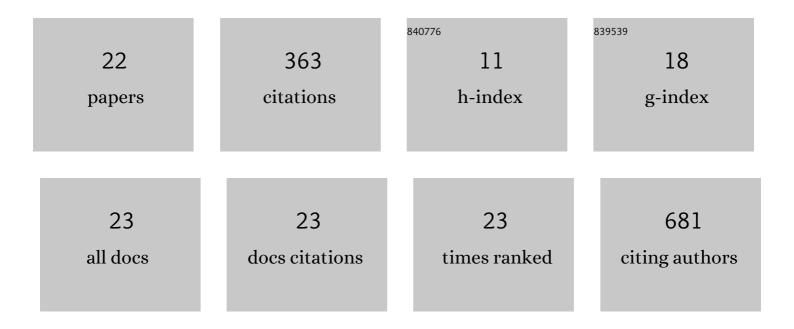
## Anne F Buckley

List of Publications by Year in descending order

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ANNE F RUCKLEY

#	Article	IF	CITATIONS
1	Somatic variants in diverse genes leads to a spectrum of focal cortical malformations. Brain, 2022, 145, 2704-2720.	7.6	33
2	Case Report: Unusual Aggregation of Different Glomerulopathies in a Family Resolved by Genetic Testing and Reverse Phenotyping. Frontiers in Pediatrics, 2022, 10, 826330.	1.9	1
3	Resolution of radiation necrosis with bevacizumab following radiation therapy for primary CNS lymphoma. Oncotarget, 2022, 13, 576-582.	1.8	4
4	Blockade of the natriuretic peptide clearance receptor attenuates proteinuria in a mouse model of focal segmental glomerulosclerosis. Physiological Reports, 2021, 9, e15095.	1.7	1
5	lrgm1-deficiency leads to myeloid dysfunction in colon lamina propria and susceptibility to the intestinal pathogen Citrobacter rodentium. PLoS Pathogens, 2020, 16, e1008553.	4.7	14
6	Skeletal Muscle Is an Antigen Reservoir in Integrase-Defective Lentiviral Vector-Induced Long-Term Immunity. Molecular Therapy - Methods and Clinical Development, 2020, 17, 532-544.	4.1	18
7	Knockout of TRPC6 promotes insulin resistance and exacerbates glomerular injury in Akita mice. Kidney International, 2019, 95, 321-332.	5.2	41
8	PATH-60. PRIMARY CNS LYMPHOMA PRECEDED BY ACUTE DEMYELINATION. Neuro-Oncology, 2019, 21, vi156-vi157.	1.2	0
9	Sudden Unexpected Death in a Child From an Anaplastic Ependymoma. American Journal of Forensic Medicine and Pathology, 2019, 40, 275-278.	0.8	3
10	Regulation of cofilin phosphorylation in glomerular podocytes by testis specific kinase 1 (TESK1). Scientific Reports, 2018, 8, 12286.	3.3	9
11	A Diffuse Leptomeningeal Glioneuronal Tumor Without Diffuse Leptomeningeal Involvement: Detailed Molecular and Clinical Characterization. Journal of Neuropathology and Experimental Neurology, 2018, 77, 751-756.	1.7	25
12	Podocyte-specific knockout of cyclooxygenase 2 exacerbates diabetic kidney disease. American Journal of Physiology - Renal Physiology, 2017, 313, F430-F439.	2.7	10
13	Mitochondrial disorders: The need for precision ultrastructural analysis in clinical diagnosis. Ultrastructural Pathology, 2017, 41, 76-76.	0.9	0
14	The importance of managing the patient and not the gene: expanded phenotype of <i>GLE1</i> -associated arthrogryposis. Journal of Physical Education and Sports Management, 2017, 3, a002063.	1.2	4
15	<i>PRKAG2</i> mutations presenting in infancy. Journal of Inherited Metabolic Disease, 2017, 40, 823-830.	3.6	6
16	Severe Cardiomyopathy as the Isolated Presenting Feature in an Adult with Late-Onset Pompe Disease: A Case Report. JIMD Reports, 2016, 31, 79-83.	1.5	20
17	The fornix provides multiple biomarkers to characterize circuit disruption in a mouse model of Alzheimer's disease. Neurolmage, 2016, 142, 498-511.	4.2	30
18	C-C Motif Chemokine 5 Attenuates Angiotensin Il–Dependent Kidney Injury by Limiting Renal Macrophage Infiltration. American Journal of Pathology, 2016, 186, 2846-2856.	3.8	41

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19	Rhabdomyomatous mesenchymal hamartoma presenting as a sacral skin tag in two neonates with spinal dysraphism. Journal of Cutaneous Pathology, 2015, 42, 774-778.	1.3	15
20	Radioprotection of the Brain White Matter by Mn(III) <i>N</i> -Butoxyethylpyridylporphyrin–Based Superoxide Dismutase Mimic MnTnBuOE-2-PyP5+. Molecular Cancer Therapeutics, 2015, 14, 70-79.	4.1	60
21	Augmenting podocyte injury promotes advanced diabetic kidney disease in Akita mice. Biochemical and Biophysical Research Communications, 2014, 444, 622-627.	2.1	11
22	Skeletal Muscle Microvasculature in the Diagnosis of Neuromuscular Disease. Journal of Neuropathology and Experimental Neurology, 2013, 72, 906-918.	1.7	17