Marijn F Stokman

List of Publications by Year
in descending order

13	404	933447 10	1199594 12
papers	citations	h-index	g-index

13	13	13	1004
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Next-generation sequencing for research and diagnostics in kidney disease. Nature Reviews Nephrology, 2014, 10, 433-444.	9.6	88
2	Nephronophthisis-Associated CEP164 Regulates Cell Cycle Progression, Apoptosis and Epithelial-to-Mesenchymal Transition. PLoS Genetics, 2014, 10, e1004594.	3.5	73
3	The expanding phenotypic spectra of kidney diseases: insights from genetic studies. Nature Reviews Nephrology, 2016, 12, 472-483.	9.6	61
4	Non-invasive sources of cells with primary cilia from pediatric and adult patients. Cilia, 2015, 4, 8.	1.8	34
5	Loss-of-function mutations inKIF14cause severe microcephaly and kidney development defects in humans and zebrafish. Human Molecular Genetics, 2019, 28, 778-795.	2.9	33
6	Preimplantation Genetic Testing for Monogenic Kidney Disease. Clinical Journal of the American Society of Nephrology: CJASN, 2020, 15, 1279-1286.	4.5	27
7	Changes in the urinary extracellular vesicle proteome are associated with nephronophthisis-related ciliopathies. Journal of Proteomics, 2019, 192, 27-36.	2.4	22
8	Renal Ciliopathies: Sorting Out Therapeutic Approaches for Nephronophthisis. Frontiers in Cell and Developmental Biology, 2021, 9, 653138.	3.7	22
9	Clinical and genetic analyses of a Dutch cohort of 40 patients with a nephronophthisis-related ciliopathy. Pediatric Nephrology, 2018, 33, 1701-1712.	1.7	20
10	Compound heterozygous NEK1 variants in two siblings with oral-facial-digital syndrome type II (Mohr) Tj ETQq0 0 0 rgBT /Overlock 10 Tf	2.8	13
11	Congenital Amegakaryocytic Thrombocytopenia Type II Presenting with Multiple Central Nervous System Anomalies. Neuropediatrics, 2016, 47, 128-131.	0.6	8
12	De novo 14q24.2q24.3 microdeletion including IFT43 is associated with intellectual disability, skeletal anomalies, cardiac anomalies, and myopia. American Journal of Medical Genetics, Part A, 2016, 170, 1566-1569.	1.2	3
13	P0050PRE-IMPLANTATION GENETIC TESTING FOR MONOGENIC KIDNEY DISEASE: TWENTY-FIVE YEAR EXPERIENCE IN THE NETHERLANDS. Nephrology Dialysis Transplantation, 2020, 35, .	0.7	0