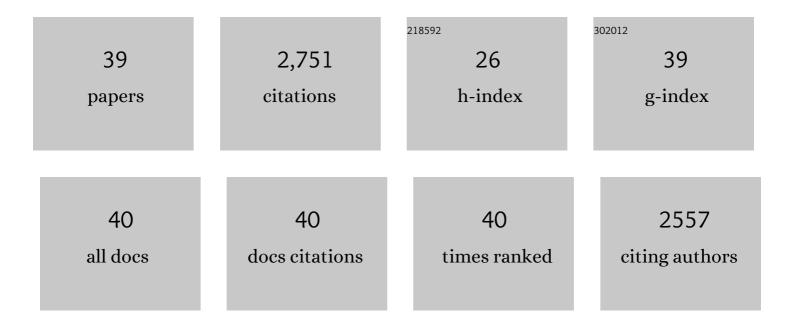
Kathleen J Sweadner

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Misfolding, altered membrane distributions, and the unfolded protein response contribute to pathogenicity differences in Na,K-ATPase ATP1A3 mutations. Journal of Biological Chemistry, 2021, 296, 100019.	1.6	15
2	Rapid-Onset Dystonia-Parkinsonism Phenotype Consistency for a Novel Variant of ATP1A3 in Patients Across 3 Global Populations. Neurology: Genetics, 2021, 7, e562.	0.9	2
3	Revising rapidâ€onset dystonia–parkinsonism: Broadening indications for <i>ATP1A3</i> testing. Movement Disorders, 2019, 34, 1528-1536.	2.2	34
4	Factors in the disease severity of ATP1A3 mutations: Impairment, misfolding, and allele competition. Neurobiology of Disease, 2019, 132, 104577.	2.1	31
5	Genotype-structure-phenotype relationships diverge in paralogs <i>ATP1A1</i> , <i>ATP1A2</i> , and <i>ATP1A3</i> . Neurology: Genetics, 2019, 5, e303.	0.9	33
6	De novo ATP1A3 and compound heterozygous NLRP3 mutations in a child with autism spectrum disorder, episodic fatigue and somnolence, and muckle-wells syndrome. Molecular Genetics and Metabolism Reports, 2018, 16, 23-29.	0.4	12
7	An ion-transport enzyme that rocks. Nature, 2017, 545, 162-164.	13.7	5
8	Impaired AQP2 trafficking in Fxyd1 knockout mice: A role for FXYD1 in regulated vesicular transport. PLoS ONE, 2017, 12, e0188006.	1.1	11
9	ATP1A3 Mutation in Adult Rapid-Onset Ataxia. PLoS ONE, 2016, 11, e0151429.	1.1	34
10	Colorimetric Assays of Na,K-ATPase. Methods in Molecular Biology, 2016, 1377, 89-104.	0.4	4
11	Functional Studies of Na+,K+-ATPase Using Transfected Cell Cultures. Methods in Molecular Biology, 2016, 1377, 321-332.	0.4	2
12	A dystonia-like movement disorder with brain and spinal neuronal defects is caused by mutation of the mouse laminin β1 subunit, Lamb1. ELife, 2015, 4, .	2.8	21
13	Novel mutations in <i>ATP1A3</i> associated with catastrophic early life epilepsy, episodic prolonged apnea, and postnatal microcephaly. Epilepsia, 2015, 56, 422-430.	2.6	107
14	Rapid-onset dystonia-parkinsonism associated with the I758S mutation of the ATP1A3 gene: a neuropathologic and neuroanatomical study of four siblings. Acta Neuropathologica, 2014, 128, 81-98.	3.9	37
15	Distinct neurological disorders with ATP1A3 mutations. Lancet Neurology, The, 2014, 13, 503-514.	4.9	206
16	Paradoxical activation of the sodium chloride cotransporter (NCC) without hypertension in kidney deficient in a regulatory subunit of Na,K-ATPase, FXYD2. Physiological Reports, 2014, 2, e12226.	0.7	8
17	Oxidative inhibition of the vascular Na+-K+ pump via NADPH oxidase-dependent β1-subunit glutathionylation: Implications for angiotensin Il-induced vascular dysfunction. Free Radical Biology and Medicine, 2013, 65, 563-572.	1.3	31
18	Hyperplasia of Pancreatic Beta Cells and Improved Glucose Tolerance in Mice Deficient in the FXYD2 Subunit of Na.K-ATPase. Journal of Biological Chemistry, 2013, 288, 7077-7085.	1.6	28

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19	FXYD Proteins Reverse Inhibition of the Na+-K+ Pump Mediated by Glutathionylation of Its β1 Subunit. Journal of Biological Chemistry, 2011, 286, 18562-18572.	1.6	79
20	Hypertrophy, increased ejection fraction, and reduced Na-K-ATPase activity in phospholemman-deficient mice. American Journal of Physiology - Heart and Circulatory Physiology, 2005, 288, H1982-H1988.	1.5	66
21	Na,K-ATPase from Mice Lacking the Î ³ Subunit (FXYD2) Exhibits Altered Na+ Affinity and Decreased Thermal Stability. Journal of Biological Chemistry, 2005, 280, 19003-19011.	1.6	77
22	Mutations in the Na+/K+-ATPase α3 Gene ATP1A3 Are Associated with Rapid-Onset Dystonia Parkinsonism. Neuron, 2004, 43, 169-175.	3.8	466
23	FXYD Proteins as Regulators of the Na,Kâ€ATPase in the Kidney. Annals of the New York Academy of Sciences, 2003, 986, 382-387.	1.8	45
24	Phospholemman, a Single-Span Membrane Protein, Is an Accessory Protein of Na,K-ATPase in Cerebellum and Choroid Plexus. Journal of Neuroscience, 2003, 23, 2161-2169.	1.7	96
25	Phospholemman expression in extraglomerular mesangium and afferent arteriole of the juxtaglomerular apparatus. American Journal of Physiology - Renal Physiology, 2003, 285, F121-F129.	1.3	28
26	Distribution and oligomeric association of splice forms of Na ⁺ -K ⁺ -ATPase regulatory γ-subunit in rat kidney. American Journal of Physiology - Renal Physiology, 2002, 282, F393-F407.	1.3	73
27	Predicted location and limited accessibility of protein kinase A phosphorylation site on Na-K-ATPase. American Journal of Physiology - Cell Physiology, 2001, 280, C1017-C1026.	2.1	41
28	Structural similarities of Na,K-ATPase and SERCA, the Ca2+-ATPase of the sarcoplasmic reticulum. Biochemical Journal, 2001, 356, 685.	1.7	116
29	Structural similarities of Na,K-ATPase and SERCA, the Ca2+-ATPase of the sarcoplasmic reticulum. Biochemical Journal, 2001, 356, 685-704.	1.7	180
30	Oligodendrocytes in brain and optic nerve express the ?3 subunit isoform of Na,K-ATPase. Glia, 2000, 31, 206-218.	2.5	30
31	Carbachol inhibits Na ⁺ -K ⁺ -ATPase activity in choroid plexus via stimulation of the NO/cGMP pathway. American Journal of Physiology - Cell Physiology, 2000, 279, C1685-C1693.	2.1	72
32	Genomic Organization of the Human FXYD2 Gene Encoding the Î ³ Subunit of the Na,K-ATPase. Biochemical and Biophysical Research Communications, 2000, 279, 196-201.	1.0	44
33	The FXYD Gene Family of Small Ion Transport Regulators or Channels: cDNA Sequence, Protein Signature Sequence, and Expression. Genomics, 2000, 68, 41-56.	1.3	382
34	Cellular and Subcellular Specification of Na,K-ATPase α and β Isoforms in the Postnatal Development of Mouse Retina. Journal of Neuroscience, 1999, 19, 9878-9889.	1.7	101
35	Rat skeletal muscle in culture expresses the ?1 but not the ?2 protein subunit isoform of the Na+/K+ pump. Journal of Cellular Physiology, 1999, 180, 236-244.	2.0	25
36	Phosphorylation of Na,K-ATPase by Protein Kinases. Annals of the New York Academy of Sciences, 1997, 834, 479-488.	1.8	29

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37	Epitope and mimotope for an antibody to the Na, Kâ€ATPase. Protein Science, 1997, 6, 1537-1548.	3.1	18
38	Constraints on Models for the Folding of the Na,K-ATPase. Annals of the New York Academy of Sciences, 1992, 671, 217-227.	1.8	20
39	Multiple tubulin forms are expressed by a single neurone. Nature, 1981, 294, 477-480.	13.7	142