## Jenny Aj Van Der Wijst

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Impaired basolateral sorting of pro-EGF causes isolated recessive renal hypomagnesemia. Journal of Clinical Investigation, 2007, 117, 2260-2267.	8.2	307
2	A missense mutation in the Kv1.1 voltage-gated potassium channel–encoding gene KCNA1 is linked to human autosomal dominant hypomagnesemia. Journal of Clinical Investigation, 2009, 119, 936-942.	8.2	138
3	Targeted Next-Generation Sequencing of a 12.5 Mb Homozygous Region Reveals ANO10 Mutations in Patients with Autosomal-Recessive Cerebellar Ataxia. American Journal of Human Genetics, 2010, 87, 813-819.	6.2	125
4	Structural insight into TRPV5 channel function and modulation. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 8869-8878.	7.1	78
5	Methionine Sulfoxide Reductase B1 (MsrB1) Recovers TRPM6 Channel Activity during Oxidative Stress. Journal of Biological Chemistry, 2010, 285, 26081-26087.	3.4	71
6	New TRPM6 missense mutations linked to hypomagnesemia with secondary hypocalcemia. European Journal of Human Genetics, 2014, 22, 497-504.	2.8	70
7	RACK1 Inhibits TRPM6 Activity via Phosphorylation of the Fused α-Kinase Domain. Current Biology, 2008, 18, 168-176.	3.9	67
8	TRP channels in calcium homeostasis: from hormonal control to structure-function relationship of TRPV5 and TRPV6. Biochimica Et Biophysica Acta - Molecular Cell Research, 2017, 1864, 883-893.	4.1	65
9	Regulation of the Epithelial Mg2+ Channel TRPM6 by Estrogen and the Associated Repressor Protein of Estrogen Receptor Activity (REA). Journal of Biological Chemistry, 2009, 284, 14788-14795.	3.4	63
10	Learning Physiology from Inherited Kidney Disorders. Physiological Reviews, 2019, 99, 1575-1653.	28.8	60
11	Functional Analysis of the Kv1.1 N255D Mutation Associated with Autosomal Dominant Hypomagnesemia. Journal of Biological Chemistry, 2010, 285, 171-178.	3.4	50
12	Renal Handling of Circulating and Renal-Synthesized Hepcidin and Its Protective Effects against Hemoglobin–Mediated Kidney Injury. Journal of the American Society of Nephrology: JASN, 2016, 27, 2720-2732.	6.1	50
13	Effects of the EGFR Inhibitor Erlotinib on Magnesium Handling. Journal of the American Society of Nephrology: JASN, 2010, 21, 1309-1316.	6.1	47
14	Regulation of Two Renal Chloride Transporters, AE1 and Pendrin, by Electrolytes and Aldosterone. PLoS ONE, 2013, 8, e55286.	2.5	36
15	Mg2+ homeostasis. Current Opinion in Nephrology and Hypertension, 2014, 23, 361-369.	2.0	35
16	Low gut microbiota diversity and dietary magnesium intake are associated with the development of PPlâ€induced hypomagnesemia. FASEB Journal, 2019, 33, 11235-11246.	0.5	32
17	Epithelial Mg2+ channel TRPM6: insight into theÂmolecular regulation. Magnesium Research, 2009, 22, 127-132.	0.5	31
18	Effects of a high-sodium/low-potassium diet on renal calcium, magnesium, and phosphate handling. American Journal of Physiology - Renal Physiology, 2018, 315, F110-F122.	2.7	27

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19	A Gate Hinge Controls the Epithelial Calcium Channel TRPV5. Scientific Reports, 2017, 7, 45489.	3.3	23
20	A de novo <b><i>KCNA1</i></b> Mutation in a Patient with Tetany and Hypomagnesemia. Nephron, 2018, 139, 359-366.	1.8	22
21	Highâ€resolution structures of <scp>transient receptor potential vanilloid</scp> channels: Unveiling a functionally diverse group of ion channels. Protein Science, 2020, 29, 1569-1580.	7.6	20
22	TRPV5 in renal tubular calcium handling and its potential relevance for nephrolithiasis. Kidney International, 2019, 96, 1283-1291.	5.2	17
23	Kinase and channel activity of TRPM6 are co-ordinated by a dimerization motif and pocket interaction. Biochemical Journal, 2014, 460, 165-175.	3.7	15
24	Epithelial Mg2+ channel TRPM6: insight into the molecular regulation. Magnesium Research, 2009, 22, 127-32.	0.5	14
25	TRPC5 inhibition to treat progressive kidney disease. Nature Reviews Nephrology, 2018, 14, 145-146.	9.6	9
26	Renal sodium and magnesium reabsorption are not coupled in a mouse model of Gordon syndrome. Physiological Reports, 2018, 6, e13728.	1.7	8
27	Interspecies differences in PTH-mediated PKA phosphorylation of the epithelial calcium channel TRPV5. Pflugers Archiv European Journal of Physiology, 2017, 469, 1301-1311.	2.8	7
28	Urinary β-galactosidase stimulates Ca <sup>2+</sup> transport by stabilizing TRPV5 at the plasma membrane. Glycobiology, 2016, 26, 472-481.	2.5	6
29	Possible role for rare <i>TRPM7</i> variants in patients with hypomagnesaemia with secondary hypocalcaemia. Nephrology Dialysis Transplantation, 2023, 38, 679-690.	0.7	6
30	Dominant functional role of the novel phosphorylation site S811 in the human renal NaCl cotransporter. FASEB Journal, 2018, 32, 4482-4493.	0.5	5
31	Role of the alternative splice variant of NCC in blood pressure control. Channels, 2018, 12, 346-355.	2.8	4
32	Colonic expression of calcium transporter TRPV6 is regulated by dietary sodium butyrate. Pflugers Archiv European Journal of Physiology, 2022, 474, 293-302.	2.8	3
33	Modeling Distal Convoluted Tubule (Patho)Physiology: An Overview of Past Developments and an Outlook Toward the Future. Tissue Engineering - Part C: Methods, 2021, 27, 200-212.	2.1	2
34	Effects of flow and polycystinâ€1 dysfunction on ATP release in the inner medullary collecting duct of the kidney. FASEB Journal, 2022, 36, .	0.5	0
35	Title: Jealous protons sour another happy marriage; the story of how TRPV5 and PI(4,5)P2 split up Cell Calcium, 2022, , 102609.	2.4	0