

Christine Gast

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7276565/publications.pdf>

Version: 2024-02-01

6
papers

277
citations

1936888

4
h-index

2272555

4
g-index

6
all docs

6
docs citations

6
times ranked

499
citing authors

#	ARTICLE	IF	CITATIONS
1	MO048PATHOGENIC VARIANTS IN CHLORIDE VOLTAGE-GATED CHANNEL 5 (CLCN5), ASSOCIATED WITH DENT DISEASE TYPE 1, SHOULD BE CONSIDERED IN END-STAGE KIDNEY DISEASE OF UNKNOWN AETIOLOGY. Nephrology Dialysis Transplantation, 2021, 36, .	0.4	0
2	Genetic and Clinical Predictors of Age of ESKD in Individuals With Autosomal Dominant Tubulointerstitial Kidney Disease Due to UMOD Mutations. Kidney International Reports, 2020, 5, 1472-1485.	0.4	30
3	Autosomal dominant tubulointerstitial kidney disease-UMOD is the most frequent non polycystic genetic kidney disease. BMC Nephrology, 2018, 19, 301.	0.8	39
4	From juvenile hyperuricaemia to dysfunctional uromodulin: an ongoing metamorphosis. Pediatric Nephrology, 2016, 31, 2035-2042.	0.9	9
5	Collagen (<i>COL4A</i>) mutations are the most frequent mutations underlying adult focal segmental glomerulosclerosis. Nephrology Dialysis Transplantation, 2016, 31, 961-970.	0.4	199
6	Hemizygous loss of function mutations in <i>CLCN5</i> causing end-stage kidney disease without Dent disease phenotype. CKJ: Clinical Kidney Journal, 0, , .	1.4	0