Patricia Outeda

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Monoallelic IFT140 pathogenic variants are an important cause of the autosomal dominant polycystic kidney-spectrum phenotype. American Journal of Human Genetics, 2022, 109, 136-156.	6.2	62
2	TWEAK Signaling Pathway Blockade Slows Cyst Growth and Disease Progression in Autosomal Dominant Polycystic Kidney Disease. Journal of the American Society of Nephrology: JASN, 2021, 32, 1913-1932.	6.1	18
3	Polycystinâ€1 dependent regulation of polycystinâ€2 via GRP94, a member of HSP90 family that resides in the endoplasmic reticulum. FASEB Journal, 2021, 35, e21865.	0.5	4
4	GDNF drives rapid tubule morphogenesis in novel 3D in vitro model for ADPKD. Journal of Cell Science, 2020, 133, .	2.0	7
5	A genetic screen in Drosophila reveals an unexpected role for the KIP1 ubiquitination-promoting complex in male fertility. PLoS Genetics, 2020, 16, e1009217.	3.5	4
6	A cleavage product of Polycystin-1 is a mitochondrial matrix protein that affects mitochondria morphology and function when heterologously expressed. Scientific Reports, 2018, 8, 2743.	3.3	75
7	NEDD4-family E3 ligase dysfunction due to PKHD1/Pkhd1 defects suggests a mechanistic model for ARPKD pathobiology. Scientific Reports, 2017, 7, 7733.	3.3	22
8	A novel model of autosomal recessive polycystic kidney questions the role of the fibrocystin C-terminus in disease mechanism. Kidney International, 2017, 92, 1130-1144.	5.2	43
9	The polycystin complex mediates Wnt/Ca2+ signalling. Nature Cell Biology, 2016, 18, 752-764.	10.3	132
10	Inhibition of histone deacetylase 6 activity reduces cyst growth in polycystic kidney disease. Kidney International, 2016, 90, 90-99.	5.2	58
11	Ciliary membrane proteins traffic through the Golgi via a Rabep1/GGA1/Arl3-dependent mechanism. Nature Communications, 2014, 5, 5482.	12.8	101
12	Polycystin Signaling Is Required for Directed Endothelial Cell Migration and Lymphatic Development. Cell Reports, 2014, 7, 634-644.	6.4	71