Verena KlÃ¤mbt

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7745792/publications.pdf

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17	366	933264 10	940416 16
papers	citations	h-index	g-index

18	18	18	618
all docs	docs citations	times ranked	citing authors

#	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 Recipientâ€“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 NOS1AP 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