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	A case of autoimmune enteropathy with CTLA4 haploinsufficiency. Intestinal Research, 2022, 20, 144-149.	2.6	6
2	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
3	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
4	Cartilageâ€hair hypoplasia with Tâ€cell dysfunction. Pediatrics International, 2022, 64, e15080.	0.5	3
5	Case Report: Rotavirus Vaccination and Severe Combined Immunodeficiency in Japan. Frontiers in Immunology, 2022, 13, 786375.	4.8	2
6	<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
7	Preemptive hematopoietic cell transplantation for asymptomatic patients with X-linked lymphoproliferative syndrome type 1. Clinical Immunology, 2022, 237, 108993.	3.2	1
8	Skeletal dysplasia in adenosine deaminase deficiency. Pediatrics International, 2022, 64, .	0.5	1
9	Cytomegalovirus Laryngitis in Primary Combined Immunodeficiency Diseases. Journal of Clinical Immunology, 2021, 41, 243-247.	3.8	4
10	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
11	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
12	Functional analysis of novel A20 variants in patients with atypical inflammatory diseases. Arthritis Research and Therapy, 2021, 23, 52.	3.5	15
13	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
14	Epstein-Barr Virusâ€“Negative Granulomatous Disease Due to SAP Deficiency. Journal of Clinical Immunology, 2021, 41, 1372-1375.	3.8	1
15	Genomics analysis of leukaemia predisposition in Xâ€linked agammaglobulinaemia. British Journal of Haematology, 2021, 193, 1277-1281.	2.5	1
16	A variant in human AIOLOS impairs adaptive immunity by interfering with IKAROS. Nature Immunology, 2021, 22, 893-903.	14.5	33
17	Heterozygous OAS1 gain-of-function variants cause an autoinflammatory immunodeficiency. Science Immunology, 2021, 6, .	11.9	36
18	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

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19	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
20	Hematopoietic Cell Transplantation Ameliorates Autoinflammation in A20 Haploinsufficiency. Journal of Clinical Immunology, 2021, 41, 1954-1956.	3.8	9
21	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
22	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
23	Endocrinopathies in Inborn Errors of Immunity. Frontiers in Immunology, 2021, 12, 786241.	4.8	3
24	Clinical Courses of IKAROS and CTLA4 Deficiencies: A Systematic Literature Review and Retrospective Longitudinal Study. Frontiers in Immunology, 2021, 12, 784901.	4.8	4
25	Dysregulation of the Intestinal Microbiome in Patients With Haploinsufficiency of A20. Frontiers in Cellular and Infection Microbiology, 2021, 11, 787667.	3.9	0
26	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
27	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
28	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
29	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
30	Influenzaâ€induced hemolytic crisis in glucoseâ€6â€phosphate dehydrogenase deficiency. Pediatrics International, 2020, 62, 1003-1004.	0.5	0
31	Current Perspectives and Unmet Needs of Primary Immunodeficiency Care in Asia Pacific. Frontiers in Immunology, 2020, 11, 1605.	4.8	13
32	Impaired B-Cell Differentiation in a Patient With STAT1 Gain-of-Function Mutation. Frontiers in Immunology, 2020, 11, 557521.	4.8	9
33	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
34	Fatal Progressive Meningoencephalitis Diagnosed in Two Members of a Family With X-Linked Agammaglobulinemia. Frontiers in Pediatrics, 2020, 8, 579.	1.9	1
35	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
36	Helicobacter cinaedi-Associated Refractory Cellulitis in Patients with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2020, 40, 1132-1137.	3.8	8

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37	A Cry for the Development of Newborn Screening for Familial Hemophagocytic Lymphohistiocytosis. Journal of Clinical Immunology, 2020, 40, 1196-1198.	3.8	3
38	DNA Ligase IV Deficiency Identified by Chance Following Vaccine-Derived Rubella Virus Infection. Journal of Clinical Immunology, 2020, 40, 1187-1190.	3.8	4
39	Inherited chromosomally integrated human herpesvirusâ€6 in a patient with XIAP deficiency. Transplant Infectious Disease, 2020, 22, e13331.	1.7	6
40	Longâ€term outcomes of children with extracutaneous juvenile xanthogranulomas in Japan. Pediatric Blood and Cancer, 2020, 67, e28381.	1.5	12
41	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
42	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
43	Immunophenotyping of A20 haploinsufficiency by multicolor flow cytometry. Clinical Immunology, 2020, 216, 108441.	3.2	5
44	APRIL-dependent lifelong plasmacyte maintenance and immunoglobulin production in humans. Journal of Allergy and Clinical Immunology, 2020, 146, 1109-1120.e4.	2.9	33
45	A Novel Homozygous Mutation Destabilizes IKKÎ² and Leads to Human Combined Immunodeficiency. Frontiers in Immunology, 2020, 11, 517544.	4.8	8
46	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
47	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
48	A synonymous splice site mutation in IL2RG gene causes late-onset combined immunodeficiency. International Journal of Hematology, 2019, 109, 603-611.	1.6	11
49	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
50	Epstein-Barr Virus-Associated Î³Î´ T-Cell Lymphoproliferative Disorder Associated With Hypomorphic IL2RG Mutation. Frontiers in Pediatrics, 2019, 7, 15.	1.9	12
51	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
52	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
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	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

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55	Outcomes in children with hemophagocytic lymphohistiocytosis treated using HLH-2004 protocol in Japan. International Journal of Hematology, 2019, 109, 206-213.	1.6	32
56	B-lymphoblastic lymphoma with TCF3-PBX1 fusion gene. Haematologica, 2019, 104, e35-e37.	3.5	9
57	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
58	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
59	Droplet Digital PCR-Based Chimerism Analysis for Primary Immunodeficiency Diseases. Journal of Clinical Immunology, 2018, 38, 300-306.	3.8	14
60	Dysregulation of Epstein-Barr Virus Infection in Hypomorphic ZAP70 Mutation. Journal of Infectious Diseases, 2018, 218, 825-834.	4.0	22
61	Heterozygous Mutations in OAS1 Cause Infantile-Onset Pulmonary Alveolar Proteinosis with Hypogammaglobulinemia. American Journal of Human Genetics, 2018, 102, 480-486.	6.2	26
62	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
63	Haploinsufficiency of A20 causes autoinflammatory and autoimmune disorders. Journal of Allergy and Clinical Immunology, 2018, 141, 1485-1488.e11.	2.9	100
64	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
65	Flow cytometry-based diagnosis of primary immunodeficiency diseases. Allergology International, 2018, 67, 43-54.	3.3	97
66	Genetic heterogeneity of uncharacterized childhood autoimmune diseases with lymphoproliferation. Pediatric Blood and Cancer, 2018, 65, e26831.	1.5	18
67	Hematopoietic stem cell transplantation for pulmonary alveolar proteinosis associated with primary immunodeficiency disease. International Journal of Hematology, 2018, 107, 610-614.	1.6	10
68	Population Pharmacokinetics of Intravenous Busulfan in Japanese Pediatric Patients With Primary Immunodeficiency Diseases. Journal of Clinical Pharmacology, 2018, 58, 327-331.	2.0	13
69	Hematopoietic cell transplantation for asymptomatic X-linked lymphoproliferative syndrome type 1. Allergy, Asthma and Clinical Immunology, 2018, 14, 82.	2.0	2
70	Inflammatory bowel diseases and primary immunodeficiency diseases. Immunological Medicine, 2018, 41, 154-161.	2.6	6
71	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
72	Clinical and Immunological Characterization of ICF Syndrome in Japan. Journal of Clinical Immunology, 2018, 38, 927-937.	3.8	29

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73	Gain-of-function IKBKB mutation causes human combined immune deficiency. Journal of Experimental Medicine, 2018, 215, 2715-2724.	8.5	69
74	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
75	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
76	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
77	IKBA S32 Mutations Underlie Ectodermal Dysplasia with Immunodeficiency and Severe Noninfectious Systemic Inflammation. Journal of Clinical Immunology, 2018, 38, 543-545.	3.8	10
78	Reactive peripheral blood plasmacytosis in Kawasaki disease. Pediatrics International, 2018, 60, 884-885.	0.5	5
79	Dominant-negative IKZF1 mutations cause a T, B, and myeloid cell combined immunodeficiency. Journal of Clinical Investigation, 2018, 128, 3071-3087.	8.2	133
80	Hematopoietic Stem Cell Transplantation in Children with Refractory Langerhans Cell Histiocytosis. Blood, 2018, 132, 4657-4657.	1.4	1
81	Clinical and Genetic Characteristics of Patients with Shwachman-Diamond Syndrome in Japan. Blood, 2018, 132, 3862-3862.	1.4	0
82	Multicolor Flow Cytometry for the Diagnosis of Primary Immunodeficiency Diseases. Journal of Clinical Immunology, 2017, 37, 486-495.	3.8	42
83	Abnormal hematopoiesis and autoimmunity in human subjects with germline IKZF1 mutations. Journal of Allergy and Clinical Immunology, 2017, 140, 223-231.	2.9	99
84	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
85	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
86	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
87	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
88	Hematopoietic Stem Cell Transplantation for XIAP Deficiency in Japan. Journal of Clinical Immunology, 2017, 37, 85-91.	3.8	63
89	Novel compound heterozygous mutations in a Japanese girl with Janus kinase 3 deficiency. Pediatrics International, 2016, 58, 1076-1080.	0.5	6
90	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

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91	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
92	Neonatal acute megakaryoblastic leukemia mimicking congenital neuroblastoma. Clinical Case Reports (discontinued), 2015, 3, 145-149.	0.5	9
93	X-Linked Agammaglobulinemia Associated with B-Precursor Acute Lymphoblastic Leukemia. Journal of Clinical Immunology, 2015, 35, 108-111.	3.8	20
94	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
95	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
96	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
97	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
98	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
99	Identification of novel fusion genes with 28S ribosomal DNA in hematologic malignancies. International Journal of Oncology, 2014, 44, 1193-1198.	3.3	7
100	Cost-minimization Analysis of IgPro20, a Subcutaneous Immunoglobulin, in Japanese Patients With Primary Immunodeficiency. Clinical Therapeutics, 2014, 36, 1616-1624.	2.5	22
101	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
102	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
103	Sustained elevation of serum interleukin-18 and its association with hemophagocytic lymphohistiocytosis in XIAP deficiency. Cytokine, 2014, 65, 74-78.	3.2	112
104	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
105	Extensive serum biomarker analysis in patients with enterohemorrhagic Escherichia coli O111-induced hemolytic-uremic syndrome. Cytokine, 2014, 66, 1-6.	3.2	18
106	XIAP Restricts TNF- and RIP3-Dependent Cell Death and Inflammasome Activation. Cell Reports, 2014, 7, 1796-1808.	6.4	210
107	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
108	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

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109	Allogeneic hematopoietic cell transplantation for XIAP deficiency: an international survey reveals poor outcomes. Blood, 2013, 121, 877-883.	1.4	132
110	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
111	SAP and XIAP deficiency in hemophagocytic lymphohistiocytosis. Pediatrics International, 2012, 54, 447-454.	0.5	35
112	Intracranial calcification in a uremic infant with Wilmsâ€™ tumor in a solitary kidney. CEN Case Reports, 2012, 1, 86-89.	0.9	1
113	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
114	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
115	Clinical and Genetic Characteristics of XIAP Deficiency in Japan. Journal of Clinical Immunology, 2012, 32, 411-420.	3.8	84
116	Genetic Basis of Myeloid Proliferation Related to Down Syndrome. Blood, 2012, 120, 535-535.	1.4	1
117	Identification of Novel Fusion Genes with 28S Ribosomal DNA in Hematologic Malignancies. Blood, 2012, 120, 4418-4418.	1.4	7
118	Knockdown of Shwachman-Diamond Syndrome Gene, SBDS, Induces Increased Expression of Galectin-1 and Impaired Cell Growth.. Blood, 2012, 120, 2360-2360.	1.4	0
119	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
120	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
121	Successful treatment of very large congenital infantile fibrosarcoma. Pediatrics International, 2011, 53, 768-770.	0.5	6
122	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
123	Nationwide Survey of Patients with Primary Immunodeficiency Diseases in Japan. Journal of Clinical Immunology, 2011, 31, 968-976.	3.8	77
124	Autoimmune lymphoproliferative syndrome mimicking chronic active Epsteinâ€“Barr virus infection. International Journal of Hematology, 2011, 93, 760-764.	1.6	16
125	Mislocalization or low expression of mutated Shwachmanâ€“Bodianâ€“Diamond syndrome protein. International Journal of Hematology, 2011, 94, 54-62.	1.6	3
126	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

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127	Early and rapid detection of X-linked lymphoproliferative syndrome with SH2D1A mutations by flow cytometry. , 2011, 80B, 8-13.		19
128	GATA1 Mutants Lacking Rb-Binding Motif Observed in Transient Abnormal Myelopoiesis in Down Syndrome. Blood, 2011, 118, 1491-1491.	1.4	0
129	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
130	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
131	Three brothers of X-linked agammaglobulinemia: the relation between phenotype and neutropenia. International Journal of Hematology, 2009, 90, 117-119.	1.6	4
132	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
133	Early lineage switch in an infant acute lymphoblastic leukemia. International Journal of Hematology, 2009, 90, 653-655.	1.6	30
134	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
135	A CIAS1 mutation in a Japanese girl with familial cold autoinflammatory syndrome. European Journal of Pediatrics, 2008, 167, 245-247.	2.7	1
136	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
137	Identification of DKC1 gene mutations in Japanese patients with Xâ€linked dyskeratosis congenita. British Journal of Haematology, 2005, 129, 432-434.	2.5	22
138	Genetic analysis of patients with defects in early Bâ€cell development. Immunological Reviews, 2005, 203, 216-234.	6.0	170
139	X-linked lymphoproliferative syndrome presenting with systemic lymphocytic vasculitis. American Journal of Hematology, 2005, 78, 130-133.	4.1	36
140	Severe Neutropenia in Japanese Patients with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2005, 25, 491-495.	3.8	34
141	Female agammaglobulinemia due to the Bruton tyrosine kinase deficiency caused by extremely skewed X-chromosome inactivation. Blood, 2004, 103, 185-187.	1.4	59
142	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
143	Autoimmune lymphoproliferative syndrome presenting with glomerulonephritis. Pediatric Nephrology, 2003, 18, 454-456.	1.7	27
144	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

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145	Lymphoproliferative disorders caused by hereditary genetic defects. Japanese Journal of Clinical Immunology, 2003, 26, 311-322.	0.0	0
146	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
147	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
148	ï¼¸â€ï½Œï½‰ï½Žï½‹ï½…ï½„â€ï½Œï½™ï½ï½ï½ˆï½ï½ï½’ï½ï½Œï½‰ï½†ï½…ï½’ï½ï½”ï½‰ï½–ï½…â€ï½“ï½™ï½Žï½„ï		
149	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
150	Maternal germinal mosaicism of Xâ€linked agammaglobulinemia. American Journal of Medical Genetics Part A, 2001, 99, 234-237.	2.4	9
151	Bruton tyrosine kinase gene mutations in Turkish patients with presumed X-linked agammaglobulinemia. Human Mutation, 2001, 18, 356-356.	2.5	15
152	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
153	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
154	Point mutation in intron 11 of Bruton's tyrosine kinase in atypical X-linked agammaglobulinemia. Pediatrics International, 2000, 42, 689-692.	0.5	4
155	X-linked thrombocytopenia identified by flow cytometric demonstration of defective Wiskott-Aldrich syndrome protein in lymphocytes. Blood, 2000, 95, 1110-1111.	1.4	30
156	Agammaglobulinemia as early B cell defects. Japanese Journal of Clinical Immunology, 2000, 23, 435-444.	0.0	0
157	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
158	Atypical X-linked Agammaglobulinemia Diagnosed in Three Adults.. Internal Medicine, 1999, 38, 722-725.	0.7	39
159	EBV-NK Cells Interactions and Lymphoproliferative Disorders. Leukemia and Lymphoma, 1998, 29, 491-498.	1.3	38
160	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
161	A case of influenza A infection followed by multiple organ failure (MOF). Journal of the Japanese Society of Intensive Care Medicine, 1997, 4, 375-380.	0.0	0
162	Mononucleosisâ€like illness in an infant associated with human herpesvirus 6 infection. Pediatrics International, 1995, 37, 227-229.	0.5	6

#	Article	IF	Citations
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