

Marijn F Stokman

List of Publications by Year in descending order

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Version: 2024-02-01

13
papers

404
citations

933447

10
h-index

1199594

12
g-index

13
all docs

13
docs citations

13
times ranked

1004
citing authors

#	ARTICLE	IF	CITATIONS
1	Renal Ciliopathies: Sorting Out Therapeutic Approaches for Nephronophthisis. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 653138.	3.7	22
2	P0050PRE-IMPLANTATION GENETIC TESTING FOR MONOGENIC KIDNEY DISEASE: TWENTY-FIVE YEAR EXPERIENCE IN THE NETHERLANDS. <i>Nephrology Dialysis Transplantation</i> , 2020, 35, .	0.7	0
3	Preimplantation Genetic Testing for Monogenic Kidney Disease. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2020, 15, 1279-1286.	4.5	27
4	Changes in the urinary extracellular vesicle proteome are associated with nephronophthisis-related ciliopathies. <i>Journal of Proteomics</i> , 2019, 192, 27-36.	2.4	22
5	Loss-of-function mutations in <i>KIF14</i> cause severe microcephaly and kidney development defects in humans and zebrafish. <i>Human Molecular Genetics</i> , 2019, 28, 778-795.	2.9	33
6	Clinical and genetic analyses of a Dutch cohort of 40 patients with a nephronophthisis-related ciliopathy. <i>Pediatric Nephrology</i> , 2018, 33, 1701-1712.	1.7	20
7	The expanding phenotypic spectra of kidney diseases: insights from genetic studies. <i>Nature Reviews Nephrology</i> , 2016, 12, 472-483.	9.6	61
8	Congenital Amegakaryocytic Thrombocytopenia Type II Presenting with Multiple Central Nervous System Anomalies. <i>Neuropediatrics</i> , 2016, 47, 128-131.	0.6	8
9	Compound heterozygous <i>NEK1</i> variants in two siblings with oral-facial-digital syndrome type II (Mohr) Tj ETQq1 1 0,784314 rgBT /Over	2.8	13
10	De novo 14q24.2q24.3 microdeletion including <i>IFT43</i> is associated with intellectual disability, skeletal anomalies, cardiac anomalies, and myopia. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 1566-1569.	1.2	3
11	Non-invasive sources of cells with primary cilia from pediatric and adult patients. <i>Cilia</i> , 2015, 4, 8.	1.8	34
12	Nephronophthisis-Associated CEP164 Regulates Cell Cycle Progression, Apoptosis and Epithelial-to-Mesenchymal Transition. <i>PLoS Genetics</i> , 2014, 10, e1004594.	3.5	73
13	Next-generation sequencing for research and diagnostics in kidney disease. <i>Nature Reviews Nephrology</i> , 2014, 10, 433-444.	9.6	88