

Glucocerebrosidase mutations in subjects with parkinson

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Parkinsonism among Gaucher disease carriers. <i>Journal of Medical Genetics</i> , 2004, 41, 937-940.	1.5	320
2	Gaucher disease: complexity in a "simple" disorder. <i>Molecular Genetics and Metabolism</i> , 2004, 83, 6-15.	0.5	350
3	Molecular genetic pathways in Parkinson's disease: a review. <i>Clinical Science</i> , 2005, 109, 355-364.	1.8	37
4	Gaucher disease: pathological mechanisms and modern management. <i>British Journal of Haematology</i> , 2005, 129, 178-188.	1.2	240
5	The emerging field of lipidomics. <i>Nature Reviews Drug Discovery</i> , 2005, 4, 594-610.	21.5	1,104
6	Analysis of the glucocerebrosidase gene in Parkinson's disease. <i>Movement Disorders</i> , 2005, 20, 367-370.	2.2	107
7	Pilot association study of the Î²-glucocerebrosidase N370S allele and Parkinson's disease in subjects of Jewish ethnicity. <i>Movement Disorders</i> , 2005, 20, 100-103.	2.2	93
8	Neurodegenerative Disorders: The Role of Genetic Factors in Their Origin and the Efficiency of Treatment. <i>Human Physiology</i> , 2005, 31, 472-482.	0.1	3
9	Study of Multimodal Evoked Potentials in Patients With Type 1 Gaucher's Disease. <i>Journal of Child Neurology</i> , 2005, 20, 124-128.	0.7	15
10	The Glucocerebrosidase Gene and Parkinson's Disease in Ashkenazi Jews. <i>New England Journal of Medicine</i> , 2005, 352, 728-731.	13.9	90
11	LRRK2 gene in Parkinson disease: Mutation analysis and case control association study. <i>Neurology</i> , 2005, 65, 696-700.	1.5	160
12	Mutations in the glucocerebrosidase gene and Parkinson disease: Phenotype "genotype" correlation. <i>Neurology</i> , 2005, 65, 1460-1461.	1.5	74
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14	Gaucher disease and parkinsonism. <i>Molecular Genetics and Metabolism</i> , 2005, 84, 302-304.	0.5	75
15	Nosology of Parkinson's Disease: Looking for the Way Out of a Quackmire. <i>Neuron</i> , 2005, 47, 479-482.	3.8	87
16	A Chaperone-Mediated Approach to Enzyme Enhancement as a Therapeutic Option for the Lysosomal Storage Disorders. <i>Drugs in R and D</i> , 2006, 7, 339-348.	1.1	19
17	Glucocerebrosidase mutations are an important risk factor for Lewy body disorders. <i>Neurology</i> , 2006, 67, 908-910.	1.5	204
19	From genes to systems: New global strategies for the characterization of NCL biology. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2006, 1762, 934-944.	1.8	14

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20	Increased incidence of Parkinson disease among relatives of patients with Gaucher disease. <i>Blood Cells, Molecules, and Diseases</i> , 2006, 36, 426-428.	0.6	118
21	Detection of 12 new mutations in Gaucher disease Brazilian patients. <i>Blood Cells, Molecules, and Diseases</i> , 2006, 37, 204-209.	0.6	22
22	Glucocerebrosidase mutations are not found in association with LRRK2 G2019S in subjects with parkinsonism. <i>Neuroscience Letters</i> , 2006, 404, 163-165.	1.0	14
23	Heterozygosity for a Mendelian disorder as a risk factor for complex disease. <i>Clinical Genetics</i> , 2006, 70, 275-282.	1.0	53
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25	Self-stimulatory behavior associated with deep brain stimulation in Parkinson's disease. <i>Movement Disorders</i> , 2006, 21, 283-285.	2.2	34
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29	Parkinsons Disease: Genetics and Beyond. <i>Current Neuropharmacology</i> , 2007, 5, 99-113.	1.4	28
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34	Glucocerebrosidase mutations in Chinese subjects from Taiwan with sporadic Parkinson disease. <i>Molecular Genetics and Metabolism</i> , 2007, 91, 195-200.	0.5	111
36	GM1 Specifically Interacts with α -Synuclein and Inhibits Fibrillation. <i>Biochemistry</i> , 2007, 46, 1868-1877.	1.2	239
37	Lysosomal hydrolases in cerebrospinal fluid from subjects with Parkinson's disease. <i>Movement Disorders</i> , 2007, 22, 1481-1484.	2.2	103
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39	Movement and mood disorder in two brothers with Gaucher disease. <i>Clinical Genetics</i> , 2007, 72, 357-361.	1.0	10

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41	Glucocerebrosidase gene mutations are associated with Parkinson's disease in southern Italy. <i>Movement Disorders</i> , 2008, 23, 460-463.	2.2	83
42	Emerging pathways in genetic Parkinson's disease: Potential role of ceramide metabolism in Lewy body disease. <i>FEBS Journal</i> , 2008, 275, 5767-5773.	2.2	121
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56	MUTATIONS IN <i>GBA</i> ARE ASSOCIATED WITH FAMILIAL PARKINSON DISEASE SUSCEPTIBILITY AND AGE AT ONSET. <i>Neurology</i> , 2009, 73, 1424-1426.	1.5	142
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58	Recommendations on Diagnosis, Treatment, and Monitoring for Gaucher Disease. <i>Journal of Pediatrics</i> , 2009, 155, S10-S18.	0.9	45

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60	The association between mutations in the lysosomal protein glucocerebrosidase and parkinsonism. <i>Movement Disorders</i> , 2009, 24, 1571-1578.	2.2	71
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67	From Genes to Proteins in Mendelian Parkinson's Disease: An Overview. <i>Anatomical Record</i> , 2009, 292, 1893-1901.	0.8	14
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73	Glucosidase-beta variations and Lewy body disorders. <i>Parkinsonism and Related Disorders</i> , 2009, 15, 414-416.	1.1	36
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95	Facial tic associated with lamotrigine in adults. Movement Disorders, 2010, 25, 1512-1513.	2.2	10
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99	Changes in apraxia after deep brain stimulation of the nucleus basalis Meynert in a patient with Parkinson dementia syndrome. <i>Movement Disorders</i> , 2010, 25, 1519-1520.	2.2	43
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142	A DNA resequencing array for genes involved in Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2012, 18, 386-390.	1.1	7
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146	Glucocerebrosidase Involvement in Parkinson Disease and Other Synucleinopathies. <i>Frontiers in Neurology</i> , 2012, 3, 65.	1.1	15
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158	Unfolded protein response in Gaucher disease: from human to <i>Drosophila</i> . <i>Orphanet Journal of Rare Diseases</i> , 2013, 8, 140.	1.2	88
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165	Factors Influencing the Measurement of Lysosomal Enzymes Activity in Human Cerebrospinal Fluid. <i>PLoS ONE</i> , 2014, 9, e101453.	1.1	23
166	Glucocerebrosidase Gene Mutations Associated with Parkinson's Disease: A Meta-Analysis in a Chinese population. <i>PLoS ONE</i> , 2014, 9, e115747.	1.1	32
167	Original article The key role of sphingosine kinases in the molecular mechanism of neuronal cell survival and death in an experimental model of Parkinson's disease. <i>Folia Neuropathologica</i> , 2014, 3, 260-269.	0.5	51
168	The genetics of Parkinson's disease: review of current and emerging candidates. <i>Journal of Parkinsonism and Restless Legs Syndrome</i> , 2014, , 63.	0.8	1
169	Multiple System Atrophy: Genetic or Epigenetic?. <i>Experimental Neurobiology</i> , 2014, 23, 277-291.	0.7	28
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