

論文発表者 Authors	論文名 Title	Journal	Volume	Issue	Pages
Fujimura J, Nozu K, Yamamura T, Minamikawa S, Nakanishi K, Horinouchi T, Nagano C, Sakakibara N, Nakanishi K, Shima Y, Miyako K, Nozu Y, Morisada N, Nagase H, Ninchoji T, Kaito H, Iijima K	Clinical and Genetic Characteristics in Patients With Gitelman Syndrome.	Kidney Int Rep	4		119–125
Hashikami K, Asahina M, Nozu K, Iijima K, Nagata M, Takeyama M	Establishment of X-linked Alport syndrome model mice with a Col4a5 R471X mutation	Biochem Biophys Rep	17		81–86
Horinouchi T, Morisada N, Uemura H, Kobayashi D, Nozu K, Okamoto N, Iijima K:	Male CDPX2 patient with EBP mosaicism and asymmetrically lateralized skin lesions with strict midline demarcation.	Am J Med Genet A	179		1315–1318
Horinouchi T, Nozu K, Yamamura T, Minamikawa S, Nagano C, Sakakibara N, Nakanishi K, Shima Y, Morisada N, Ishiko S, Aoto Y, Nagase H, Takeda H, Rossanti R, Kaito H, Matsuo M, Iijima K	Determination of the pathogenicity of known COL4A5 intronic variants by in vitro splicing assay.	Sci Rep	9		12696
Ishimori S, Kaito H, Shima Y, Kamioka I, Hamahira K, Nozu K, Nakanishi K, Tanaka R, Yoshikawa N, Iijima K	Clinicopathological characteristics and renal outcomes of childhood-onset lupus nephritis with acute kidney injury: A multicenter study.	Mod Rheumatol	29		970–976
Kyono Y, Nozu K, Nakagawa T, Takami Y, Fujita H, Ioroi T, Kugo M, Iijima K, Kamiyoshi N	Combination of furosemide and fludrocortisone as a loading test for diagnosis of distal renal tubular acidosis in a pediatric case.	CEN Case Rep			
Lee JM, Nozu K, Choi DE, Kang HG, Ha IS, Cheong HI	Features of Autosomal Recessive Alport Syndrome: A Systematic Review.	J Clin Med	8		
Nagano C, Morisada N, Nozu K, Kamei K, Tanaka R, Kanda S, Shiona S, Araki Y, Ohara S, Matsumura C, Kasahara K, Mori Y, Seo A, Miura K, Washiyama M, Sugimoto K, Harada R, Tazoe S, Kourakata H, Enseki M, Aotani D, Yamada T, Sakakibara N, Yamamura T, Minamikawa S, Ishikura K, Ito S, Hattori M, Iijima K	Clinical characteristics of HNF1B-related disorders in a Japanese population.	Clin Exp Nephrol	23		1119–1129
Nagano C, Nozu K, Yamamura T, Minamikawa S, Fujimura J, Sakakibara N, Nakanishi K, Horinouchi T, Iwafuchi Y, Kusuhara S, Matsumiya W, Yoshikawa N, Iijima K	TGFBI-associated corneal dystrophy and nephropathy: a novel syndrome?	CEN Case Rep	8		14–17
Nagano C, Sako M, Kamei K, Ishikura K, Nakamura H, Nakanishi K, Omori T, Nozu K, Iijima K	Study protocol: multicenter double-blind, randomized, placebo-controlled trial of rituximab for the treatment of childhood-onset early-stage uncomplicated frequently relapsing or steroid-dependent nephrotic syndrome (JSKDC10 trial)	BMC Nephrol	20		293
Nakanishi K, Okamoto T, Nozu K, Hara S, Sato Y, Hayashi A, Takahashi T, Nagano C, Sakakibara N, Horinouchi T, Fujimura J, Minamikawa S, Yamamura T, Rossanti R, Nagase H, Kaito H, Ariga T, Iijima K	Pair analysis and custom array CGH can detect a small copy number variation in COQ6 gene.	Clin Exp Nephrol	23		669–675
Nozu K, Nakanishi K, Abe Y, Udagawa T, Okada S, Okamoto T, Kaito H, Kanemoto K, Kobayashi A, Tanaka E, Tanaka K, Hama T, Fujimaru R, Miwa S, Yamamura T, Yamamura N, Horinouchi T, Minamikawa S, Nagata M, Iijima K	A review of clinical characteristics and genetic backgrounds in Alport syndrome.	Clin Exp Nephrol,	23		158–168
Okamoto T, Nozu K, Iijima K, Ariga T	Germline mosaicism is a pitfall in the diagnosis of “sporadic” X-linked Alport syndrome.	J Nephrol	32		155–159
Rossanti R, Shono A, Miura K, Hattori M, Yamamura T, Nakanishi K, Minamikawa S, Fujimura J, Horinouchi T, Nagano C, Sakakibara N, Kaito H, Nagase H, Morisada N, Asanuma K, Matsuo M, Nozu K, Iijima K	Molecular assay for an intronic variant in NUP93 that causes steroid resistant nephrotic syndrome.	J Hum Genet	64		673–679
Sakakibara N, Morisada N, Nozu K, Nagatani K, Ohta T, Shimizu J, Wada T, Shima Y, Yamamura T, Minamikawa S, Fujimura J, Horinouchi T, Nagano C, Shono A, Ye MJ, Nozu Y, Nakanishi K, Iijima K	Clinical spectrum of male patients with OFD1 mutations.	J Hum Genet	64		3–9
Shima Y, Nakanishi K, Sako M, Saito-Oba M, Hamasaki Y, Hataya H, Honda M, Kamei K, Ishikura K, Ito S, Kaito H, Tanaka R, Nozu K, Nakamura H, Ohashi Y, Iijima K, Yoshikawa N	apanese Study Group of Kidney Disease in C: Lisinopril versus lisinopril and losartan for mild childhood IgA nephropathy: a randomized controlled trial (JSKDC01 study).	Pediatr Nephrol	34		837–846
Yamamura T, Nozu K, Minamikawa S, Horinouchi T, Sakakibara N, Nagano C, Aoto Y, Ishiko S, Nakanishi K, Shima Y, Nagase H, Rossanti R, Ye MJ, Nozu Y, Ishimori S, Morisada N, Kaito H, Iijima K	Comparison between conventional and comprehensive sequencing approaches for genetic diagnosis of Alport syndrome.	Mol Genet Genomic Med	7		e883
Uemura S, Ishida T, Mon Thwin KK, Yamamoto N, Tamura A, Kishimoto K, Hasegawa D, Kosaka Y, Nino N, San Lin K, Takafuji S, Mori T, Iijima K, Nishimura N	Dynamics of Minimal Residual Disease in Neuroblastoma Patients	Front Oncol	9	455	
Nakano Y, Hasegawa D, Stewart DR, Schultz KAP, Harris AK, Hirato J, Uemura S, Tamura A, Saito A, Kawamura A, Yoshida M, Yamasaki K, Yamashita S, Ushijima T, Kosaka Y, Ichimura K, Dehner LP, Hill AD	Presacral Malignant Teratoid Neoplasm in Association With Pathogenic DICER1 Variation	Mod Pathol	32	12	1744–1750
Mon Thwin KK, Ishida T, Uemura S, Yamamoto N, San Lin K, Tamura A, Kozaki A, Saito A, Kishimoto K, Mori T, Hasegawa D, Kosaka Y, Nino N, Takafuji S, Iijima K, Nishimura N	Level of Seven Neuroblastoma-Associated mRNAs Detected by Droplet Digital PCR Is Associated With Tumor Relapse/Regrowth of High-Risk Neuroblastoma Patients	J Mol Diagn			
Uemura S, Mori T, Nino N, Sakakibara N, Takafuji T, Myojin S, Takami Y, Morioka I, Nishimura N, Kugo M, Iijima K	An infant with refractory cytomegalovirus-induced thrombocytopenia	Clin Case Rep			
Tamura A, Yamamoto N, Nino N, Ichikawa T, Nakatani N, Nakamura S, Saito A, Kozaki A, Kishimoto K, Ishida T, Yoshida M, Akasaka Y, Hasegawa D, Kosaka Y	Pazopanib maintenance therapy after tandem high-dose chemotherapy for disseminated Ewing sarcoma	Int Cancer Conf J	8	3	95–100
Kishimoto K, Hasegawa D, Kawasaki K, Tamura A, Yamamoto N, Saito A, Kozaki A, Ishida T, Kosaka Y.	Early posttransplant plasma ADAMTS13 activity reduction in stem cell transplantation: a prospective study of 46 pediatric patients	Bone Marrow Transplant	54	12	1926–1929

論文発表者 Authors	論文名 Title	Journal	Volume	Issue	Pages
Uchida A, Tanimura K, Morizane M, Fujioka K, Morioka I, Oohashi M, Minematsu T, Yamada H.	Clinical factors associated with congenital cytomegalovirus infection: A cohort study of pregnant women and newborns.	Clin Infect Dis.			
Nakasuji Y, Tanimura K, Sasagawa Y, Imafuku H, Morizane M, Fujioka K, Ohji G, Yamada H.	Case report of eight pregnant women with syphilis.	J Infect Chemother.			
Yamamoto K, Fukushima S, Mishima Y, Hashimoto M, Yamakawa K, Fujioka K, Iijima K, Yano I.	Pharmacokinetic assessment of alprazolam-induced neonatal abstinence syndrome using physiologically based pharmacokinetic model.	Drug Metab Pharmacokinet.	34	6	400-402
Matsumoto M, Nagano N, Awano H, Ohyama S, Fujioka K, Iwatani S, Urakami T, Iijima K, Morioka I.	Incidence and Neonatal Risk factors of Short Stature and Growth Hormone treatment in Japanese Preterm Infants Born Small for Gestational Age.	Sci Rep.	9	1	12238
Ohyama S, Fujioka K, Fukushima S, Abe S, Ashina M, Ikuta T, Nishida K, Matsumoto H, Nakamachi Y, Tanimura K, Yamada H, Iijima K.	Diagnostic Value of Cytomegalovirus IgM Antibodies at Birth in PCR-Confirmed Congenital Cytomegalovirus Infection.	Int J Mol Sci.	20	13	
Fukushima S, Morioka I, Ohyama S, Nishida K, Iwatani S, Fujioka K, Mandai T, Matsumoto H, Nakamachi Y, Deguchi M, Tanimura K, Iijima K, Yamada H.	Prediction of poor neurological development in patients with symptomatic congenital cytomegalovirus diseases after oral valganciclovir treatment.	Brain Dev.	41	9	743-750
Abe S, Fujioka K.	Mesenchymal stem cell therapy for neonatal intraventricular hemorrhage: a long way to go?	Ann Transl Med.	7	5	79
Ohyama S, Morioka I, Fukushima S, Yamana K, Nishida K, Iwatani S, Fujioka K, Matsumoto H, Imanishi T, Nakamachi Y, Deguchi M, Tanimura K, Iijima K, Yamada H.	Efficacy of Valganciclovir Treatment Depends on the Severity of Hearing Dysfunction in Symptomatic Infants with Congenital Cytomegalovirus Infection.	Int J Mol Sci.	20	6	
Htun Y, Nakamura S, Nakao Y, Mitsue T, Nakamura M, Yamato S, Jinnai W, Koyano K, Ohta K, Morimoto A, Wakabayashi T, Sugino M, Fujioka K, Kato I, Kondo S, Yasuda S, Miki T, Ueno M, Kusaka T.	Hydrogen ventilation combined with mild hypothermia improves short-term neurological outcomes in a 5-day neonatal hypoxia-ischaemia piglet model.	Sci Rep.	9	1	4088
Iwatani S, Yoshida M, Yamana K, Kurokawa D, Kuroda J, Thwin KKM, Uemura S, Takafuji S, Nino N, Koda T, Mizobuchi M, Nishiyama M, Fujioka K, Nagase H, Morioka I, Iijima K, Nishimura N.	Isolation and Characterization of Human Umbilical Cord-derived Mesenchymal Stem Cells from Preterm and Term Infants.	J Vis Exp.		143	
Fukushima S, Fujioka K, Ashina M, Ohyama S, Ikuta T, Nishida K, Miyauchi H, Okata Y, Bitoh Y, Tanimura K, Deguchi M, Yamada H, Iijima K.	Fetal Primary Small Bowel Volvulus Associated with Acute Gastric Dilatation Detected by Ultrasonography.	Kobe J Med Sci.	64	4	E157-E159
Fujioka K, Nishida K, Ashina M, Abe S, Fukushima S, Ikuta T, Ohyama S, Morioka I, Iijima K.	DNA methylation of the Rtl1 promoter in the placentas with fetal growth restriction.	Pediatr Neonatol.	60	5	512-516
Ashina M, Fujioka K, Totsu S, Shoji H, Miyazawa T, Wada K, Iijima K, Morioka I.	Feeding interval and use of donor breast milk for very low birthweight infants: A nationwide survey in Japan.	Pediatr Neonatol.	60	3	245-251
Suga S, Hoshina T, Ichikawa S, Araki S, Kusuvara K.	A survey of the implementation status of selected infection control strategies in neonatal intensive care units in Japan.	J Hosp Infect.			
Miyauchi H, Fujioka K, Okubo S, Nishida K, Ashina M, Ikuta T, Okata Y, Maeda K, Iijima K, Bitoh Y.	Insulin Therapy for Hyperglycemia in Neonatal Sepsis Using a Preterm Mouse Model.	Pediatr Int.			
Nishiyama M, Nagase H, Tomioka K, Tanaka T, Yamaguchi H, Ishida Y, Toyoshima D, Fujita K, Maruyama A, Sasaki K, Oyazato Y, Nakagawa T, Takami Y, Nozu K, Nishimura N, Nakashima I, Iijima K	Clinical time course of pediatric acute disseminated encephalomyelitis	Brain Dev	41	6	531-537
Tanaka T, Nagase H, Yamaguchi H, Ishida Y, Tomioka K, Nishiyama M, Toyoshima D, Maruyama A, Fujita K, Nozu K, Nishimura N, Kurosawa H, Tanaka R, Iijima K	Predicting the outcomes of targeted temperature management for children with seizures and/or impaired consciousness accompanied by fever without known etiology	Brain Dev	41	7	604-613
Yamaguchi H, Nagase H, Nishiyama M, Tokumoto S, Toyoshima D, Akasaka Y, Maruyama A, Iijima K	Fibrocartilaginous Embolism of the Spinal Cord in Children: A Case Report and Review of Literature	Pediatr Neurol	99		3-6
Awano H, Nagai M, Bo R, Muraio M, Ishida Y, Tanaka T, Tomioka K, Nishiyama M, Nagase H, Iijima K	Preliminary Effectiveness and Safety of High Frequency Oscillation in Addition to Mechanical Insufflation and Exsufflation for Intratracheal Mucus Removal in Patients With Neuromuscular Disease: Protocol for a Prospective Study	JMIR Res Protoc	8	6	e12102
Tomioka K, Nishiyama M, Nagase H, Ishida Y, Tanaka T, Tokumoto S, Yamaguchi H, Toyoshima D, Maruyama A, Fujita K, Aoki K, Seino Y, Nozu K, Nishimura N, Kurosawa H, Iijima K	Detailed clinical course of fatal acute encephalopathy in children	Brain Dev	41	8	691-698
Nishimura A, Yamaguchi H, Ito Y, Tokumoto S, Toyoshima D, Kasai M, Maruyama A	Empyema necessitatis due to Pseudomonas aeruginosa in a child with cerebral palsy	J Infect Public Health			
Yamada Y, Yamaguchi H, Ito Y, Takeuchi N, Kasai M	Pyogenic sacroiliitis caused by pneumococcal serotype 16F in a child	Pediatr Int			
Yamaguchi H, Maruyama A	Reply to the comments on "Spontaneous spinal epidural hematoma mimicking Guillain-Barre Syndrome".	Brain Dev	41	9	827-828
Yamaguchi H, Nagase H, Yoshida S, Tokumoto S, Hayashi K, Toyoshima D, Kurosawa H, Tanaka T, Maruyama A, Iijima K.	Acute encephalopathy with biphasic seizures and late reduced diffusion accompanied by Takotsubo cardiomyopathy	Brain Dev	41	3	305-309

論文発表者 Authors	論文名 Title	Journal	Volume	Issue	Pages
Nagao H, Yamaguchi, H, Ito Y, Kasai M	Infective Endocarditis Due to Streptococcus gallolyticus subsp. pasteurianus with Pulmonary Vein Obstruction	Indian J Pediatr	86	2	192
Yamaguchi H, Nagase H, Nishiyama M, Tokumoto S, Ishida Y, Tomioka K, Tanaka T, Fujita K, Toyoshima D, Nishimura N, Kurosawa H, Nozu K, Maruyama A, Tanaka R, Iijima K	Nonconvulsive Seizure Detection by Reduced-Lead Electroencephalography in Children with Altered Mental Status in the Emergency Department	J Pediatr	207		213-219
Kato H, Yamaguchi H, Ito Y, Imuta N, Nishi J, Kasai M	Escherichia coli O157 Enterocolitis Followed by Non-diarrheagenic Escherichia coli Bacteremia	Indian J Pediatr	86	8	750
Kondo A, Yamaguchi H, Ishida Y, Toyoshima D, Azumi M, Akutsu N, Koyama J, Kurosawa H, Kawamura A, Maruyama A	Spontaneous spinal epidural hematoma mimicking Guillain-Barre Syndrome	Brain Dev	41	4	392-395
Rani AQM, Farea MA, Maeta K, Kawaguchi T, Awano H, Nagai M, Nishio H, Matsuo M	Identification of the shortest splice variant of Dp71, together with five known variants, in glioblastoma cells	Biochemical and Biophysical Research Communications	508	2	640-645
Matsuo M, Awano H, Nishio H	Can urinary titin be used for predicting Duchenne muscular dystrophy?	Clinica Chimica Acta	490		162
Matsumoto M, Awano H, Hirota Y, Nagai M, Bo R, Matsuoka A, Hamaguchi T, Takeuchi T, Nakagawa Y, Ogawa W, Iijima K	The prescription rates of glucagon for hypoglycemia by pediatricians and physicians are low in Japan	Endocrine	64	2	233-238
Shimomura H, Lee T, Tanaka Y, Awano H, Itoh K, Nishino I, Takeshima Y	Two closely spaced mutations in cis result in Ullrich congenital muscular dystrophy	Hum Genome Var	6	21	
Matsuo M, Awano H, Maruyama N, Nishio H	Titin fragment in urine: A nonvasive biomarker of muscle degradation	Advances in Clinical Chemistry	90	6	1-23
Katsuyama A, Kusuhara S, Awano F, Nagase H, Matsumiya W, Nakamura M	A case of probable Vogt-Koyanagi-Harada disease in a 3-year-old girl	BMC Ophthalmology	19	1	179
Rani AQM, Maeta K, Kawaguchi T, Awano H, Nagai M, Nishio H, Matsuo M	Schwann cell-specific Dp116 is expressed in glioblastoma cells, revealing two novel DMD gene splicing patterns	Biochemistry and Biophysics Reports	20		
Takeuchi A, Tode C, Nishino M, Wijaya YOS, Niba ETE, Awano H, Takeshima Y, Saito T, Saito K, Lai PS, Bouike Y, Nishio H, Shinohara M	Newborn screening for spinal muscular atrophy: DNA preparation from dried blood spot and DNA polymerase selection in PCR	Kobe J Med Sci	65	6	E95-99
Komaki H, Maegaki Y, Matsumura T, Shiraishi K, Awano H, Nakamura A, Kinoshita S, Ogata K, Ishigaki K, Satio S, Funato M, Kuru S, Nakayama, T, Iwata Y, Yajima H, Takeda S	Early phase 2 trial of TAS-205 in patients with Duchenne muscular dystrophy	Annals of Clinical and Translational Neurology			
Yamaguchi H, Taniguchi-Ikeda M, Nagase H, Ito Y, Tokumoto S, Toyoshima D, Enkhjargal S, Nishiyama M, Awano H, Kurosawa H, Kasai M, Maruyama A, Iijima K	Acute Rhabdomyolysis Following Viral Infection with Coxsackie A4 in a 50-Day-Old Infant with Fukuyama Congenital Muscular Dystrophy	journal of infection and chemotherapy			