Nahid Tayebi

List of Publications by Year in Descending Order

20 papers	556 citations	10 h-index	21 g-index
21 ext. papers	777 ext. citations	6.4 avg, IF	3.3 L-index

#	Paper	IF	Citations
20	No Evidence That Glucosylsphingosine Is a Biomarker for Parkinson's Disease: Statistical Differences Do Not Necessarily Indicate Biological Significance.. *Movement Disorders*, **2022**,	7	2
19	Next-Generation Sequencing Analysis of : The Challenge of Detecting Complex Recombinant Alleles. *Frontiers in Genetics*, **2021**, 12, 684067	4.5	2
18	Genome sequencing analysis identifies new loci associated with Lewy body dementia and provides insights into its genetic architecture. *Nature Genetics*, **2021**, 53, 294-303	36.3	31
17	Clinical evaluation of sibling pairs with gaucher disease discordant for parkinsonism. *Movement Disorders*, **2020**, 35, 359-365	7	6
16	The natural history of type 2 Gaucher disease in the 21st century: A retrospective study. *Neurology*, **2020**, 95, e2119-e2130	6.5	8
15	EEG abnormalities in patients with chronic neuronopathic Gaucher disease: A retrospective review. *Molecular Genetics and Metabolism*, **2020**, 131, 358-363	3.7	4
14	C-terminal Esynuclein truncations are linked to cysteine cathepsin activity in Parkinson's disease. *Journal of Biological Chemistry*, **2019**, 294, 9973-9984	5.4	23
13	Can GBA1-Associated Parkinson Disease Be Modeled in the Mouse?. *Trends in Neurosciences*, **2019**, 42, 631-643	13.3	6
12	Variation in cognitive function over time in Gaucher disease type 3. *Neurology*, **2019**, 93, e2272-e2283	6.5	10
11	ACE phenotyping in Gaucher disease. *Molecular Genetics and Metabolism*, **2018**, 123, 501-510	3.7	16
10	Exploring genetic modifiers of Gaucher disease: The next horizon. *Human Mutation*, **2018**, 39, 1739-1751	4.7	28
9	The role of epigenetics in lysosomal storage disorders: Uncharted territory. *Molecular Genetics and Metabolism*, **2017**, 122, 10-18	3.7	26
8	Glucocerebrosidase haploinsufficiency in A53T Esynuclein mice impacts disease onset and course. *Molecular Genetics and Metabolism*, **2017**, 122, 198-208	3.7	18
7	Type 2 Gaucher disease in an infant despite a normal maternal glucocerebrosidase gene. *American Journal of Medical Genetics, Part A*, **2017**, 173, 3211-3215	2.5	4
6	A new glucocerebrosidase-deficient neuronal cell model provides a tool to probe pathophysiology and therapeutics for Gaucher disease. *DMM Disease Models and Mechanisms*, **2016**, 9, 769-78	4.1	14
5	Clinical course and prognosis in patients with Gaucher disease and parkinsonism. *Neurology: Genetics*, **2016**, 2, e57	3.8	23
4	Lysosomal integral membrane protein-2: a new player in lysosome-related pathology. *Molecular Genetics and Metabolism*, **2014**, 111, 84-91	3.7	46

3	A multicenter study of glucocerebrosidase mutations in dementia with Lewy bodies. *JAMA Neurology*, **2013**, 70, 727-35	17.2	285
2	Localization of Non-Factor VIII Sequence Involved in the Factor VIII Gene Inversion in Hemophilia A Dogs.. *Blood*, **2007**, 110, 1146-1146	2.2	
1	A novel alteration in metaxin 1, F202L, is associated with N370S in Gaucher disease. *Journal of Human Genetics*, **2004**, 49, 220-222	4.3	4