

Franz Schaefer

List of Publications by Year in descending order

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Version: 2024-02-01

280
papers

21,949
citations

8181

76
h-index

10158

140
g-index

295
all docs

295
docs citations

295
times ranked

17770
citing authors

#	ARTICLE	IF	CITATIONS
1	Treatment and long-term outcome in primary nephrogenic diabetes insipidus. <i>Nephrology Dialysis Transplantation</i> , 2023, 38, 2120-2130.	0.7	9
2	Genetic testing in the diagnosis of chronic kidney disease: recommendations for clinical practice. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 239-254.	0.7	63
3	Impact of COVID-19 pandemic on use of rituximab among children with difficult nephrotic syndrome. <i>Pediatric Research</i> , 2022, 92, 3-5.	2.3	7
4	Acute paediatric kidney replacement therapies in Europe: demographic results from the EurAKid Registry. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 770-780.	0.7	3
5	Persistence of behavioral abnormalities following corticosteroid therapy in children with initial episode of idiopathic nephrotic syndrome: a prospective longitudinal observation. <i>Jornal Brasileiro De Nefrologia: Orgao Oficial De Sociedades Brasileira E Latino-Americana De Nefrologia</i> , 2022, 44, 58-67.	0.9	1
6	Dialysis disequilibrium syndrome (DDS) in pediatric patients on dialysis: systematic review and clinical practice recommendations. <i>Pediatric Nephrology</i> , 2022, 37, 263-274.	1.7	8
7	An update on the use of tolvaptan for autosomal dominant polycystic kidney disease: consensus statement on behalf of the ERA Working Group on Inherited Kidney Disorders, the European Rare Kidney Disease Reference Network and Polycystic Kidney Disease International. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 825-839.	0.7	44
8	Domain-Specific Common Data Elements for Rare Disease Registration: Conceptual Approach of a European Joint Initiative Toward Semantic Interoperability in Rare Disease Research. <i>JMIR Medical Informatics</i> , 2022, 10, e32158.	2.6	8
9	Polycystic Kidney Diseaseâ€™Related Disease Burden in Adolescents With Autosomal Dominant Polycystic Kidney Disease: An International Qualitative Study. <i>Kidney Medicine</i> , 2022, 4, 100415.	2.0	7
10	Definition, diagnosis and management of fetal lower urinary tract obstruction: consensus of the ERKNet CAKUT-Obstructive Uropathy Work Group. <i>Nature Reviews Urology</i> , 2022, 19, 295-303.	3.8	27
11	Meta-GWAS Reveals Novel Genetic Variants Associated with Urinary Excretion of Uromodulin. <i>Journal of the American Society of Nephrology: JASN</i> , 2022, 33, 511-529.	6.1	14
12	Findings from 4C-T Study demonstrate an increased cardiovascular burden in girls with end stage kidney disease and kidney transplantation. <i>Kidney International</i> , 2022, 101, 585-596.	5.2	16
13	Variation of the clinical spectrum and genotype-phenotype associations in Coenzyme Q10 deficiency associated glomerulopathy. <i>Kidney International</i> , 2022, 102, 592-603.	5.2	12
14	Phenotypic Variability in Siblings With Autosomal Recessive Polycystic Kidney Disease. <i>Kidney International Reports</i> , 2022, 7, 1643-1652.	0.8	6
15	Inactivation of Osteoblast PKC Signaling Reduces Cortical Bone Mass and Density and Aggravates Renal Osteodystrophy in Mice with Chronic Kidney Disease on High Phosphate Diet. <i>International Journal of Molecular Sciences</i> , 2022, 23, 6404.	4.1	4
16	Definition, diagnosis and clinical management of non-obstructive kidney dysplasia: a consensus statement by the ERKNet Working Group on Kidney Malformations. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 2351-2362.	0.7	6
17	COVID-19 in children treated with immunosuppressive medication for kidney diseases. <i>Archives of Disease in Childhood</i> , 2021, 106, 798-801.	1.9	46
18	Targeting optimal PD management in children: what have we learned from the IPPN registry?. <i>Pediatric Nephrology</i> , 2021, 36, 1053-1063.	1.7	10

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19	Pathophysiology and consequences of arterial stiffness in children with chronic kidney disease. <i>Pediatric Nephrology</i> , 2021, 36, 1683-1695.	1.7	20
20	Infectious Complications of Peritoneal Dialysis in Children. , 2021, , 265-290.		1
21	Management of congenital nephrotic syndrome: consensus recommendations of the ERKNet-ESPN Working Group. <i>Nature Reviews Nephrology</i> , 2021, 17, 277-289.	9.6	41
22	Hemodiafiltration maintains a sustained improvement in blood pressure compared to conventional hemodialysis in children—the HDF, heart and height (3H) study. <i>Pediatric Nephrology</i> , 2021, 36, 2393-2403.	1.7	9
23	Differential assessment of fluid compartments by bioimpedance in pediatric patients with kidney diseases. <i>Pediatric Nephrology</i> , 2021, 36, 1843-1850.	1.7	7
24	FC 109GLUCOSE DERIVATIVE INDUCED VASCULOPATHY IN CHILDREN ON PERITONEAL DIALYSIS. <i>Nephrology Dialysis Transplantation</i> , 2021, 36, .	0.7	0
25	MO107CLINICAL CHARACTERISTICS OF A PATIENT POPULATION WITH ATYPICAL HAEMOLYTIC URAEMIC SYNDROME AND MALIGNANT HYPERTENSION: THE GLOBAL AHUS REGISTRY ANALYSIS. <i>Nephrology Dialysis Transplantation</i> , 2021, 36, .	0.7	0
26	MO001THE EUROPEAN DRTA REGISTRY: AN INITIAL DATA ANALYSIS*. <i>Nephrology Dialysis Transplantation</i> , 2021, 36, .	0.7	0
27	The European Rare Kidney Disease Registry (ERKReg): objectives, design and initial results. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 251.	2.7	26
28	Low-Dose Antibiotic Prophylaxis Induces Rapid Modifications of the Gut Microbiota in Infants With Vesicoureteral Reflux. <i>Frontiers in Pediatrics</i> , 2021, 9, 674716.	1.9	11
29	CDH12 as a Candidate Gene for Kidney Injury in Posterior Urethral Valve Cases: A Genome-wide Association Study Among Patients with Obstructive Uropathies. <i>European Urology Open Science</i> , 2021, 28, 26-35.	0.4	7
30	An Experimental Workflow for Studying Barrier Integrity, Permeability, and Tight Junction Composition and Localization in a Single Endothelial Cell Monolayer: Proof of Concept. <i>International Journal of Molecular Sciences</i> , 2021, 22, 8178.	4.1	7
31	Systematic review on outcomes used in clinical research on autosomal recessive polycystic kidney disease—are patient-centered outcomes our blind spot?. <i>Pediatric Nephrology</i> , 2021, 36, 3841-3851.	1.7	3
32	Glucose Derivative Induced Vasculopathy in Children on Chronic Peritoneal Dialysis. <i>Circulation Research</i> , 2021, 129, e102-e118.	4.5	17
33	Mortality in Children Treated With Maintenance Peritoneal Dialysis: Findings From the International Pediatric Peritoneal Dialysis Network Registry. <i>American Journal of Kidney Diseases</i> , 2021, 78, 380-390.	1.9	13
34	Refining genotype–phenotype correlations in 304 patients with autosomal recessive polycystic kidney disease and PKHD1 gene variants. <i>Kidney International</i> , 2021, 100, 650-659.	5.2	38
35	Generation of an induced pluripotent stem cell line (DHMCi006-A) from a patient with autosomal recessive polycystic kidney disease (ARPKD) carrying a compound heterozygous missense mutation in the fibrocystin encoding PKHD1 gene. <i>Stem Cell Research</i> , 2021, 57, 102579.	0.7	1
36	Generation of an induced pluripotent stem cell line (DHMCi007-A) from a patient with autosomal recessive polycystic kidney disease (ARPKD) carrying a homozygous missense mutation in the fibrocystin-encoding PKHD1 gene. <i>Stem Cell Research</i> , 2021, 57, 102573.	0.7	0

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37	Early childhood height-adjusted total kidney volume as a risk marker of kidney survival in ARPKD. <i>Scientific Reports</i> , 2021, 11, 21677.	3.3	12
38	Cardiovascular risk factors in children on dialysis: an update. <i>Pediatric Nephrology</i> , 2020, 35, 41-57.	1.7	20
39	Prenatal alcohol exposure affects renal function in overweight schoolchildren: birth cohort analysis. <i>Pediatric Nephrology</i> , 2020, 35, 695-702.	1.7	3
40	Severe neurological outcomes after very early bilateral nephrectomies in patients with autosomal recessive polycystic kidney disease (ARPKD). <i>Scientific Reports</i> , 2020, 10, 16025.	3.3	14
41	Randomized clinical trial to compare efficacy and safety of repeated courses of rituximab to single-course rituximab followed by maintenance mycophenolate-mofetil in children with steroid dependent nephrotic syndrome. <i>BMC Nephrology</i> , 2020, 21, 520.	1.8	5
42	Renal developmental genes are differentially regulated after unilateral ureteral obstruction in neonatal and adult mice. <i>Scientific Reports</i> , 2020, 10, 19302.	3.3	6
43	Patient- and parent proxy-reported outcome measures for life participation in children with chronic kidney disease: a systematic review. <i>Nephrology Dialysis Transplantation</i> , 2020, 35, 1924-1937.	0.7	10
44	MO026 TREATMENT WITH ACTIVE VITAMIN D DOES NOT IMPROVE LEFT VENTRICULAR HYPERTROPHY BUT FURTHER INCREASES FGF23 AND ACCELERATES CKD PROGRESSION IN CHILDREN. <i>Nephrology Dialysis Transplantation</i> , 2020, 35, .	0.7	0
45	Cinacalcet studies in pediatric subjects with secondary hyperparathyroidism receiving dialysis. <i>Pediatric Nephrology</i> , 2020, 35, 1679-1697.	1.7	12
46	IPNA clinical practice recommendations for the diagnosis and management of children with steroid-resistant nephrotic syndrome. <i>Pediatric Nephrology</i> , 2020, 35, 1529-1561.	1.7	179
47	Genetic aspects of congenital nephrotic syndrome: a consensus statement from the ERKNetâ€“ESPN inherited glomerulopathy working group. <i>European Journal of Human Genetics</i> , 2020, 28, 1368-1378.	2.8	28
48	The severity of COVID-19 in children on immunosuppressive medication. <i>The Lancet Child and Adolescent Health</i> , 2020, 4, e17-e18.	5.6	87
49	Implications of early diagnosis of autosomal dominant polycystic kidney disease: A post hoc analysis of the TEMPO 3:4 trial. <i>Scientific Reports</i> , 2020, 10, 4294.	3.3	2
50	Nomenclature for kidney function and disease: report of a Kidney Disease: Improving Global Outcomes (KDIGO) Consensus Conference. <i>Kidney International</i> , 2020, 97, 1117-1129.	5.2	407
51	Maintenance Peritoneal Dialysis in Children With Autosomal Recessive Polycystic Kidney Disease: A Comparative Cohort Study of the International Pediatric Peritoneal Dialysis Network Registry. <i>American Journal of Kidney Diseases</i> , 2020, 75, 460-464.	1.9	8
52	Clinical Interventions and All-Cause Mortality of Patients with Chronic Kidney Disease: An Umbrella Systematic Review of Meta-Analyses. <i>Journal of Clinical Medicine</i> , 2020, 9, 394.	2.4	5
53	Discontinuation of RAAS Inhibition in Children with Advanced CKD. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2020, 15, 625-632.	4.5	19
54	Consensus guidelines for management of hyperammonaemia in paediatric patients receiving continuous kidney replacement therapy. <i>Nature Reviews Nephrology</i> , 2020, 16, 471-482.	9.6	52

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55	Targeting Tubulointerstitium to Predict Kidney Outcomes in Childhood Nephrotic Syndrome. <i>Kidney International Reports</i> , 2020, 5, 383-385.	0.8	0
56	Genes In Your Genes. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2020, 15, 10-12.	4.5	2
57	Fiji plugins for qualitative image annotations: routine analysis and application to image classification. <i>Research</i> , 2020, 9, 1248.	1.6	3
58	Serum indoxyl sulfate concentrations associate with progression of chronic kidney disease in children. <i>PLoS ONE</i> , 2020, 15, e0240446.	2.5	19
59	Hämolytisch-urämisches Syndrom. <i>Springer Reference Medizin</i> , 2020, , 2389-2393.	0.0	0
60	Chronische Niereninsuffizienz. <i>Springer Reference Medizin</i> , 2020, , 2401-2405.	0.0	0
61	Fiji plugins for qualitative image annotations: routine analysis and application to image classification. <i>Research</i> , 2020, 9, 1248.	1.6	4
62	COVID-19 in children treated with immunosuppressive medication for kidney diseases. , 2020, ,		1
63	Genetic associations of hemoglobin in children with chronic kidney disease in the PediGFR Consortium. <i>Pediatric Research</i> , 2019, 85, 324-328.	2.3	1
64	Indoxyl sulfate associates with cardiovascular phenotype in children with chronic kidney disease. <i>Pediatric Nephrology</i> , 2019, 34, 2571-2582.	1.7	27
65	Impaired Systolic and Diastolic Left Ventricular Function in Children with Chronic Kidney Disease - Results from the 4C Study. <i>Scientific Reports</i> , 2019, 9, 11462.	3.3	20
66	Current management of transition of young people affected by rare renal conditions in the ERKNet. <i>European Journal of Human Genetics</i> , 2019, 27, 1783-1790.	2.8	14
67	Arterial tissue transcriptional profiles associate with tissue remodeling and cardiovascular phenotype in children with end-stage kidney disease. <i>Scientific Reports</i> , 2019, 9, 10316.	3.3	12
68	Treatment of Hypertension in Chronic Kidney Disease. <i>Updates in Hypertension and Cardiovascular Protection</i> , 2019, , 239-255.	0.1	1
69	Determinants of Statural Growth in European Children With Chronic Kidney Disease: Findings From the Cardiovascular Comorbidity in Children With Chronic Kidney Disease (4C) Study. <i>Frontiers in Pediatrics</i> , 2019, 7, 278.	1.9	19
70	Hemodialysis vascular access and subsequent transplantation: a report from the ESPN/ERA-EDTA Registry. <i>Pediatric Nephrology</i> , 2019, 34, 713-721.	1.7	10
71	Clinical courses and complications of young adults with Autosomal Recessive Polycystic Kidney Disease (ARPKD). <i>Scientific Reports</i> , 2019, 9, 7919.	3.3	50
72	Tolvaptan use in children and adolescents with autosomal dominant polycystic kidney disease: rationale and design of a two-part, randomized, double-blind, placebo-controlled trial. <i>European Journal of Pediatrics</i> , 2019, 178, 1013-1021.	2.7	29

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73	A Smart Imaging Workflow for Organ-Specific Screening in a Cystic Kidney Zebrafish Disease Model. <i>International Journal of Molecular Sciences</i> , 2019, 20, 1290.	4.1	26
74	Uremic Toxin Concentrations are Related to Residual Kidney Function in the Pediatric Hemodialysis Population. <i>Toxins</i> , 2019, 11, 235.	3.4	20
75	Effects of Hemodiafiltration versus Conventional Hemodialysis in Children with ESKD: The HDF, Heart and Height Study. <i>Journal of the American Society of Nephrology: JASN</i> , 2019, 30, 678-691.	6.1	60
76	NUP Nephropathy: When Defective Pores Cause Leaky Glomeruli. <i>American Journal of Kidney Diseases</i> , 2019, 73, 890-892.	1.9	1
77	Global Variation of Nutritional Status in Children Undergoing Chronic Peritoneal Dialysis: A Longitudinal Study of the International Pediatric Peritoneal Dialysis Network. <i>Scientific Reports</i> , 2019, 9, 4886.	3.3	36
78	Low levels of urinary epidermal growth factor predict chronic kidney disease progression in children. <i>Kidney International</i> , 2019, 96, 214-221.	5.2	43
79	Acute dialysis in children: results of a European survey. <i>Journal of Nephrology</i> , 2019, 32, 445-451.	2.0	26
80	Peritoneal Dialysis Vintage and Glucose Exposure but Not Peritonitis Episodes Drive Peritoneal Membrane Transformation During the First Years of PD. <i>Frontiers in Physiology</i> , 2019, 10, 356.	2.8	27
81	Pediatric intradialytic hypotension: recommendations from the Pediatric Continuous Renal Replacement Therapy (PCRRT) Workgroup. <i>Pediatric Nephrology</i> , 2019, 34, 925-941.	1.7	13
82	Urinary acute kidney injury biomarkers in very low-birth-weight infants on indomethacin for patent ductus arteriosus. <i>Pediatric Research</i> , 2019, 85, 678-686.	2.3	15
83	Methods of Computational Analysis in Kidney Development. <i>Methods in Molecular Biology</i> , 2019, 1926, 235-246.	0.9	0
84	Urinary proteome signature of Renal Cysts and Diabetes syndrome in children. <i>Scientific Reports</i> , 2019, 9, 2225.	3.3	15
85	Isolated nocturnal and isolated daytime hypertension associate with altered cardiovascular morphology and function in children with chronic kidney disease. <i>Journal of Hypertension</i> , 2019, 37, 2247-2255.	0.5	45
86	Eculizumab Use for Kidney Transplantation in Patients With a Diagnosis of Atypical Hemolytic Uremic Syndrome. <i>Kidney International Reports</i> , 2019, 4, 434-446.	0.8	59
87	A randomized, double-blind, placebo-controlled study to assess the efficacy and safety of cinacalcet in pediatric patients with chronic kidney disease and secondary hyperparathyroidism receiving dialysis. <i>Pediatric Nephrology</i> , 2019, 34, 475-486.	1.7	28
88	Chronische Niereninsuffizienz bei Kindern und Jugendlichen. <i>Springer Reference Medizin</i> , 2019, , 1-5.	0.0	0
89	Simultaneous sequencing of 37 genes identified causative mutations in the majority of children with renal tubulopathies. <i>Kidney International</i> , 2018, 93, 961-967.	5.2	77
90	Hypertension in End-Stage Renal Disease: Dialysis. , 2018, , 473-485.		0

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91	RD-Connect, NeurOmics and EURenOmics: collaborative European initiative for rare diseases. <i>European Journal of Human Genetics</i> , 2018, 26, 778-785.	2.8	55
92	Prevalence of Hypertension in Children with Early-Stage ADPKD. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2018, 13, 874-883.	4.5	65
93	pH-mediated upregulation of AQP1 gene expression through the Spi-B transcription factor. <i>BMC Molecular Biology</i> , 2018, 19, 4.	3.0	3
94	Early Effects of Renal Replacement Therapy on Cardiovascular Comorbidity in Children With End-Stage Kidney Disease. <i>Transplantation</i> , 2018, 102, 484-492.	1.0	31
95	Perinatal Diagnosis, Management, and Follow-up of Cystic Renal Diseases. <i>JAMA Pediatrics</i> , 2018, 172, 74.	6.2	64
96	Efficacy and Long-Term Safety of C.E.R.A. Maintenance in Pediatric Hemodialysis Patients with Anemia of CKD. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2018, 13, 81-90.	4.5	16
97	Barriers for implementation of intensified hemodialysis: survey results from the International Pediatric Dialysis Network. <i>Pediatric Nephrology</i> , 2018, 33, 705-712.	1.7	5
98	Outcomes of renal replacement therapy in boys with prune belly syndrome: findings from the ESPN/ERA-EDTA Registry. <i>Pediatric Nephrology</i> , 2018, 33, 117-124.	1.7	18
99	Unmet needs and challenges for follow-up and treatment of autosomal dominant polycystic kidney disease: the paediatric perspective. <i>CKJ: Clinical Kidney Journal</i> , 2018, 11, i14-i26.	2.9	16
100	SuO018AN AUTOMATED HIGH CONTENT SCREENING PLATFORM FOR IDENTIFICATION OF CYSTIC KIDNEY DISEASE-MODIFYING SUBSTANCES IN ZEBRAFISH. <i>Nephrology Dialysis Transplantation</i> , 2018, 33, i623-i623.	0.7	0
101	Exploring the Clinical and Genetic Spectrum of Steroid Resistant Nephrotic Syndrome: The PodoNet Registry. <i>Frontiers in Pediatrics</i> , 2018, 6, 200.	1.9	77
102	Gastrostomy Tube Insertion in Pediatric Patients With Autosomal Recessive Polycystic Kidney Disease (ARPKD): Current Practice. <i>Frontiers in Pediatrics</i> , 2018, 6, 164.	1.9	16
103	Outrageous prices of orphan drugs: a call for collaboration. <i>Lancet</i> , 2018, 392, 791-794.	13.7	132
104	Neutral pH and low glucose degradation product dialysis fluids induce major early alterations of the peritoneal membrane in children on peritoneal dialysis. <i>Kidney International</i> , 2018, 94, 419-429.	5.2	84
105	Risk Factors for Early Dialysis Dependency in Autosomal Recessive Polycystic Kidney Disease. <i>Journal of Pediatrics</i> , 2018, 199, 22-28.e6.	1.8	39
106	Effect of haemodiafiltration vs conventional haemodialysis on growth and cardiovascular outcomes in children – the HDF, heart and height (3H) study. <i>BMC Nephrology</i> , 2018, 19, 199.	1.8	22
107	Validating the use of bioimpedance spectroscopy for assessment of fluid status in children. <i>Pediatric Nephrology</i> , 2018, 33, 1601-1607.	1.7	31
108	Intimal and medial arterial changes defined by ultra-high-frequency ultrasound: Response to changing risk factors in children with chronic kidney disease. <i>PLoS ONE</i> , 2018, 13, e0198547.	2.5	18

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109	Efficacy of Rituximab vs Tacrolimus in Pediatric Corticosteroid-Dependent Nephrotic Syndrome. <i>JAMA Pediatrics</i> , 2018, 172, 757.	6.2	94
110	Clinical and genetic predictors of atypical hemolytic uremic syndrome phenotype and outcome. <i>Kidney International</i> , 2018, 94, 408-418.	5.2	117
111	Hemodiafiltration is associated with reduced inflammation, oxidative stress and improved endothelial risk profile compared to high-flux hemodialysis in children. <i>PLoS ONE</i> , 2018, 13, e0198320.	2.5	42
112	The Human Phenotype Ontology in 2017. <i>Nucleic Acids Research</i> , 2017, 45, D865-D876.	14.5	699
113	Peritoneal Dialysis Access Revision in Children: Causes, Interventions, and Outcomes. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2017, 12, 105-112.	4.5	50
114	The Phenotypic Spectrum of Nephropathies Associated with Mutations in Diacylglycerol Kinase $\hat{\mu}$. <i>Journal of the American Society of Nephrology: JASN</i> , 2017, 28, 3066-3075.	6.1	50
115	Long-Term Outcome of Steroid-Resistant Nephrotic Syndrome in Children. <i>Journal of the American Society of Nephrology: JASN</i> , 2017, 28, 3055-3065.	6.1	142
116	Longer duration of obesity is associated with a reduction in urinary angiotensinogen in prepubertal children. <i>Pediatric Nephrology</i> , 2017, 32, 1411-1422.	1.7	3
117	Mortality risk disparities in children receiving chronic renal replacement therapy for the treatment of end-stage renal disease across Europe: an ESPN-ERA/EDTA registry analysis. <i>Lancet, The</i> , 2017, 389, 2128-2137.	13.7	48
118	Infants Requiring Maintenance Dialysis: Outcomes of Hemodialysis and Peritoneal Dialysis. <i>American Journal of Kidney Diseases</i> , 2017, 69, 617-625.	1.9	53
119	The association of donor and recipient age with graft survival in paediatric renal transplant recipients in a European Society for Paediatric Nephrology/European Renal Association "European Dialysis and Transplantation Association Registry study. <i>Nephrology Dialysis Transplantation</i> , 2017, 32, 1949-1956.	0.7	35
120	Association of Serum Soluble Urokinase Receptor Levels With Progression of Kidney Disease in Children. <i>JAMA Pediatrics</i> , 2017, 171, e172914.	6.2	46
121	Chronic dialysis in children and adolescents: challenges and outcomes. <i>The Lancet Child and Adolescent Health</i> , 2017, 1, 68-77.	5.6	55
122	Metabolic acidosis is common and associates with disease progression in children with chronic kidney disease. <i>Kidney International</i> , 2017, 92, 1507-1514.	5.2	66
123	Cardiovascular Phenotypes in Children with CKD: The 4C Study. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2017, 12, 19-28.	4.5	138
124	Low renal but high extrarenal phenotype variability in Schimke immuno-osseous dysplasia. <i>PLoS ONE</i> , 2017, 12, e0180926.	2.5	25
125	Mutations in sphingosine-1-phosphate lyase cause nephrosis with ichthyosis and adrenal insufficiency. <i>Journal of Clinical Investigation</i> , 2017, 127, 912-928.	8.2	160
126	An inducible mouse model of podocin-mutation-related nephrotic syndrome. <i>PLoS ONE</i> , 2017, 12, e0186574.	2.5	15

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127	Hypertension in End-Stage Renal Disease: Dialysis. , 2017, , 1-13.		0
128	SP701 EFFICACY OF CONTINUOUS PERITONEAL DIALYSIS VERSUS DAILY HAEMODIALYSIS IN MANAGING PEDIATRIC ACUTE KIDNEY INJURY. Nephrology Dialysis Transplantation, 2016, 31, i330-i330.	0.7	0
129	The expanding phenotypic spectra of kidney diseases: insights from genetic studies. Nature Reviews Nephrology, 2016, 12, 472-483.	9.6	61
130	Quantitative Histomorphometry of the Healthy Peritoneum. Scientific Reports, 2016, 6, 21344.	3.3	77
131	2016 European Society of Hypertension guidelines for the management of high blood pressure in children and adolescents. Journal of Hypertension, 2016, 34, 1887-1920.	0.5	898
132	Determinants of carotid-femoral pulse wave velocity in prepubertal children. International Journal of Cardiology, 2016, 218, 37-42.	1.7	31
133	Averting the Legacy of Kidney Disease - Focus on Childhood. Kidney Diseases (Basel, Switzerland), 2016, 2, 46-52.	2.5	5
134	Normalization of glomerular filtration rate in obese children. Pediatric Nephrology, 2016, 31, 1321-1328.	1.7	21
135	Mortality risk in European children with end-stage renal disease on dialysis. Kidney International, 2016, 89, 1355-1362.	5.2	73
136	Efficacy and outcomes of continuous peritoneal dialysis versus daily intermittent hemodialysis in pediatric acute kidney injury. Pediatric Nephrology, 2016, 31, 1681-1689.	1.7	15
137	Association of myeloperoxidase levels with cardiometabolic factors and renal function in prepubertal children. European Journal of Clinical Investigation, 2016, 46, 50-59.	3.4	16
138	Oxidative stress and nitric oxide are increased in obese children and correlate with cardiometabolic risk and renal function. British Journal of Nutrition, 2016, 116, 805-815.	2.3	37
139	International Network of Chronic Kidney Disease cohort studies (iNET-CKD): a global network of chronic kidney disease cohorts. BMC Nephrology, 2016, 17, 121.	1.8	44
140	Timing of renal replacement therapy does not influence survival and growth in children with congenital nephrotic syndrome caused by mutations in NPHS1: data from the ESPN/ERA-EDTA Registry. Pediatric Nephrology, 2016, 31, 2317-2325.	1.7	25
141	Safety and usage of darbepoetin alfa in children with chronic kidney disease: prospective registry study. Pediatric Nephrology, 2016, 31, 443-453.	1.7	19
142	Urinary fibrogenic cytokines ET-1 and TGF- β 1 are associated with urinary angiotensinogen levels in obese children. Pediatric Nephrology, 2016, 31, 455-464.	1.7	4
143	Accelerated growth during childhood is associated with increased arterial stiffness in prepubertal children. International Journal of Cardiology, 2016, 204, 83-85.	1.7	6
144	Genome-wide association studies in pediatric chronic kidney disease. Pediatric Nephrology, 2016, 31, 1241-1252.	1.7	9

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145	Averting the Legacy of Kidney Diseaseâ€™Focus on Childhood. American Journal of Hypertension, 2016, 29, 537-541.	2.0	0
146	Averting the legacy of kidney diseaseâ€™focus on childhood. Kidney International, 2016, 89, 512-518.	5.2	22
147	Left Ventricular Mass Indexing in Infants, Children, and Adolescents: A Simplified Approach for the Identification of Left Ventricular Hypertrophy in Clinical Practice. Journal of Pediatrics, 2016, 170, 193-198.	1.8	70
148	Kidney disease in children: latest advances and remaining challenges. Nature Reviews Nephrology, 2016, 12, 182-191.	9.6	31
149	Racial Disparities in Access to and Outcomes of Kidney Transplantation in Children, Adolescents, and Young Adults: Results From the ESPN/ERA-EDTA (European Society of Pediatric Nephrology/European) Diseases. 2016, 67, 293-301.	1.9	55
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