Poster Presentation

Poster 1 Genetic abnormalities: Study

P-1

Analysis of TRPC6 variation in children with steroid-resistant nephritic syndrome

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P-2

Sphingomyelin phosphodiesterase acid-like 3b (SMPDL3b) mediates TLR3 signaling in human podocytes

Shojiro Watanabe, Takao Karasawa, Shun Hashimoto, Hidenori Umetsu, Riko Sato, Masashi Fujita, Tomomi Aizawa, Koji Tsugawa, Hiroshi Tanaka

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P-3

Association of podocyte lesions with novel CUBN gene mutations in three isolated proteinuric children

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P-4

Genotype-phenotype correlations in pediatric patients with HNF1B mutations

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P-5

The Clinical and Genetic Heterogeneity of Chinese Children with TTC21B Mutation Caused Nephronophthisis-Related Ciliopathies

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Poster 2 Genetic Abnormalities: Case

P-6

A Novel Compound Heterozygous Mutation in LAMB2 Gene in a Chinese Boy With Isolated Proteinuria

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P-7

A PKD1 and SLC36A2 mutation in pediatric patient with polycystic kidney and nephrocalcinosis

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P-8

A novel de novo truncating TRIM8 mutation associated with childhood-onset focal segmental glomerulosclerosis without epileptic encephalopathy

Yoko Shirai[1], Kenichiro Miura[1], Naoto Kaneko[1], Kiyonobu Ishizuka[1], Amane Endo[2], Taeko Hashimoto[3], Shoichiro Kanda[4], Yutaka Harita[4], Motoshi Hattori[1]

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Mutations in TTC21B in two siblings with young childhood onset cause different phenotypes in China

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P-10

Exercise-induced acute kidney injury in an adolescent boy with a mutation of SCL22A12 gene causing renal hypouricemia

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A case of focal segmental glomerulosclerosis (FSGS) in an infant in whom hepatic fibrosis were observed and mutations in TTC21B were identified

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P-12

A case of Gordon syndrome with a CUL3 mutation presented with short stature

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Two male relatives with OFD1 mutations

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A case with Proteinuria and Renal biopsy findings resembling Fabry disease

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A HNF1B-related disorders: two cases in Japanese

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A case report of hereditary renal tubule dysplasia with anemia

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Poster 3 Alport syndrome

P-17

Alport syndrome overlapping with IgA nephropathy: a case report

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A Case of Autosomal Recessive Alport Syndrome with Acute Kidney Injury on Lisinopril medication

Daisuke Nakanishi[1], Wataru Shimabukuro[1], Kazuya Hamada[1], Shougo Nakada[1], Masatsugu Uehara[2], Shigeru Fukuyama[2], Tomoo Kise[2], Kandai Nozu[3], Kazumoto Iijima[3], Noriko Kinjo[1], Koichi Nakanishi[1]

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P-19

Study on the timing of renal biopsy in Alport syndrome

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Poster 4 Congenital Disease

P-20

A case with familial short - rib thoracic dysplasia with or without polydactyly

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Wilson's disease in a child with nephrotic syndrome: a case report and literature review

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Withdrawn

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A case of primary hyperoxaluria diagnosed by urine gas chromatography / mass spectrometry

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A case of Russell-Silver syndrome born in the phenotype of Potter sequence and renal malformations

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A case of cyclic vomiting syndrome with persistent strong hypertension

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Long-term outcome of Bartter syndrome

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Poster 5 IgA nephropathy/HSPN

P-27

Involvement of activated cytotoxic T lymphocytes and natural killer cells in Henoch–Schönlein purpura nephritis

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MYH9 associated disorders with the onset of Henoch- Schönlein purpuric nephritis: a case report

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A case of glomerulonephritis with the glomerular basement membrane abnormality and mesangial IgA deposition

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P-30

The role of school urinalysis in the discovery of IgA nephropathy in the Kyushu / Okinawa area

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Poster 6 C3 nephropathy/aHUS

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Discrepancies between CH50 and Free C5 concentrations in 4 cases of aHUS treated with ravulizumab

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C3 Glomerulonephritis: A Entity of Complement Driven-Renal Disease in Children with Unsatisfactory Outcome

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A case of crescent C3 glomerulonephritis (C3GN) difficult to distinguish from monoclonal immunoglobulin (Ig)G deposits (PGNMID)

Takashi Honda[1,2], Riku Hamada[1,2], Atsuko Anno[2], Kantaro Kobayashi[2], Keiji Akamine[1], Chikako Terano[1], Ryoko Harada[1], Hideyo Oguchi[3], Takatoshi Tsuchihashi[2], Kazutoshi Shibuya[4], Ken Sakai[3], Hideo Cho[2], Yutaka Yamaguchi[5], Masataka Honda[1], Hiroshi Hataya[1]

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WT1 disorder associated with C3 glomerulonephritis? Difficulty in differential diagnosis: A case report

Lin Zhilang, Rong Liping, Pei Yuxin, Qiu Yuanquan, Yue Zhihui, Mo Ying, Jiang Xiaoyun

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Poster 7 Glimeruronephritis

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Periodic Fever with Pauci-immune Crescentic Glomerulonephritis in a Chinese Girl

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Focal segmental glomerulosclerosis in a patient with atypical progeroid

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A case of shunt nephritis characterized by Lupus-like syndrome

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Two cases of children who had hematuria with or without mild proteinuria with normal renal function and positive anti-neutrophil cytoplasmic antibody

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Staphylococcus associated glomerulonephritis in child: a case report and review of literature

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Withdrawn

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Successful Treatment of two cases with Streptococcus Pneumoniae-associated Hemolytic Uremic Syndrome by anti-T-antibody Negative plasma exchange

Ling Hou, Yubin Wu, Yue Du

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Case Report: Were Membranous nephropathy Associated With Evans' syndrome in Childhood?

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Poster 8 Nephrotic sindrome

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Nephrotic syndrome relapse in a boy with COVID-19

Akio Kagawa, Takuji Enya, Yuichi Morimoto, Rina Oshima, Kohei Miyazaki, Tomoki Miyazawa, Mitsuru Okada, Keisuke Sugimoto

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P-44

A case of nephrotic syndrome caused by COQ6 gene mutation with significant podocyte hypertrophy

Baige Su, Suxia Wang, Xuhui Zhong, Xiaoyu Liu, Yanqin Zhang

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A steroid-resistant nephrotic syndrome in a patient with coenzyme Q6 mutation: a case report of two sibling

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Intermediate stage of Citrin deficiency with nephrotic syndrome: a case report and literature review

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Withdrawn

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A case of thrombotic microangiopathy with nephrotic syndrome

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Clinical analysis of children having primary nephrotic syndrome complicated with posterior reversible encephalopathy syndrome on treatment of immunosuppressants

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Pathological Findings of Acute Kidney Injury in a Nephrotic Syndrome Patient Requiring Emergency Dialysis during Methylprednisolone Pulse Therapy

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Poster 9 CKD/Dialysis

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Schimke immuno-osseous dysplasia; disease progression after kidney transplantation

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Risk factors for post-nephrectomy hypotension in pediatric patients: a single-center observational study

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Calcifications in children on maintenance peritoneal dialysis

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Poster 10 Others

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A case of DICER1 syndrome presenting as Wilm's tumor and contralateral progressive cystic nephroma; Newly Known Cancer Predisposition Syndrome

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A case of a 17-month-old boy presenting abnormal pelvic mass

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Additional preventative effect of probiotics on low-dose antibiotic prophylaxis for urinary tract infection in children with vesicoureteral reflux

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Group B streptococcal renal abscess in a 17-year-old girl with type 1 diabetes mellitus

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