Jens KÃ¶nig

List of Publications by Year
in descending order

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22	712	759233 12	642732 23
papers	citations	h-index	g-index

28	28	28	1010
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	International consensus statement on the diagnosis and management of autosomal dominant polycystic kidney disease in children and young people. Nature Reviews Nephrology, 2019, 15, 713-726.	9.6	86
2	Phenotypic Spectrum of Children with Nephronophthisis and Related Ciliopathies. Clinical Journal of the American Society of Nephrology: CJASN, 2017, 12, 1974-1983.	4.5	75
3	Treatment and long-term outcome in primary distal renal tubular acidosis. Nephrology Dialysis Transplantation, 2019, 34, 981-991.	0.7	75
4	Germline De Novo Mutations in ATP1A1 Cause Renal Hypomagnesemia, Refractory Seizures, and Intellectual Disability. American Journal of Human Genetics, 2018, 103, 808-816.	6.2	74
5	Perinatal Diagnosis, Management, and Follow-up of Cystic Renal Diseases. JAMA Pediatrics, 2018, 172, 74.	6.2	64
6	Rationale, design and objectives of ARegPKD, a European ARPKD registry study. BMC Nephrology, 2015, 16, 22.	1.8	46
7	HNF1B nephropathy has a slow-progressive phenotype in childhoodâ€"with the exception of very early onset cases: results of the German Multicenter HNF1B Childhood Registry. Pediatric Nephrology, 2019, 34, 1065-1075.	1.7	41
8	Risk Factors for Early Dialysis Dependency in Autosomal Recessive Polycystic Kidney Disease. Journal of Pediatrics, 2018, 199, 22-28.e6.	1.8	39
9	Refining genotypeâ€“phenotype correlations in 304 patients with autosomal recessive polycystic kidney disease and PKHD1 gene variants. Kidney International, 2021, 100, 650-659.	5.2	38
10	Outcome of renal transplantation in small infants: a match-controlled analysis. Pediatric Nephrology, 2018, 33, 1057-1068.	1.7	27
11	mTOR-Activating Mutations in RRAGD Are Causative for Kidney Tubulopathy and Cardiomyopathy. Journal of the American Society of Nephrology: JASN, 2021, 32, 2885-2899.	6.1	24
12	Vaccination titres pre- and post-transplant in paediatric renal transplant recipients and the impact of immunosuppressive therapy. Pediatric Nephrology, 2018, 33, 897-910.	1.7	20
13	Severe neurological outcomes after very early bilateral nephrectomies in patients with autosomal recessive polycystic kidney disease (ARPKD). Scientific Reports, 2020, 10, 16025.	3.3	14
14	Cardiovascular Outcome of Pediatric Patients With Bi-Allelic (Homozygous) Familial Hypercholesterolemia Before and After Initiation of Multimodal Lipid Lowering Therapy Including Lipoprotein Apheresis. American Journal of Cardiology, 2020, 136, 38-48.	1.6	13
15	Multimodal lipid-lowering treatment in pediatric patients with homozygous familial hypercholesterolemiaâ€"target attainment requires further increase of intensity. Pediatric Nephrology, 2018, 33, 1199-1208.	1.7	12
16	Incomplete vaccination coverage in European children with end-stage kidney disease prior to renal transplantation. Pediatric Nephrology, 2018, 33, 341-350.	1.7	12
17	Early childhood height-adjusted total kidney volume as a risk marker of kidney survival in ARPKD. Scientific Reports, 2021, 11, 21677.	3.3	12
18	Systematic evaluation of olfaction in patients with hereditary cystic kidney diseases/renal ciliopathies. Journal of Medical Genetics, 2021, 58, 629-636.	3.2	7

#	Article	IF	Citations
19	Different approaches to long-term treatment of aHUS due to MCP mutations: a multicenter analysis. Pediatric Nephrology, 2021, 36, 463-471.	1.7	6
20	NPHP1 gene-associated nephronophthisis is associated with an occult retinopathy. Kidney International, 2021, 100, 1092-1100.	5.2	6
21	Primary URECs: a source to better understand the pathology of renal tubular epithelia in pediatric hereditary cystic kidney diseases. Orphanet Journal of Rare Diseases, 2022, 17, 122.	2.7	5
22	Development of an Interactive Dashboard for OSSE Rare Disease Registries. Studies in Health Technology and Informatics, 2022, 293, 187-188.	0.3	0