

# Grisel J Lopez

## List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

33  
papers

1,416  
citations

13  
h-index

34  
g-index

34  
ext. papers

1,807  
ext. citations

6.5  
avg, IF

4.52  
L-index

#	Paper	IF	Citations
33	The link between the GBA gene and parkinsonism. <i>Lancet Neurology, The</i> , <b>2012</b> , 11, 986-98	24.1	353
32	A multicenter study of glucocerebrosidase mutations in dementia with Lewy bodies. <i>JAMA Neurology</i> , <b>2013</b> , 70, 727-35	17.2	285
31	A New Glucocerebrosidase Chaperone Reduces $\beta$ Synuclein and Glycolipid Levels in iPSC-Derived Dopaminergic Neurons from Patients with Gaucher Disease and Parkinsonism. <i>Journal of Neuroscience</i> , <b>2016</b> , 36, 7441-52	6.6	150
30	The spectrum of parkinsonian manifestations associated with glucocerebrosidase mutations. <i>Archives of Neurology</i> , <b>2008</b> , 65, 1353-7		142
29	Gaucher disease: Progress and ongoing challenges. <i>Molecular Genetics and Metabolism</i> , <b>2017</b> , 120, 8-21	3.7	72
28	Macrophage models of Gaucher disease for evaluating disease pathogenesis and candidate drugs. <i>Science Translational Medicine</i> , <b>2014</b> , 6, 240ra73	17.5	72
27	The clinical management of Type 2 Gaucher disease. <i>Molecular Genetics and Metabolism</i> , <b>2015</b> , 114, 110-122	3.7	71
26	The neurobiology of glucocerebrosidase-associated parkinsonism: a positron emission tomography study of dopamine synthesis and regional cerebral blood flow. <i>Brain</i> , <b>2012</b> , 135, 2440-8	11.2	67
25	Genome sequencing analysis identifies new loci associated with Lewy body dementia and provides insights into its genetic architecture. <i>Nature Genetics</i> , <b>2021</b> , 53, 294-303	36.3	31
24	Cerebrospinal fluid biomarkers of central dopamine deficiency predict Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , <b>2018</b> , 50, 108-112	3.6	28
23	Clinical course and prognosis in patients with Gaucher disease and parkinsonism. <i>Neurology: Genetics</i> , <b>2016</b> , 2, e57	3.8	23
22	Cardiac sympathetic denervation predicts PD in at-risk individuals. <i>Parkinsonism and Related Disorders</i> , <b>2018</b> , 52, 90-93	3.6	20
21	Gaucher disease and SARS-CoV-2 infection: Emerging management challenges. <i>Molecular Genetics and Metabolism</i> , <b>2020</b> , 130, 164-169	3.7	14
20	Autosomal recessive mutations in the development of Parkinson's disease. <i>Biomarkers in Medicine</i> , <b>2010</b> , 4, 713-21	2.3	11
19	Variation in cognitive function over time in Gaucher disease type 3. <i>Neurology</i> , <b>2019</b> , 93, e2272-e2283	6.5	10
18	Efferocytosis is impaired in Gaucher macrophages. <i>Haematologica</i> , <b>2017</b> , 102, 656-665	6.6	8
17	The natural history of type 2 Gaucher disease in the 21st century: A retrospective study. <i>Neurology</i> , <b>2020</b> , 95, e2119-e2130	6.5	8

16	Alleles with more than one mutation can complicate genotype/phenotype studies in Mendelian disorders: Lessons from Gaucher disease. <i>Molecular Genetics and Metabolism</i> , <b>2018</b> , 125, 1-3	3.7	8
15	Substrate reduction therapy for GBA1-associated Parkinsonism: Are we betting on the wrong mouse?. <i>Movement Disorders</i> , <b>2020</b> , 35, 228-230	7	6
14	Clinical evaluation of sibling pairs with gaucher disease discordant for parkinsonism. <i>Movement Disorders</i> , <b>2020</b> , 35, 359-365	7	6
13	Pro-cathepsin D, Prosaposin, and Progranulin: Lysosomal Networks in Parkinsonism. <i>Trends in Molecular Medicine</i> , <b>2020</b> , 26, 913-923	11.5	6
12	Five-parameter evaluation of dysphagia: A novel prognostic scale for assessing neurological decline in Gaucher disease type 2. <i>Molecular Genetics and Metabolism</i> , <b>2019</b> , 127, 191-199	3.7	5
11	Type 2 Gaucher disease in an infant despite a normal maternal glucocerebrosidase gene. <i>American Journal of Medical Genetics, Part A</i> , <b>2017</b> , 173, 3211-3215	2.5	4
10	EEG abnormalities in patients with chronic neuronopathic Gaucher disease: A retrospective review. <i>Molecular Genetics and Metabolism</i> , <b>2020</b> , 131, 358-363	3.7	4
9	Longitudinal Positron Emission Tomography of Dopamine Synthesis in Subjects with GBA1 Mutations. <i>Annals of Neurology</i> , <b>2020</b> , 87, 652-657	9.4	3
8	Bilateral Femoral Osteolytic Lesions in a Patient with Type 3 Gaucher Disease. <i>Molecular Genetics and Metabolism Reports</i> , <b>2015</b> , 5, 107-109	1.8	3
7	Glucocerebrosidase mutations and parkinsonism: how much does the mutation matter?. <i>Journal of Xiangya Medicine</i> , <b>2018</b> , 3, 1-1	0.1	3
6	No Evidence That Glucosylsphingosine Is a Biomarker for Parkinson's Disease: Statistical Differences Do Not Necessarily Indicate Biological Significance.. <i>Movement Disorders</i> , <b>2022</b> ,	7	2
5	Understanding and managing parkinsonism in patients with glucocerebrosidase mutations. <i>Expert Opinion on Orphan Drugs</i> , <b>2015</b> , 3, 549-562	1.1	1
4	Neuropathological Features of Gaucher Disease and Gaucher Disease with Parkinsonism. <i>International Journal of Molecular Sciences</i> , <b>2022</b> , 23, 5842	6.3	0
3	Studies of glucocerebrosidase provide new therapeutic targets for parkinsonism. <i>Future Neurology</i> , <b>2014</b> , 9, 407-409	1.5	
2	Gaucher Disease and Heart Failure of Unknown Origin. <i>American Journal of Medicine</i> , <b>2021</b> , 134, 745-748.	2.4	
1	Imaging and genetics in Parkinson's disease: assessment of the GBA1 mutation. <i>Journal of Neurology</i> ,	5.5	