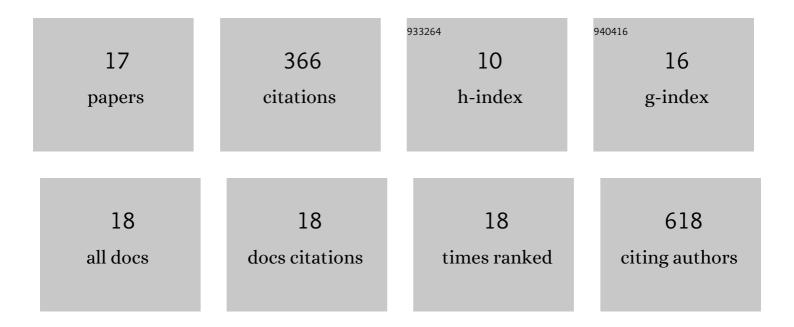
Verena Klämbt

List of Publications by Year in descending order

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VEDENA KIÃMIRT

| # | Article | IF | CITATIONS |
|----|---|------|-----------|
| 1 | A CRISPR-based assay for the detection of opportunistic infections post-transplantation and for the monitoring of transplant rejection. Nature Biomedical Engineering, 2020, 4, 601-609. | 11.6 | 80 |
| 2 | A Novel Function for P2Y2 in Myeloid Recipient–Derived Cells during Graft-versus-Host Disease. Journal of Immunology, 2015, 195, 5795-5804. | 0.4 | 51 |
| 3 | Whole exome sequencing identified ATP6V1C2 as a novel candidate gene for recessive distal renal tubular acidosis. Kidney International, 2020, 97, 567-579. | 2.6 | 42 |
| 4 | Multisystem inflammation and susceptibility to viral infections in human ZNFX1 deficiency. Journal of Allergy and Clinical Immunology, 2021, 148, 381-393. | 1.5 | 40 |
| 5 | Mutations of the Transcriptional Corepressor ZMYM2 Cause Syndromic Urinary Tract Malformations. American Journal of Human Genetics, 2020, 107, 727-742. | 2.6 | 25 |
| 6 | Recessive <i>NOS1AP</i> variants impair actin remodeling and cause glomerulopathy in humans and mice. Science Advances, 2021, 7, . | 4.7 | 21 |
| 7 | CAKUT and Autonomic Dysfunction Caused by Acetylcholine Receptor Mutations. American Journal of Human Genetics, 2019, 105, 1286-1293. | 2.6 | 18 |
| 8 | Phenotype expansion of heterozygous FOXC1 pathogenic variants toward involvement of congenital anomalies of the kidneys and urinary tract (CAKUT). Genetics in Medicine, 2020, 22, 1673-1681. | 1.1 | 16 |
| 9 | Mutations in PRDM15 Are a Novel Cause of Galloway-Mowat Syndrome. Journal of the American Society of Nephrology: JASN, 2021, 32, 580-596. | 3.0 | 15 |
| 10 | De novo TRIM8 variants impair its protein localization to nuclear bodies and cause developmental delay, epilepsy, and focal segmental glomerulosclerosis. American Journal of Human Genetics, 2021, 108, 357-367. | 2.6 | 14 |
| 11 | Reverse phenotyping facilitates disease allele calling in exome sequencing of patients with CAKUT. Genetics in Medicine, 2022, 24, 307-318. | 1.1 | 13 |
| 12 | DAAM2 Variants Cause Nephrotic Syndrome via Actin Dysregulation. American Journal of Human Genetics, 2020, 107, 1113-1128. | 2.6 | 12 |
| 13 | Recessive Mutations in SYNPO2 as a Candidate of Monogenic Nephrotic Syndrome. Kidney International Reports, 2021, 6, 472-483. | 0.4 | 7 |
| 14 | Different approaches to long-term treatment of aHUS due to MCP mutations: a multicenter analysis. Pediatric Nephrology, 2021, 36, 463-471. | 0.9 | 6 |
| 15 | Ribavirin therapy of hepatitis E infection may cause hyporegenerative anemia in pediatric renal transplant patients. Pediatric Transplantation, 2018, 22, e13195. | 0.5 | 4 |
| 16 | Generation of Monogenic Candidate Genes for Human Nephrotic Syndrome Using 3 Independent Approaches. Kidney International Reports, 2021, 6, 460-471. | 0.4 | 2 |
| 17 | Mutations in transcription factor CP2-like 1 may cause a novel syndrome with distal renal tubulopathy in humans. Nephrology Dialysis Transplantation, 2021, 36, 237-246. | 0.4 | 0 |