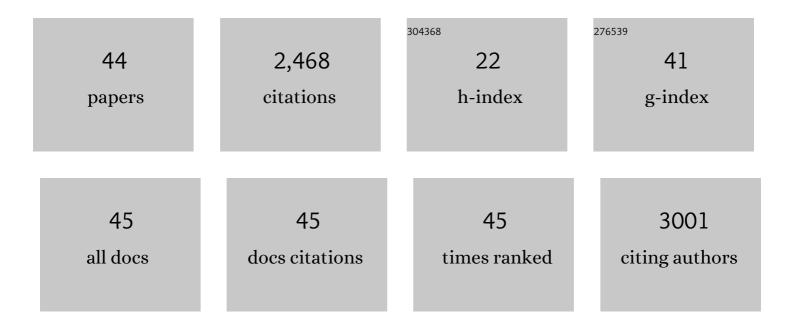
## Jaaakko Patrakka

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Hereditary Proteinuria Syndromes and Mechanisms of Proteinuria. New England Journal of Medicine, 2006, 354, 1387-1401.	13.9	492
2	Large-scale identification of genes implicated in kidney glomerulus development and function. EMBO Journal, 2006, 25, 1160-1174.	3.5	196
3	New insights into the role of podocytes in proteinuria. Nature Reviews Nephrology, 2009, 5, 463-468.	4.1	159
4	RECURRENCE OF NEPHROTIC SYNDROME IN KIDNEY GRAFTS OF PATIENTS WITH CONGENITAL NEPHROTIC SYNDROME OF THE FINNISH TYPE. Transplantation, 2002, 73, 394-403.	0.5	134
5	Nephrin – a unique structural and signaling protein of the kidney filter. Trends in Molecular Medicine, 2007, 13, 396-403.	3.5	126
6	CD2AP in mouse and human podocytes controls a proteolytic program that regulates cytoskeletal structure and cellular survival. Journal of Clinical Investigation, 2011, 121, 3965-3980.	3.9	124
7	Reducing VEGF-B Signaling Ameliorates Renal Lipotoxicity and Protects against Diabetic Kidney Disease. Cell Metabolism, 2017, 25, 713-726.	7.2	115
8	Molecular make-up of the glomerular filtration barrier. Biochemical and Biophysical Research Communications, 2010, 396, 164-169.	1.0	103
9	Role of Nephrin in Cell Junction Formation in Human Nephrogenesis. American Journal of Pathology, 2000, 157, 1905-1916.	1.9	100
10	Expression of Nephrin in Pediatric Kidney Diseases. Journal of the American Society of Nephrology: JASN, 2001, 12, 289-296.	3.0	75
11	Expression and Subcellular Distribution of Novel Glomerulus-Associated Proteins Dendrin, Ehd3, Sh2d4a, Plekhh2, and 2310066E14Rik. Journal of the American Society of Nephrology: JASN, 2007, 18, 689-697.	3.0	72
12	Proteinuria and prenatal diagnosis of congenital nephrosis in fetal carriers of nephrin gene mutations. Lancet, The, 2002, 359, 1575-1577.	6.3	69
13	A flexible, multilayered protein scaffold maintains the slit in between glomerular podocytes. JCI Insight, 2016, 1, .	2.3	69
14	A CRISP(e)R view on kidney organoids allows generation of an induced pluripotent stem cell–derived kidney model for drug discovery. Kidney International, 2018, 94, 1099-1110.	2.6	60
15	Single-cell RNA sequencing reveals the mesangial identity and species diversity of glomerular cell transcriptomes. Nature Communications, 2021, 12, 2141.	5.8	55
16	Neuronal proteins are novel components of podocyte major processes and their expression in glomerular crescents supports their role in crescent formation. Kidney International, 2013, 83, 63-71.	2.6	47
17	Confocal super-resolution imaging of the glomerular filtration barrier enabled by tissueÂexpansion. Kidney International, 2018, 93, 1008-1013.	2.6	47
18	The Number of Podocyte Slit Diaphragms Is Decreased in Minimal Change Nephrotic Syndrome. Pediatric Research, 2002, 52, 349-355.	1.1	39

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#	Article	IF	CITATIONS
19	Genetic kidney diseases disclose the pathogenesis of proteinuria. Annals of Medicine, 2001, 33, 526-533.	1.5	34
20	Understanding Podocyte Biology to Develop Novel Kidney Therapeutics. Frontiers in Endocrinology, 2018, 9, 409.	1.5	29
21	Novel insights into the disease transcriptome of human diabetic glomeruli and tubulointerstitium. Nephrology Dialysis Transplantation, 2020, 35, 2059-2072.	0.4	28
22	Glomerular Transcriptome Changes Associated with Lipopolysaccharide-Induced Proteinuria. American Journal of Nephrology, 2009, 29, 558-570.	1.4	27
23	Pdlim2 is a novel actin-regulating protein of podocyte foot processes. Kidney International, 2011, 80, 1045-1054.	2.6	24
24	Molecular insights into the early stage of glomerular injury in IgA nephropathy using single-cell RNA sequencing. Kidney International, 2022, 101, 752-765.	2.6	23
25	Novel INF2 mutation p. L77P in a family with glomerulopathy and Charcot-Marie-Tooth neuropathy. Pediatric Nephrology, 2013, 28, 339-343.	0.9	21
26	Myo1e Impairment Results in Actin Reorganization, Podocyte Dysfunction, and Proteinuria in Zebrafish and Cultured Podocytes. PLoS ONE, 2013, 8, e72750.	1.1	21
27	Plekhh2, a novel podocyte protein downregulated in human focal segmental glomerulosclerosis, is involved in matrix adhesion and actin dynamics. Kidney International, 2012, 82, 1071-1083.	2.6	20
28	GPRC5b Modulates Inflammatory Response in Glomerular Diseases via NF-κB Pathway. Journal of the American Society of Nephrology: JASN, 2019, 30, 1573-1586.	3.0	18
29	A fast and simple clearing and swelling protocol for 3D in-situ imaging of the kidney across scales. Kidney International, 2021, 99, 1010-1020.	2.6	18
30	Dendrin Ablation Prolongs Life Span by Delaying Kidney Failure. American Journal of Pathology, 2015, 185, 2143-2157.	1.9	17
31	Depletion of Gprc5a Promotes Development of Diabetic Nephropathy. Journal of the American Society of Nephrology: JASN, 2018, 29, 1679-1689.	3.0	16
32	Retinoic acid receptor responder1 promotes development of glomerular diseases via the Nuclear Factor-κB signaling pathway. Kidney International, 2021, 100, 809-823.	2.6	16
33	Schip1 Is a Novel Podocyte Foot Process Protein that Mediates Actin Cytoskeleton Rearrangements and Forms a Complex with Nherf2 and Ezrin. PLoS ONE, 2015, 10, e0122067.	1.1	14
34	Neph1 Is Reduced in Primary Focal Segmental Glomerulosclerosis, Minimal Change Nephrotic Syndrome, and Corresponding Experimental Animal Models of Adriamycin-Induced Nephropathy and Puromycin Aminonucleoside Nephrosis. Nephron Extra, 2014, 4, 146-154.	1.1	13
35	Coro2b, a podocyte protein downregulated in human diabetic nephropathy, is involved in the development of protamine sulphate-induced foot process effacement. Scientific Reports, 2019, 9, 8888.	1.6	8
36	Wtip- and Gadd45a-Interacting Protein Dendrin Is Not Crucial for the Development or Maintenance of the Glomerular Filtration Barrier. PLoS ONE, 2013, 8, e83133.	1.1	7

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#	Article	IF	CITATIONS
37	Novel NPHS2 variant in patients with familial steroid-resistant nephrotic syndrome with early onset, slow progression and dominant inheritance pattern. Clinical and Experimental Nephrology, 2017, 21, 677-684.	0.7	7
38	FYVE domain-containing protein ZFYVE28 regulates EGFR-signaling in podocytes but is not critical for the function of filtration barrier in mice. Scientific Reports, 2018, 8, 4712.	1.6	6
39	Association of crumbs homolog-2 with mTORC1 in developing podocyte. PLoS ONE, 2018, 13, e0202400.	1.1	6
40	Knockdown of Tmem234 in zebrafish results in proteinuria. American Journal of Physiology - Renal Physiology, 2015, 309, F955-F966.	1.3	5
41	The Number of Podocyte Slit Diaphragms Is Decreased in Minimal Change Nephrotic Syndrome. , 0, .		4
42	The role of Dendrin in IgA Nephropathy. Nephrology Dialysis Transplantation, 0, , .	0.4	3
43	Inactivation of mediator complex protein 22 in podocytes results in intracellular vacuole formation, podocyte loss and premature death. Scientific Reports, 2020, 10, 20037.	1.6	1
44	MO263KLOTHO IS PROTECTIVE IN THE CONTEXT OF ACUTE GLOMERULAR INJURY BUT IS NOT EXPRESSED IN PODOCYTES. Nephrology Dialysis Transplantation, 2021, 36, .	0.4	0