Yo Han Ahn

List of Publications by Year
in descending order

53	474	686830 13	752256 20
papers	citations	h-index	g-index

55	55	55	824
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Efficacy and safety of rituximab in childhood-onset, difficult-to-treat nephrotic syndrome. Medicine (United States), 2018, 97, e13157.	0.4	41
2	COQ6 Mutations in Children With Steroid-Resistant Focal Segmental Glomerulosclerosis and Sensorineural Hearing Loss. American Journal of Kidney Diseases, 2017, 70, 139-144.	2.1	40
3	Long-term renal outcome in children withOCRLmutations: retrospective analysis of a large international cohort. Nephrology Dialysis Transplantation, 2016, 33, gfw350.	0.4	27
4	Loss of podocalyxin causes a novel syndromic type of congenital nephrotic syndrome. Experimental and Molecular Medicine, 2017, 49, e414-e414.	3.2	27
5	Targeted exome sequencing resolves allelic and the genetic heterogeneity in the genetic diagnosis of nephronophthisis-related ciliopathy. Experimental and Molecular Medicine, 2016, 48, e251-e251.	3.2	26
6	Genotypeâ€“phenotype analysis of pediatric patients with WT1 glomerulopathy. Pediatric Nephrology, 2017, 32, 81-89.	0.9	25
7	Measurement of Fluid Status Using Bioimpedance Methods in Korean Pediatric Patients on Hemodialysis. Journal of Korean Medical Science, 2017, 32, 1828.	1.1	25
8	Mental health and psychosocial adjustment in pediatric chronic kidney disease derived from the KNOW-Ped CKD study. Pediatric Nephrology, 2019, 34, 1753-1764.	0.9	23
9	NUP107mutations in children with steroid-resistant nephrotic syndrome. Nephrology Dialysis Transplantation, 2017, 32, gfw103.	0.4	22
10	Genetic Study in Korean Pediatric Patients with Steroid-Resistant Nephrotic Syndrome or Focal Segmental Glomerulosclerosis. Journal of Clinical Medicine, 2020, 9, 2013.	1.0	20
11	Genotype and Phenotype Analyses in Pediatric Patients with HNF1B Mutations. Journal of Clinical Medicine, 2020, 9, 2320.	1.0	18
12	Targeted Exome Sequencing Provided Comprehensive Genetic Diagnosis of Congenital Anomalies of the Kidney and Urinary Tract. Journal of Clinical Medicine, 2020, 9, 751.	1.0	18
13	Long-term repeated rituximab treatment for childhood steroid-dependent nephrotic syndrome. Kidney Research and Clinical Practice, 2017, 36, 257-263.	0.9	15
14	Tumour lysis syndrome in children: experience of last decade. Hematological Oncology, 2011, 29, 196-201.	0.8	14
15	Posttransplantation lymphoproliferative disorder after pediatric solid organ transplantation: experiences of 20 years in a single center. Korean Journal of Pediatrics, 2017, 60, 86.	1.9	13
16	Post-Transplant Lymphoproliferative Diseases in Pediatric Kidney Allograft Recipients with Epstein-Barr Virus Viremia. Journal of Korean Medical Science, 2019, 34, e203.	1.1	13
17	Three cases of Gordon syndrome with dominant KLHL3 mutations. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 361-364.	0.4	12
18	A familial case of Blau syndrome caused by a novelNOD2genetic mutation. Korean Journal of Pediatrics, 2016, 59, S5.	1.9	11

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19	Dyslipidemia in pediatric CKD patients: results from KNOW-PedCKD (KoreaN cohort study for) Tj ETQq1 1 0.784314 rgBT /Overlock 1	0.9	10
20	Two Korean Infants with Genetically Confirmed Congenital Nephrotic Syndrome of Finnish Type. Journal of Korean Medical Science, 2009, 24, S210.	1.1	9
21	Long-Term Outcomes of Pediatric Renovascular Hypertension. Kidney and Blood Pressure Research, 2017, 42, 617-627.	0.9	7
22	Maternal antibiotic exposure during pregnancy is a risk factor for community-acquired urinary tract infection caused by extended-spectrum beta-lactamase-producing bacteria in infants. Pediatric Nephrology, 2022, 37, 163-170.	0.9	7
23	Reninoma: a rare cause of curable hypertension. Korean Journal of Pediatrics, 2019, 62, 144-147.	1.9	7
24	Impact of Antibiotic Prescribing Patterns on Susceptibilities of Uropathogens in Children below 24 Months Old. Antibiotics, 2020, 9, 915.	1.5	6
25	Urine biomarkers for monitoring acute kidney injury in premature infants. Kidney Research and Clinical Practice, 2020, 39, 284-294.	0.9	6
26	Rasburicase improves the outcome of acute kidney injury from typical hemolytic uremic syndrome. Pediatric Nephrology, 2020, 35, 2183-2189.	0.9	5
27	Case of catastrophic antiphospholipid syndrome presenting as neuroretinitis and vaso-occlusive retinopathy. BMC Ophthalmology, 2020, 20, 482.	0.6	4
28	Clinical Relevance of Fluid Volume Status Assessment by Bioimpedance Spectroscopy in Children Receiving Maintenance Hemodialysis or Peritoneal Dialysis. Journal of Clinical Medicine, 2021, 10, 79.	1.0	4
29	Renal Syndromic Hearing Loss is Common in Childhood-onset Chronic Kidney Disease. Journal of Korean Medical Science, 2020, 35, e364.	1.1	4
30	Higher Incidence of BK Virus Nephropathy in Pediatric Kidney Allograft Recipients with Alport Syndrome. Journal of Clinical Medicine, 2019, 8, 491.	1.0	3
31	Intellectual Functioning of Pediatric Patients with Chronic Kidney Disease: Results from the KNOW-Ped CKD. Journal of Korean Medical Science, 2021, 36, e138.	1.1	3
32	Baseline characteristics of participants enrolled in the KoreaN cohort study for Outcomes in patients With Pediatric Chronic Kidney Disease (KNOW-Ped CKD). Pediatric Nephrology, 2022, 37, 3177-3187.	0.9	2
33	Cystic dysplasia of the kidneys in extremely preterm infants following acute kidney injury. Pediatric Nephrology, 2020, 35, 2369-2372.	0.9	1
34	Risk Factors for the Progression of Chronic Kidney Disease in Children. Childhood Kidney Diseases, 2021, 25, 1-7.	0.1	1
35	Severe Anemia Due to Parvovirus Infection Following Treatment with Rituximab in a Pediatric Kidney Transplant Recipient : Anemia after Treatment of Rituximab in Kidney Recipient Patient. Childhood Kidney Diseases, 2015, 19, 176-179.	0.1	1
36	Peritoneal Protein Loss in Nephrotic Syndrome on Peritoneal Dialysis. Journal of the Korean Society of Pediatric Nephrology, 2009, 13, 189.	0.1	1

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37	Chronic Renal Failure in Russell-Silver Syndrome. Journal of the Korean Society of Pediatric Nephrology, 2009, 13, 256.	0.1	1
38	Concurrent cytomegalovirus enteritis and atypical hemolytic uremic syndrome with gastrointestinal tract involvement: a case report. Korean Journal of Transplantation, 2021, , .	0.0	1
39	Genetic analysis using whole-exome sequencing in pediatric chronic kidney disease: a single center's experience. Childhood Kidney Diseases, 0, , .	0.1	1
40	Pharmacodynamic Monitoring of Calcineurin Inhibitor in Pediatric Kidney Transplantation. The Journal of the Korean Society for Transplantation, 2015, 29, 16.	0.2	0
41	Current usage and effects of steroids in the management of childhood mycoplasma pneumonia in a secondary hospital. Allergy Asthma & Respiratory Disease, 2018, 6, 122.	0.3	0
42	SP013GENOTYPE-PHENOTYPE ANALYSES OF PEDIATRIC PATIENTS WITH PAX2 MUTATIONS. Nephrology Dialysis Transplantation, 2019, 34, .	0.4	0
43	Clinical Outcomes of Non-carbapenem Treatment for Urinary Tract Infections Caused by Extended-spectrum Î²-lactamase-producing Escherichia coli. Childhood Kidney Diseases, 2021, 25, 22-28.	0.1	0
44	Renal replacement therapy is an alarm sign of survival outcome in pediatric liver transplantation. Korean Journal of Transplantation, 2021, 35, S116-S116.	0.0	0
45	Posterior Reversible Encephalopathy Syndrome Accompanied by a Cerebral Hemorrhage and Restricted Diffusion in a Child with Uncontrolled Nephrotic Syndrome. Journal of the Korean Child Neurology Society, 2017, 25, 174-178.	0.0	0
46	Two cases of children presenting with polydipsia, polyuria, and malignant hypertension: Answers. Pediatric Nephrology, 2022, 37, 559-561.	0.9	0
47	Two cases of children presenting with polydipsia, polyuria, and malignant hypertension: Questions. Pediatric Nephrology, 2021, , 1.	0.9	0
48	Gorham-Stout Syndrome with Focal Segmental Glomerulosclerosis: A Case Report. Childhood Kidney Diseases, 2020, 24, 120-125.	0.1	0
49	Response to Chan. Pediatric Nephrology, 2022, , .	0.9	0
50	A patient with multiple arterial stenosis diagnosed with Alagille syndrome: A case report. Journal of Genetic Medicine, 2021, 18, 142-146.	0.1	0
51	Extraskeletal Calcifications in Children with Maintenance Peritoneal Dialysis. Childhood Kidney Diseases, 2021, 25, 117-121.	0.1	0
52	BK polyomavirus associated nephropathy. Childhood Kidney Diseases, 0, , .	0.1	0
53	A case of recurrent hemolytic uremic syndrome caused by <i>DGKE</i> gene mutation. Childhood Kidney Diseases, 0, , .	0.1	0