Kenji Ihara

List of Publications by Year
in descending order

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	SAMD9 mutations cause a novel multisystem disorder, MIRAGE syndrome, and are associated with loss of chromosome 7. Nature Genetics, 2016, 48, 792-797.	21.4	243
2	Epsteinâ€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
3	MLL2 and KDM6A mutations in patients with Kabuki syndrome. American Journal of Medical Genetics, Part A, 2013, 161, 2234-2243.	1.2	148
4	Three-Dimensional Structure of a DNA Repair Enzyme, 3-Methyladenine DNA Glycosylase II, from Escherichia coli. Cell, 1996, 86, 311-319.	28.9	147
5	A functional polymorphism in the promoter/enhancer region of the FOXP3/Scurfin gene associated with typeÂ1 diabetes. Immunogenetics, 2003, 55, 149-156.	2.4	130
6	Association of Vascular Endothelial Growth Factor (VEGF) and VEGF Receptor Gene Polymorphisms with Coronary Artery Lesions of Kawasaki Disease. Pediatric Research, 2004, 56, 953-959.	2.3	109
7	Moyamoya disease susceptibility gene RNF213 links inflammatory and angiogenic signals in endothelial cells. Scientific Reports, 2015, 5, 13191.	3.3	105
8	Newborn screening for Fabry disease in Japan: prevalence and genotypes of Fabry disease in a pilot study. Journal of Human Genetics, 2013, 58, 548-552.	2.3	102
9	Oversecretion of IL-18 in haemophagocytic lymphohistiocytosis: a novel marker of disease activity. British Journal of Haematology, 1999, 106, 182-189.	2.5	97
10	Characteristic expression of aryl hydrocarbon receptor repressor gene in human tissues: Organ-specific distribution and variable induction patterns in mononuclear cells. Life Sciences, 2004, 74, 1039-1049.	4.3	91
11	Association of IFN-Î³ and IFN regulatory factor 1 polymorphisms with childhood atopic asthma. Journal of Allergy and Clinical Immunology, 2001, 107, 499-504.	2.9	85
12	Mutations in genes encoding polycomb repressive complex 2 subunits cause Weaver syndrome. Human Mutation, 2017, 38, 637-648.	2.5	80
13	Pulmonary Hypertension in Patients With Congenital Portosystemic Venous Shunt: A Previously Unrecognized Association. Pediatrics, 2008, 121, e892-e899.	2.1	79
14	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
15	Perforin defects of primary haemophagocytic lymphohistiocytosis in Japan. British Journal of Haematology, 2002, 116, 346-349.	2.5	71
16	In vivo hepatogenic capacity and therapeutic potential of stem cells from human exfoliated deciduous teeth in liver fibrosis in mice. Stem Cell Research and Therapy, 2015, 6, 171.	5.5	67
17	Childhood Atopic Asthma: Positive Association with a Polymorphism of IL-4 Receptor Î± Gene but Not with That of IL-4 Promoter or Fc Îµ Receptor I Î² Gene. Experimental and Clinical Immunogenetics, 2000, 17, 63-70.	1.2	63
18	PD-1 gene haplotype is associated with the development of type 1 diabetes mellitus in Japanese children. Human Genetics, 2007, 121, 223-232.	3.8	63

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19	Female agammaglobulinemia due to the Bruton tyrosine kinase deficiency caused by extremely skewed X-chromosome inactivation. Blood, 2004, 103, 185-187.	1.4	59
20	Inflammatory Bowel Disease-Like Colitis in Glycogen Storage Disease Type 1b. Inflammatory Bowel Diseases, 2001, 7, 128-132.	1.9	56
21	Physical Growth and Retinopathy in Preterm Infants: Involvement of IGF-I and GH. Pediatric Research, 2001, 50, 732-736.	2.3	53
22	PD1 as a common candidate susceptibility gene of subacute sclerosing panencephalitis. Human Genetics, 2010, 127, 411-419.	3.8	45
23	High expression of platelet-derived growth factor and transforming growth factor-Î²1 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
24	Contribution of the Interleukin 4 Gene to Susceptibility to Subacute Sclerosing Panencephalitis. Archives of Neurology, 2002, 59, 822-7.	4.5	42
25	Identification of a novel type 1 diabetes susceptibility gene, T-bet. Human Genetics, 2004, 115, 177-84.	3.8	40
26	Quick quantitative analysis of gene dosages associated with prognosis in neuroblastoma. Cancer Letters, 2001, 166, 89-94.	7.2	36
27	Mouse methyltransferase for repair of O6-methylguanine and O4-methylthymine in DNA. Carcinogenesis, 1995, 16, 1595-1602.	2.8	35
28	TYK2 Promoter Variant and Diabetes Mellitus in the Japanese. EBioMedicine, 2015, 2, 744-749.	6.1	35
29	Activation of an Innate Immune Receptor, Nod1, Accelerates Atherogenesis in Apoeâ^'/â^' Mice. Journal of Immunology, 2015, 194, 773-780.	0.8	35
30	De novo missense mutations in NALCN cause developmental and intellectual impairment with hypotonia. Journal of Human Genetics, 2016, 61, 451-455.	2.3	35
31	Genetic Analysis of MMP Gene Polymorphisms in Patients With Kawasaki Disease. Pediatric Research, 2008, 63, 182-185.	2.3	34
32	Nonsense mutation in exon 4 of human complement C9 gene is the major cause of Japanese complement C9 deficiency. Human Genetics, 1998, 102, 605-610.	3.8	29
33	Dominant expression of interleukin 10 but not interferon Î³ in CD4- CD8- Î±Î²T cells of autoimmune lymphoproliferative syndrome. British Journal of Haematology, 2002, 119, 535-538.	2.5	29
34	HRAS mutants identified in Costello syndrome patients can induce cellular senescence: possible implications for the pathogenesis of Costello syndrome. Journal of Human Genetics, 2011, 56, 707-715.	2.3	29
35	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
36	Neurotrophin-3 Levels in Cerebrospinal Fluid From Children With Bacterial Meningitis, Viral Meningitis, or Encephalitis. Journal of Child Neurology, 2000, 15, 19-21.	1.4	27

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37	Five novelSLC7A7 variants and y+L gene-expression pattern in cultured lymphoblasts from Japanese patients with lysinuric protein intolerance. Human Mutation, 2002, 20, 375-381.	2.5	27
38	Successful bone marrow transplantation in a patient withc-mpl-mutated congenital amegakaryocytic thrombocytopenia from a carrier donor. Pediatric Transplantation, 2005, 9, 101-103.	1.0	26
39	HLA-class II and class I genotypes among Japanese children with Type 1A diabetes and their families. Pediatric Diabetes, 2012, 13, 33-44.	2.9	26
40	Survival and Neurodevelopmental Outcome of Preterm Infants Born at 22-24 Weeks of Gestational Age. Neonatology, 2014, 105, 79-84.	2.0	26
41	Cytokine imbalance in hyperâ€IgE syndrome: reduced expression of transforming growth factor Î² and interferon Î³ genes in circulating activated T cells. British Journal of Haematology, 2003, 121, 324-331.	2.5	25
42	Clinical significance of a highly sensitive analysis for gene dosage and the expression level of MYCN in neuroblastoma. Journal of Pediatric Surgery, 2004, 39, 63-68.	1.6	25
43	A Case of Congenital Afibrinogenemia: Fibrinogen Hakata, a Novel Nonsense Mutation of the Fibrinogen Î³-Chain Gene. Thrombosis and Haemostasis, 2000, 84, 49-53.	3.4	24
44	Congenital amegakaryocytic thrombocytopenia in three siblings: molecular analysis of atypical clinical presentation. Experimental Hematology, 2005, 33, 1215-1221.	0.4	24
45	Association study of the NRAMP1 gene promoter polymorphism and early-onset type 1 diabetes. Immunogenetics, 2002, 54, 282-285.	2.4	23
46	Genetic susceptibility to Kawasaki disease: Analysis of pattern recognition receptor genes. Human Immunology, 2012, 73, 654-660.	2.4	23
47	Two neonatal cholestasis patients with mutations in the SRD5B1 (AKR1D1) gene: diagnosis and bile acid profiles during chenodeoxycholic acid treatment. Journal of Inherited Metabolic Disease, 2013, 36, 565-573.	3.6	22
48	Hyperinsulinemic hypoglycemia of infancy in Sotos syndrome. American Journal of Medical Genetics, Part A, 2013, 161, 34-37.	1.2	22
49	Founder effect of the C9 R95X mutation in Orientals. Human Genetics, 2003, 112, 244-248.	3.8	21
50	Inducible and endothelial constitutive nitric oxide synthase gene polymorphisms in Kawasaki disease. Pediatrics International, 2003, 45, 130-134.	0.5	21
51	Therapeutic potential of hepatocyte-like-cells converted from stem cells from human exfoliated deciduous teeth in fulminant Wilsonâ€™s disease. Scientific Reports, 2019, 9, 1535.	3.3	21
52	Molecular epidemiology of C9 deficiency heterozygotes with an Arg95Stop mutation of the C9 gene in Japan. Journal of Human Genetics, 1999, 44, 109-111.	2.3	20
53	Novel Polymorphism in the Coding Region of the IL-13 Receptor Alphaâ€™ Gene: Association Study with Atopic Asthma in the Japanese Population. Experimental and Clinical Immunogenetics, 2000, 17, 18-22.	1.2	20
54	In Vivo Selection of Genetically Modified Erythroid Cells Using a Jak2-Based Cell Growth Switch. Molecular Therapy, 2004, 10, 456-468.	8.2	20

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55	Serum chemokine levels and developmental outcome in preterm infants. Early Human Development, 2011, 87, 439-443.	1.8	20
56	Efficacy of bosentan therapy for segmental pulmonary artery hypertension due to major aortopulmonary collateral arteries in children. International Journal of Cardiology, 2012, 161, e1-e3.	1.7	20
57	Long-term liposteroid therapy for idiopathic pulmonary hemosiderosis. European Journal of Pediatrics, 2013, 172, 1475-1481.	2.7	20
58	Hyperinsulinemic hypoglycemia in Beckwithâ€“Wiedemann, Sotos, and Kabuki syndromes: A nationwide survey in Japan. American Journal of Medical Genetics, Part A, 2017, 173, 360-367.	1.2	20
59	HIGH EXPRESSION BUT NO INTERNAL TANDEM DUPLICATION OF FLT3 IN NORMAL HEMATOPOIETIC CELLS. Pediatric Hematology and Oncology, 1999, 16, 437-441.	0.8	19
60	Rubinstein-Taybi syndrome: A girl with a history of neuroblastoma and premature thelarche. , 1999, 83, 365-366.		19
61	Fatigue and quality of life in citrin deficiency during adaptation and compensation stage. Molecular Genetics and Metabolism, 2013, 109, 9-13.	1.1	19
62	Novel mutations in five Japanese patients with 3-methylcrotonyl-CoA carboxylase deficiency. Journal of Human Genetics, 2007, 52, 1040-1043.	2.3	18
63	Malignant transformation of phosphaturic mesenchymal tumor: a case report and literature review. Clinical Pediatric Endocrinology, 2020, 29, 69-75.	0.8	18
64	Blood Reference Intervals for Preterm Low-Birth-Weight Infants: A Multicenter Cohort Study in Japan. PLoS ONE, 2016, 11, e0161439.	2.5	17
65	The First Nationwide Survey and Genetic Analyses of Bardet-Biedl Syndrome in Japan. PLoS ONE, 2015, 10, e0136317.	2.5	16
66	Tyrosine pre-transfer RNA fragments are linked to p53-dependent neuronal cell death via PKM2. Biochemical and Biophysical Research Communications, 2020, 525, 726-732.	2.1	16
67	<scp>FUT</scp>2 nonâ€secretor status is associated with Type 1 diabetes susceptibility in Japanese children. Diabetic Medicine, 2017, 34, 586-589.	2.3	15
68	Immune checkpoint inhibitor therapy for pediatric cancers: A mini review of endocrine adverse events. Clinical Pediatric Endocrinology, 2019, 28, 59-68.	0.8	15
69	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
70	Linear nevus sebaceous syndrome with hypophosphatemic rickets with elevated FGF-23. Pediatric Nephrology, 2012, 27, 861-863.	1.7	13
71	Tracheal aspirate gene expression in preterm newborns and development of bronchopulmonary dysplasia. Pediatrics International, 2012, 54, 208-214.	0.5	13
72	Hypothalamic pituitary complications in Kabuki syndrome. Pituitary, 2013, 16, 133-138.	2.9	13

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73	Association study between interleukin-12 receptor Î²1/Î²2 genes and type 1 diabetes or asthma in the Japanese population. Immunogenetics, 2003, 55, 189-192.	2.4	12
74	The lipocalin 24p3, which is an essential molecule in IL-3 withdrawal-induced apoptosis, is not involved in the G-CSF withdrawal-induced apoptosis. European Journal of Haematology, 2003, 71, 412-417.	2.2	12
75	Association Study of Human MTH1 Gene Polymorphisms with Type 1 Diabetes Mellitus. Endocrine Journal, 2004, 51, 493-498.	1.6	12
76	Left ventricular efficiency after ligation of patent ductus arteriosus for premature infants. Journal of Thoracic and Cardiovascular Surgery, 2013, 146, 1353-1358.	0.8	12
77	Circulating tricarboxylic acid cycle metabolite levels in citrin-deficient children with metabolic adaptation, with and without sodium pyruvate treatment. Molecular Genetics and Metabolism, 2017, 120, 207-212.	1.1	12
78	Incidence rate and characteristics of symptomatic vitamin D deficiency in children: a nationwide survey in Japan. Endocrine Journal, 2018, 65, 593-599.	1.6	12
79	A severe pulmonary complication in a patient with COL4A1 -related disorder: A case report. European Journal of Medical Genetics, 2017, 60, 169-171.	1.3	11
80	The signal transducer and activator of transcription 5B gene polymorphism contributes to the cholesterol metabolism in Japanese children with growth hormone deficiency. Clinical Endocrinology, 2011, 74, 611-617.	2.4	10
81	Comprehensive screening for monogenic diabetes in 89 Japanese children with insulin-requiring antibody-negative type 1 diabetes. Pediatric Diabetes, 2018, 19, 243-250.	2.9	10
82	Analysis of Measles Virus Binding Sites of the CD46 Gene in Patients with Subacute Sclerosing Panencephalitis. Journal of Infectious Diseases, 2000, 181, 1447-1449.	4.0	9
83	Lack of association between E148Q MEFV variant and Kawasaki disease. Human Immunology, 2009, 70, 468-471.	2.4	9
84	Gonadal macrophage infiltration in congenital lipoid adrenal hyperplasia. European Journal of Endocrinology, 2016, 175, 127-132.	3.7	9
85	The clinical characteristics of Asian patients with classical-type Hutchinsonâ€“Gilford progeria syndrome. Journal of Human Genetics, 2017, 62, 1031-1035.	2.3	9
86	Serum Amino Acid Profiling in Citrin-Deficient Children Exhibiting Normal Liver Function During the Apparently Healthy Period. JIMD Reports, 2018, 43, 53-61.	1.5	9
87	Diagnostic potential of stored dried blood spots for inborn errors of metabolism: a metabolic autopsy of medium-chain acyl-CoA dehydrogenase deficiency. Journal of Clinical Pathology, 2018, 71, 885-889.	2.0	9
88	The association between brain morphological development and the quality of general movements. Brain and Development, 2019, 41, 490-500.	1.1	9
89	Coagulopathy in Patients With Late-Onset Ornithine Transcarbamylase Deficiency in Remission State: A Previously Unrecognized Complication. Pediatrics, 2013, 131, e327-e330.	2.1	8
90	Increased levels of anti-phosphatidylcholine and anti-phosphatidylethanolamine antibodies in pediatric patients with cerebral infarction. Brain and Development, 2017, 39, 542-546.	1.1	8

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91	Genomic imprinting of insulin-like growth factor-2 in infant leukemia and childhood neuroblastoma. , 2000, 88, 2372-2377.		7
92	Bacterial pericarditis caused by Lactobacillus iners in an infant. Diagnostic Microbiology and Infectious Disease, 2012, 74, 181-182.	1.8	7
93	Unique cell tropism of HHVâ€6B in an infantile autopsy case of primary HHVâ€6B encephalitis. Neuropathology, 2018, 38, 400-406.	1.2	7
94	Case Report: Adjuvant Therapy with a High Dose of Mitotane for Adrenocortical Carcinoma in a 4-year-old Boy. Clinical Pediatric Endocrinology, 2008, 17, 71-74.	0.8	7
95	Thrombocytosis in preterm infants: a possible involvement of thrombopoietin receptor gene expression. Journal of Molecular Medicine, 2005, 83, 316-320.	3.9	6
96	Biotin Deficiency in a Glycogen Storage Disease Type 1b Girl Fed Only with Glycogen Storage Diseaseâ€Related Formula. Pediatric Dermatology, 2011, 28, 339-341.	0.9	6
97	Acute Liver Failure as the Initial Manifestation of Wilson Disease Triggered by Human Parvovirus B19 Infection. Pediatric Infectious Disease Journal, 2012, 31, 103-104.	2.0	6
98	Endocrine complications in primary immunodeficiency diseases in Japan. Clinical Endocrinology, 2012, 77, 628-634.	2.4	6
99	Hypothyroidism and Levothyroxine-Responsive Liver Dysfunction in a Patient with Ring Chromosome 18 Syndrome. Thyroid, 2012, 22, 1080-1083.	4.5	6
100	Neuroendocrine phenotypes in a boy with 5q14 deletion syndrome implicate the regulatory roles of myocyte-specific enhancer factor 2C in the postnatal hypothalamus. European Journal of Medical Genetics, 2013, 56, 475-483.	1.3	6
101	Autosomal dominant distal renal tubular acidosis caused by a mutation in the anion exchanger 1 gene in a Japanese family. CEN Case Reports, 2015, 4, 218-222.	0.9	6
102	The long-term management of congenital generalized lipodystrophy (Berardinelli-Seip syndrome): the clinical manifestations of Japanese siblings for approximately 20 years. Clinical Pediatric Endocrinology, 2019, 28, 139-145.	0.8	6
103	Macroglossia in Beckwith-Wiedemann Syndrome Is Attributed to Skeletal Muscle Hyperplasia. Case Reports in Dentistry, 2020, 2020, 1-6.	0.5	6
104	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
105	Mutant alleles associated with late-onset ornithine transcarbamylase deficiency in male patients have recurrently arisen and have been retained in some populations. Journal of Human Genetics, 2010, 55, 18-22.	2.3	5
106	A girl with Hajdu-Cheney syndrome and premature ovarian failure. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 171-3.	0.9	5
107	Reversible cerebral vasoconstriction syndrome manifesting as focal seizures without a thunderclap headache: A pediatric case report. Brain and Development, 2016, 38, 880-883.	1.1	5
108	Three-Quarters Adrenalectomy for Infantile-Onset Cushing Syndrome due to Bilateral Adrenal Hyperplasia in McCune-Albright Syndrome. Hormone Research in Paediatrics, 2017, 88, 285-290.	1.8	5

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109	A case of CHARGE syndrome associated with hyperinsulinemic hypoglycemia in infancy. European Journal of Medical Genetics, 2018, 61, 312-314.	1.3	5
110	Cholesterol Metabolism Is Enhanced in the Liver and Brain of Children With Citrin Deficiency. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 2488-2497.	3.6	5
111	Respiratory assessment in a spinal muscular atrophy infant treated with nusinersen. Pediatrics International, 2019, 61, 1051-1053.	0.5	5
112	Present status of prophylactic thyroidectomy in pediatric multiple endocrine neoplasia 2: a nationwide survey in Japan 1997â€“2017. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 585-595.	0.9	5
113	Balloonâ€occluded retrograde transvenous obliteration for congenital portosystemic venous shunt: Report of two cases. Pediatrics International, 2012, 54, 419-421.	0.5	4
114	Bilateral stenosis of carotid siphon in Hutchinson-Gilford progeria syndrome. Brain and Development, 2013, 35, 690-693.	1.1	4
115	Overexpression of p53 but not <scp>R</scp>b in the cytoplasm of neurons and small vessels in an autopsy of a patient with <scp>C</scp>ockayne syndrome. Neuropathology, 2015, 35, 266-272.	1.2	4
116	Toxic epidermal necrolysis in a child 6 months postâ€hematopoietic stem cell transplantation. Pediatric Transplantation, 2017, 21, e12931.	1.0	4
117	Auto-immune disorders in a child with PIK3CD variant and 22q13 deletion. European Journal of Medical Genetics, 2018, 61, 631-633.	1.3	4
118	Leucyl-tRNA synthetase deficiency systemically induces excessive autophagy in zebrafish. Scientific Reports, 2021, 11, 8392.	3.3	4
119	Aminophylline-associated irritable behaviour in preterm neonates. Early Human Development, 2016, 99, 37-41.	1.8	3
120	Metabolic and immunological assessment of small-for-gestational-age children during one-year treatment with growth hormone: the clinical impact of apolipoproteins. Endocrine Journal, 2018, 65, 449-459.	1.6	3
121	Pediatric procedural sedation in Japan: A singleâ€facility study of 1,436 cases. Pediatrics International, 2020, 62, 1346-1350.	0.5	3
122	Endocrine and Metabolic Abnormalities in a Girl with Childhood <scp>W</scp>erner Syndrome: Case Report. Journal of the American Geriatrics Society, 2014, 62, 1404-1405.	2.6	2
123	Analysis of Death Due to Infectious Diseases in Patients Hospitalized in the Pediatric Ward of a Single Japanese Tertiary Medical Facility. Japanese Journal of Infectious Diseases, 2016, 69, 464-470.	1.2	2
124	First mixoploid infant with full triploidy in blood cells. Pediatrics International, 2016, 58, 1354-1355.	0.5	2
125	Vitamin D deficiency associated with dilated cardiomyopathy in early infancy caused by maternal cholestasis. Clinical Pediatric Endocrinology, 2018, 27, 187-192.	0.8	2
126	Language delay and developmental catch-up would be a clinical feature of pseudohypoparathyroidism type 1A during childhood. Endocrine Journal, 2019, 66, 215-221.	1.6	2

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127	Clinical features of infantile subcutaneous panniculitisâ€like Tâ€cell lymphoma. Pediatrics International, 2019, 61, 1261-1262.	0.5	2
128	Early Recovery of Height Velocity in Prepubertal Children With Acute Lymphoblastic Leukemia Treated by a Short Intensive Phase Without Cranial Radiation Therapy. Journal of Pediatric Hematology/Oncology, 2020, 42, 271-274.	0.6	2
129	A Child with Prostaglandin I₂-associated Thyrotoxicosis: Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 207-210.	0.9	2
130	Stability of amino acids, free and acylâ€carnitine in stored dried blood spots. Pediatrics International, 2022, 64, .	0.5	2
131	Rothmundâ€Thomson syndrome investigated by two nationwide surveys in Japan. Pediatrics International, 2022, 64, .	0.5	2
132	Preoperative Management With Nitrogen Inhalation Therapy for a Low-Birth Weight Infant With Tetralogy of Fallot and Absent Pulmonary Valve. Pediatric Cardiology, 2011, 32, 685-688.	1.3	1
133	Thyroid Follicular Carcinoma in a Fourteen-year-old Girl with Gravesâ€™ Disease. Clinical Pediatric Endocrinology, 2014, 23, 59-64.	0.8	1
134	Management of congenital diaphragmatic hernia with transposition of the great arteries. Pediatrics International, 2016, 58, 516-518.	0.5	1
135	Critical association of Pallisterâ€“Hall syndrome and congenital heart disease. Pediatrics International, 2019, 61, 827-828.	0.5	1
136	Nonâ€immune hydrops fetalis neonate born to a mother with yellow nail syndrome. Pediatrics International, 2019, 61, 313-315.	0.5	1
137	The autophagy reaction in the human umbilical cord: a potential marker for estimating fetal nutrition and neonatal growth. Journal of Maternal-Fetal and Neonatal Medicine, 2022, 35, 625-629.	1.5	1
138	A case of HDR syndrome coexisting with tetralogy of Fallot, with a novel GATA3 mutation, which manifested as a renal abscess. CEN Case Reports, 2021, 10, 241-243.	0.9	1
139	Hydrocortisone improved dexamethasoneâ€induced neuropsychological adverse effects. Pediatrics International, 2021, 63, 339-342.	0.5	1
140	Topoisomerase IIÎ² 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. Brain Tumor Pathology, 2021, 38, 109-121.	1.7	1
141	Metabolome Characteristics of Liver Autophagy Deficiency under Starvation Conditions in Infancy. Nutrients, 2021, 13, 3026.	4.1	1
142	Renal lesions mimicking acute focal bacterial nephritis in pediatric leukemia. Pediatrics International, 2022, 64, .	0.5	1
143	Fidgety Movements Assessment Accuracy Survey in Japan. International Journal of Environmental Research and Public Health, 2021, 18, 13428.	2.6	1
144	Inflammatory bowel diseaseâ€“like complication in anhidrotic ectodermal dysplasia. American Journal of Gastroenterology, 2000, 95, 3651-3652.	0.4	0

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145	<scp>NUP</scp>98â€“<scp>HOXC</scp>13 fusion gene in acute myeloid leukemia: Pediatric case. Pediatrics International, 2017, 59, 1105-1106.	0.5	0
146	Caffeine not associated with irritable behaviour in very low-birth-weight infants. Early Human Development, 2019, 137, 104835.	1.8	0
147	Gastric Carcinoma as Second Malignant Neoplasm in a Survivor From High-risk Neuroblastoma. Journal of Pediatric Hematology/Oncology, 2020, 42, 160-162.	0.6	0
148	A case of septo-optic dysplasia with hereditary hemorrhagic telangiectasia: a previously unrecognized combination of malformations. Clinical Dysmorphology, 2020, 29, 49-52.	0.3	0
149	Immune escape by loss of heterozygosis of HLA genes facilitated the transmission of diffuse large B cell lymphoma from mother to offspring. British Journal of Haematology, 2020, 189, e234-e237.	2.5	0
150	A Japanese infant presenting with hypocalcemic seizures resulting from hypovitaminosis D induced by non-celiac gluten sensitivity. Clinical Pediatric Endocrinology, 2021, 30, 105-110.	0.8	0
151	Atypical food proteinâ€induced enterocolitis syndrome after vaccinations. Pediatrics International, 2022, 64, e14891.	0.5	0
152	A longitudinal ulcer due to Yersinia pseudotuberculosis infection in a girl. Pediatrics International, 2022, 64, e15036.	0.5	0
153	Simultaneous monitoring of oxygen and carbon dioxide for Pittâ€Hopkins syndrome. Pediatrics International, 2022, 64, e15180.	0.5	0
154	Fetoâ€maternal hemorrhage with placental chorioangioma: Two case reports. Pediatrics International, 2022, 64, .	0.5	0