

Kenji Ihara

List of Publications by Year in descending order

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Version: 2024-02-01

154
papers

3,731
citations

159585

30
h-index

155660

55
g-index

156
all docs

156
docs citations

156
times ranked

5842
citing authors

#	ARTICLE	IF	CITATIONS
1	The autophagy reaction in the human umbilical cord: a potential marker for estimating fetal nutrition and neonatal growth. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2022, 35, 625-629.	1.5	1
2	Renal lesions mimicking acute focal bacterial nephritis in pediatric leukemia. <i>Pediatrics International</i> , 2022, 64, .	0.5	1
3	Stability of amino acids, free and acylâ€carnitine in stored dried blood spots. <i>Pediatrics International</i> , 2022, 64, .	0.5	2
4	Rothmundâ€Thomson syndrome investigated by two nationwide surveys in Japan. <i>Pediatrics International</i> , 2022, 64, .	0.5	2
5	Atypical food proteinâ€induced enterocolitis syndrome after vaccinations. <i>Pediatrics International</i> , 2022, 64, e14891.	0.5	0
6	A longitudinal ulcer due to <i>Yersinia pseudotuberculosis</i> infection in a girl. <i>Pediatrics International</i> , 2022, 64, e15036.	0.5	0
7	Simultaneous monitoring of oxygen and carbon dioxide for Pittâ€Hopkins syndrome. <i>Pediatrics International</i> , 2022, 64, e15180.	0.5	0
8	Fetoâ€maternal hemorrhage with placental chorioangioma: Two case reports. <i>Pediatrics International</i> , 2022, 64, .	0.5	0
9	A case of HDR syndrome coexisting with tetralogy of Fallot, with a novel GATA3 mutation, which manifested as a renal abscess. <i>CEN Case Reports</i> , 2021, 10, 241-243.	0.9	1
10	A Japanese infant presenting with hypocalcemic seizures resulting from hypovitaminosis D induced by non-celiac gluten sensitivity. <i>Clinical Pediatric Endocrinology</i> , 2021, 30, 105-110.	0.8	0
11	Hydrocortisone improved dexamethasoneâ€induced neuropsychological adverse effects. <i>Pediatrics International</i> , 2021, 63, 339-342.	0.5	1
12	Topoisomerase III ² immunoreactivity (IR) co-localizes with neuronal marker-IR but not glial fibrillary acidic protein-IR in GLI3-positive medulloblastomas: an immunohistochemical analysis of 124 medulloblastomas from the Japan Childrenâ€™s Cancer Group. <i>Brain Tumor Pathology</i> , 2021, 38, 109-121.	1.7	1
13	Leucyl-tRNA synthetase deficiency systemically induces excessive autophagy in zebrafish. <i>Scientific Reports</i> , 2021, 11, 8392.	3.3	4
14	Metabolome Characteristics of Liver Autophagy Deficiency under Starvation Conditions in Infancy. <i>Nutrients</i> , 2021, 13, 3026.	4.1	1
15	Fidgety Movements Assessment Accuracy Survey in Japan. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 13428.	2.6	1
16	Gastric Carcinoma as Second Malignant Neoplasm in a Survivor From High-risk Neuroblastoma. <i>Journal of Pediatric Hematology/Oncology</i> , 2020, 42, 160-162.	0.6	0
17	Macroglossia in Beckwith-Wiedemann Syndrome Is Attributed to Skeletal Muscle Hyperplasia. <i>Case Reports in Dentistry</i> , 2020, 2020, 1-6.	0.5	6
18	Early Recovery of Height Velocity in Prepubertal Children With Acute Lymphoblastic Leukemia Treated by a Short Intensive Phase Without Cranial Radiation Therapy. <i>Journal of Pediatric Hematology/Oncology</i> , 2020, 42, 271-274.	0.6	2

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19	Pediatric procedural sedation in Japan: A single-center facility study of 1,436 cases. <i>Pediatrics International</i> , 2020, 62, 1346-1350.	0.5	3
20	Tyrosine pre-transfer RNA fragments are linked to p53-dependent neuronal cell death via PKM2. <i>Biochemical and Biophysical Research Communications</i> , 2020, 525, 726-732.	2.1	16
21	A case of septo-optic dysplasia with hereditary hemorrhagic telangiectasia: a previously unrecognized combination of malformations. <i>Clinical Dysmorphology</i> , 2020, 29, 49-52.	0.3	0
22	Immune escape by loss of heterozygosity of HLA genes facilitated the transmission of diffuse large B cell lymphoma from mother to offspring. <i>British Journal of Haematology</i> , 2020, 189, e234-e237.	2.5	0
23	Malignant transformation of phosphaturic mesenchymal tumor: a case report and literature review. <i>Clinical Pediatric Endocrinology</i> , 2020, 29, 69-75.	0.8	18
24	Respiratory assessment in a spinal muscular atrophy infant treated with nusinersen. <i>Pediatrics International</i> , 2019, 61, 1051-1053.	0.5	5
25	The long-term management of congenital generalized lipodystrophy (Berardinelli-Seip syndrome): the clinical manifestations of Japanese siblings for approximately 20 years. <i>Clinical Pediatric Endocrinology</i> , 2019, 28, 139-145.	0.8	6
26	Critical association of Pallister-Hall syndrome and congenital heart disease. <i>Pediatrics International</i> , 2019, 61, 827-828.	0.5	1
27	Caffeine not associated with irritable behaviour in very low-birth-weight infants. <i>Early Human Development</i> , 2019, 137, 104835.	1.8	0
28	Immune checkpoint inhibitor therapy for pediatric cancers: A mini review of endocrine adverse events. <i>Clinical Pediatric Endocrinology</i> , 2019, 28, 59-68.	0.8	15
29	Language delay and developmental catch-up would be a clinical feature of pseudohypoparathyroidism type 1A during childhood. <i>Endocrine Journal</i> , 2019, 66, 215-221.	1.6	2
30	Present status of prophylactic thyroidectomy in pediatric multiple endocrine neoplasia 2: a nationwide survey in Japan 1997-2017. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2019, 32, 585-595.	0.9	5
31	Non-immune hydrops fetalis neonate born to a mother with yellow nail syndrome. <i>Pediatrics International</i> , 2019, 61, 313-315.	0.5	1
32	Therapeutic potential of hepatocyte-like-cells converted from stem cells from human exfoliated deciduous teeth in fulminant Wilson's disease. <i>Scientific Reports</i> , 2019, 9, 1535.	3.3	21
33	The association between brain morphological development and the quality of general movements. <i>Brain and Development</i> , 2019, 41, 490-500.	1.1	9
34	Clinical features of infantile subcutaneous panniculitis-like T-cell lymphoma. <i>Pediatrics International</i> , 2019, 61, 1261-1262.	0.5	2
35	A Child with Prostaglandin I ₂ -associated Thyrotoxicosis: Case Report. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2019, 11, 207-210.	0.9	2
36	Auto-immune disorders in a child with PIK3CD variant and 22q13 deletion. <i>European Journal of Medical Genetics</i> , 2018, 61, 631-633.	1.3	4

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37	Serum Amino Acid Profiling in Citrin-Deficient Children Exhibiting Normal Liver Function During the Apparently Healthy Period. <i>JIMD Reports</i> , 2018, 43, 53-61.	1.5	9
38	Unique cell tropism of HHV-6B in an infantile autopsy case of primary HHV-6B encephalitis. <i>Neuropathology</i> , 2018, 38, 400-406.	1.2	7
39	A case of CHARGE syndrome associated with hyperinsulinemic hypoglycemia in infancy. <i>European Journal of Medical Genetics</i> , 2018, 61, 312-314.	1.3	5
40	Diagnostic potential of stored dried blood spots for inborn errors of metabolism: a metabolic autopsy of medium-chain acyl-CoA dehydrogenase deficiency. <i>Journal of Clinical Pathology</i> , 2018, 71, 885-889.	2.0	9
41	Comprehensive screening for monogenic diabetes in 89 Japanese children with insulin-requiring antibody-negative type 1 diabetes. <i>Pediatric Diabetes</i> , 2018, 19, 243-250.	2.9	10
42	Vitamin D deficiency associated with dilated cardiomyopathy in early infancy caused by maternal cholestasis. <i>Clinical Pediatric Endocrinology</i> , 2018, 27, 187-192.	0.8	2
43	Cholesterol Metabolism Is Enhanced in the Liver and Brain of Children With Citrin Deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 2488-2497.	3.6	5
44	Incidence rate and characteristics of symptomatic vitamin D deficiency in children: a nationwide survey in Japan. <i>Endocrine Journal</i> , 2018, 65, 593-599.	1.6	12
45	Metabolic and immunological assessment of small-for-gestational-age children during one-year treatment with growth hormone: the clinical impact of apolipoproteins. <i>Endocrine Journal</i> , 2018, 65, 449-459.	1.6	3
46	Mutations in genes encoding polycomb repressive complex 2 subunits cause Weaver syndrome. <i>Human Mutation</i> , 2017, 38, 637-648.	2.5	80
47	Increased levels of anti-phosphatidylcholine and anti-phosphatidylethanolamine antibodies in pediatric patients with cerebral infarction. <i>Brain and Development</i> , 2017, 39, 542-546.	1.1	8
48	Toxic epidermal necrolysis in a child 6 months post-â€“hematopoietic stem cell transplantation. <i>Pediatric Transplantation</i> , 2017, 21, e12931.	1.0	4
49	A severe pulmonary complication in a patient with COL4A1 -related disorder: A case report. <i>European Journal of Medical Genetics</i> , 2017, 60, 169-171.	1.3	11
50	Circulating tricarboxylic acid cycle metabolite levels in citrin-deficient children with metabolic adaptation, with and without sodium pyruvate treatment. <i>Molecular Genetics and Metabolism</i> , 2017, 120, 207-212.	1.1	12
51	<i><sc>NUP</sc>98^{â€“}<sc>HOXC</sc>13</i> fusion gene in acute myeloid leukemia: Pediatric case. <i>Pediatrics International</i> , 2017, 59, 1105-1106.	0.5	0
52	Three-Quarters Adrenalectomy for Infantile-Onset Cushing Syndrome due to Bilateral Adrenal Hyperplasia in McCune-Albright Syndrome. <i>Hormone Research in Paediatrics</i> , 2017, 88, 285-290.	1.8	5
53	The clinical characteristics of Asian patients with classical-type Hutchinsonâ€“Gilford progeria syndrome. <i>Journal of Human Genetics</i> , 2017, 62, 1031-1035.	2.3	9
54	Hyperinsulinemic hypoglycemia in Beckwithâ€“Wiedemann, Sotos, and Kabuki syndromes: A nationwide survey in Japan. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 360-367.	1.2	20

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55	<i> FUT</i> non- ϵ secretor status is associated with Type 1 diabetes susceptibility in Japanese children. <i>Diabetic Medicine</i> , 2017, 34, 586-589.	2.3	15
56	Blood Reference Intervals for Preterm Low-Birth-Weight Infants: A Multicenter Cohort Study in Japan. <i>PLoS ONE</i> , 2016, 11, e0161439.	2.5	17
57	Analysis of Death Due to Infectious Diseases in Patients Hospitalized in the Pediatric Ward of a Single Japanese Tertiary Medical Facility. <i>Japanese Journal of Infectious Diseases</i> , 2016, 69, 464-470.	1.2	2
58	Management of congenital diaphragmatic hernia with transposition of the great arteries. <i>Pediatrics International</i> , 2016, 58, 516-518.	0.5	1
59	First mixoploid infant with full triploidy in blood cells. <i>Pediatrics International</i> , 2016, 58, 1354-1355.	0.5	2
60	SAMD9 mutations cause a novel multisystem disorder, MIRAGE syndrome, and are associated with loss of chromosome 7. <i>Nature Genetics</i> , 2016, 48, 792-797.	21.4	243
61	Gonadal macrophage infiltration in congenital lipid adrenal hyperplasia. <i>European Journal of Endocrinology</i> , 2016, 175, 127-132.	3.7	9
62	Reversible cerebral vasoconstriction syndrome manifesting as focal seizures without a thunderclap headache: A pediatric case report. <i>Brain and Development</i> , 2016, 38, 880-883.	1.1	5
63	Aminophylline-associated irritable behaviour in preterm neonates. <i>Early Human Development</i> , 2016, 99, 37-41.	1.8	3
64	De novo missense mutations in NALCN cause developmental and intellectual impairment with hypotonia. <i>Journal of Human Genetics</i> , 2016, 61, 451-455.	2.3	35
65	Moyamoya disease susceptibility gene RNF213 links inflammatory and angiogenic signals in endothelial cells. <i>Scientific Reports</i> , 2015, 5, 13191.	3.3	105
66	In vivo hepatogenic capacity and therapeutic potential of stem cells from human exfoliated deciduous teeth in liver fibrosis in mice. <i>Stem Cell Research and Therapy</i> , 2015, 6, 171.	5.5	67
67	The First Nationwide Survey and Genetic Analyses of Bardet-Biedl Syndrome in Japan. <i>PLoS ONE</i> , 2015, 10, e0136317.	2.5	16
68	TYK2 Promoter Variant and Diabetes Mellitus in the Japanese. <i>EBioMedicine</i> , 2015, 2, 744-749.	6.1	35
69	Overexpression of p53 but not <i>Rb</i> in the cytoplasm of neurons and small vessels in an autopsy of a patient with Cockayne syndrome. <i>Neuropathology</i> , 2015, 35, 266-272.	1.2	4
70	Autosomal dominant distal renal tubular acidosis caused by a mutation in the anion exchanger 1 gene in a Japanese family. <i>CEN Case Reports</i> , 2015, 4, 218-222.	0.9	6
71	Activation of an Innate Immune Receptor, Nod1, Accelerates Atherogenesis in <i>Apoe</i> ^{-/-} Mice. <i>Journal of Immunology</i> , 2015, 194, 773-780.	0.8	35
72	Thyroid Follicular Carcinoma in a Fourteen-year-old Girl with Graves' Disease. <i>Clinical Pediatric Endocrinology</i> , 2014, 23, 59-64.	0.8	1

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73	Survival and Neurodevelopmental Outcome of Preterm Infants Born at 22-24 Weeks of Gestational Age. <i>Neonatology</i> , 2014, 105, 79-84.	2.0	26
74	Endocrine and Metabolic Abnormalities in a Girl with Childhood Werner Syndrome: Case Report. <i>Journal of the American Geriatrics Society</i> , 2014, 62, 1404-1405.	2.6	2
75	Hypothalamic pituitary complications in Kabuki syndrome. <i>Pituitary</i> , 2013, 16, 133-138.	2.9	13
76	Long-term liposteroid therapy for idiopathic pulmonary hemosiderosis. <i>European Journal of Pediatrics</i> , 2013, 172, 1475-1481.	2.7	20
77	Bilateral stenosis of carotid siphon in Hutchinson-Gilford progeria syndrome. <i>Brain and Development</i> , 2013, 35, 690-693.	1.1	4
78	Two neonatal cholestasis patients with mutations in the <i>SRD5B1</i> (<i>AKR1D1</i>) gene: diagnosis and bile acid profiles during chenodeoxycholic acid treatment. <i>Journal of Inherited Metabolic Disease</i> , 2013, 36, 565-573.	3.6	22
79	Newborn screening for Fabry disease in Japan: prevalence and genotypes of Fabry disease in a pilot study. <i>Journal of Human Genetics</i> , 2013, 58, 548-552.	2.3	102
80	Coagulopathy in Patients With Late-Onset Ornithine Transcarbamylase Deficiency in Remission State: A Previously Unrecognized Complication. <i>Pediatrics</i> , 2013, 131, e327-e330.	2.1	8
81	Hyperinsulinemic hypoglycemia of infancy in Sotos syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 34-37.	1.2	22
82	Left ventricular efficiency after ligation of patent ductus arteriosus for premature infants. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2013, 146, 1353-1358.	0.8	12
83	Fatigue and quality of life in citrin deficiency during adaptation and compensation stage. <i>Molecular Genetics and Metabolism</i> , 2013, 109, 9-13.	1.1	19
84	Neuroendocrine phenotypes in a boy with 5q14 deletion syndrome implicate the regulatory roles of myocyte-specific enhancer factor 2C in the postnatal hypothalamus. <i>European Journal of Medical Genetics</i> , 2013, 56, 475-483.	1.3	6
85	<i>MLL2</i> and <i>KDM6A</i> mutations in patients with Kabuki syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 2234-2243.	1.2	148
86	A girl with Hajdu-Cheney syndrome and premature ovarian failure. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2012, 25, 171-3.	0.9	5
87	Acute Liver Failure as the Initial Manifestation of Wilson Disease Triggered by Human Parvovirus B19 Infection. <i>Pediatric Infectious Disease Journal</i> , 2012, 31, 103-104.	2.0	6
88	Endocrine complications in primary immunodeficiency diseases in Japan. <i>Clinical Endocrinology</i> , 2012, 77, 628-634.	2.4	6
89	Genetic susceptibility to Kawasaki disease: Analysis of pattern recognition receptor genes. <i>Human Immunology</i> , 2012, 73, 654-660.	2.4	23
90	Efficacy of bosentan therapy for segmental pulmonary artery hypertension due to major aortopulmonary collateral arteries in children. <i>International Journal of Cardiology</i> , 2012, 161, e1-e3.	1.7	20

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91	Bacterial pericarditis caused by <i>Lactobacillus iners</i> in an infant. <i>Diagnostic Microbiology and Infectious Disease</i> , 2012, 74, 181-182.	1.8	7
92	Hypothyroidism and Levothyroxine-Responsive Liver Dysfunction in a Patient with Ring Chromosome 18 Syndrome. <i>Thyroid</i> , 2012, 22, 1080-1083.	4.5	6
93	Linear nevus sebaceous syndrome with hypophosphatemic rickets with elevated FGF-23. <i>Pediatric Nephrology</i> , 2012, 27, 861-863.	1.7	13
94	HLA-class II and class I genotypes among Japanese children with Type 1A diabetes and their families. <i>Pediatric Diabetes</i> , 2012, 13, 33-44.	2.9	26
95	Balloon-occluded retrograde transvenous obliteration for congenital portosystemic venous shunt: Report of two cases. <i>Pediatrics International</i> , 2012, 54, 419-421.	0.5	4
96	Tracheal aspirate gene expression in preterm newborns and development of bronchopulmonary dysplasia. <i>Pediatrics International</i> , 2012, 54, 208-214.	0.5	13
97	The signal transducer and activator of transcription 5B gene polymorphism contributes to the cholesterol metabolism in Japanese children with growth hormone deficiency. <i>Clinical Endocrinology</i> , 2011, 74, 611-617.	2.4	10
98	Biotin Deficiency in a Glycogen Storage Disease Type 1b Girl Fed Only with Glycogen Storage Disease-Related Formula. <i>Pediatric Dermatology</i> , 2011, 28, 339-341.	0.9	6
99	Serum chemokine levels and developmental outcome in preterm infants. <i>Early Human Development</i> , 2011, 87, 439-443.	1.8	20
100	Preoperative Management With Nitrogen Inhalation Therapy for a Low-Birth Weight Infant With Tetralogy of Fallot and Absent Pulmonary Valve. <i>Pediatric Cardiology</i> , 2011, 32, 685-688.	1.3	1
101	HRAS mutants identified in Costello syndrome patients can induce cellular senescence: possible implications for the pathogenesis of Costello syndrome. <i>Journal of Human Genetics</i> , 2011, 56, 707-715.	2.3	29
102	PD1 as a common candidate susceptibility gene of subacute sclerosing panencephalitis. <i>Human Genetics</i> , 2010, 127, 411-419.	3.8	45
103	Mutant alleles associated with late-onset ornithine transcarbamylase deficiency in male patients have recurrently arisen and have been retained in some populations. <i>Journal of Human Genetics</i> , 2010, 55, 18-22.	2.3	5
104	Lack of association between E148Q MEFV variant and Kawasaki disease. <i>Human Immunology</i> , 2009, 70, 468-471.	2.4	9
105	Genetic Analysis of MMP Gene Polymorphisms in Patients With Kawasaki Disease. <i>Pediatric Research</i> , 2008, 63, 182-185.	2.3	34
106	Pulmonary Hypertension in Patients With Congenital Portosystemic Venous Shunt: A Previously Unrecognized Association. <i>Pediatrics</i> , 2008, 121, e892-e899.	2.1	79
107	Case Report: Adjuvant Therapy with a High Dose of Mitotane for Adrenocortical Carcinoma in a 4-year-old Boy. <i>Clinical Pediatric Endocrinology</i> , 2008, 17, 71-74.	0.8	7
108	PD-1 gene haplotype is associated with the development of type 1 diabetes mellitus in Japanese children. <i>Human Genetics</i> , 2007, 121, 223-232.	3.8	63

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109	Novel mutations in five Japanese patients with 3-methylcrotonyl-CoA carboxylase deficiency. <i>Journal of Human Genetics</i> , 2007, 52, 1040-1043.	2.3	18
110	Successful bone marrow transplantation in a patient withc-mpl-mutated congenital amegakaryocytic thrombocytopenia from a carrier donor. <i>Pediatric Transplantation</i> , 2005, 9, 101-103.	1.0	26
111	Congenital amegakaryocytic thrombocytopenia in three siblings: molecular analysis of atypical clinical presentation. <i>Experimental Hematology</i> , 2005, 33, 1215-1221.	0.4	24
112	Thrombocytosis in preterm infants: a possible involvement of thrombopoietin receptor gene expression. <i>Journal of Molecular Medicine</i> , 2005, 83, 316-320.	3.9	6
113	Association of Vascular Endothelial Growth Factor (VEGF) and VEGF Receptor Gene Polymorphisms with Coronary Artery Lesions of Kawasaki Disease. <i>Pediatric Research</i> , 2004, 56, 953-959.	2.3	109
114	In Vivo Selection of Genetically Modified Erythroid Cells Using a Jak2-Based Cell Growth Switch. <i>Molecular Therapy</i> , 2004, 10, 456-468.	8.2	20
115	Identification of a novel type 1 diabetes susceptibility gene, T-bet. <i>Human Genetics</i> , 2004, 115, 177-84.	3.8	40
116	Characteristic expression of aryl hydrocarbon receptor repressor gene in human tissues: Organ-specific distribution and variable induction patterns in mononuclear cells. <i>Life Sciences</i> , 2004, 74, 1039-1049.	4.3	91
117	Clinical significance of a highly sensitive analysis for gene dosage and the expression level of MYCN in neuroblastoma. <i>Journal of Pediatric Surgery</i> , 2004, 39, 63-68.	1.6	25
118	Female agammaglobulinemia due to the Bruton tyrosine kinase deficiency caused by extremely skewed X-chromosome inactivation. <i>Blood</i> , 2004, 103, 185-187.	1.4	59
119	Association Study of Human MTH1 Gene Polymorphisms with Type 1 Diabetes Mellitus. <i>Endocrine Journal</i> , 2004, 51, 493-498.	1.6	12
120	Founder effect of the C9 R95X mutation in Orientals. <i>Human Genetics</i> , 2003, 112, 244-248.	3.8	21
121	A functional polymorphism in the promoter/enhancer region of the FOXP3/Scurfin gene associated with type 1 diabetes. <i>Immunogenetics</i> , 2003, 55, 149-156.	2.4	130
122	Association study between interleukin-12 receptor $\beta 1/\beta 2$ genes and type 1 diabetes or asthma in the Japanese population. <i>Immunogenetics</i> , 2003, 55, 189-192.	2.4	12
123	The lipocalin 24p3, which is an essential molecule in IL-3 withdrawal-induced apoptosis, is not involved in the G-CSF withdrawal-induced apoptosis. <i>European Journal of Haematology</i> , 2003, 71, 412-417.	2.2	12
124	Inducible and endothelial constitutive nitric oxide synthase gene polymorphisms in Kawasaki disease. <i>Pediatrics International</i> , 2003, 45, 130-134.	0.5	21
125	Cytokine imbalance in hyper-IgE syndrome: reduced expression of transforming growth factor $\beta 2$ and interferon $\beta 3$ genes in circulating activated T cells. <i>British Journal of Haematology</i> , 2003, 121, 324-331.	2.5	25
126	Contribution of the Interleukin 4 Gene to Susceptibility to Subacute Sclerosing Panencephalitis. <i>Archives of Neurology</i> , 2002, 59, 822-7.	4.5	42

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127	A Case of Growth Hormone and Gonadotropin Deficiency Associated with Unilateral Anophthalmia, Microphallus, Cryptorchidism, and Mental Retardation.. Endocrine Journal, 2002, 49, 15-20.	1.6	5
128	Five novel SLC7A7 variants and γ L gene-expression pattern in cultured lymphoblasts from Japanese patients with lysinuric protein intolerance. Human Mutation, 2002, 20, 375-381.	2.5	27
129	Association study of the NRAMP1 gene promoter polymorphism and early-onset type 1 diabetes. Immunogenetics, 2002, 54, 282-285.	2.4	23
130	Perforin defects of primary haemophagocytic lymphohistiocytosis in Japan. British Journal of Haematology, 2002, 116, 346-349.	2.5	71
131	Dominant expression of interleukin 10 but not interferon γ in CD4- CD8- \hat{I}^2 T cells of autoimmune lymphoproliferative syndrome. British Journal of Haematology, 2002, 119, 535-538.	2.5	29
132	Association of IFN- \hat{I}^3 and IFN regulatory factor 1 polymorphisms with childhood atopic asthma. Journal of Allergy and Clinical Immunology, 2001, 107, 499-504.	2.9	85
133	Human Biotin-Containing Subunit of 3-Methylcrotonyl-CoA Carboxylase Gene (MCCA): cDNA Sequence, Genomic Organization, Localization to Chromosomal Band 3q27, and Expression. Genomics, 2001, 72, 145-152.	2.9	13
134	Quick quantitative analysis of gene dosages associated with prognosis in neuroblastoma. Cancer Letters, 2001, 166, 89-94.	7.2	36
135	Association studies of CTLA-4, CD28, and ICOS gene polymorphisms with type 1 diabetes in the Japanese population. Immunogenetics, 2001, 53, 447-454.	2.4	75
136	High expression of platelet-derived growth factor and transforming growth factor- \hat{I}^2 1 in blast cells from patients with Down Syndrome suffering from transient myeloproliferative disorder and organ fibrosis. British Journal of Haematology, 2001, 115, 472-475.	2.5	44
137	Inflammatory Bowel Disease-Like Colitis in Glycogen Storage Disease Type 1b. Inflammatory Bowel Diseases, 2001, 7, 128-132.	1.9	56
138	Epstein-Barr Virus (EBV) Load and Cytokine Gene Expression in Activated T Cells of Chronic Active EBV Infection. Journal of Infectious Diseases, 2001, 183, 1-7.	4.0	186
139	Physical Growth and Retinopathy in Preterm Infants: Involvement of IGF-I and GH. Pediatric Research, 2001, 50, 732-736.	2.3	53
140	Childhood Atopic Asthma: Positive Association with a Polymorphism of IL-4 Receptor \hat{I}^2 Gene but Not with That of IL-4 Promoter or Fc \hat{I}^2 Receptor \hat{I}^2 Gene. Experimental and Clinical Immunogenetics, 2000, 17, 63-70.	1.2	63
141	Genomic imprinting of insulin-like growth factor-2 in infant leukemia and childhood neuroblastoma. , 2000, 88, 2372-2377.		7
142	A Case of Congenital Afibrinogenemia: Fibrinogen Hakata, a Novel Nonsense Mutation of the Fibrinogen \hat{I}^3 -Chain Gene. Thrombosis and Haemostasis, 2000, 84, 49-53.	3.4	24
143	Novel Polymorphism in the Coding Region of the IL-13 Receptor Alpha \hat{I}^2 Gene: Association Study with Atopic Asthma in the Japanese Population. Experimental and Clinical Immunogenetics, 2000, 17, 18-22.	1.2	20
144	Lack of Association between CD28/CTLA-4 Gene Polymorphisms and Atopic Asthma in the Japanese Population. Experimental and Clinical Immunogenetics, 2000, 17, 179-184.	1.2	28

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145	Neurotrophin-3 Levels in Cerebrospinal Fluid From Children With Bacterial Meningitis, Viral Meningitis, or Encephalitis. <i>Journal of Child Neurology</i> , 2000, 15, 19-21.	1.4	27
146	Analysis of Measles Virus Binding Sites of the CD46 Gene in Patients with Subacute Sclerosing Panencephalitis. <i>Journal of Infectious Diseases</i> , 2000, 181, 1447-1449.	4.0	9
147	Inflammatory bowel disease-like complication in anhidrotic ectodermal dysplasia. <i>American Journal of Gastroenterology</i> , 2000, 95, 3651-3652.	0.4	0
148	HIGH EXPRESSION BUT NO INTERNAL TANDEM DUPLICATION OF FLT3 IN NORMAL HEMATOPOIETIC CELLS. <i>Pediatric Hematology and Oncology</i> , 1999, 16, 437-441.	0.8	19
149	Oversecretion of IL-18 in haemophagocytic lymphohistiocytosis: a novel marker of disease activity. <i>British Journal of Haematology</i> , 1999, 106, 182-189.	2.5	97
150	Molecular epidemiology of C9 deficiency heterozygotes with an Arg95Stop mutation of the C9 gene in Japan. <i>Journal of Human Genetics</i> , 1999, 44, 109-111.	2.3	20
151	Rubinstein-Taybi syndrome: A girl with a history of neuroblastoma and premature thelarche. , 1999, 83, 365-366.		19
152	Nonsense mutation in exon 4 of human complement C9 gene is the major cause of Japanese complement C9 deficiency. <i>Human Genetics</i> , 1998, 102, 605-610.	3.8	29
153	Three-Dimensional Structure of a DNA Repair Enzyme, 3-Methyladenine DNA Glycosylase II, from <i>Escherichia coli</i> . <i>Cell</i> , 1996, 86, 311-319.	28.9	147
154	Mouse methyltransferase for repair of O6-methylguanine and O4-methylthymine in DNA. <i>Carcinogenesis</i> , 1995, 16, 1595-1602.	2.8	35