

David K Simon

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

74
papers

5,373
citations

34
h-index

73
g-index

76
ext. papers

6,252
ext. citations

6.9
avg, IF

5.32
L-index

#	Paper	IF	Citations
74	Associations between exercise classes and self-reported exercise by people with Parkinson's disease at Parkinson's foundation centers of excellence.. <i>Clinical Parkinsonism & Related Disorders</i> , 2022 , 6, 100137	0.9	
73	Efficacy of Nilotinib in Patients With Moderately Advanced Parkinson Disease: A Randomized Clinical Trial. <i>JAMA Neurology</i> , 2021 , 78, 312-320	17.2	34
72	A New Approach to the Development of Disease-Modifying Therapies for PD. <i>Movement Disorders</i> , 2021 , 36, 1281	7	
71	Differences in the Presentation and Progression of Parkinson's Disease by Sex. <i>Movement Disorders</i> , 2021 , 36, 106-117	7	16
70	The Relationship Between Olfactory Dysfunction and Constipation in Early Parkinson's Disease. <i>Movement Disorders</i> , 2021 , 36, 781-782	7	1
69	Effect of Urate-Elevating Inosine on Early Parkinson Disease Progression: The SURE-PD3 Randomized Clinical Trial. <i>JAMA - Journal of the American Medical Association</i> , 2021 , 326, 926-939	27.4	16
68	Acute readmission following deep brain stimulation surgery for Parkinson's disease: A nationwide analysis. <i>Parkinsonism and Related Disorders</i> , 2020 , 70, 96-102	3.6	3
67	Parkinson Disease Epidemiology, Pathology, Genetics, and Pathophysiology. <i>Clinics in Geriatric Medicine</i> , 2020 , 36, 1-12	3.8	143
66	VPS35 and the mitochondria: Connecting the dots in Parkinson's disease pathophysiology. <i>Neurobiology of Disease</i> , 2020 , 145, 105056	7.5	9
65	Boxing Exercises as Therapy for Parkinson Disease. <i>Topics in Geriatric Rehabilitation</i> , 2020 , 36, 160-165	0.7	3
64	Altered muscle electrical tissue properties in a mouse model of premature aging. <i>Muscle and Nerve</i> , 2019 , 60, 801-810	3.4	6
63	Genetic risk of Parkinson disease and progression:: An analysis of 13 longitudinal cohorts. <i>Neurology: Genetics</i> , 2019 , 5, e348	3.8	57
62	Transportation innovation to aid Parkinson disease trial recruitment. <i>Contemporary Clinical Trials Communications</i> , 2019 , 16, 100449	1.8	4
61	Revisiting protein aggregation as pathogenic in sporadic Parkinson and Alzheimer diseases. <i>Neurology</i> , 2019 , 92, 329-337	6.5	144
60	Efficacy of Deep Brain Stimulation in a Patient with Genetically Confirmed Chorea-Acanthocytosis. <i>Case Reports in Neurology</i> , 2019 , 11, 199-204	1	2
59	Genomewide association study of Parkinson's disease clinical biomarkers in 12 longitudinal patients' cohorts. <i>Movement Disorders</i> , 2019 , 34, 1839-1850	7	69
58	Inverse probability weighted Cox regression for doubly truncated data. <i>Biometrics</i> , 2018 , 74, 481-487	1.8	13

57	Biomarker-driven phenotyping in Parkinson's disease: A translational missing link in disease-modifying clinical trials. <i>Movement Disorders</i> , 2017 , 32, 319-324	7	111
56	Caffeine, creatine, GRIN2A and Parkinson's disease progression. <i>Journal of the Neurological Sciences</i> , 2017 , 375, 355-359	3.2	17
55	Autonomic and electrocardiographic findings in Parkinson's disease. <i>Autonomic Neuroscience: Basic and Clinical</i> , 2017 , 205, 93-98	2.4	11
54	Factors associated with falling in early, treated Parkinson's disease: The NET-PD LS1 cohort. <i>Journal of the Neurological Sciences</i> , 2017 , 377, 137-143	3.2	19
53	Association of metabolic syndrome and change in Unified Parkinson's Disease Rating Scale scores. <i>Neurology</i> , 2017 , 89, 1789-1794	6.5	15
52	Mitochondrial DNA mutations in Parkinson's disease brain. <i>Acta Neuropathologica Communications</i> , 2017 , 5, 33	7.3	7
51	Clinical Impact of 123I-Ioflupane SPECT (DaTscan) in a Movement Disorder Center. <i>Neurodegenerative Diseases</i> , 2017 , 17, 38-43	2.3	8
50	Head injury at early ages is associated with risk of Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2016 , 23, 57-61	3.6	33
49	Complicated spontaneous intracranial hypotension treated with intrathecal saline infusion. <i>Practical Neurology</i> , 2016 , 16, 146-9	2.4	8
48	Metabolomic analysis of exercise effects in the POLG mitochondrial DNA mutator mouse brain. <i>Neurobiology of Aging</i> , 2015 , 36, 2972-2983	5.6	23
47	Peripheral Biomarkers of Parkinson's Disease Progression and Pioglitazone Effects. <i>Journal of Parkinsons Disease</i> , 2015 , 5, 731-6	5.3	20
46	Effect of creatine monohydrate on clinical progression in patients with Parkinson disease: a randomized clinical trial. <i>JAMA - Journal of the American Medical Association</i> , 2015 , 313, 584-93	27.4	153
45	Caffeine and Progression of Parkinson Disease: A Deleterious Interaction With Creatine. <i>Clinical Neuropharmacology</i> , 2015 , 38, 163-9	1.4	18
44	Somatic mitochondrial DNA mutations do not increase neuronal vulnerability to MPTP in young POLG mutator mice. <i>Neurotoxicology and Teratology</i> , 2014 , 46, 62-7	3.9	11
43	Rapamycin drives selection against a pathogenic heteroplasmic mitochondrial DNA mutation. <i>Human Molecular Genetics</i> , 2014 , 23, 637-47	5.6	63
42	A randomized clinical trial of high-dosage coenzyme Q10 in early Parkinson disease: no evidence of benefit. <i>JAMA Neurology</i> , 2014 , 71, 543-52	17.2	239
41	Computationally simple estimation and improved efficiency for special cases of double truncation. <i>Lifetime Data Analysis</i> , 2014 , 20, 335-54	1.3	8
40	No sex differences in use of dopaminergic medication in early Parkinson disease in the US and Canada - baseline findings of a multicenter trial. <i>PLoS ONE</i> , 2014 , 9, e112287	3.7	9

39	The inverse association of cancer and Alzheimer's: a bioenergetic mechanism. <i>Journal of the Royal Society Interface</i> , 2013 , 10, 20130006	4.1	38
38	Behavioral and metabolic characterization of heterozygous and homozygous POLG mutator mice. <i>Mitochondrion</i> , 2013 , 13, 282-91	4.9	23
37	Genetic risk factors in Parkinson's disease: single gene effects and interactions of genotypes. <i>Journal of Neurology</i> , 2012 , 259, 2503-5	5.5	6
36	An inverse-Warburg effect and the origin of Alzheimer's disease. <i>Biogerontology</i> , 2012 , 13, 583-94	4.5	59
35	The utility of laser-generated visual-cueing in Parkinsonian patients with gait freezing. <i>Parkinsonism and Related Disorders</i> , 2012 , 18, 401	3.6	
34	Singing in groups for Parkinson's disease (SING-PD): a pilot study of group singing therapy for PD-related voice/speech disorders. <i>Parkinsonism and Related Disorders</i> , 2012 , 18, 548-52	3.6	36
33	Meta-analysis of Parkinson's disease: identification of a novel locus, RIT2. <i>Annals of Neurology</i> , 2012 , 71, 370-84	9.4	214
32	Somatic mitochondrial DNA mutations in early Parkinson and incidental Lewy body disease. <i>Annals of Neurology</i> , 2012 , 71, 850-4	9.4	79
31	Frequency of the D620N mutation in VPS35 in Parkinson disease. <i>Archives of Neurology</i> , 2012 , 69, 1360-4		66
30	Pgc-1 β overexpression downregulates Pitx3 and increases susceptibility to MPTP toxicity associated with decreased Bdnf. <i>PLoS ONE</i> , 2012 , 7, e48925	3.7	52
29	Mitochondria and Parkinson's disease. <i>Parkinson's Disease</i> , 2011 , 2011, 261791	2.6	2
28	Do somatic mitochondrial DNA mutations contribute to Parkinson's disease?. <i>Parkinson's Disease</i> , 2011 , 2011, 659694	2.6	10
27	The c.-237_236GA>TT THAP1 sequence variant does not increase risk for primary dystonia. <i>Movement Disorders</i> , 2011 , 26, 549-52	7	15
26	Association of cumulative lead exposure with Parkinson's disease. <i>Environmental Health Perspectives</i> , 2010 , 118, 1609-13	8.4	97
25	MELAS syndrome, cardiomyopathy, rhabdomyolysis, and autism associated with the A3260G mitochondrial DNA mutation. <i>Biochemical and Biophysical Research Communications</i> , 2010 , 402, 443-7	3.4	19
24	Maternal inheritance and mitochondrial DNA variants in familial Parkinson's disease. <i>BMC Medical Genetics</i> , 2010 , 11, 53	2.1	26
23	Oral N-acetyl-cysteine attenuates loss of dopaminergic terminals in alpha-synuclein overexpressing mice. <i>PLoS ONE</i> , 2010 , 5, e12333	3.7	80
22	Do mtDNA deletions drive premature aging in mtDNA mutator mice?. <i>Aging Cell</i> , 2009 , 8, 502-6	9.9	37

21	Transcribe to survive: transcriptional control of antioxidant defense programs for neuroprotection in Parkinson's disease. <i>Antioxidants and Redox Signaling</i> , 2009 , 11, 509-28	8.4	77
20	Caffeine and progression of Parkinson disease. <i>Clinical Neuropharmacology</i> , 2008 , 31, 189-96	1.4	30
19	Noninvasive brain stimulation for Parkinson's disease and dystonia. <i>Neurotherapeutics</i> , 2008 , 5, 345-61	6.4	103
18	Mitochondrial complex I gene variant associated with early age at onset in spinocerebellar ataxia type 2. <i>Archives of Neurology</i> , 2007 , 64, 1042-4		29
17	Dystonia. <i>New England Journal of Medicine</i> , 2006 , 355, 818-29	59.2	205
16	Suppression of reactive oxygen species and neurodegeneration by the PGC-1 transcriptional coactivators. <i>Cell</i> , 2006 , 127, 397-408	56.2	1689
15	Somatic mitochondrial DNA mutations in single neurons and glia. <i>Neurobiology of Aging</i> , 2005 , 26, 1343-55	5.5	69
14	Mitochondrial cyclic AMP response element-binding protein (CREB) mediates mitochondrial gene expression and neuronal survival. <i>Journal of Biological Chemistry</i> , 2005 , 280, 40398-401	5.4	155
13	Mistaken diagnosis of psychogenic gait disorder in a man with status cataplecticus ("limp man syndrome"). <i>Movement Disorders</i> , 2004 , 19, 838-840	7	15
12	Attenuation of free radical production and paracrystalline inclusions by creatine supplementation in a patient with a novel cytochrome b mutation. <i>Muscle and Nerve</i> , 2004 , 29, 537-47	3.4	41
11	Somatic mitochondrial DNA mutations in cortex and substantia nigra in aging and Parkinson's disease. <i>Neurobiology of Aging</i> , 2004 , 25, 71-81	5.6	95
10	A common NURR1 polymorphism associated with Parkinson disease and diffuse Lewy body disease. <i>Archives of Neurology</i> , 2003 , 60, 722-5		79
9	A heteroplasmic mitochondrial complex I gene mutation in adult-onset dystonia. <i>Neurogenetics</i> , 2003 , 4, 199-205	3	32
8	High aggregate burden of somatic mtDNA point mutations in aging and Alzheimer's disease brain. <i>Human Molecular Genetics</i> , 2002 , 11, 133-45	5.6	262
7	A frameshift mitochondrial complex I gene mutation in a patient with dystonia and cataracts: is the mutation pathogenic?. <i>Journal of Medical Genetics</i> , 2001 , 38, 58-61	5.8	7
6	Plasticity in the development of topographic order in the mammalian retinocollicular projection. <i>Developmental Biology</i> , 1994 , 162, 384-93	3.1	47
5	Responses of retinal axons in vivo and in vitro to position-encoding molecules in the embryonic superior colliculus. <i>Neuron</i> , 1992 , 9, 977-89	13.9	90
4	Influence of position along the medial-lateral axis of the superior colliculus on the topographic targeting and survival of retinal axons. <i>Developmental Brain Research</i> , 1992 , 69, 167-72		40

3	Relationship of retinotopic ordering of axons in the optic pathway to the formation of visual maps in central targets. <i>Journal of Comparative Neurology</i> , 1991 , 307, 393-404	3.4	62
2	Limited topographic specificity in the targeting and branching of mammalian retinal axons. <i>Developmental Biology</i> , 1990 , 137, 125-34	3.1	87
1	Nilotinib in Patients with Advanced Parkinson Disease: A Randomized Phase 2A Study (NILO-PD)		6