function variants cause an autoinflammatory immunodeficiency. Science Immunology, 2021, 6, .	11.9	36
35	SAP and XIAP deficiency in hemophagocytic lymphohistiocytosis. Pediatrics International, 2012, 54, 447-454.	0.5	35
36	Severe Neutropenia in Japanese Patients with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2005, 25, 491-495.	3.8	34

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37	APRIL-dependent lifelong plasmacyte maintenance and immunoglobulin production in humans. Journal of Allergy and Clinical Immunology, 2020, 146, 1109-1120.e4.	2.9	33
38	A variant in human AIOLOS impairs adaptive immunity by interfering with IKAROS. Nature Immunology, 2021, 22, 893-903.	14.5	33
39	Outcomes in children with hemophagocytic lymphohistiocytosis treated using HLH-2004 protocol in Japan. International Journal of Hematology, 2019, 109, 206-213.	1.6	32
40	X-linked thrombocytopenia identified by flow cytometric demonstration of defective Wiskott-Aldrich syndrome protein in lymphocytes. Blood, 2000, 95, 1110-1111.	1.4	30
41	Early lineage switch in an infant acute lymphoblastic leukemia. International Journal of Hematology, 2009, 90, 653-655.	1.6	30
42	Somatic Mosaicism for a NRAS Mutation Associates with Disparate Clinical Features in RAS-associated Leukoproliferative Disease: a Report of Two Cases. Journal of Clinical Immunology, 2015, 35, 454-458.	3.8	30
43	Detailed analysis of Japanese patients with adenosine deaminase 2 deficiency reveals characteristic elevation of type II interferon signature and STAT1 hyperactivation. Journal of Allergy and Clinical Immunology, 2021, 148, 550-562.	2.9	30
44	Clinical features and outcome of Xâ€linked lymphoproliferative syndrome type 1 (SAP deficiency) in Japan identified by the combination of flow cytometric assay and genetic analysis. Pediatric Allergy and Immunology, 2012, 23, 488-493.	2.6	29
45	Efficacy and Safety of IgPro20, a Subcutaneous Immunoglobulin, in Japanese Patients with Primary Immunodeficiency Diseases. Journal of Clinical Immunology, 2014, 34, 204-211.	3.8	29
46	Clinical and Immunological Characterization of ICF Syndrome in Japan. Journal of Clinical Immunology, 2018, 38, 927-937.	3.8	29
47	A Female Patient with Incomplete Hemophagocytic Lymphohistiocytosis Caused by a Heterozygous XIAP Mutation Associated with Non-Random X-Chromosome Inactivation Skewed Towards the Wild-Type XIAP Allele. Journal of Clinical Immunology, 2015, 35, 244-248.	3.8	28
48	Robust and highly efficient hiPSC generation from patient non-mobilized peripheral blood-derived CD34+ cells using the auto-erasable Sendai virus vector. Stem Cell Research and Therapy, 2019, 10, 185.	5.5	28
49	Autoimmune lymphoproliferative syndrome presenting with glomerulonephritis. Pediatric Nephrology, 2003, 18, 454-456.	1.7	27
50	Allogeneic stem cell transplantation for X-linked agammaglobulinemia using reduced intensity conditioning as a model of the reconstitution of humoral immunity. Journal of Hematology and Oncology, 2016, 9, 9.	17.0	27
51	Late-Onset Combined Immunodeficiency with a Novel IL2RG Mutation and Probable Revertant Somatic Mosaicism. Journal of Clinical Immunology, 2015, 35, 610-614.	3.8	26
52	Heterozygous Mutations in OAS1 Cause Infantile-Onset Pulmonary Alveolar Proteinosis with Hypogammaglobulinemia. American Journal of Human Genetics, 2018, 102, 480-486.	6.2	26
53	S100A4 Protein Is Essential for the Development of Mature Microfold Cells in Peyerâ€™s Patches. Cell Reports, 2019, 29, 2823-2834.e7.	6.4	25
54	Identification of DKC1 gene mutations in Japanese patients with Xâ€linked dyskeratosis congenita. British Journal of Haematology, 2005, 129, 432-434.	2.5	22

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55	Cost-minimization Analysis of IgPro20, a Subcutaneous Immunoglobulin, in Japanese Patients With Primary Immunodeficiency. Clinical Therapeutics, 2014, 36, 1616-1624.	2.5	22
56	Dysregulation of Epstein-Barr Virus Infection in Hypomorphic ZAP70 Mutation. Journal of Infectious Diseases, 2018, 218, 825-834.	4.0	22
57	Mechanisms of genotype-phenotype correlation in autosomal dominant anhidrotic ectodermal dysplasia with immune deficiency. Journal of Allergy and Clinical Immunology, 2018, 141, 1060-1073.e3.	2.9	22
58	Atypical SIFD with novel TRNT1 mutations: a case study on the pathogenesis of B-cell deficiency. International Journal of Hematology, 2019, 109, 382-389.	1.6	22
59	Kawasaki disease with a concomitant primary Epstein-Barr virus infection. Pediatrics International, 1994, 36, 713-716.	0.5	21
60	Infectious mononucleosis as a disease of early childhood in Japan caused by primary Epsteinâ€Barr virus infection. Pediatrics International, 1997, 39, 166-171.	0.5	20
61	X-Linked Agammaglobulinemia Associated with B-Precursor Acute Lymphoblastic Leukemia. Journal of Clinical Immunology, 2015, 35, 108-111.	3.8	20
62	Early and rapid detection of X-linked lymphoproliferative syndrome with SH2D1A mutations by flow cytometry. , 2011, 80B, 8-13.		19
63	Extensive serum biomarker analysis in patients with enterohemorrhagic Escherichia coli O111-induced hemolytic-uremic syndrome. Cytokine, 2014, 66, 1-6.	3.2	18
64	Genetic heterogeneity of uncharacterized childhood autoimmune diseases with lymphoproliferation. Pediatric Blood and Cancer, 2018, 65, e26831.	1.5	18
65	High frequencies of asymptomatic Epstein-Barr virus viremia in affected and unaffected individuals with CTLA4 mutations. Clinical Immunology, 2018, 195, 45-48.	3.2	18
66	X-linked lymphoproliferative syndrome in mainland China: review of clinical, genetic, and immunological characteristic. European Journal of Pediatrics, 2020, 179, 327-338.	2.7	18
67	Treatment Satisfaction with Subcutaneous Immunoglobulin Replacement Therapy in Patients with Primary Immunodeficiency: a Pooled Analysis of Six HizentraÂ® Studies. Journal of Clinical Immunology, 2018, 38, 886-897.	3.8	17
68	XIAP restrains TNF-driven intestinal inflammation and dysbiosis by promoting innate immune responses of Paneth and dendritic cells. Science Immunology, 2021, 6, eabf7235.	11.9	17
69	Autoimmune lymphoproliferative syndrome mimicking chronic active Epsteinâ€“Barr virus infection. International Journal of Hematology, 2011, 93, 760-764.	1.6	16
70	Bruton tyrosine kinase gene mutations in Turkish patients with presumed X-linked agammaglobulinemia. Human Mutation, 2001, 18, 356-356.	2.5	15
71	Identification of mutations in the Bruton's tyrosine kinase gene, including a novel genomic rearrangements resulting in large deletion, in Korean X-linked agammaglobulinemia patients. Journal of Human Genetics, 2003, 48, 322-326.	2.3	15
72	Functional analysis of novel A20 variants in patients with atypical inflammatory diseases. Arthritis Research and Therapy, 2021, 23, 52.	3.5	15

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73	Hematopoietic Cell Transplantation Rescues Inflammatory Bowel Disease and Dysbiosis of Gut Microbiota in XIAP Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 3767-3780.	3.8	15
74	Spontaneous regression of aleukemic leukemia cutis harboring a NPM/RARA fusion gene in an infant with cutaneous mastosytosis. International Journal of Hematology, 2009, 89, 86-90.	1.6	14
75	Droplet Digital PCR-Based Chimerism Analysis for Primary Immunodeficiency Diseases. Journal of Clinical Immunology, 2018, 38, 300-306.	3.8	14
76	Prospective Study of Allogeneic Hematopoietic Stem Cell Transplantation with Post-Transplantation Cyclophosphamide and Antithymocyte Globulin from HLA-Mismatched Related Donors for Nonmalignant Diseases. Biology of Blood and Marrow Transplantation, 2020, 26, e286-e291.	2.0	14
77	Population Pharmacokinetics of Intravenous Busulfan in Japanese Pediatric Patients With Primary Immunodeficiency Diseases. Journal of Clinical Pharmacology, 2018, 58, 327-331.	2.0	13
78	Current Perspectives and Unmet Needs of Primary Immunodeficiency Care in Asia Pacific. Frontiers in Immunology, 2020, 11, 1605.	4.8	13
79	Epstein-Barr Virus-Associated Î³Î´ T-Cell Lymphoproliferative Disorder Associated With Hypomorphic IL2RG Mutation. Frontiers in Pediatrics, 2019, 7, 15.	1.9	12
80	Comprehensive Targeted Sequencing Identifies Monogenic Disorders in Patients With Earlyâ€onset Refractory Diarrhea. Journal of Pediatric Gastroenterology and Nutrition, 2020, 71, 333-339.	1.8	12
81	Longâ€term outcomes of children with extracutaneous juvenile xanthogranulomas in Japan. Pediatric Blood and Cancer, 2020, 67, e28381.	1.5	12
82	Development of EBV-Positive T-cell Lymphoma Following Infection of Peripheral Blood T Cells with EBV. Leukemia and Lymphoma, 1999, 34, 603-607.	1.3	11
83	Serum tau protein as a marker of disease activity in enterohemorrhagic Escherichia coli O111-induced hemolytic uremic syndrome. Neurochemistry International, 2015, 85-86, 24-30.	3.8	11
84	A synonymous splice site mutation in IL2RG gene causes late-onset combined immunodeficiency. International Journal of Hematology, 2019, 109, 603-611.	1.6	11
85	Hematopoietic stem cell transplantation for pulmonary alveolar proteinosis associated with primary immunodeficiency disease. International Journal of Hematology, 2018, 107, 610-614.	1.6	10
86	IKBA S32 Mutations Underlie Ectodermal Dysplasia with Immunodeficiency and Severe Noninfectious Systemic Inflammation. Journal of Clinical Immunology, 2018, 38, 543-545.	3.8	10
87	Clinical and Immunological Heterogeneity in Japanese Patients with Gain-of-Function Variants in STAT3. Journal of Clinical Immunology, 2021, 41, 780-790.	3.8	10
88	Maternal germinal mosaicism of Xâ€linked agammaglobulinemia. American Journal of Medical Genetics Part A, 2001, 99, 234-237.	2.4	9
89	Hematopoietic stem cell transplantation with reduced intensity conditioning from a family haploidentical donor in an infant with familial hemophagocytic lymphohistocytosis. International Journal of Hematology, 2011, 94, 285-290.	1.6	9
90	Neonatal acute megakaryoblastic leukemia mimicking congenital neuroblastoma. Clinical Case Reports (discontinued), 2015, 3, 145-149.	0.5	9

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91	Maternal T and B cell engraftment in two cases of X-linked severe combined immunodeficiency with IgG1 gammopathy. Clinical Immunology, 2017, 183, 112-120.	3.2	9
92	B-lymphoblastic lymphoma with TCF3-PBX1 fusion gene. Haematologica, 2019, 104, e35-e37.	3.5	9
93	Nationwide retrospective review of hematopoietic stem cell transplantation in children with refractory Langerhans cell histiocytosis. International Journal of Hematology, 2020, 111, 137-148.	1.6	9
94	Novel AP3B1 compound heterozygous mutations in a Japanese patient with Hermanskyâ€“Pudlak syndrome type 2. Journal of Dermatology, 2020, 47, 185-189.	1.2	9
95	Impaired B-Cell Differentiation in a Patient With STAT1 Gain-of-Function Mutation. Frontiers in Immunology, 2020, 11, 557521.	4.8	9
96	Hematopoietic Cell Transplantation with Reduced Intensity Conditioning Using Fludarabine/Busulfan or Fludarabine/Melphalan for Primary Immunodeficiency Diseases. Journal of Clinical Immunology, 2021, 41, 944-957.	3.8	9
97	Hematopoietic Cell Transplantation Ameliorates Autoinflammation in A20 Haploinsufficiency. Journal of Clinical Immunology, 2021, 41, 1954-1956.	3.8	9
98	Effect of reducedâ€intensity conditioning and the risk of lateâ€onset nonâ€infectious pulmonary complications in pediatric patients. European Journal of Haematology, 2017, 99, 525-531.	2.2	8
99	Factors predicting the recurrence of Epsteinâ€“Barr virus-associated hemophagocytic lymphohistiocytosis in children after treatment using the HLH-2004 protocol. International Journal of Hematology, 2019, 109, 612-617.	1.6	8
100	Modification of cellular and humoral immunity by somatically reverted T cells in X-linked lymphoproliferative syndrome type 1. Journal of Allergy and Clinical Immunology, 2019, 143, 421-424.e11.	2.9	8
101	Helicobacter cinaedi-Associated Refractory Cellulitis in Patients with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2020, 40, 1132-1137.	3.8	8
102	A Novel Homozygous Mutation Destabilizes IKKÎ² and Leads to Human Combined Immunodeficiency. Frontiers in Immunology, 2020, 11, 517544.	4.8	8
103	Acute thrombocytopenic purpura associated with primary Epsteinâ€Barr virus infection. Pediatrics International, 1994, 36, 423-426.	0.5	7
104	Non-Hodgkin's Lymphoma of the Ascending Colon in a Patient with Becker Muscular Dystrophy: Report of a Case. Surgery Today, 2001, 31, 1016-1019.	1.5	7
105	Identification of novel fusion genes with 28S ribosomal DNA in hematologic malignancies. International Journal of Oncology, 2014, 44, 1193-1198.	3.3	7
106	Identification of Novel Fusion Genes with 28S Ribosomal DNA in Hematologic Malignancies. Blood, 2012, 120, 4418-4418.	1.4	7
107	Mononucleosisâ€like illness in an infant associated with human herpesvirus 6 infection. Pediatrics International, 1995, 37, 227-229.	0.5	6
108	Successful treatment of very large congenital infantile fibrosarcoma. Pediatrics International, 2011, 53, 768-770.	0.5	6

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109	Novel compound heterozygous mutations in a Japanese girl with Janus kinase 3 deficiency. Pediatrics International, 2016, 58, 1076-1080.	0.5	6
110	Inflammatory bowel diseases and primary immunodeficiency diseases. Immunological Medicine, 2018, 41, 154-161.	2.6	6
111	Comprehensive molecular diagnosis of Epsteinâ€“Barr virus-associated lymphoproliferative diseases using next-generation sequencing. International Journal of Hematology, 2018, 108, 319-328.	1.6	6
112	Inherited chromosomally integrated human herpesvirusâ€6 in a patient with XIAP deficiency. Transplant Infectious Disease, 2020, 22, e13331.	1.7	6
113	A case of autoimmune enteropathy with CTLA4 haploinsufficiency. Intestinal Research, 2022, 20, 144-149.	2.6	6
114	Identification of Germline Non-coding Deletions in XIAP Gene Causing XIAP Deficiency Reveals a Key Promoter Sequence. Journal of Clinical Immunology, 2022, 42, 559-571.	3.8	6
115	Reactive peripheral blood plasmacytosis in Kawasaki disease. Pediatrics International, 2018, 60, 884-885.	0.5	5
116	Immunophenotyping of A20 haploinsufficiency by multicolor flow cytometry. Clinical Immunology, 2020, 216, 108441.	3.2	5
117	<i>BRAF</i> V600E-positive cells as molecular markers of bone marrow disease in pediatric Langerhans cell histiocytosis. Haematologica, 2022, 107, 1719-1725.	3.5	5
118	Distinct Clones are Associated with the Development of Transient Myeloproliferative Disorder and Acute Megakaryocytic Leukemia in a Patient with Down Syndrome. International Journal of Hematology, 2007, 86, 250-252.	1.6	5
119	Point mutation in intron 11 of Bruton's tyrosine kinase in atypical X-linked agammaglobulinemia. Pediatrics International, 2000, 42, 689-692.	0.5	4
120	Three brothers of X-linked agammaglobulinemia: the relation between phenotype and neutropenia. International Journal of Hematology, 2009, 90, 117-119.	1.6	4
121	Successful bone marrow transplantation with reduced intensity conditioning in a patient with delayedâ€onset adenosine deaminase deficiency. Pediatric Transplantation, 2013, 17, E29-32.	1.0	4
122	Intravenous immunoglobulin (IVIG) efficiency in women with common variable immunodeficiency (CVID) decreases significantly during pregnancy. Journal of Maternal-Fetal and Neonatal Medicine, 2019, 32, 3092-3096.	1.5	4
123	Mycobacterium genavense Infection Presenting as an Endobronchial Polyp and Upper Lobe Atelectasis. American Journal of Respiratory and Critical Care Medicine, 2020, 202, e144-e145.	5.6	4
124	DNA Ligase IV Deficiency Identified by Chance Following Vaccine-Derived Rubella Virus Infection. Journal of Clinical Immunology, 2020, 40, 1187-1190.	3.8	4
125	Cytomegalovirus Laryngitis in Primary Combined Immunodeficiency Diseases. Journal of Clinical Immunology, 2021, 41, 243-247.	3.8	4
126	Clinical Courses of IKAROS and CTLA4 Deficiencies: A Systematic Literature Review and Retrospective Longitudinal Study. Frontiers in Immunology, 2021, 12, 784901.	4.8	4

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127	Advances in Understanding the Pathogenesis of Epstein-Barr Virus-Associated Lymphoproliferative Disorders. Iranian Journal of Allergy, Asthma and Immunology, 2015, 14, 462-71.	0.4	4
128	Acute Epsteinâ€"Barr virus infection presenting as severe gastroenteritis without infectious mononucleosis-like manifestations. Clinical Journal of Gastroenterology, 2009, 2, 398-403.	0.8	3
129	Mislocalization or low expression of mutated Shwachmanâ€"Bodianâ€"Diamond syndrome protein. International Journal of Hematology, 2011, 94, 54-62.	1.6	3
130	A Cry for the Development of Newborn Screening for Familial Hemophagocytic Lymphohistiocytosis. Journal of Clinical Immunology, 2020, 40, 1196-1198.	3.8	3
131	Endocrinopathies in Inborn Errors of Immunity. Frontiers in Immunology, 2021, 12, 786241.	4.8	3
132	Cartilageâ€hair hypoplasia with Tâ€cell dysfunction. Pediatrics International, 2022, 64, e15080.	0.5	3
133	Hematopoietic cell transplantation for asymptomatic X-linked lymphoproliferative syndrome type 1. Allergy, Asthma and Clinical Immunology, 2018, 14, 82.	2.0	2
134	Early Surgery Is Feasible for a Very Large Congenital Infantile Fibrosarcoma Associated With Life Threatening Coagulopathy: A Case Report and Literature Review. Frontiers in Pediatrics, 2019, 7, 529.	1.9	2
135	Highâ€throughput analysis revealed the unique immunoglobulin gene rearrangements in plasmacytomaâ€like postâ€transplant lymphoproliferative disorder. British Journal of Haematology, 2020, 189, e164-e168.	2.5	2
136	Case Report: Infantile-Onset Fulminant Type 1 Diabetes Mellitus Caused by Novel Compound Heterozygous LRBA Variants. Frontiers in Immunology, 2021, 12, 677572.	4.8	2
137	Wiskott-Aldrich Syndrome: A Retrospective Study on 575 Patients Analyzing the Impact of Splenectomy, Stem Cell Transplantation, or No Definitive Treatment on Frequency of Disease-Related Complications and Physician-Perceived Quality of Life. Blood, 2016, 128, 366-366.	1.4	2
138	Case Report: Rotavirus Vaccination and Severe Combined Immunodeficiency in Japan. Frontiers in Immunology, 2022, 13, 786375.	4.8	2
139	A CIAS1 mutation in a Japanese girl with familial cold autoinflammatory syndrome. European Journal of Pediatrics, 2008, 167, 245-247.	2.7	1
140	M-protein-positive chronic active Epsteinâ€"Barr virus infection: features mimicking HIV-1 infection. International Journal of Hematology, 2009, 90, 235-238.	1.6	1
141	Intracranial calcification in a uremic infant with Wilmsâ€™ tumor in a solitary kidney. CEN Case Reports, 2012, 1, 86-89.	0.9	1
142	Hematopoietic cell transplantation with reduced intensity conditioning using fludarabine and busulfan for X-linked hyper IgM syndrome. Journal of Hematopoietic Cell Transplantation, 2019, 8, 43-49.	0.1	1
143	Epstein-Barr Virus (EBV)-induced B-cell Lymphoproliferative Disorder Mimicking the Recurrence of EBV-associated Hemophagocytic Lymphohistiocytosis. Journal of Pediatric Hematology/Oncology, 2019, 41, e44-e46.	0.6	1
144	Fatal Progressive Meningoencephalitis Diagnosed in Two Members of a Family With X-Linked Agammaglobulinemia. Frontiers in Pediatrics, 2020, 8, 579.	1.9	1

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145	Successful Artery Embolization in a Patient with Autoimmune Lymphoproliferative Syndrome Associated with Splenic Rupture. Journal of Clinical Immunology, 2020, 40, 780-782.	3.8	1
146	Epstein-Barr Virusâ€“Negative Granulomatous Disease Due to SAP Deficiency. Journal of Clinical Immunology, 2021, 41, 1372-1375.	3.8	1
147	Genomics analysis of leukaemia predisposition in Xâ€linked agammaglobulinaemia. British Journal of Haematology, 2021, 193, 1277-1281.	2.5	1
148	Hematopoietic Stem Cell Transplantation in Children with Refractory Langerhans Cell Histiocytosis. Blood, 2018, 132, 4657-4657.	1.4	1
149	Genetic Basis of Myeloid Proliferation Related to Down Syndrome. Blood, 2012, 120, 535-535.	1.4	1
150	Graft versus tumor effect against neuroblastoma: a case report with long-term survival and a review of the literature. Journal of Hematopoietic Cell Transplantation, 2014, 3, 97-101.	0.1	1
151	A sporadic case of CTLA4 haploinsufficiency manifesting as Epsteinâ€“Barr virus-positive diffuse large B-cell lymphoma. Journal of Clinical and Experimental Hematopathology: JCEH, 2021, , .	0.8	1
152	Early diagnosis of partial interferon-Î³ receptor 1 deficiency prevents the development of Bacille de Calmette et GuÃ©rin osteomyelitis. Clinical Immunology, 2022, 235, 108933.	3.2	1
153	Preemptive hematopoietic cell transplantation for asymptomatic patients with X-linked lymphoproliferative syndrome type 1. Clinical Immunology, 2022, 237, 108993.	3.2	1
154	Skeletal dysplasia in adenosine deaminase deficiency. Pediatrics International, 2022, 64, .	0.5	1
155	Peripheral blood lymphocyte subpopulations in three infants with hepatosplenomegaly caused by cytomegalovirus infection. Pediatrics International, 1995, 37, 370-373.	0.5	0
156	Gastritis and colitis can be associated with <scp>XLPâ€1</scp> (<scp>SAP</scp> deficiency): Reply. Pediatrics International, 2012, 54, 964-965.	0.5	0
157	Influenzaâ€induced hemolytic crisis in glucoseâ€6â€phosphate dehydrogenase deficiency. Pediatrics International, 2020, 62, 1003-1004.	0.5	0
158	Agammaglobulinemia as early B cell defects. Japanese Journal of Clinical Immunology, 2000, 23, 435-444.	0.0	0
159	ï¼‚â€ï½Œï½‰ï½Žï½‹ï½…ã€€ï½Œï½™ï½ï½ï½˜ï½ï½’ï½ï½Œï½‰ï½†ï½…ï½’ï½ï½”ï½‰ï½–ï½…ã€€ï½“ï½™ï½Žï½		
160	Lymphoproliferative disorders caused by hereditary genetic defects. Japanese Journal of Clinical Immunology, 2003, 26, 311-322.	0.0	0
161	Rapid Detection of SAP Deficiency in Cytotoxic Lymphocytes from Patients with X-Linked Lymphoproliferative Disease and Their Family Members.. Blood, 2004, 104, 3846-3846.	1.4	0
162	GATA1 Mutants Lacking Rb-Binding Motif Observed in Transient Abnormal Myelopoiesis in Down Syndrome. Blood, 2011, 118, 1491-1491.	1.4	0

#	Article	IF	Citations
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