

# Joel M Henderson

## List of Publications by Year in descending order

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31  
papers

2,511  
citations

489802

18  
h-index

536525

29  
g-index

32  
all docs

32  
docs citations

32  
times ranked

4502  
citing authors

#	ARTICLE	IF	CITATIONS
1	PODO: Trial Design: Phase 2 Study of PF-06730512 in Focal Segmental Glomerulosclerosis. <i>Kidney International Reports</i> , 2021, 6, 1629-1633.	0.4	4
2	KIM-1 mediates fatty acid uptake by renal tubular cells to promote progressive diabetic kidney disease. <i>Cell Metabolism</i> , 2021, 33, 1042-1061.e7.	7.2	103
3	A conceptual framework linking immunology, pathology, and clinical features in primary membranous nephropathy. <i>Kidney International</i> , 2021, 100, 289-300.	2.6	25
4	T95 nucleophosmin phosphorylation as a novel mediator and marker of regulated cell death in acute kidney injury. <i>American Journal of Physiology - Renal Physiology</i> , 2020, 319, F552-F561.	1.3	3
5	Loss of Roundabout Guidance Receptor 2 (Robo2) in Podocytes Protects Adult Mice from Glomerular Injury by Maintaining Podocyte Foot Process Structure. <i>American Journal of Pathology</i> , 2020, 190, 799-816.	1.9	10
6	Control of Podocyte and Glomerular Capillary Wall Structure and Elasticity by WNK1 Kinase. <i>Frontiers in Cell and Developmental Biology</i> , 2020, 8, 618898.	1.8	5
7	Monoclonal IgG4/2 <sup>+</sup> Deposition Following Eculizumab Therapy for Recurrent Atypical Hemolytic Uremic Syndrome in Kidney Transplantation. <i>Kidney Medicine</i> , 2019, 1, 139-143.	1.0	0
8	Segmentation of Glomeruli Within Trichrome Images Using Deep Learning. <i>Kidney International Reports</i> , 2019, 4, 955-962.	0.4	126
9	Association of Pathological Fibrosis With Renal Survival Using Deep Neural Networks. <i>Kidney International Reports</i> , 2018, 3, 464-475.	0.4	114
10	Similar Biophysical Abnormalities in Glomeruli and Podocytes from Two Distinct Models. <i>Journal of the American Society of Nephrology: JASN</i> , 2018, 29, 1501-1512.	3.0	23
11	Concurrent Presentation of Thrombotic Thrombocytopenic Purpura and Membranous Nephropathy. <i>Kidney International Reports</i> , 2018, 3, 476-481.	0.4	3
12	TMIGD1 acts as a tumor suppressor through regulation of p21Cip1/p27Kip1 in renal cancer. <i>Oncotarget</i> , 2018, 9, 9672-9684.	0.8	20
13	Targeting STUB1-tissue factor axis normalizes hyperthrombotic uremic phenotype without increasing bleeding risk. <i>Science Translational Medicine</i> , 2017, 9, .	5.8	38
14	SLIT2/ROBO2 signaling pathway inhibits nonmuscle myosin IIA activity and destabilizes kidney podocyte adhesion. <i>JCI Insight</i> , 2016, 1, e86934.	2.3	34
15	Nonmuscle myosin IIA is critical for podocyte actin organization, contractility, and attenuation of cell motility. <i>Cytoskeleton</i> , 2016, 73, 377-395.	1.0	20
16	Pregnancy in a Patient With Primary Membranous Nephropathy and Circulating Anti-PLA2R Antibodies: A Case Report. <i>American Journal of Kidney Diseases</i> , 2016, 67, 775-778.	2.1	26
17	<sc>KIM</sc> mediated phagocytosis links <sc>ATG</sc> dependent clearance of apoptotic cells to antigen presentation. <i>EMBO Journal</i> , 2015, 34, 2441-2464.	3.5	76
18	Perivascular Gli1+ Progenitors Are Key Contributors to Injury-Induced Organ Fibrosis. <i>Cell Stem Cell</i> , 2015, 16, 51-66.	5.2	738

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19	Distinct Renal Pathology and a Chemotactic Phenotype after Enterohemorrhagic Escherichia coli Shiga Toxins in Non-Human Primate Models of Hemolytic Uremic Syndrome. American Journal of Pathology, 2013, 182, 1227-1238.	1.9	35
20	Gl $\alpha$ 12 activation in podocytes leads to cumulative changes in glomerular collagen expression, proteinuria and glomerulosclerosis. Laboratory Investigation, 2012, 92, 662-675.	1.7	19
21	Biophysical properties of normal and diseased renal glomeruli. American Journal of Physiology - Cell Physiology, 2011, 300, C397-C405.	2.1	91
22	Mutations in the formin gene INF2 cause focal segmental glomerulosclerosis. Nature Genetics, 2010, 42, 72-76.	9.4	381
23	Patients with ACTN4 Mutations Demonstrate Distinctive Features of Glomerular Injury. Journal of the American Society of Nephrology: JASN, 2009, 20, 961-968.	3.0	52
24	A Case of Familial Kidney Disease. Clinical Journal of the American Society of Nephrology: CJASN, 2007, 2, 1367-1374.	2.2	18
25	Novel Mutations in NPHP4 in a Consanguineous Family With Histological Findings of Focal Segmental Glomerulosclerosis. American Journal of Kidney Diseases, 2007, 50, 855-864.	2.1	16
26	Targeted Overexpression of the Transcription Factor XBP-1 in B Cells Promotes Plasma Cell and Lymphoplasmacytic Neoplasms in Transgenic Mice.. Blood, 2005, 106, 359-359.	0.6	4
27	$\alpha$ -Actinin-4-Mediated FSGS: An Inherited Kidney Disease Caused by an Aggregated and Rapidly Degraded Cytoskeletal Protein. PLoS Biology, 2004, 2, e167.	2.6	129
28	Traumatic calcinosis cutis in a dialysis patient. American Journal of Kidney Diseases, 2004, 44, e18-e21.	2.1	13
29	Gemcitabine-associated thrombotic microangiopathy. Cancer, 2004, 100, 2664-2670.	2.0	175
30	Mice deficient in $\alpha$ -actinin-4 have severe glomerular disease. Journal of Clinical Investigation, 2003, 111, 1683-1690.	3.9	210
31	Kidney Glomerulonephritis and Renal Ischemia. , 0, , 304-316.		0