# Jens KÃ¶nig

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7052821/publications.pdf

Version: 2024-02-01

| 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 | 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 |
| 2 | Development of an Interactive Dashboard for OSSE Rare Disease Registries. Studies in Health Technology and Informatics, 2022, 293, 187-188. | 0.3 | 0 |
| 3 | Systematic evaluation of olfaction in patients with hereditary cystic kidney diseases/renal ciliopathies. Journal of Medical Genetics, 2021, 58, 629-636. | 3.2 | 7 |
| 4 | Different approaches to long-term treatment of aHUS due to MCP mutations: a multicenter analysis. Pediatric Nephrology, 2021, 36, 463-471. | 1.7 | 6 |
| 5 | NPHP1 gene-associated nephronophthisis is associated with an occult retinopathy. Kidney International, 2021, 100, 1092-1100. | 5.2 | 6 |
| 6 | 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 |
| 7 | 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 |
| 8 | Early childhood height-adjusted total kidney volume as a risk marker of kidney survival in ARPKD. Scientific Reports, 2021, 11, 21677. | 3.3 | 12 |
| 9 | 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 |
| 10 | 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 |
| 11 | 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 |
| 12 | 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 |
| 13 | Treatment and long-term outcome in primary distal renal tubular acidosis. Nephrology Dialysis Transplantation, 2019, 34, 981-991. | 0.7 | 75 |
| 14 | 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 |
| 15 | 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 |
| 16 | Outcome of renal transplantation in small infants: a match-controlled analysis. Pediatric Nephrology, 2018, 33, 1057-1068. | 1.7 | 27 |
| 17 | Incomplete vaccination coverage in European children with end-stage kidney disease prior to renal transplantation. Pediatric Nephrology, 2018, 33, 341-350. | 1.7 | 12 |
| 18 | Perinatal Diagnosis, Management, and Follow-up of Cystic Renal Diseases. JAMA Pediatrics, 2018, 172, 74. | 6.2 | 64 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | 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 |
| 20 | Risk Factors for Early Dialysis Dependency in Autosomal Recessive Polycystic Kidney Disease. Journal of Pediatrics, 2018, 199, 22-28.e6. | 1.8 | 39 |
| 21 | 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 |
| 22 | Rationale, design and objectives of ARegPKD, a European ARPKD registry study. BMC Nephrology, 2015, 16, 22. | 1.8 | 46 |