

The ubiquitin pathway in Parkinson's disease

Nature

395, 451-452

DOI: [10.1038/26652](https://doi.org/10.1038/26652)

Citation Report

#	ARTICLE	IF	CITATIONS
1	Maladie de Parkinson Vers un mécanisme de mort neuronale. Journal of Engineering and Technology Management - JET-M, 1997, 14, 25-45.	1.4	65
2	Intron-exon Structure of Ubiquitin C-terminal Hydrolase-L1. DNA Research, 1998, 5, 397-400.	1.5	25
3	Gene therapy for Parkinson's disease: review and update. Expert Opinion on Investigational Drugs, 1999, 8, 1551-1564.	1.9	1
4	Cluster headaches. Neurology, 1999, 53, 543-543.	1.5	31
5	Etiology of Parkinson's Disease. Canadian Journal of Neurological Sciences, 1999, 26, S5-S12.	0.3	23
6	Aberrant Protein Deposition and Neurological Disease. Journal of Biological Chemistry, 1999, 274, 37507-37510.	1.6	107
7	A chromosome 4p haplotype segregating with Parkinson's disease and postural tremor. Human Molecular Genetics, 1999, 8, 81-85.	1.4	229
8	Role of Mitochondria in Parkinson Disease. Biological Chemistry, 1999, 380, 865-70.	1.2	63
9	Evidence for Proteasome Involvement in Polyglutamine Disease: Localization to Nuclear Inclusions in SCA3/MJD and Suppression of Polyglutamine Aggregation in vitro. Human Molecular Genetics, 1999, 8, 673-682.	1.4	386
10	Candidate genes and Parkinson's disease. Neurology, 1999, 53, 1382-1382.	1.5	24
11	Axonal transport of synucleins is mediated by all rate components. European Journal of Neuroscience, 1999, 11, 3369-3376.	1.2	94
12	Intragenic deletion in the gene encoding ubiquitin carboxy-terminal hydrolase in gad mice. Nature Genetics, 1999, 23, 47-51.	9.4	467
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17	Rapid-onset dystonia-parkinsonism: Linkage to chromosome 19q13. Annals of Neurology, 1999, 46, 176-182.	2.8	97
18	Tremor arrest with thalamic microinjections of muscimol in patients with essential tremor. Annals of Neurology, 1999, 46, 249-252.	2.8	93

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19	Lymphocyte migration and multiple sclerosis: Relation with disease course and therapy. <i>Annals of Neurology</i> , 1999, 46, 253-256.	2.8	56
20	Phosphorus and proton magnetic resonance spectroscopy in episodic ataxia type 2. <i>Annals of Neurology</i> , 1999, 46, 256-259.	2.8	49
21	Increased tissue copper and manganese content in the lentiform nucleus in primary adult-onset dystonia. <i>Annals of Neurology</i> , 1999, 46, 260-263.	2.8	91
22	Selective suppression of cerebellar GABAergic transmission by an autoantibody to glutamic acid decarboxylase. <i>Annals of Neurology</i> , 1999, 46, 263-267.	2.8	109
23	Cytomegalovirus is not associated with IgM anti-myelin-associated glycoprotein/sulphate-3-glucuronyl paragloboside antibody-associated neuropathy. <i>Annals of Neurology</i> , 1999, 46, 267-270.	2.8	7
24	Ataxin 1 and ataxin 3 in neuronal intranuclear inclusion disease. <i>Annals of Neurology</i> , 1999, 46, 271-273.	2.8	46
25	Muscle phosphoglycerate mutase deficiency with tubular aggregates: Effect of dantrolene. <i>Annals of Neurology</i> , 1999, 46, 274-277.	2.8	29
26	Reduced expression of the G209A α -synuclein allele in familial parkinsonism. <i>Annals of Neurology</i> , 1999, 46, 374-381.	2.8	89
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61	Subhypnotic doses of zolpidem oppose dopaminergic-induced dyskinesia in Parkinson's disease. <i>Movement Disorders</i> , 2000, 15, 734-735.	2.2	38
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149	Expression of A53T Mutant But Not Wild-Type α -Synuclein in PC12 Cells Induces Alterations of the Ubiquitin-Dependent Degradation System, Loss of Dopamine Release, and Autophagic Cell Death. <i>Journal of Neuroscience</i> , 2001, 21, 9549-9560.	1.7	540
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151	No genetic association of the Ubiquitin Carboxy-terminal Hydrolase-L1 gene S18Y polymorphism with familial Parkinson's disease. <i>Journal of Neural Transmission</i> , 2001, 108, 979-984.	1.4	39
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