Rik Westland

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9555944/publications.pdf

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23 papers	1,559 citations	430754 18 h-index	610775 24 g-index
26 all docs	26 docs citations	26 times ranked	1966 citing authors

#	Article	IF	Citations
1	Unilateral renal agenesis: a systematic review on associated anomalies and renal injury. Nephrology Dialysis Transplantation, 2013, 28, 1844-1855.	0.4	206
2	Unilateral multicystic dysplastic kidney: a meta-analysis of observational studies on the incidence, associated urinary tract malformations and the contralateral kidney. Nephrology Dialysis Transplantation, 2009, 24, 1810-1818.	0.4	165
3	Whole-Exome Sequencing in Adults With Chronic Kidney Disease. Annals of Internal Medicine, 2018, 168, 100.	2.0	154
4	The copy number variation landscape of congenital anomalies of the kidney and urinary tract. Nature Genetics, 2019, 51, 117-127.	9.4	144
5	Genetic Drivers of Kidney Defects in the DiGeorge Syndrome. New England Journal of Medicine, 2017, 376, 742-754.	13.9	120
6	Risk Factors for Renal Injury in Children With a Solitary Functioning Kidney. Pediatrics, 2013, 131, e478-e485.	1.0	113
7	Genetic basis of human congenital anomalies of the kidney and urinary tract. Journal of Clinical Investigation, 2018, 128, 4-15.	3.9	91
8	Clinical Implications of the Solitary Functioning Kidney. Clinical Journal of the American Society of Nephrology: CJASN, 2014, 9, 978-986.	2.2	81
9	Copy number variation analysis identifies novel CAKUT candidate genes in children with a solitary functioning kidney. Kidney International, 2015, 88, 1402-1410.	2.6	65
10	Exome-wide Association Study Identifies GREB1L Mutations in Congenital Kidney Malformations. American Journal of Human Genetics, 2017, 101, 789-802.	2.6	63
11	The Burden of Candidate Pathogenic Variants for Kidney and Genitourinary Disorders Emerging From Exome Sequencing. Annals of Internal Medicine, 2019, 170, 11.	2.0	60
12	Phenotypic Expansion of DGKE-Associated Diseases. Journal of the American Society of Nephrology: JASN, 2014, 25, 1408-1414.	3.0	59
13	Precision of Estimating Equations for GFR in Children with a Solitary Functioning Kidney. Clinical Journal of the American Society of Nephrology: CJASN, 2013, 8, 764-772.	2.2	38
14	Clinical Integration of Genome Diagnostics for Congenital Anomalies of the Kidney and Urinary Tract. Clinical Journal of the American Society of Nephrology: CJASN, 2021, 16, 128-137.	2.2	37
15	Ambulatory blood pressure monitoring is recommended in the clinical management of children with a solitary functioning kidney. Pediatric Nephrology, 2014, 29, 2205-2211.	0.9	28
16	Preimplantation Genetic Testing for Monogenic Kidney Disease. Clinical Journal of the American Society of Nephrology: CJASN, 2020, 15, 1279-1286.	2.2	27
17	Long-term follow-up of blood pressure and glomerular filtration rate in patients with a solitary functioning kidney: a comparison between Wilms tumor survivors and nephrectomy for other reasons. Pediatric Nephrology, 2016, 31, 435-441.	0.9	26
18	Clinical Management of Children with a Congenital Solitary Functioning Kidney: Overview and Recommendations. European Urology Open Science, 2021, 25, 11-20.	0.2	25

#	Article	IF	Citations
19	Copy Number Variant Analysis and Genome-wide Association Study Identify Loci with Large Effect for Vesicoureteral Reflux. Journal of the American Society of Nephrology: JASN, 2021, 32, 805-820.	3.0	17
20	Multidisciplinary approaches for elucidating genetics and molecular pathogenesis of urinary tract malformations. Kidney International, 2022, 101, 473-484.	2.6	16
21	Gender differences in solitary functioning kidney: do they affect renal outcome?. Pediatric Nephrology, 2014, 29, 2243-2244.	0.9	8
22	Bartter syndrome Type III and congenital anomalies of the kidney and urinary tract: an antenatal presentation. Clinical Nephrology, 2012, 78, 492-496.	0.4	7
23	Recessive mutations in CAKUT and VACTERL association. Kidney International, 2014, 85, 1253-1255.	2.6	4