Shuichi Ito

List of Publications by Year in descending order

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Снисни Іто

#	Article	IF	CITATIONS
1	Eculizumab for paediatric patients with atypical haemolytic uraemic syndrome: full dataset analysis of post-marketing surveillance in Japan. Nephrology Dialysis Transplantation, 2023, 38, 414-424.	0.7	6
2	Effectiveness of Pediatric Teleconsultation to Prevent Skin Conditions in Infants and Reduce Parenting Stress in Mothers: Randomized Controlled Trial. JMIR Pediatrics and Parenting, 2022, 5, e27615.	1.6	2
3	Defining renal remission in an international cohort of 248 children and adolescents with lupus nephritis. Rheumatology, 2022, 61, 2563-2571.	1.9	8
4	Relations of mold, stove, and fragrance products on childhood wheezing and asthma: A prospective cohort study from the Japan Environment and Children's Study. Indoor Air, 2022, 32, .	4.3	7
5	New-onset pediatric nephrotic syndrome following Pfizer-BioNTech SARS-CoV-2 vaccination: a case report and literature review. CEN Case Reports, 2022, 11, 242-246.	0.9	20
6	Glucocorticoid discontinuation in pediatric-onset systemic lupus erythematosus: a single-center experience. Pediatric Nephrology, 2022, 37, 2131-2139.	1.7	4
7	Association between a single mother family and childhood undervaccination, and mediating effect of household income: a nationwide, prospective birth cohort from the Japan Environment and Children's Study (JECS). BMC Public Health, 2022, 22, 117.	2.9	5
8	Epigenetic insights into the pathogenesis of Kawasaki disease. Pediatric Research, 2022, 92, 347-348.	2.3	1
9	Association of maternal heavy metal exposure during pregnancy with isolated cleft lip and palate in offspring: Japan Environment and Children's Study (JECS) cohort study. PLoS ONE, 2022, 17, e0265648.	2.5	4
10	Impaired Interleukinâ€18 Signaling in Natural Killer Cells From Patients With Systemic Juvenile Idiopathic Arthritis. ACR Open Rheumatology, 2022, 4, 503-510.	2.1	6
11	Mycophenolate Mofetil after Rituximab for Childhood-Onset Complicated Frequently-Relapsing or Steroid-Dependent Nephrotic Syndrome. Journal of the American Society of Nephrology: JASN, 2022, 33, 401-419.	6.1	24
12	Persistence of Robust Humoral Immune Response in Coronavirus Disease 2019 Convalescent Individuals Over 12 Months After Infection. Open Forum Infectious Diseases, 2022, 9, ofab626.	0.9	6
13	Lipschütz ulcer induced by acute Epstein–Barr virus infection in a young girl. Pediatrics International, 2022, 64, e15022.	0.5	1
14	Clinical, Pathological, and Genetic Characteristics in Patients with Focal Segmental Glomerulosclerosis. Kidney360, 2022, 3, 1384-1393.	2.1	3
15	Neonatal wearable device for colorimetry-based real-time detection of jaundice with simultaneous sensing of vitals. Science Advances, 2021, 7, .	10.3	32
16	A case of multisystem inflammatory syndrome in children in a Japanese boy: with discussion of cytokine profile. Modern Rheumatology Case Reports, 2021, 5, 442-447.	0.7	16
17	Intravenous immunoglobulin for the treatment of Kawasaki disease. The Cochrane Library, 2021, 2021, .	2.8	1
18	Exposures associated with the onset of Kawasaki disease in infancy from the Japan Environment and Children's Study. Scientific Reports, 2021, 11, 13309.	3.3	8

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19	Genetic features of precursor Bâ€cell phenotype Burkitt leukemia with IGH―MYC rearrangement. Cancer Reports, 2021, , e1545.	1.4	1
20	A prospective cohort study of the association between the Apgar score and developmental status at 3Ayears of age: the Japan Environment and Children's Study (JECS). European Journal of Pediatrics, 2021, , 1.	2.7	2
21	<i>NUDT15</i> variants confer high incidence of second malignancies in children with acute lymphoblastic leukemia. Blood Advances, 2021, 5, 5420-5428.	5.2	4
22	Safety and efficacy of azilsartan in paediatric patients with hypertension: a phase 3, single-arm, open-label, prospective study. Clinical and Experimental Nephrology, 2021, , 1.	1.6	1
23	Establishing clinical remission criteria and the framework of a treat-to-targetÂalgorithm for Takayasu arteritis: Results of a Delphi exercise carried out by an expert panel of the Japan Research Committee of the Ministry of Health, Labour and Welfare for intractable vasculitis. Modern Rheumatology, 2021,	1.8	5
24	Urate-lowering therapy for gout and asymptomatic hyperuricemia in the pediatric population: a cross-sectional study of a Japanese health insurance database. BMC Pediatrics, 2021, 21, 581.	1.7	0
25	Influenza virus vaccination in pediatric nephrotic syndrome significantly reduces rate of relapse and influenza virus infection as assessed in a nationwide survey. Scientific Reports, 2021, 11, 23305.	3.3	4
26	Rituximab therapy for refractory steroid-resistant nephrotic syndrome in children. Pediatric Nephrology, 2020, 35, 17-24.	1.7	41
27	A novel truncating PAX2 mutation in a boy with renal coloboma syndrome with focal segmental glomerulosclerosis causing rapid progression to end-stage kidney disease. CEN Case Reports, 2020, 9, 19-23.	0.9	8
28	Allogeneic Bone Marrow Transplantation versus Peripheral Blood Stem Cell Transplantation for Hematologic Malignancies in Children: A Systematic Review and Meta-Analysis. Biology of Blood and Marrow Transplantation, 2020, 26, 88-93.	2.0	15
29	Prevalence of gout and asymptomatic hyperuricemia in the pediatric population: a cross-sectional study of a Japanese health insurance database. BMC Pediatrics, 2020, 20, 481.	1.7	6
30	Prevalence of germline <i>GATA2</i> and <i>SAMD9/9L</i> variants in paediatric haematological disorders with monosomy 7. British Journal of Haematology, 2020, 191, 835-843.	2.5	17
31	Influenza virus vaccination in children with nephrotic syndrome: insignificant risk of relapse. Clinical and Experimental Nephrology, 2020, 24, 1069-1076.	1.6	9
32	A 2â€yearâ€old patient with a diffuse intrinsic pontine glioma and radiationâ€induced moyamoya syndrome. Pediatric Blood and Cancer, 2020, 67, e28618.	1.5	0
33	IL-33 Is Essential for Adjuvant Effect of Hydroxypropyl-β-Cyclodexrin on the Protective Intranasal Influenza Vaccination. Frontiers in Immunology, 2020, 11, 360.	4.8	12
34	Successful Resolution of Recurrent Vaginal Pinworm Infection With Intermittent Albendazole Administration. Pediatric Infectious Disease Journal, 2020, 39, 254-255.	2.0	0
35	Clinical and genetic variability of PAX2-related disorder in the Japanese population. Journal of Human Genetics, 2020, 65, 541-549.	2.3	15
36	Clinical characteristics and treatment of 50 cases of Blau syndrome in Japan confirmed by genetic analysis of the <i>NOD2</i> mutation. Annals of the Rheumatic Diseases, 2020, 79, 1492-1499.	0.9	47

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37	JCS 2017 Guideline on Management of Vasculitis Syndrome ― Digest Version ―. Circulation Journal, 2020, 84, 299-359.	1.6	59
38	High prevalence of SMARCB1 constitutional abnormalities including mosaicism in malignant rhabdoid tumors. European Journal of Human Genetics, 2020, 28, 1124-1128.	2.8	0
39	Wearable Multi Vital Monitor for Newborns. , 2020, , .		2
40	Maternal multivitamin intake and orofacial clefts in offspring: Japan Environment and Children's Study (JECS) cohort study. BMJ Open, 2020, 10, e035817.	1.9	12
41	ATRT-11. PREVALENCE OF GERMLINE VARIANTS IN SMARCB1 INCLUDING SOMATIC MOSAICISM IN AT/RT AND OTHER RHABDOID TUMORS. Neuro-Oncology, 2020, 22, iii277-iii278.	1.2	0
42	Safety and effectiveness of eculizumab for pediatric patients with atypical hemolytic–uremic syndrome in Japan: interim analysis of post-marketing surveillance. Clinical and Experimental Nephrology, 2019, 23, 112-121.	1.6	31
43	Safety and effectiveness of eculizumab for adult patients with atypical hemolytic–uremic syndrome in Japan: interim analysis of post-marketing surveillance. Clinical and Experimental Nephrology, 2019, 23, 65-75.	1.6	15
44	Clinical significance of subcutaneous fat and fascial involvement in juvenile dermatomyositis. Modern Rheumatology, 2019, 29, 808-813.	1.8	10
45	Wearable Optical Device for Real-Time Monitoring of Newborn Jaundice. , 2019, , .		2
46	<i>Clostridium botulinum</i> infection in an exclusively breastâ€fed infant. Pediatrics International, 2019, 61, 1050-1051.	0.5	1
47	Clinical characteristics of HNF1B-related disorders in a Japanese population. Clinical and Experimental Nephrology, 2019, 23, 1119-1129.	1.6	31
48	Epidemiology and clinical features of childhood-onset anti-neutrophil cytoplasmic antibody–associated vasculitis: a clinicopathological analysis. Pediatric Nephrology, 2019, 34, 1425-1433.	1.7	15
49	Genetic landscape of Rett syndrome-like phenotypes revealed by whole exome sequencing. Journal of Medical Genetics, 2019, 56, 396-407.	3.2	30
50	Association between the clinical presentation of congenital anomalies of the kidney and urinary tract (CAKUT) and gene mutations: an analysis of 66 patients at a single institution. Pediatric Nephrology, 2019, 34, 1457-1464.	1.7	25
51	Antenatal Administration of Betamethasone Contributes to Intimal Thickening of the Rat Ductus Arteriosus. Circulation Journal, 2019, 83, 654-661.	1.6	2
52	Lisinopril versus lisinopril and losartan for mild childhood IgA nephropathy: a randomized controlled trial (JSKDC01 study). Pediatric Nephrology, 2019, 34, 837-846.	1.7	23
53	The Detection of Minor Clones with Somatic KIT D816V Mutations Using Droplet Digital PCR in Pediatric De Novo AML: AML-05 Trial from the Japanese Pediatric Leukemia/Lymphoma Study Group. Blood, 2019, 134, 1419-1419.	1.4	0
54	Clinically diverse phenotypes and genotypes of patients with branchio-oto-renal syndrome. Journal of Human Genetics, 2018, 63, 647-656.	2.3	31

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55	Prospective Study of Live Attenuated Vaccines for Patients with Nephrotic Syndrome Receiving Immunosuppressive Agents. Journal of Pediatrics, 2018, 196, 217-222.e1.	1.8	29
56	Novel recessive mutations in MSTO1 cause cerebellar atrophy with pigmentary retinopathy. Journal of Human Genetics, 2018, 63, 263-270.	2.3	19
57	Haploinsufficiency of A20 causes autoinflammatory and autoimmune disorders. Journal of Allergy and Clinical Immunology, 2018, 141, 1485-1488.e11.	2.9	100
58	Relapse of nephrotic syndrome during post-rituximab peripheral blood B-lymphocyte depletion. Clinical and Experimental Nephrology, 2018, 22, 110-116.	1.6	20
59	Coagulopathy as a complication of kidney biopsies in paediatric systemic lupus erythematosus patients with antiphospholipid syndrome. Nephrology, 2018, 23, 592-596.	1.6	2
60	A novel mutation in SLC1A3 causes episodic ataxia. Journal of Human Genetics, 2018, 63, 207-211.	2.3	42
61	Infliximab for the Treatment of Refractory Kawasaki Disease: A Nationwide Survey in Japan. Journal of Pediatrics, 2018, 195, 115-120.e3.	1.8	32
62	Study protocol: mycophenolate mofetil as maintenance therapy after rituximab treatment for childhood-onset, complicated, frequently-relapsing nephrotic syndrome or steroid-dependent nephrotic syndrome: a multicenter double-blind, randomized, placebo-controlled trial (JSKDC07). BMC Nephrology, 2018, 19, 302.	1.8	10
63	Effects of physical activity during pregnancy on preterm delivery and mode of delivery: The Japan Environment and Children's Study, birth cohort study. PLoS ONE, 2018, 13, e0206160.	2.5	32
64	Changes in Hemoglobin Concentrations Post-immunoglobulin Therapy in Patients with Kawasaki Disease: A Population-Based Study Using a Claims Database in Japan. Paediatric Drugs, 2018, 20, 585-591.	3.1	5
65	Study protocol: high-dose mizoribine with prednisolone therapy in short-term relapsing steroid-sensitive nephrotic syndrome to prevent frequent relapse (JSKDC05 trial). BMC Nephrology, 2018, 19, 223.	1.8	6
66	Diplotype analysis of NUDT15 variants and 6-mercaptopurine sensitivity in pediatric lymphoid neoplasms. Leukemia, 2018, 32, 2710-2714.	7.2	26
67	A novel SLC9A1 mutation causes cerebellar ataxia. Journal of Human Genetics, 2018, 63, 1049-1054.	2.3	28
68	Tissue-type plasminogen activator contributes to remodeling of the rat ductus arteriosus. PLoS ONE, 2018, 13, e0190871.	2.5	11
69	A consensus statement on health-care transition of patients with childhood-onset chronic kidney diseases: providing adequate medical care in adolescence and young adulthood. Clinical and Experimental Nephrology, 2018, 22, 743-751.	1.6	12
70	Is There Difference between the Survival Rate and Treatment Related Complications between Peripheral Blood Stem Cell Transplantation and Bone Marrow Transplantation for Pediatric Hematological Malignancy?: Systematic Review and Meta-Analysis. Blood, 2018, 132, 3378-3378.	1.4	0
71	Identification of novel <i><scp>SNORD118</scp></i> mutations in seven patients with leukoencephalopathy with brain calcifications and cysts. Clinical Genetics, 2017, 92, 180-187.	2.0	28
72	Successful therapy switch from eculizumab to mycophenolate mofetil in a girl with DEAP-HUS. Pediatric Nephrology, 2017, 32, 1997-1998.	1.7	3

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73	Simultaneous development of Kawasaki disease following acute human adenovirus infection in monozygotic twins: A case report. Pediatric Rheumatology, 2017, 15, 39.	2.1	15
74	Long-term outcome of childhood-onset complicated nephrotic syndrome after a multicenter, double-blind, randomized, placebo-controlled trial of rituximab. Pediatric Nephrology, 2017, 32, 2071-2078.	1.7	35
75	Diagnostic challenge in a patient with nephropathic juvenile cystinosis: a case report. BMC Nephrology, 2017, 18, 300.	1.8	6
76	Childhood-onset inflammatory bowel diseases associated with mutation of Wiskott-Aldrich syndrome protein gene. World Journal of Gastroenterology, 2017, 23, 8544-8552.	3.3	14
77	Infliximab for Intravenous Immunoglobulin-Resistant Patients with Kawasaki Disease: A Single-Institute Study. Nihon Shoni Junkanki Gakkai Zasshi = Pediatric Cardiology and Cardiac Surgery, 2017, 33, 43-49.	0.0	0
78	Significant improvement in Fabry disease podocytopathy after 3Âyears of treatment with agalsidase beta. Pediatric Nephrology, 2016, 31, 1369-1373.	1.7	16
79	Clinical guides for atypical hemolytic uremic syndrome in Japan. Pediatrics International, 2016, 58, 549-555.	0.5	19
80	Clinical guides for atypical hemolytic uremic syndrome in Japan. Clinical and Experimental Nephrology, 2016, 20, 536-543.	1.6	41
81	Milder progressive cerebellar atrophy caused by biallelic SEPSECS mutations. Journal of Human Genetics, 2016, 61, 527-531.	2.3	30
82	Evolution of IgA nephropathy into anaphylactoid purpura in six cases—further evidence that IgA nephropathy and Henoch–Schonlein purpura nephritis share common pathogenesis. Pediatric Nephrology, 2016, 31, 779-785.	1.7	18
83	Risk factors for relapse and long-term outcome in steroid-dependent nephrotic syndrome treated with rituximab. Pediatric Nephrology, 2016, 31, 89-95.	1.7	24
84	Efficacy and safety of eculizumab in childhood atypical hemolytic uremic syndrome in Japan. Clinical and Experimental Nephrology, 2016, 20, 265-272.	1.6	22
85	Proteinuria during Follow-Up Period and Long-Term Renal Survival of Childhood IgA Nephropathy. PLoS ONE, 2016, 11, e0150885.	2.5	26
86	Childhood SLE and lupus nephritis. Japanese Journal of Pediatric Nephrology, 2016, 29, 1-7.	0.0	0
87	Early discharge after the initial treatment for pediatric idiopathic nephrotic syndrome. Japanese Journal of Pediatric Nephrology, 2016, 29, 161-165.	0.0	0
88	Growth impairment in children with pre-dialysis chronic kidney disease in Japan. Clinical and Experimental Nephrology, 2015, 19, 1142-1148.	1.6	15
89	Creatinine-based equation to estimate the glomerular filtration rate in Japanese children and adolescents with chronic kidney disease. Clinical and Experimental Nephrology, 2014, 18, 626-633.	1.6	194
90	Methotrexate therapy for juvenile rheumatoid arthritis in Japan — surveillance with a questionnaire at seven main facilities. Japanese Journal of Rheumatology, 1999, 9, 229-237.	0.0	2

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91	Methotrexate therapy for juvenile rheumatoid arthritis in Japan — surveillance with a questionnaire at seven main facilities. Japanese Journal of Rheumatology, 1999, 9, 229-237.	0.0	Ο