

# Daniela D Berg

## List of Publications by Citations

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557  
papers

34,080  
citations

91  
h-index

169  
g-index

589  
ext. papers

40,971  
ext. citations

5.9  
avg, IF

6.93  
L-index

#	Paper	IF	Citations
557	MDS clinical diagnostic criteria for Parkinson's disease. <i>Movement Disorders</i> , <b>2015</b> , 30, 1591-601	7	2371
556	Genome-wide association study reveals genetic risk underlying Parkinson's disease. <i>Nature Genetics</i> , <b>2009</b> , 41, 1308-12	36.3	1469
555	Multicenter analysis of glucocerebrosidase mutations in Parkinson's disease. <i>New England Journal of Medicine</i> , <b>2009</b> , 361, 1651-61	59.2	1351
554	The Parkinson Progression Marker Initiative (PPMI). <i>Progress in Neurobiology</i> , <b>2011</b> , 95, 629-35	10.9	793
553	Imputation of sequence variants for identification of genetic risks for Parkinson's disease: a meta-analysis of genome-wide association studies. <i>Lancet, The</i> , <b>2011</b> , 377, 641-9	40	733
552	MDS research criteria for prodromal Parkinson's disease. <i>Movement Disorders</i> , <b>2015</b> , 30, 1600-11	7	645
551	Differentiation between intracerebral hemorrhage and ischemic stroke by transcranial color-coded duplex-sonography. <i>Stroke</i> , <b>1998</b> , 29, 2563-7	6.7	583
550	Mutations in the gene encoding epsilon-sarcoglycan cause myoclonus-dystonia syndrome. <i>Nature Genetics</i> , <b>2001</b> , 29, 66-9	36.3	458
549	Loss of function mutations in the gene encoding Omi/HtrA2 in Parkinson's disease. <i>Human Molecular Genetics</i> , <b>2005</b> , 14, 2099-111	5.6	446
548	Loss-of-function of human PINK1 results in mitochondrial pathology and can be rescued by parkin. <i>Journal of Neuroscience</i> , <b>2007</b> , 27, 12413-8	6.6	431
547	Comprehensive research synopsis and systematic meta-analyses in Parkinson's disease genetics: The PDGene database. <i>PLoS Genetics</i> , <b>2012</b> , 8, e1002548	6	420
546	Identification of common variants influencing risk of the tauopathy progressive supranuclear palsy. <i>Nature Genetics</i> , <b>2011</b> , 43, 699-705	36.3	386
545	Genetic correction of a LRRK2 mutation in human iPSCs links parkinsonian neurodegeneration to ERK-dependent changes in gene expression. <i>Cell Stem Cell</i> , <b>2013</b> , 12, 354-67	18	382
544	Past, present, and future of Parkinson's disease: A special essay on the 200th Anniversary of the Shaking Palsy. <i>Movement Disorders</i> , <b>2017</b> , 32, 1264-1310	7	375
543	Echogenicity of the substantia nigra: association with increased iron content and marker for susceptibility to nigrostriatal injury. <i>Archives of Neurology</i> , <b>2002</b> , 59, 999-1005		366
542	14-3-3 proteins in the nervous system. <i>Nature Reviews Neuroscience</i> , <b>2003</b> , 4, 752-62	13.5	350
541	iPSC-derived neurons from GBA1-associated Parkinson's disease patients show autophagic defects and impaired calcium homeostasis. <i>Nature Communications</i> , <b>2014</b> , 5, 4028	17.4	324

540	EFNS/MDS-ES/ENS [corrected] recommendations for the diagnosis of Parkinson's disease. <i>European Journal of Neurology</i> , <b>2013</b> , 20, 16-34	6	299
539	Time to redefine PD? Introductory statement of the MDS Task Force on the definition of Parkinson's disease. <i>Movement Disorders</i> , <b>2014</b> , 29, 454-62	7	290
538	Transcranial sonography in movement disorders. <i>Lancet Neurology</i> , <b>2008</b> , 7, 1044-55	24.1	290
537	A multicenter study of glucocerebrosidase mutations in dementia with Lewy bodies. <i>JAMA Neurology</i> , <b>2013</b> , 70, 727-35	17.2	285
536	Unbiased screen for interactors of leucine-rich repeat kinase 2 supports a common pathway for sporadic and familial Parkinson disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2014</b> , 111, 2626-31	11.5	282
535	Brain iron pathways and their relevance to Parkinson's disease. <i>Journal of Neurochemistry</i> , <b>2001</b> , 79, 225-36	6	275
534	Echogenicity of the substantia nigra in Parkinson's disease and its relation to clinical findings. <i>Journal of Neurology</i> , <b>2001</b> , 248, 684-9	5.5	256
533	Increased susceptibility to sporadic Parkinson's disease by a certain combined alpha-synuclein/apolipoprotein E genotype. <i>Annals of Neurology</i> , <b>1999</b> , 45, 611-7	9.4	245
532	Association of LRRK2 exonic variants with susceptibility to Parkinson's disease: a case-control study. <i>Lancet Neurology</i> , <b>2011</b> , 10, 898-908	24.1	237
531	Parkin is transcriptionally regulated by ATF4: evidence for an interconnection between mitochondrial stress and ER stress. <i>Cell Death and Differentiation</i> , <b>2011</b> , 18, 769-82	12.7	235
530	Genome-wide association study of intracranial aneurysm identifies three new risk loci. <i>Nature Genetics</i> , <b>2010</b> , 42, 420-5	36.3	234
529	SNCA variants are associated with increased risk for multiple system atrophy. <i>Annals of Neurology</i> , <b>2009</b> , 65, 610-4	9.4	232
528	Vulnerability of the nigrostriatal system as detected by transcranial ultrasound. <i>Neurology</i> , <b>1999</b> , 53, 1026-31	6.5	227
527	Loss of VPS13C Function in Autosomal-Recessive Parkinsonism Causes Mitochondrial Dysfunction and Increases PINK1/Parkin-Dependent Mitophagy. <i>American Journal of Human Genetics</i> , <b>2016</b> , 98, 500-513	11.3	225
526	Type and frequency of mutations in the LRRK2 gene in familial and sporadic Parkinson's disease*. <i>Brain</i> , <b>2005</b> , 128, 3000-11	11.2	217
525	Excessive burden of lysosomal storage disorder gene variants in Parkinson's disease. <i>Brain</i> , <b>2017</b> , 140, 3191-3203	11.2	209
524	A two-stage meta-analysis identifies several new loci for Parkinson's disease. <i>PLoS Genetics</i> , <b>2011</b> , 7, e1002142	6	209
523	Genetic variability in the SNCA gene influences alpha-synuclein levels in the blood and brain. <i>FASEB Journal</i> , <b>2008</b> , 22, 1327-34	0.9	209

522	Progression of Parkinson's disease in the clinical phase: potential markers. <i>Lancet Neurology, The</i> , <b>2009</b> , 8, 1158-71	24.1	208
521	Multiple regions of alpha-synuclein are associated with Parkinson's disease. <i>Annals of Neurology</i> , <b>2005</b> , 57, 535-41	9.4	206
520	A phase 2 trial of the GSK-3 inhibitor tideglusib in progressive supranuclear palsy. <i>Movement Disorders</i> , <b>2014</b> , 29, 470-8	7	201
519	Five-year follow-up study of hyperechogenicity of the substantia nigra in Parkinson's disease. <i>Movement Disorders</i> , <b>2005</b> , 20, 383-5	7	199
518	The specificity and sensitivity of transcranial ultrasound in the differential diagnosis of Parkinson's disease: a prospective blinded study. <i>Lancet Neurology, The</i> , <b>2008</b> , 7, 417-24	24.1	195
517	Transcranial brain parenchyma sonography in movement disorders: state of the art. <i>Ultrasound in Medicine and Biology</i> , <b>2007</b> , 33, 15-25	3.5	195
516	Advances in markers of prodromal Parkinson disease. <i>Nature Reviews Neurology</i> , <b>2016</b> , 12, 622-634	15	189
515	GBA-associated PD presents with nonmotor characteristics. <i>Neurology</i> , <b>2011</b> , 77, 276-80	6.5	182
514	Phosphorylated $\beta$ synuclein in Parkinson's disease. <i>Science Translational Medicine</i> , <b>2012</b> , 4, 121ra20	17.5	182
513	Role of iron in neurodegenerative disorders. <i>Topics in Magnetic Resonance Imaging</i> , <b>2006</b> , 17, 5-17	2.3	178
512	In vivo detection of iron and neuromelanin by transcranial sonography: a new approach for early detection of substantia nigra damage. <i>Movement Disorders</i> , <b>2005</b> , 20, 1278-85	7	178
511	Update of the MDS research criteria for prodromal Parkinson's disease. <i>Movement Disorders</i> , <b>2019</b> , 34, 1464-1470	7	177
510	Higher frequency of regulatory T cells in the elderly and increased suppressive activity in neurodegeneration. <i>Journal of Neuroimmunology</i> , <b>2007</b> , 188, 117-27	3.5	170
509	Relationship of substantia nigra echogenicity and motor function in elderly subjects. <i>Neurology</i> , <b>2001</b> , 56, 13-7	6.5	167
508	Redox imbalance. <i>Cell and Tissue Research</i> , <b>2004</b> , 318, 201-13	4.2	160
507	Cortical PIB binding in Lewy body disease is associated with Alzheimer-like characteristics. <i>Neurobiology of Disease</i> , <b>2009</b> , 34, 107-12	7.5	158
506	Ubiquitylation of synphilin-1 and alpha-synuclein by SIAH and its presence in cellular inclusions and Lewy bodies imply a role in Parkinson's disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2004</b> , 101, 5500-5	11.5	157
505	Genetic analysis implicates APOE, SNCA and suggests lysosomal dysfunction in the etiology of dementia with Lewy bodies. <i>Human Molecular Genetics</i> , <b>2014</b> , 23, 6139-46	5.6	152

504	Iron accumulation in the substantia nigra in rats visualized by ultrasound. <i>Ultrasound in Medicine and Biology</i> , <b>1999</b> , 25, 901-4	3.5	150
503	AFQ056 treatment of levodopa-induced dyskinesias: results of 2 randomized controlled trials. <i>Movement Disorders</i> , <b>2011</b> , 26, 1243-50	7	147
502	Using genome-wide complex trait analysis to quantify 'missing heritability' in Parkinson's disease. <i>Human Molecular Genetics</i> , <b>2012</b> , 21, 4996-5009	5.6	145
501	Angiogenin variants in Parkinson disease and amyotrophic lateral sclerosis. <i>Annals of Neurology</i> , <b>2011</b> , 70, 964-73	9.4	144
500	Iron metabolism in Parkinsonian syndromes. <i>Movement Disorders</i> , <b>2006</b> , 21, 1299-310	7	144
499	The Parkinson's progression markers initiative (PPMI) - establishing a PD biomarker cohort. <i>Annals of Clinical and Translational Neurology</i> , <b>2018</b> , 5, 1460-1477	5.3	142
498	GBA-associated Parkinson's disease: reduced survival and more rapid progression in a prospective longitudinal study. <i>Movement Disorders</i> , <b>2015</b> , 30, 407-11	7	135
497	Enlarged substantia nigra hyperechogenicity and risk for Parkinson disease: a 37-month 3-center study of 1847 older persons. <i>Archives of Neurology</i> , <b>2011</b> , 68, 932-7		131
496	Parkin protects mitochondrial genome integrity and supports mitochondrial DNA repair. <i>Human Molecular Genetics</i> , <b>2009</b> , 18, 3832-50	5.6	130
495	Brain-first versus body-first Parkinson's disease: a multimodal imaging case-control study. <i>Brain</i> , <b>2020</b> , 143, 3077-3088	11.2	130
494	Mutation analysis and association studies of the UCHL1 gene in German Parkinson's disease patients. <i>NeuroReport</i> , <b>2000</b> , 11, 2079-82	1.7	128
493	Plasma ceramide and glucosylceramide metabolism is altered in sporadic Parkinson's disease and associated with cognitive impairment: a pilot study. <i>PLoS ONE</i> , <b>2013</b> , 8, e73094	3.7	124
492	[11C]PIB binding in Parkinson's disease dementia. <i>NeuroImage</i> , <b>2008</b> , 39, 1027-33	7.9	120
491	Differentiation of Parkinson's disease and atypical parkinsonian syndromes by transcranial ultrasound. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2005</b> , 76, 423-5	5.5	120
490	S100B is increased in Parkinson's disease and ablation protects against MPTP-induced toxicity through the RAGE and TNF- $\alpha$ pathway. <i>Brain</i> , <b>2012</b> , 135, 3336-47	11.2	118
489	Cerebral activation patterns in patients with writer's cramp: a functional magnetic resonance imaging study. <i>Journal of Neurology</i> , <b>2001</b> , 248, 10-7	5.5	112
488	Substantia nigra hypoechogenicity: definition and findings in restless legs syndrome. <i>Movement Disorders</i> , <b>2007</b> , 22, 187-92	7	111
487	Lrrk2 R1441C parkinsonism is clinically similar to sporadic Parkinson disease. <i>Neurology</i> , <b>2008</b> , 70, 1456-60	6.5	109

486	Changing the research criteria for the diagnosis of Parkinson's disease: obstacles and opportunities. <i>Lancet Neurology, The</i> , <b>2013</b> , 12, 514-24	24.1	108
485	Multiregional brain iron deficiency in restless legs syndrome. <i>Movement Disorders</i> , <b>2008</b> , 23, 1184-7	7	108
484	The patients' perception of prodromal symptoms before the initial diagnosis of Parkinson's disease. <i>Movement Disorders</i> , <b>2011</b> , 26, 653-8	7	107
483	Tideglusib reduces progression of brain atrophy in progressive supranuclear palsy in a randomized trial. <i>Movement Disorders</i> , <b>2014</b> , 29, 479-87	7	106
482	Large-scale replication and heterogeneity in Parkinson disease genetic loci. <i>Neurology</i> , <b>2012</b> , 79, 659-67	6.5	106
481	Novel homozygous p.E64D mutation in DJ1 in early onset Parkinson disease (PARK7). <i>Human Mutation</i> , <b>2004</b> , 24, 321-9	4.7	104
480	Validation of the MDS clinical diagnostic criteria for Parkinson's disease. <i>Movement Disorders</i> , <b>2018</b> , 33, 1601-1608	7	101
479	Ceruloplasmin gene variations and substantia nigra hyperechogenicity in Parkinson disease. <i>Neurology</i> , <b>2004</b> , 63, 1912-7	6.5	99
478	Neuroimaging in basal ganglia disorders: perspectives for transcranial ultrasound. <i>Movement Disorders</i> , <b>2001</b> , 16, 23-32	7	98
477	Depression in Parkinson's disease: brainstem midline alteration on transcranial sonography and magnetic resonance imaging. <i>Journal of Neurology</i> , <b>1999</b> , 246, 1186-93	5.5	98
476	Defining at-risk populations for Parkinson's disease: lessons from ongoing studies. <i>Movement Disorders</i> , <b>2012</b> , 27, 656-65	7	96
475	A pathway-based analysis provides additional support for an immune-related genetic susceptibility to Parkinson's disease. <i>Human Molecular Genetics</i> , <b>2013</b> , 22, 1039-49	5.6	96
474	Health-Related Quality of Life in patients with Parkinson's disease--A systematic review based on the ICF model. <i>Neuroscience and Biobehavioral Reviews</i> , <b>2016</b> , 61, 26-34	9	95
473	Echogenicity of substantia nigra determined by transcranial ultrasound correlates with severity of parkinsonian symptoms induced by neuroleptic therapy. <i>Biological Psychiatry</i> , <b>2001</b> , 50, 463-7	7.9	95
472	To rise and to fall: functional connectivity in cognitively normal and cognitively impaired patients with Parkinson's disease. <i>Neurobiology of Aging</i> , <b>2015</b> , 36, 1727-1735	5.6	94
471	The transcription factor PITX3 is associated with sporadic Parkinson's disease. <i>Neurobiology of Aging</i> , <b>2009</b> , 30, 731-8	5.6	92
470	The PRIPS study: screening battery for subjects at risk for Parkinson's disease. <i>European Journal of Neurology</i> , <b>2013</b> , 20, 102-8	6	91
469	Osteopontin is elevated in Parkinson's disease and its absence leads to reduced neurodegeneration in the MPTP model. <i>Neurobiology of Disease</i> , <b>2007</b> , 25, 473-82	7.5	91

468	Disturbance of iron metabolism in Parkinson's disease -- ultrasonography as a biomarker. <i>Neurotoxicity Research</i> , <b>2006</b> , 9, 1-13	4.3	91
467	Echogenicity of the substantia nigra in relatives of patients with sporadic Parkinson's disease. <i>NeuroImage</i> , <b>2003</b> , 18, 416-22	7.9	91
466	Enlarged hyperechogenic substantia nigra as a risk marker for Parkinson's disease. <i>Movement Disorders</i> , <b>2013</b> , 28, 216-9	7	90
465	Increased tissue copper and manganese content in the lentiform nucleus in primary adult-onset dystonia. <i>Annals of Neurology</i> , <b>1999</b> , 46, 260-3	9.4	86
464	Longitudinal CSF biomarkers in patients with early Parkinson disease and healthy controls. <i>Neurology</i> , <b>2017</b> , 89, 1959-1969	6.5	84
463	Genetic analysis of immunomodulating factors in sporadic Parkinson's disease. <i>Journal of Neural Transmission</i> , <b>2000</b> , 107, 553-62	4.3	82
462	Arm swing as a potential new prodromal marker of Parkinson's disease. <i>Movement Disorders</i> , <b>2016</b> , 31, 1527-1534	7	80
461	Serum iron levels and the risk of Parkinson disease: a Mendelian randomization study. <i>PLoS Medicine</i> , <b>2013</b> , 10, e1001462	11.6	80
460	Urinary LRRK2 phosphorylation predicts parkinsonian phenotypes in G2019S LRRK2 carriers. <i>Neurology</i> , <b>2016</b> , 86, 994-9	6.5	78
459	Prodromal features for Parkinson's disease--baseline data from the TREND study. <i>European Journal of Neurology</i> , <b>2014</b> , 21, 766-72	6	76
458	Independent and joint effects of the MAPT and SNCA genes in Parkinson disease. <i>Annals of Neurology</i> , <b>2011</b> , 69, 778-92	9.4	76
457	Loss of nocturnal blood pressure fall in various extrapyramidal syndromes. <i>Movement Disorders</i> , <b>2009</b> , 24, 2136-42	7	76
456	Penetrance estimate of LRRK2 p.G2019S mutation in individuals of non-Ashkenazi Jewish ancestry. <i>Movement Disorders</i> , <b>2017</b> , 32, 1432-1438	7	74
455	The basal ganglia in haemochromatosis. <i>Neuroradiology</i> , <b>2000</b> , 42, 9-13	3.2	74
454	Biomarkers for the early detection of Parkinson's and Alzheimer's disease. <i>Neurodegenerative Diseases</i> , <b>2008</b> , 5, 133-6	2.3	72
453	NeuroX, a fast and efficient genotyping platform for investigation of neurodegenerative diseases. <i>Neurobiology of Aging</i> , <b>2015</b> , 36, 1605.e7-12	5.6	70
452	Olfactory dysfunction: common in later life and early warning of neurodegenerative disease. <i>Deutsches A&amp;#x0308;rzteblatt International</i> , <b>2013</b> , 110, 1-7, e1	2.5	70
451	Validation of the MDS research criteria for prodromal Parkinson's disease: Longitudinal assessment in a REM sleep behavior disorder (RBD) cohort. <i>Movement Disorders</i> , <b>2017</b> , 32, 865-873	7	69



450	Efficacy of rasagiline in patients with the parkinsonian variant of multiple system atrophy: a randomised, placebo-controlled trial. <i>Lancet Neurology, The</i> , <b>2015</b> , 14, 145-52	24.1	69
449	Functional relevance of ceruloplasmin mutations in Parkinson's disease. <i>FASEB Journal</i> , <b>2005</b> , 19, 1851-30.9		68
448	Movement disorder society criteria for clinically established early Parkinson's disease. <i>Movement Disorders</i> , <b>2018</b> , 33, 1643-1646	7	67
447	The role of high-field magnetic resonance imaging in parkinsonian disorders: Pushing the boundaries forward. <i>Movement Disorders</i> , <b>2017</b> , 32, 510-525	7	65
446	Iron as a therapeutic target for Parkinson's disease. <i>Movement Disorders</i> , <b>2018</b> , 33, 568-574	7	65
445	Microglia activation is related to substantia nigra echogenicity. <i>Journal of Neural Transmission</i> , <b>2010</b> , 117, 1287-92	4.3	64
444	Aging-related cortical reorganization of verbal fluency processing: a functional near-infrared spectroscopy study. <i>Neurobiology of Aging</i> , <b>2013</b> , 34, 439-50	5.6	63
443	Alpha-synuclein and Parkinson's disease: implications from the screening of more than 1,900 patients. <i>Movement Disorders</i> , <b>2005</b> , 20, 1191-4	7	63
442	Man-in-the-barrel syndrome caused by cervical spinal cord infarction. <i>Acta Neurologica Scandinavica</i> , <b>1998</b> , 97, 417-9	3.8	62
441	The modulation of Amyotrophic Lateral Sclerosis risk by ataxin-2 intermediate polyglutamine expansions is a specific effect. <i>Neurobiology of Disease</i> , <b>2012</b> , 45, 356-61	7.5	61
440	The new definition and diagnostic criteria of Parkinson's disease. <i>Lancet Neurology, The</i> , <b>2016</b> , 15, 546-8	24.1	61
439	Prodromal Parkinson's Disease: The Decade Past, the Decade to Come. <i>Movement Disorders</i> , <b>2019</b> , 34, 665-675	7	59
438	L-dopa increases $\beta$ synuclein DNA methylation in Parkinson's disease patients in vivo and in vitro. <i>Movement Disorders</i> , <b>2015</b> , 30, 1794-801	7	59
437	miRNA-based signatures in cerebrospinal fluid as potential diagnostic tools for early stage Parkinson's disease. <i>Oncotarget</i> , <b>2018</b> , 9, 17455-17465	3.3	59
436	Poor trail making test performance is directly associated with altered dual task prioritization in the elderly--baseline results from the TREND study. <i>PLoS ONE</i> , <b>2011</b> , 6, e27831	3.7	58
435	Impaired trunk stability in individuals at high risk for Parkinson's disease. <i>PLoS ONE</i> , <b>2012</b> , 7, e32240	3.7	58
434	Serum neuronal exosomes predict and differentiate Parkinson's disease from atypical parkinsonism. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2020</b> , 91, 720-729	5.5	57
433	Genome-wide association study in essential tremor identifies three new loci. <i>Brain</i> , <b>2016</b> , 139, 3163-3169	1.2	57



432	Gait analysis with wearables predicts conversion to parkinson disease. <i>Annals of Neurology</i> , <b>2019</b> , 86, 357-367	9.4	57
431	Reproducibility and diagnostic accuracy of substantia nigra sonography for the diagnosis of Parkinson's disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2010</b> , 81, 1087-92	5.5	57
430	Basal limbic system alteration in major depression: a hypothesis supported by transcranial sonography and MRI findings. <i>International Journal of Neuropsychopharmacology</i> , <b>2001</b> , 4, 21-31	5.8	57
429	Transcranial ultrasound in different monogenetic subtypes of Parkinson's disease. <i>Journal of Neurology</i> , <b>2007</b> , 254, 613-6	5.5	56
428	Subtypes of mild cognitive impairment in patients with Parkinson's disease: evidence from the LANDSCAPE study. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2016</b> , 87, 1099-105	5.5	56
427	Clinical and brain imaging characteristics in leucine-rich repeat kinase 2-associated PD and asymptomatic mutation carriers. <i>Movement Disorders</i> , <b>2011</b> , 26, 2335-42	7	55
426	Biomarker candidates of neurodegeneration in Parkinson's disease for the evaluation of disease-modifying therapeutics. <i>Journal of Neural Transmission</i> , <b>2012</b> , 119, 39-52	4.3	54
425	ATP13A2 variants in early-onset Parkinson's disease patients and controls. <i>Movement Disorders</i> , <b>2009</b> , 24, 2104-11	7	54
424	Assessment of idiopathic rapid-eye-movement sleep behavior disorder by transcranial sonography, olfactory function test, and FP-CIT-SPECT. <i>Movement Disorders</i> , <b>2008</b> , 23, 596-9	7	54
423	In vivo detection of iron and neuromelanin by transcranial sonography--a new approach for early detection of substantia nigra damage. <i>Journal of Neural Transmission</i> , <b>2006</b> , 113, 775-80	4.3	54
422	The correlation between ventricular diameter measured by transcranial sonography and clinical disability and cognitive dysfunction in patients with multiple sclerosis. <i>Archives of Neurology</i> , <b>2000</b> , 57, 1289-92		54
421	Application of the movement disorder society prodromal Parkinson's disease research criteria in 2 independent prospective cohorts. <i>Movement Disorders</i> , <b>2017</b> , 32, 1025-1034	7	53
420	Cognitive changes in prodromal Parkinson's disease: A review. <i>Movement Disorders</i> , <b>2017</b> , 32, 1655-1666		53
419	αSynuclein in Parkinson's disease: causal or bystander?. <i>Journal of Neural Transmission</i> , <b>2019</b> , 126, 815-840	4.3	53
418	Distinct metabolomic signature in cerebrospinal fluid in early parkinson's disease. <i>Movement Disorders</i> , <b>2017</b> , 32, 1401-1408	7	53
417	Characterizing POLG ataxia: clinics, electrophysiology and imaging. <i>Cerebellum</i> , <b>2012</b> , 11, 1002-11	4.3	53
416	Genetics and iron in the systems biology of Parkinson's disease and some related disorders. <i>Neurochemistry International</i> , <b>2013</b> , 62, 637-52	4.4	53
415	Identification of novel Angiogenin (ANG) gene missense variants in German patients with amyotrophic lateral sclerosis. <i>Journal of Neurology</i> , <b>2009</b> , 256, 1337-42	5.5	53

414	Motor dual-tasking deficits predict falls in Parkinson's disease: A prospective study. <i>Parkinsonism and Related Disorders</i> , <b>2016</b> , 26, 73-7	3.6	53
413	In-vivo evidence that high mobility group box 1 exerts deleterious effects in the 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine model and Parkinson's disease which can be attenuated by glycyrrhizin. <i>Neurobiology of Disease</i> , <b>2016</b> , 91, 59-68	7.5	52
412	Sonographic abnormalities of brainstem structures in restless legs syndrome. <i>Sleep Medicine</i> , <b>2008</b> , 9, 782-9	4.6	51
411	Predictive value of transcranial sonography in the diagnosis of Parkinson's disease. <i>Movement Disorders</i> , <b>2006</b> , 21, 1763-5	7	51
410	MicroRNA-101 Regulates Multiple Developmental Programs to Constrain Excitation in Adult Neural Networks. <i>Neuron</i> , <b>2016</b> , 92, 1337-1351	13.9	50
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406	Reduction of dyskinesia and induction of akinesia induced by morphine in two parkinsonian patients with severe sciatica. <i>Journal of Neural Transmission</i> , <b>1999</b> , 106, 725-8	4.3	49
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