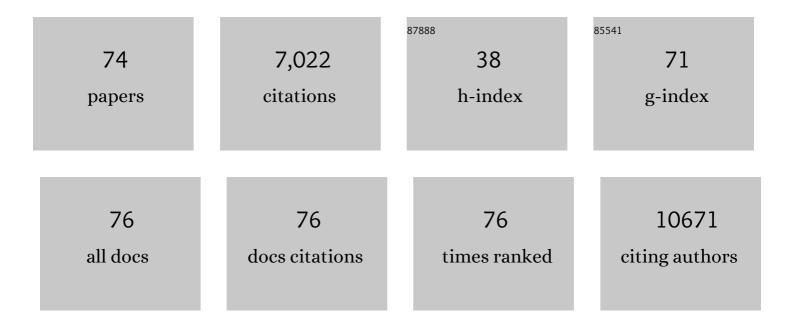
## David K Simon

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Suppression of Reactive Oxygen Species and Neurodegeneration by the PGC-1 Transcriptional Coactivators. Cell, 2006, 127, 397-408.	28.9	1,948
2	Parkinson Disease Epidemiology, Pathology, Genetics, and Pathophysiology. Clinics in Geriatric Medicine, 2020, 36, 1-12.	2.6	487
3	High aggregate burden of somatic mtDNA point mutations in aging and Alzheimer's disease brain. Human Molecular Genetics, 2002, 11, 133-145.	2.9	318
4	A Randomized Clinical Trial of High-Dosage Coenzyme Q10 in Early Parkinson Disease. JAMA Neurology, 2014, 71, 543.	9.0	312
5	Metaâ€analysis of Parkinson's Disease: Identification of a novel locus, <i>RIT2</i> . Annals of Neurology, 2012, 71, 370-384.	5.3	264
6	Dystonia. New England Journal of Medicine, 2006, 355, 818-829.	27.0	243
7	Revisiting protein aggregation as pathogenic in sporadic Parkinson and Alzheimer diseases. Neurology, 2019, 92, 329-337.	1.1	194
8	Effect of Creatine Monohydrate on Clinical Progression in Patients With Parkinson Disease. JAMA - Journal of the American Medical Association, 2015, 313, 584.	7.4	192
9	Mitochondrial Cyclic AMP Response Element-binding Protein (CREB) Mediates Mitochondrial Gene Expression and Neuronal Survival. Journal of Biological Chemistry, 2005, 280, 40398-40401.	3.4	187
10	Biomarkerâ€driven phenotyping in Parkinson's disease: A translational missing link in diseaseâ€modifying clinical trials. Movement Disorders, 2017, 32, 319-324.	3.9	145
11	Association of Cumulative Lead Exposure with Parkinson's Disease. Environmental Health Perspectives, 2010, 118, 1609-1613.	6.0	137
12	Genomewide association study of Parkinson's disease clinical biomarkers in 12 longitudinal patients' cohorts. Movement Disorders, 2019, 34, 1839-1850.	3.9	122
13	Noninvasive Brain Stimulation for Parkinson's Disease and Dystonia. Neurotherapeutics, 2008, 5, 345-361.	4.4	121
14	Somatic mitochondrial DNA mutations in cortex and substantia nigra in aging and Parkinson's disease. Neurobiology of Aging, 2004, 25, 71-81.	3.1	110
15	Genetic risk of Parkinson disease and progression:. Neurology: Genetics, 2019, 5, e348.	1.9	109
16	Somatic mitochondrial DNA mutations in early parkinson and incidental lewy body disease. Annals of Neurology, 2012, 71, 850-854.	5.3	103
17	Responses of retinal axons in vivo and in vitro to position-encoding molecules in the embryonic superior colliculus. Neuron, 1992, 9, 977-989.	8.1	99
18	Oral N-Acetyl-Cysteine Attenuates Loss of Dopaminergic Terminals in α-Synuclein Overexpressing Mice. PLoS ONE, 2010, 5, e12333.	2.5	97

DAVID K SIMON

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19	Limited topographic specificity in the targeting and branching of mammalian retinal axons. Developmental Biology, 1990, 137, 125-134.	2.0	96
20	A Common NURR1 Polymorphism Associated With Parkinson Disease and Diffuse Lewy Body Disease. Archives of Neurology, 2003, 60, 722.	4.5	90
21	Transcribe to Survive: Transcriptional Control of Antioxidant Defense Programs for Neuroprotection in Parkinson's Disease. Antioxidants and Redox Signaling, 2009, 11, 509-528.	5.4	88
22	Efficacy of Nilotinib in Patients With Moderately Advanced Parkinson Disease. JAMA Neurology, 2021, 78, 312.	9.0	83
23	Effect of Urate-Elevating Inosine on Early Parkinson Disease Progression. JAMA - Journal of the American Medical Association, 2021, 326, 926.	7.4	80
24	Rapamycin drives selection against a pathogenic heteroplasmic mitochondrial DNA mutation. Human Molecular Genetics, 2014, 23, 637-647.	2.9	79
25	Somatic mitochondrial DNA mutations in single neurons and glia. Neurobiology of Aging, 2005, 26, 1343-1355.	3.1	77
26	Frequency of the D620N Mutation in VPS35 in Parkinson Disease. Archives of Neurology, 2012, 69, 1360.	4.5	76
27	An inverse-Warburg effect and the origin of Alzheimer's disease. Biogerontology, 2012, 13, 583-594.	3.9	73
28	Relationship of retinotopic ordering of axons in the optic pathway to the formation of visual maps in central targets. Journal of Comparative Neurology, 1991, 307, 393-404.	1.6	67
29	Pgc-1α Overexpression Downregulates Pitx3 and Increases Susceptibility to MPTP Toxicity Associated with Decreased Bdnf. PLoS ONE, 2012, 7, e48925.	2.5	63
30	Differences in the Presentation and Progression of Parkinson's Disease by Sex. Movement Disorders, 2021, 36, 106-117.	3.9	54
31	Head injury at early ages is associated with risk of Parkinson's disease. Parkinsonism and Related Disorders, 2016, 23, 57-61.	2.2	50
32	Plasticity in the Development of Topographic Order in the Mammalian Retinocollicular Projection. Developmental Biology, 1994, 162, 384-393.	2.0	48
33	Singing in groups for Parkinson's disease (SING-PD): A pilot study of group singing therapy for PD-related voice/speech disorders. Parkinsonism and Related Disorders, 2012, 18, 548-552.	2.2	48
34	Attenuation of free radical production and paracrystalline inclusions by creatine supplementation in a patient with a novel cytochromeb mutation. Muscle and Nerve, 2004, 29, 537-547.	2.2	43
35	Influence of position along the medial-lateral axis of the superior colliculus on the topographic targeting and survival of retinal axons. Developmental Brain Research, 1992, 69, 167-172.	1.7	42
36	Do mtDNA deletions drive premature aging in mtDNA mutator mice?. Aging Cell, 2009, 8, 502-506.	6.7	40

DAVID K SIMON

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37	The inverse association of cancer and Alzheimer's: a bioenergetic mechanism. Journal of the Royal Society Interface, 2013, 10, 20130006.	3.4	39
38	A heteroplasmic mitochondrial complex I gene mutation in adult-onset dystonia. Neurogenetics, 2003, 4, 199-205.	1.4	38
39	Caffeine and Progression of Parkinson Disease. Clinical Neuropharmacology, 2008, 31, 189-196.	0.7	35
40	Metabolomic analysis of exercise effects in the POLG mitochondrial DNA mutator mouse brain. Neurobiology of Aging, 2015, 36, 2972-2983.	3.1	34
41	Behavioral and metabolic characterization of heterozygous and homozygous POLG mutator mice. Mitochondrion, 2013, 13, 282-291.	3.4	33
42	Mitochondrial Complex I Gene Variant Associated With Early Age at Onset in Spinocerebellar Ataxia Type 2. Archives of Neurology, 2007, 64, 1042.	4.5	30
43	MELAS syndrome, cardiomyopathy, rhabdomyolysis, and autism associated with the A3260G mitochondrial DNA mutation. Biochemical and Biophysical Research Communications, 2010, 402, 443-447.	2.1	28
44	Factors associated with falling in early, treated Parkinson's disease: The NET-PD LS1 cohort. Journal of the Neurological Sciences, 2017, 377, 137-143.	0.6	27
45	Maternal inheritance and mitochondrial DNA variants in familial Parkinson's disease. BMC Medical Genetics, 2010, 11, 53.	2.1	26
46	Peripheral Biomarkers of Parkinson's Disease Progression and Pioglitazone Effects. Journal of Parkinson's Disease, 2015, 5, 731-736.	2.8	25
47	Caffeine and Progression of Parkinson Disease. Clinical Neuropharmacology, 2015, 38, 163-169.	0.7	25
48	The c.â€237_236CA>TT <i>THAP1</i> sequence variant does not increase risk for primary dystonia. Movement Disorders, 2011, 26, 549-553.	3.9	24
49	Association of metabolic syndrome and change in Unified Parkinson's Disease Rating Scale scores. Neurology, 2017, 89, 1789-1794.	1.1	24
50	Caffeine, creatine, GRIN2A and Parkinson's disease progression. Journal of the Neurological Sciences, 2017, 375, 355-359.	0.6	23
51	Inverse Probability Weighted Cox Regression for Doubly Truncated Data. Biometrics, 2018, 74, 481-487.	1.4	21
52	Mistaken diagnosis of psychogenic gait disorder in a man with status cataplecticus ("Limp Man) Tj ETQq0 0 0 rg	BT /Overlo	ock 10 Tf 50 1
53	Autonomic and electrocardiographic findings in Parkinson's disease. Autonomic Neuroscience: Basic and Clinical, 2017, 205, 93-98.	2.8	19

DAVID K SIMON

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55	Somatic mitochondrial DNA mutations do not increase neuronal vulnerability to MPTP in young POLG mutator mice. Neurotoxicology and Teratology, 2014, 46, 62-67.	2.4	14
56	Clinical Impact of <sup>123</sup> I-Ioflupane SPECT (DaTscan) in a Movement Disorder Center. Neurodegenerative Diseases, 2017, 17, 38-43.	1.4	13
5 <b>7</b>	Do Somatic Mitochondrial DNA Mutations Contribute to Parkinson's Disease?. Parkinson's Disease, 2011, 2011, 1-9.	1.1	12
58	Computationally simple estimation and improved efficiency for special cases of double truncation. Lifetime Data Analysis, 2014, 20, 335-354.	0.9	12
59	Complicated spontaneous intracranial hypotension treated with intrathecal saline infusion. Practical Neurology, 2016, 16, 146-149.	1.1	12
60	A frameshift mitochondrial complex I gene mutation in a patient with dystonia and cataracts: is the mutation pathogenic?. Journal of Medical Genetics, 2001, 38, 58-61.	3.2	12
61	No Sex Differences in Use of Dopaminergic Medication in Early Parkinson Disease in the US and Canada - Baseline Findings of a Multicenter Trial. PLoS ONE, 2014, 9, e112287.	2.5	12
62	Altered muscle electrical tissue properties in a mouse model of premature aging. Muscle and Nerve, 2019, 60, 801-810.	2.2	11
63	Genetic risk factors in Parkinson's disease: single gene effects and interactions of genotypes. Journal of Neurology, 2012, 259, 2503-2505.	3.6	7
64	Mitochondrial DNA mutations in Parkinson's disease brain. Acta Neuropathologica Communications, 2017, 5, 33.	5.2	7
65	Transportation innovation to aid Parkinson disease trial recruitment. Contemporary Clinical Trials Communications, 2019, 16, 100449.	1.1	6
66	Acute readmission following deep brain stimulation surgery for Parkinson's disease: A nationwide analysis. Parkinsonism and Related Disorders, 2020, 70, 96-102.	2.2	6
67	Efficacy of Deep Brain Stimulation in a Patient with Genetically Confirmed Chorea-Acanthocytosis. Case Reports in Neurology, 2019, 11, 199-204.	0.7	5
68	Boxing Exercises as Therapy for Parkinson Disease. Topics in Geriatric Rehabilitation, 2020, 36, 160-165.	0.4	4
69	Mitochondria and Parkinson's Disease. Parkinson's Disease, 2011, 2011, 1-2.	1.1	3
70	The Relationship Between Olfactory Dysfunction and Constipation in Early Parkinson's Disease. Movement Disorders, 2021, 36, 781-782.	3.9	3
71	Novel human pathological mutations. Gene symbol: THAP1. Disease: dystonia 6. Human Genetics, 2010, 127, 470.	3.8	1
72	The utility of laser-generated visual-cueing in Parkinsonian patients with gait freezing. Parkinsonism and Related Disorders, 2012, 18, 401.	2.2	0

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73	A New Approach to the Development of <scp>Diseaseâ€Modifying</scp> Therapies for <scp>PD</scp> . Movement Disorders, 2021, 36, 1281-1281.	3.9	0
74	Associations between exercise classes and self-reported exercise by people with Parkinson's disease at Parkinson's foundation centers of excellence. Clinical Parkinsonism & Related Disorders, 2022, 6, 100137.	0.9	0