## Max Werth

## List of Publications by Year in descending order

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1163117 1474206 1,556 9 8 9 citations h-index g-index papers 12 12 12 3607 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	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.	0.7	O
2	The copy number variation landscape of congenital anomalies of the kidney and urinary tract. Nature Genetics, 2019, 51, 117-127.	21.4	144
3	Single-cell transcriptomics of the mouse kidney reveals potential cellular targets of kidney disease. Science, 2018, 360, 758-763.	12.6	797
4	Genetic Drivers of Kidney Defects in the DiGeorge Syndrome. New England Journal of Medicine, 2017, 376, 742-754.	27.0	120
5	Transcription factor TFCP2L1 patterns cells in the mouse kidney collecting ducts. ELife, 2017, 6, .	6.0	58
6	A <i>Grhl2</i> -dependent gene network controls trophoblast branching morphogenesis. Development (Cambridge), 2015, 142, 1125-1136.	2.5	61
7	A Grainyhead-Like 2/Ovo-Like 2 Pathway Regulates Renal Epithelial Barrier Function and Lumen Expansion. Journal of the American Society of Nephrology: JASN, 2015, 26, 2704-2715.	6.1	69
8	α–Intercalated cells defend the urinary system from bacterial infection. Journal of Clinical Investigation, 2014, 124, 2963-2976.	8.2	127
9	The transcription factor grainyhead-like 2 regulates the molecular composition of the epithelial apical junctional complex. Development (Cambridge), 2010, 137, 3835-3845.	2.5	169