

Glucocerebrosidase mutations in subjects with parkinson

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Parkinsonism among Gaucher disease carriers. Journal of Medical Genetics, 2004, 41, 937-940.	3.2	320
2	Gaucher disease: complexity in a "simple" disorder. Molecular Genetics and Metabolism, 2004, 83, 6-15.	1.1	350
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4	Gaucher disease: pathological mechanisms and modern management. British Journal of Haematology, 2005, 129, 178-188.	2.5	240
5	The emerging field of lipidomics. Nature Reviews Drug Discovery, 2005, 4, 594-610.	46.4	1,104
6	Analysis of the glucocerebrosidase gene in Parkinson's disease. Movement Disorders, 2005, 20, 367-370.	3.9	107
7	Pilot association study of the Î²-glucocerebrosidase N370S allele and Parkinson's disease in subjects of Jewish ethnicity. Movement Disorders, 2005, 20, 100-103.	3.9	93
8	Neurodegenerative Disorders: The Role of Genetic Factors in Their Origin and the Efficiency of Treatment. Human Physiology, 2005, 31, 472-482.	0.4	3
9	Study of Multimodal Evoked Potentials in Patients With Type 1 Gaucher's Disease. Journal of Child Neurology, 2005, 20, 124-128.	1.4	15
10	The Glucocerebrosidase Gene and Parkinson's Disease in Ashkenazi Jews. New England Journal of Medicine, 2005, 352, 728-731.	27.0	90
11	<i>LRRK2</i> gene in Parkinson disease. Neurology, 2005, 65, 696-700.	1.1	160
12	Mutations in the glucocerebrosidase gene and Parkinson disease: Phenotype "genotype correlation". Neurology, 2005, 65, 1460-1461.	1.1	74
13	Hematologically important mutations: Gaucher disease. Blood Cells, Molecules, and Diseases, 2005, 35, 355-364.	1.4	154
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19	From genes to systems: New global strategies for the characterization of NCL biology. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2006, 1762, 934-944.	3.8	14

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21	Detection of 12 new mutations in Gaucher disease Brazilian patients. Blood Cells, Molecules, and Diseases, 2006, 37, 204-209.	1.4	22
22	Glucocerebrosidase mutations are not found in association with LRRK2 G2019S in subjects with parkinsonism. Neuroscience Letters, 2006, 404, 163-165.	2.1	14
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24	Glucocerebrosidase mutations are also found in subjects with early-onset parkinsonism from Venezuela. Movement Disorders, 2006, 21, 282-283.	3.9	64
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28	Lewy bodies. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 1661-1668.	7.1	401
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60	The association between mutations in the lysosomal protein glucocerebrosidase and parkinsonism. <i>Movement Disorders</i> , 2009, 24, 1571-1578.	3.9	71
61	Parkinsonism in Gaucher's disease type 1: Ten new cases and a review of the literature. <i>Movement Disorders</i> , 2009, 24, 1524-1530.	3.9	28
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65	Tremor on smiling. <i>Movement Disorders</i> , 2009, 24, 1542-1545.	3.9	9
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73	Glucosidase-beta variations and Lewy body disorders. <i>Parkinsonism and Related Disorders</i> , 2009, 15, 414-416.	2.2	36
74	Complete screening for glucocerebrosidase mutations in Parkinson disease patients from Greece. <i>Neuroscience Letters</i> , 2009, 452, 87-89.	2.1	94
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78	Etiology and Pathogenesis of Parkinson Disease. <i>Neurologic Clinics</i> , 2009, 27, 583-603.	1.8	105

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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.	3.9	43
100	Congenital mirror movements in Parkinson's disease: Clinical and neurophysiological observations. <i>Movement Disorders</i> , 2010, 25, 1520-1523.	3.9	6
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103	Association of the glucocerebrosidase N370S allele with Parkinson's disease in two separate Chinese Han populations of mainland China. <i>European Journal of Neurology</i> , 2010, 17, 1476-1478.	3.3	37
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136	Genetics of Parkinson's Disease. Cold Spring Harbor Perspectives in Medicine, 2012, 2, a008888-a008888.	6.2	1,026
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148	Glucocerebrosidase mutations confer a greater risk of dementia during Parkinson's disease course. Movement Disorders, 2012, 27, 393-399.	3.9	144
149	The Association Between α -Glucocerebrosidase Mutations and Parkinsonism. Current Neurology and Neuroscience Reports, 2013, 13, 368.	4.2	32
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162	Glucocerebrosidase, a new player changing the old rules in Lewy body diseases. <i>Biological Chemistry</i> , 2013, 394, 807-818.	2.5	14
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166	Glucocerebrosidase Gene Mutations Associated with Parkinson's Disease: A Meta-Analysis in a Chinese population. <i>PLoS ONE</i> , 2014, 9, e115747.	2.5	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.	1.2	51
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177	Neuroinflammation and α -synuclein accumulation in response to glucocerebrosidase deficiency are accompanied by synaptic dysfunction. <i>Molecular Genetics and Metabolism</i> , 2014, 111, 152-162.	1.1	94
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