

# CHCHD2 mutations in autosomal dominant late-onset linkage and sequencing study

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Citation Report

#	ARTICLE	IF	CITATIONS
1	II. Familial Parkinson's Disease Causative Genes Shed Light on the Molecular Pathogenesis. The Journal of the Japanese Society of Internal Medicine, 2015, 104, 1552-1557.	0.0	0
2	Î±-Synuclein and Lewy pathology in Parkinson's disease. Current Opinion in Neurology, 2015, 28, 375-381.	1.8	79
3	Mitochondrial Dysfunction in Parkinson's Disease. Experimental Neurobiology, 2015, 24, 103-116.	0.7	267
4	Mitophagy Regulated by the PINK1-Parkin Pathway. , 2015, , .		8
5	CHCHD2 and Parkinson's disease. Lancet Neurology, The, 2015, 14, 679-680.	4.9	13
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8	CHCHD2 and Parkinson's disease. Lancet Neurology, The, 2015, 14, 680-681.	4.9	9
9	CHCHD2 and Parkinson's disease. Lancet Neurology, The, 2015, 14, 681-682.	4.9	29
10	CHCHD2 and Parkinson's disease—Authors' reply. Lancet Neurology, The, 2015, 14, 682-683.	4.9	6
11	Mitochondrial targeting sequence variants of the <i>CHCHD2</i> gene are a risk for Lewy body disorders. Neurology, 2015, 85, 2016-2025.	1.5	51
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18	Insights from late-onset familial parkinsonism on the pathogenesis of idiopathic Parkinson's disease. Lancet Neurology, The, 2015, 14, 1054-1064.	4.9	56

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19	The mTOR Signaling Pathway in Neurodegenerative Diseases. , 2016, , 85-104.		3
20	Genetics of movement disorders in the next generation sequencing era. Movement Disorders, 2016, 31, 458-470.	2.2	34
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23	Mutation analysis of the CHCHD2 gene in Chinese Han patients with Parkinson's disease. Parkinsonism and Related Disorders, 2016, 29, 143-144.	1.1	9
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25	Mutational analysis of CHCHD2 in Chinese patients with multiple system atrophy and amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 2016, 368, 389-391.	0.3	8
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38	Association of GCH1 and MIR4697 , but not SIPA1L2 and VPS13C polymorphisms, with Parkinson's disease in Taiwan. <i>Neurobiology of Aging</i> , 2016, 39, 221.e1-221.e5.	1.5	15
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45	Discovery of a frameshift mutation in podocalyxin-like (PODXL) gene, coding for a neural adhesion molecule, as causal for autosomal-recessive juvenile Parkinsonism. <i>Journal of Medical Genetics</i> , 2016, 53, 450-456.	1.5	37
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55	Loss of Parkinson's disease-associated protein CHCHD2 affects mitochondrial crista structure and destabilizes cytochrome c. <i>Nature Communications</i> , 2017, 8, 15500.	5.8	123

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57	Genetic analysis of CHCHD2 and CHCHD10 in Italian patients with Parkinson's disease. <i>Neurobiology of Aging</i> , 2017, 53, 193.e7-193.e8.	1.5	8
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131	Identification of Disease-Associated Variants by Targeted Gene Panel Resequencing in Parkinson's Disease. <i>Frontiers in Neurology</i> , 2020, 11, 576465.	1.1	4
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165	Reduced erythrocytic CHCHD2 mRNA is associated with brain pathology of Parkinson's disease. <i>Acta Neuropathologica Communications</i> , 2021, 9, 37.	2.4	8
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