

Neuropathology provides clues to the pathophysiology

Molecular Genetics and Metabolism

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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	Mutations in the Glucocerebrosidase Gene and Parkinson's Disease in Ashkenazi Jews. <i>New England Journal of Medicine</i> , 2004, 351, 1972-1977.	13.9	559
3	New Genetic Insights into Parkinson's Disease. <i>New England Journal of Medicine</i> , 2004, 351, 1937-1940.	13.9	42
4	Gaucher disease: complexity in a "simple" disorder. <i>Molecular Genetics and Metabolism</i> , 2004, 83, 6-15.	0.5	350
5	Computerized cognitive testing in patients with type I Gaucher disease: Effects of enzyme replacement and substrate reduction. <i>Genetics in Medicine</i> , 2005, 7, 124-130.	1.1	34
6	Gaucher disease: pathological mechanisms and modern management. <i>British Journal of Haematology</i> , 2005, 129, 178-188.	1.2	240
7	Neurological Manifestations in Lysosomal Storage Disorders - From Pathology to First Therapeutic Possibilities. <i>Neuropediatrics</i> , 2005, 36, 285-289.	0.3	31
8	PERINATAL LETHAL GAUCHER DISEASE: A DISTINCT PHENOTYPE ALONG THE NEURONOPATHIC CONTINUUM. <i>Fetal and Pediatric Pathology</i> , 2005, 24, 205-222.	0.4	44
9	Mutations in the glucocerebrosidase gene and Parkinson disease: Phenotype "genotype" correlation. <i>Neurology</i> , 2005, 65, 1460-1461.	1.5	74
10	Gaucher disease and parkinsonism. <i>Molecular Genetics and Metabolism</i> , 2005, 84, 302-304.	0.5	75
11	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
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15	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
16	High frequency of mutation G377S in Brazilian type 3 Gaucher disease patients. <i>Brazilian Journal of Medical and Biological Research</i> , 2006, 39, 1171-1179.	0.7	101
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19	Glucosylceramide transfer from lysosomes "the missing link in molecular pathology of glucosylceramidase deficiency: A hypothesis based on existing data. <i>Journal of Inherited Metabolic Disease</i> , 2006, 29, 707-715.	1.7	39

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20	Glucocerebrosidase mutations are also found in subjects with early-onset parkinsonism from Venezuela. <i>Movement Disorders</i> , 2006, 21, 282-283.	2.2	64
21	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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43	Chapter 6 Molecular and Cellular Biology of Synucleins. <i>International Review of Cell and Molecular Biology</i> , 2008, 270, 225-317.	1.6	90
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55	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	The Genetics of Parkinson Disease. <i>Journal of Geriatric Psychiatry and Neurology</i> , 2010, 23, 228-242.	1.2	259
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71	The Role of Glucocerebrosidase Mutations in Parkinson Disease and Lewy Body Disorders. <i>Current Neurology and Neuroscience Reports</i> , 2010, 10, 190-198.	2.0	131
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79	Facial tic associated with lamotrigine in adults. <i>Movement Disorders</i> , 2010, 25, 1512-1513.	2.2	10
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140	Increased incidence of Parkinsonism among Chinese with β -glucosidase mutation in central Taiwan.		

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177	Quality control gone wrong: mitochondria, lysosomal storage disorders and neurodegeneration. <i>British Journal of Pharmacology</i> , 2014, 171, 1958-1972.	2.7	71
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379	Thirty-year clinical outcomes after haematopoietic stem cell transplantation in neuronopathic Gaucher disease. <i>Orphanet Journal of Rare Diseases</i> , 2022, 17, .	1.2	7
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