Cyrus P Zabetian

List of Publications by Year
in descending order

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135 papers	13,090 citations	26630 56 h-index	24258 110 g-index
141 all docs	141 docs citations	141 times ranked	16591 citing authors

#	Article	IF	Citations
1	Blood extracellular vesicles carrying synaptic functionâ€•and brainâ€•related proteins as potential biomarkers for Alzheimer's disease. Alzheimer's and Dementia, 2023, 19, 909-923.	0.8	21
2	Polygenic risk prediction and SNCA haplotype analysis in a Latino Parkinson's disease cohort. Parkinsonism and Related Disorders, 2022, 102, 7-15.	2.2	2
3	Cognition as a mediator for gait and balance impairments in GBA-related Parkinsonâ€™s disease. Npj Parkinson's Disease, 2022, 8, .	5.3	1
4	Genomeâ€Wide Analysis of Copy Number Variation in Latin American Parkinson's Disease Patients. Movement Disorders, 2021, 36, 434-441.	3.9	12
5	Development of a Sensitive Diagnostic Assay for Parkinson Disease Quantifying Î±-Synucleinâ€“Containing Extracellular Vesicles. Neurology, 2021, 96, e2332-e2345.	1.1	18
6	Semantic fluency and processing speed are reduced in non-cognitively impaired participants with Parkinsonâ€™s disease. Journal of Clinical and Experimental Neuropsychology, 2021, 43, 469-480.	1.3	10
7	Genomewide Association Studies of <scp>LRRK2</scp> Modifiers of Parkinson's Disease. Annals of Neurology, 2021, 90, 76-88.	5.3	30
8	Characterizing the Genetic Architecture of Parkinson's Disease in Latinos. Annals of Neurology, 2021, 90, 353-365.	5.3	48
9	Exploring human-genome gut-microbiome interaction in Parkinsonâ€™s disease. Npj Parkinson's Disease, 2021, 7, 74.	5.3	15
10	Relationships Between Sensorimotor Inhibition and Mobility in Older Adults With and Without Parkinsonâ€™s Disease. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2021, 76, 630-637.	3.6	6
11	Novel compound heterozygous FBXO7 mutations in a family with early onset Parkinson's disease. Parkinsonism and Related Disorders, 2020, 80, 142-147.	2.2	8
12	Sensorimotor Inhibition and Mobility in Genetic Subgroups of Parkinson's Disease. Frontiers in Neurology, 2020, 11, 893.	2.4	3
13	Multivariate prediction of dementia in Parkinsonâ€™s disease. Npj Parkinson's Disease, 2020, 6, 20.	5.3	25
14	Characterizing dysbiosis of gut microbiome in PD: evidence for overabundance of opportunistic pathogens. Npj Parkinson's Disease, 2020, 6, 11.	5.3	140
15	Erythrocytic Î±-synuclein contained in microvesicles regulates astrocytic glutamate homeostasis: a new perspective on Parkinsonâ€™s disease pathogenesis. Acta Neuropathologica Communications, 2020, 8, 102.	5.2	26
16	Diagnostic prediction model for levodopa-induced dyskinesia in Parkinsonâ€™s disease. Arquivos De Neuro-Psiquiatria, 2020, 78, 206-216.	0.8	10
17	Cognitive associations with comprehensive gait and static balance measures in Parkinson's disease. Parkinsonism and Related Disorders, 2019, 69, 104-110.	2.2	41
18	New windows into the brain: Central nervous system-derived extracellular vesicles in blood. Progress in Neurobiology, 2019, 175, 96-106.	5.7	121

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19	The distribution and risk effect of GBA variants in a large cohort of PD patients from Colombia and Peru. Parkinsonism and Related Disorders, 2019, 63, 204-208.	2.2	31
20	Sex differences in progression to mild cognitive impairment and dementia in Parkinson's disease. Parkinsonism and Related Disorders, 2018, 50, 29-36.	2.2	94
21	Stool Immune Profiles Evince Gastrointestinal Inflammation in Parkinson's Disease. Movement Disorders, 2018, 33, 793-804.	3.9	130
22	The effect of LRRK2 mutations on the cholinergic system in manifest and premanifest stages of Parkinson's disease: a cross-sectional PET study. Lancet Neurology, The, 2018, 17, 309-316.	10.2	57
23	Association of a neuronal nitric oxide synthase gene polymorphism with levodopa-induced dyskinesia in Parkinson's disease. Nitric Oxide - Biology and Chemistry, 2018, 74, 86-90.	2.7	8
24	Diagnostic Validation for Participants in the Washington State Parkinson Disease Registry. Parkinson's Disease, 2018, 2018, 1-6.	1.1	17
25	Plasticity-related gene 3 (LPPR1) and age at diagnosis of Parkinson disease. Neurology: Genetics, 2018, 4, e271.	1.9	12
26	Detecting Mild Cognitive Deficits in <scp>P</scp>arkinson's <scp>D</scp>isease: <scp>C</scp>omparison of <scp>N</scp>europsychological <scp>T</scp>ests. Movement Disorders, 2018, 33, 1750-1759.	3.9	42
27	Intrafamilial variable phenotype including corticobasal syndrome in a family with p.P301L mutation in the MAPT gene: first report in South America. Neurobiology of Aging, 2017, 53, 195.e11-195.e17.	3.1	9
28	Parkinson's disease and Parkinson's disease medications have distinct signatures of the gut microbiome. Movement Disorders, 2017, 32, 739-749.	3.9	649
29	Large-scale exploratory genetic analysis of cognitive impairment in Parkinson's disease. Neurobiology of Aging, 2017, 56, 211.e1-211.e7.	3.1	37
30	Serotonin and dopamine transporter PET changes in the premotor phase of LRRK2 parkinsonism: cross-sectional studies. Lancet Neurology, The, 2017, 16, 351-359.	10.2	96
31	Neuropathological and genetic correlates of survival and dementia onset in synucleinopathies: a retrospective analysis. Lancet Neurology, The, 2017, 16, 55-65.	10.2	394
32	Homocysteine and cognitive function in Parkinson's disease. Parkinsonism and Related Disorders, 2017, 44, 1-5.	2.2	44
33	LARGEâ€PD: Examining the genetics of Parkinson's disease in Latin America. Movement Disorders, 2017, 32, 1330-1331.	3.9	34
34	Variable frequency of LRRK2 variants in the Latin American research consortium on the genetics of Parkinsonâ€™s disease (LARGE-PD), a case of ancestry. Npj Parkinson's Disease, 2017, 3, 19.	5.3	28
35	Some aspects of the validity of the Montreal Cognitive Assessment (MoCA)for evaluating cognitive impairment in Brazilian patients with Parkinson's disease. Dementia E Neuropsychologia, 2016, 10, 333-338.	0.8	18
36	Response to the letter â€œHaptoglobin phenotype and Parkinson disease riskâ€•by Delanghe etÂal.. Parkinsonism and Related Disorders, 2016, 22, 110-111.	2.2	3

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37	CNS tau efflux via exosomes is likely increased in Parkinson's disease but not in Alzheimer's disease. Alzheimer's and Dementia, 2016, 12, 1125-1131.	0.8	154
38	The Clinical Profile of GBA-Related Lewy Body Disorders. JAMA Neurology, 2016, 73, 1403.	9.0	1
39	Arguing against the proposed definition changes of PD. Movement Disorders, 2016, 31, 1619-1622.	3.9	55
40	Association ofGBAMutations and the E326K Polymorphism With Motor and Cognitive Progression in Parkinson Disease. JAMA Neurology, 2016, 73, 1217.	9.0	185
41	Identification of genetic modifiers of age-at-onset for familial Parkinsonâ€™s disease. Human Molecular Genetics, 2016, 25, 3849-3862.	2.9	44
42	The discovery of LRRK2 p.R1441S, a novel mutation for Parkinson's disease, adds to the complexity of a mutational hotspot. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2016, 171, 925-930.	1.7	17
43	GBA Variants are associated with a distinct pattern of cognitive deficits in <scp>P</scp>arkinson's disease. Movement Disorders, 2016, 31, 95-102.	3.9	158
44	Precision Medicine. American Journal of Pathology, 2016, 186, 500-506.	3.8	49
45	Review ofRosenbergâ€™s Molecular and Genetic Basis of Neurological and Psychiatric Disease, 5th ed. JAMA Neurology, 2015, 72, 1538.	9.0	0
46	Screening of cognitive impairment in patients with Parkinson's disease: diagnostic validity of the Brazilian versions of the Montreal Cognitive Assessment and the Addenbrooke's Cognitive Examination-Revised. Arquivos De Neuro-Psiquiatria, 2015, 73, 929-933.	0.8	25
47	A Peruvian family with a novel PARK2 mutation: Clinical andÂpathological characteristics. Parkinsonism and Related Disorders, 2015, 21, 444-448.	2.2	20
48	Haptoglobin phenotype modifies serum iron levels and the effect of smoking on Parkinson disease risk. Parkinsonism and Related Disorders, 2015, 21, 1087-1092.	2.2	29
49	Phosphorylated Î±-synuclein in Parkinsonâ€™s disease: correlation depends on disease severity. Acta Neuropathologica Communications, 2015, 3, 7.	5.2	74
50	Cerebrospinal Fluid Peptides as Potential Parkinson Disease Biomarkers: A Staged Pipeline for Discovery and Validation*. Molecular and Cellular Proteomics, 2015, 14, 544-555.	3.8	51
51	Lower plasma apolipoprotein A1 levels are found in Parkinson's disease and associate with apolipoprotein A1 genotype. Movement Disorders, 2015, 30, 805-812.	3.9	37
52	Glutamate Receptor Gene GRIN2A, Coffee, and Parkinson Disease. PLoS Genetics, 2014, 10, e1004774.	3.5	7
53	People with Parkinson's disease and normal MMSE score have a broad range of cognitive performance. Movement Disorders, 2014, 29, 1258-1264.	3.9	76
54	APOE,MAPT, andSNCAGenes and Cognitive Performance in Parkinson Disease. JAMA Neurology, 2014, 71, 1405.	9.0	172

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55	Identification of a novel Parkinsonâ€™s disease locus via stratified genome-wide association study. BMC Genomics, 2014, 15, 118.	2.8	53
56	Targeted Discovery and Validation of Plasma Biomarkers of Parkinsonâ€™s Disease. Journal of Proteome Research, 2014, 13, 4535-4545.	3.7	30
57	Evaluation of mild cognitive impairment subtypes in Parkinson's disease. Movement Disorders, 2014, 29, 756-764.	3.9	53
58	Plasma exosomal Î±-synuclein is likely CNS-derived and increased in Parkinsonâ€™s disease. Acta Neuropathologica, 2014, 128, 639-650.	7.7	504
59	Association mapping of the PARK10 region for Parkinson's disease susceptibility genes. Parkinsonism and Related Disorders, 2014, 20, 93-98.	2.2	16
60	Identification of a Japanese family with LRRK2 p.R1441G-related Parkinson's disease. Neurobiology of Aging, 2014, 35, 2656.e17-2656.e23.	3.1	24
61	Cheek cellâ€“derived Î±-synuclein and DJ-1 do not differentiate Parkinson's disease from control. Neurobiology of Aging, 2014, 35, 418-420.	3.1	30
62	Clinical and Biochemical Differences in Patients Having Parkinson Disease With vs WithoutGBAMutations. JAMA Neurology, 2013, 70, 852.	9.0	115
63	C9orf72 Hexanucleotide Repeat Expansion and Guam Amyotrophic Lateral Sclerosisâ€“Parkinsonism-Dementia Complex. JAMA Neurology, 2013, 70, 742.	9.0	22
64	Association of Parkinson Disease with Structural and Regulatory Variants in the HLA Region. American Journal of Human Genetics, 2013, 93, 984-993.	6.2	145
65	APOE Ïµ4 Increases Risk for Dementia in Pure Synucleinopathies. JAMA Neurology, 2013, 70, 223.	9.0	302
66	Neuropsychologic assessment in collaborative Parkinson's disease research: A proposal from the National Institute of Neurological Disorders and Stroke Morris K. Udall Centers of Excellence for Parkinson's Disease Research at the University of Pennsylvania and theÂUniversity of Washington. Alzheimer's and Dementia, 2013, 9, 609-614.	0.8	24
67	Plasma apolipoprotein A1 as a biomarker for Parkinson disease. Annals of Neurology, 2013, 74, 119-127.	5.3	116
68	Altered splicing of ATP6AP2 causes X-linked parkinsonism with spasticity (XPDS). Human Molecular Genetics, 2013, 22, 3259-3268.	2.9	113
69	Novel <scp>Lrrk2</scp>â€p.<scp>S1761R</scp> mutation is not a common cause of Parkinson's disease in Spain. Movement Disorders, 2013, 28, 248-248.	3.9	1
70	Pacific Northwest Udall Center of Excellence Clinical Consortium: Study Design and Baseline Cohort Characteristics. Journal of Parkinson's Disease, 2013, 3, 205-214.	2.8	64
71	Comprehensive Research Synopsis and Systematic Meta-Analyses in Parkinson's Disease Genetics: The PDGene Database. PLoS Genetics, 2012, 8, e1002548.	3.5	495
72	Phosphorylated Î±-Synuclein in Parkinsonâ€™s Disease. Science Translational Medicine, 2012, 4, 121ra20.	12.4	223

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73	Common variation in the LRRK2 gene is a risk factor for Parkinson's disease. Movement Disorders, 2012, 27, 1823-1826.	3.9	14
74	DJ-1 and Î±SYN in LRRK2 CSF do not correlate with striatal dopaminergic function. Neurobiology of Aging, 2012, 33, 836.e5-836.e7.	3.1	34
75	GBA mutations increase risk for Lewy body disease with and without Alzheimer disease pathology. Neurology, 2012, 79, 1944-1950.	1.1	138
76	Metaâ€analysis of Parkinson's Disease: Identification of a novel locus, RIT2. Annals of Neurology, 2012, 71, 370-384.	5.3	264
77	Genome-Wide Gene-Environment Study Identifies Glutamate Receptor Gene GRIN2A as a Parkinson's Disease Modifier Gene via Interaction with Coffee. PLoS Genetics, 2011, 7, e1002237.	3.5	206
78	Complement 3 and Factor H in Human Cerebrospinal Fluid in Parkinson's Disease, Alzheimer's Disease, and Multiple-System Atrophy. American Journal of Pathology, 2011, 178, 1509-1516.	3.8	97
79	Cerebrospinal fluid biomarkers and cognitive performance in non-demented patients with Parkinsonâ€™s disease. Parkinsonism and Related Disorders, 2011, 17, 61-64.	2.2	64
80	The UCHL1 S18Y polymorphism and Parkinsonâ€™s disease in a Japanese population. Parkinsonism and Related Disorders, 2011, 17, 473-475.	2.2	5
81	Lrrk2 p.Q1111H substitution and Parkinsonâ€™s disease in Latin America. Parkinsonism and Related Disorders, 2011, 17, 629-631.	2.2	15
82	Evidence for More than One Parkinson's Disease-Associated Variant within the HLA Region. PLoS ONE, 2011, 6, e27109.	2.5	60
83	Replication of MAPT and SNCA, but not PARK16â€18, as susceptibility genes for Parkinson's disease. Movement Disorders, 2011, 26, 819-823.	3.9	64
84	Diseaseâ€related and genetic correlates of psychotic symptoms in Parkinson's disease. Movement Disorders, 2011, 26, 2190-2195.	3.9	61
85	Cerebrospinal fluid biomarkers for Parkinson disease diagnosis and progression. Annals of Neurology, 2011, 69, 570-580.	5.3	371
86	Salivary Î±-synuclein and DJ-1: potential biomarkers for Parkinson's disease. Brain, 2011, 134, e178-e178.	7.6	196
87	Postural instability/gait disturbance in Parkinson's disease has distinct subtypes: an exploratory analysis. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 564-568.	1.9	90
88	Visualizing disease associations: graphic analysis of frequency distributions as a function of age using moving average plots (MAP) with application to Alzheimer's and Parkinson's disease. Genetic Epidemiology, 2010, 34, 92-99.	1.3	6
89	Glycoproteomics in neurodegenerative diseases. Mass Spectrometry Reviews, 2010, 29, 79-125.	5.4	99
90	A novel Xâ€linked fourâ€repeat tauopathy with Parkinsonism and spasticity. Movement Disorders, 2010, 25, 1409-1417.	3.9	20

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91	CSF AÎ²₄₂ and tau in Parkinson's disease with cognitive impairment. Movement Disorders, 2010, 25, 2682-2685.	3.9	162
92	Common genetic variation in the HLA region is associated with late-onset sporadic Parkinson's disease. Nature Genetics, 2010, 42, 781-785.	21.4	692
93	SNCA Variant Associated With Parkinson Disease and Plasma Î±-Synuclein Level. Archives of Neurology, 2010, 67, 1350-6.	4.5	157
94	DJ-1 and Î±-synuclein in human cerebrospinal fluid as biomarkers of Parkinsonâ€™s disease. Brain, 2010, 133, 713-726.	7.6	575
95	Significance and confounders of peripheral DJ-1 and alpha-synuclein in Parkinson's disease. Neuroscience Letters, 2010, 480, 78-82.	2.1	184
96	Association Between the Ubiquitin Carboxyl-Terminal Esterase L1 Gene (UCHL1) S18Y Variant and Parkinson's Disease: A HuGE Review and Meta-Analysis. American Journal of Epidemiology, 2009, 170, 1344-1357.	3.4	67
97	LRRK2 mutations and risk variants in Japanese patients with Parkinson's disease. Movement Disorders, 2009, 24, 1034-1041.	3.9	60
98	Ashkenazi Parkinsonâ€™s disease patients with the LRRK2 G2019S mutation share a common founder dating from the second to fifth centuries. Neurogenetics, 2009, 10, 355-358.	1.4	23
99	Lrrk2 R1441G-related Parkinsonâ€™s disease: evidence of a common founding event in the seventh century in Northern Spain. Neurogenetics, 2009, 10, 347-353.	1.4	41
100	Haplotype analysis of Lrrk2 R1441H carriers with parkinsonism. Parkinsonism and Related Disorders, 2009, 15, 466-467.	2.2	31
101	LRRK2 mutations in patients with Parkinson's disease from Peru and Uruguay. Parkinsonism and Related Disorders, 2009, 15, 370-373.	2.2	45
102	Genetic Polymorphism at Codon 129 of the Prion Protein Gene Is Not Associated With Multiple Sclerosis. Archives of Neurology, 2009, 66, 280-1.	4.5	4
103	Lack of evidence for an association between UCHL1 S18Y and Parkinsonâ€™s disease. European Journal of Neurology, 2008, 15, 134-139.	3.3	25
104	Exploring gene-environment interactions in Parkinsonâ€™s disease. Human Genetics, 2008, 123, 257-265.	3.8	92
105	Combined effects of smoking, coffee, and NSAIDs on Parkinson's disease risk. Movement Disorders, 2008, 23, 88-95.	3.9	129
106	Genetic association between Î±â€synuclein and idiopathic parkinson's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 1222-1230.	1.7	60
107	Phenotype, genotype, and worldwide genetic penetrance of LRRK2-associated Parkinson's disease: a case-control study. Lancet Neurology, The, 2008, 7, 583-590.	10.2	1,340
108	Application of Targeted Quantitative Proteomics Analysis in Human Cerebrospinal Fluid Using a Liquid Chromatography Matrix-Assisted Laser Desorption/Ionization Time-of-Flight Tandem Mass Spectrometer (LC MALDI TOF/TOF) Platform. Journal of Proteome Research, 2008, 7, 720-730.	3.7	67

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109	Glucocerebrosidase Gene Mutations. Archives of Neurology, 2008, 65, 379-82.	4.5	188
110	Heterozygous parkin point mutations are as common in control subjects as in Parkinson's patients. Annals of Neurology, 2007, 61, 47-54.	5.3	105
111	DBH â^'1021Câ†'T does not modify risk or age at onset in Parkinson's disease. Annals of Neurology, 2007, 62, 99-101.	5.3	7
112	Association analysis of MAPT H1 haplotype and subhaplotypes in Parkinson's disease. Annals of Neurology, 2007, 62, 137-144.	5.3	129
113	Genotypic and haplotypic associations of the DBH gene with plasma dopamine Î²-hydroxylase activity in African Americans. European Journal of Human Genetics, 2007, 15, 878-883.	2.8	43
114	Validity and Utility of a LRRK2 G2019S Mutation Test for the Diagnosis of Parkinson's Disease. Genetic Testing and Molecular Biomarkers, 2006, 10, 221-227.	1.7	32
115	LRRK2 G2019S in Families with Parkinson Disease Who Originated from Europe and the Middle East: Evidence of Two Distinct Founding Events Beginning Two Millennia Ago. American Journal of Human Genetics, 2006, 79, 752-758.	6.2	111
116	A Single Nucleotide Polymorphism at DBH, Possibly Associated with Attention-Deficit/Hyperactivity Disorder, Associates with Lower Plasma Dopamine Î²-Hydroxylase Activity and is in Linkage Disequilibrium with Two Putative Functional Single Nucleotide Polymorphisms. Biological Psychiatry, 2006, 60, 1034-1038.	1.3	42
117	Lack of replication of thirteen single-nucleotide polymorphisms implicated in Parkinson's disease: a large-scale international study. Lancet Neurology, The, 2006, 5, 917-923.	10.2	83
118	Parkinson's disease andLRRK2: Frequency of a common mutation in U.S. movement disorder clinics. Movement Disorders, 2006, 21, 519-523.	3.9	84
119	Clinical Features of Parkinson Disease Patients With Homozygous Leucine-Rich Repeat Kinase 2 G2019S Mutations. Archives of Neurology, 2006, 63, 1250.	4.5	91
120	Analysis of the LRRK2 G2019S Mutation in Alzheimer Disease. Archives of Neurology, 2006, 63, 156.	4.5	21
121	Haplotypeâ€controlled analysis of the association of a nonâ€synonymous single nucleotide polymorphism at DBH (+â€‰1603Câ€‰â†'â€‰T) with plasma dopamine Î²â€hydroxylase activity. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 139B, 88-90.	3.7	31
122	Escaping Parkinson's disease: A neurologically healthy octogenarian with theLRRK2 G2019S mutation. Movement Disorders, 2005, 20, 1077-1078.	3.9	73
123	Variations in the dopamine ?-hydroxylase gene are not associated with the autonomic disorders, pure autonomic failure, or multiple system atrophy. American Journal of Medical Genetics Part A, 2003, 120A, 234-236.	2.4	13
124	A revised allele frequency estimate and haplotype analysis of the DBH deficiency mutation IVS1+2Tâ€‰â†'â€‰C in African- and European-Americans. , 2003, 123A, 190-192.		11
125	The Structure of Linkage Disequilibrium at the DBH Locus Strongly Influences the Magnitude of Association between Diallelic Markers and Plasma Dopamine Î²-Hydroxylase Activity. American Journal of Human Genetics, 2003, 72, 1389-1400.	6.2	81
126	Genotype-controlled analysis of plasma dopamine Î²-hydroxylase activity in psychotic unipolar major depression. Biological Psychiatry, 2002, 51, 358-364.	1.3	58

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127	A genotype-controlled analysis of plasma dopamine Î²-hydroxylase in healthy and alcoholic subjects: evidence for alcohol-related differences in noradrenergic function. Biological Psychiatry, 2002, 52, 1151-1158.	1.3	75
128	Mutations in the dopamine ?-hydroxylase gene are associated with human norepinephrine deficiency. American Journal of Medical Genetics Part A, 2002, 108, 140-147.	2.4	88
129	Molecular Genetic Analysis of Plasma Dopamine Î²-Hydroxylase in Depression. Advances in Behavioral Biology, 2002, , 423-426.	0.2	0
130	A Quantitative-Trait Analysis of Human Plasmaâ€“Dopamine Î²-Hydroxylase Activity: Evidence for a Major Functional Polymorphism at the DBH Locus. American Journal of Human Genetics, 2001, 68, 515-522.	6.2	253
131	Functional variants atCYP2A6: New genotyping methods, population genetics, and relevance to studies of tobacco dependence. American Journal of Medical Genetics Part A, 2000, 96, 638-645.	2.4	16
132	Effects of Environmental Salinity on Pituitary Growth Hormone Content and Cell Activity in the Euryhaline Tilapia, Oreochromis mossambicus. General and Comparative Endocrinology, 1994, 95, 483-494.	1.8	49
133	[3H]-(+)-pentazocine binding to sigma recognition sites in human cerebellum. Life Sciences, 1994, 55, PL389-PL395.	4.3	18
134	Sigma receptors are associated with cortical limbic areas in the primate brain. Synapse, 1992, 12, 195-205.	1.2	70
135	Osteology and Interrelationships of the Sand Lances (Teleostei: Ammodytidae). Copeia, 1990, 1990, 78.	1.3	28