Vladimir Han

List of Publications by Year
in descending order

28 papers	757 citations	759233 12 h-index	580821 25 g-index
28 all docs	28 docs citations	28 times ranked	1246 citing authors

#	Article	IF	Citations
1	Monogenic variants in dystonia: an exome-wide sequencing study. Lancet Neurology, The, 2020, 19, 908-918.	10.2	139
2	The Clinical Syndrome of Paroxysmal Exercise-Induced Dystonia: Diagnostic Outcomes and an Algorithm. Movement Disorders Clinical Practice, 2014, 1, 57-61.	1.5	100
3	Psychiatric co-morbidity is highly prevalent in idiopathic cervical dystonia and significantly influences health-related quality of life: Results of a controlled study. Parkinsonism and Related Disorders, 2016, 30, 7-12.	2.2	81
4	Differences in <scp>MDS</scp>â€<scp>UPDRS</scp> Scores Based on Hoehn and Yahr Stage and Disease Duration. Movement Disorders Clinical Practice, 2017, 4, 536-544.	1.5	65
5	Relationship between the non-motor items of the MDSâ€"UPDRS and Quality of Life in patients with Parkinson's disease. Journal of the Neurological Sciences, 2015, 353, 87-91.	0.6	58
6	Paroxysmal exercise-induced dystonia within the phenotypic spectrum ofECHS1deficiency. Movement Disorders, 2016, 31, 1041-1048.	3.9	58
7	Relationship between the MDS-UPDRS and Quality of Life: A large multicenter study of 3206 patients. Parkinsonism and Related Disorders, 2018, 52, 83-89.	2.2	46
8	Fatigue, Sleep Disturbances, and Their Influence on Quality of Life in Cervical Dystonia Patients. Movement Disorders Clinical Practice, 2017, 4, 517-523.	1.5	36
9	Prevalence of nonâ€motor symptoms and their association with quality of life in cervical dystonia. Acta Neurologica Scandinavica, 2020, 142, 613-622.	2.1	22
10	Neurodevelopmental disorder associated with IRF2BPL gene mutation: Expanding the phenotype?. Parkinsonism and Related Disorders, 2019, 62, 239-241.	2.2	20
11	Recessive variants in ZNF142 cause a complex neurodevelopmental disorder with intellectual disability, speech impairment, seizures, and dystonia. Genetics in Medicine, 2019, 21, 2532-2542.	2.4	17
12	Clinically relevant copy-number variants in exome sequencing data of patients with dystonia. Parkinsonism and Related Disorders, 2021, 84, 129-134.	2.2	15
13	The Frequency and Self-perceived Impact on Daily Life of Motor and Non-motor Symptoms in Cervical Dystonia. Movement Disorders Clinical Practice, 2017, 4, 750-754.	1.5	14
14	WARS2 mutations cause dopa-responsive early-onset parkinsonism and progressive myoclonus ataxia. Parkinsonism and Related Disorders, 2022, 94, 54-61.	2.2	13
15	Prevalence of Prodromal Parkinsonâ€™s Disease as Defined by MDS Research Criteria among Elderly Patients Undergoing Colonoscopy. Journal of Parkinson's Disease, 2017, 7, 481-489.	2.8	12
16	Î±â€Synuclein antibody 5G4 identifies manifest and prodromal Parkinson's disease in colonic mucosa. Movement Disorders, 2018, 33, 1366-1368.	3.9	12
17	Dystonia as a prominent presenting feature in developmental and epileptic encephalopathies: A case series. Parkinsonism and Related Disorders, 2021, 90, 73-78.	2.2	9
18	A Recurrent <scp>VPS16</scp> p.Arg187* Nonsense Variant in Earlyâ€Onset Generalized Dystonia. Movement Disorders, 2021, 36, 1984-1985.	3.9	7

#	Article	IF	Citations
19	Scoring Algorithmâ€Based Genomic Testing in Dystonia: A Prospective Validation Study. Movement Disorders, 2021, 36, 1959-1964.	3.9	7
20	Whole exome sequencing identifies a homozygous POLG2 missense variant in an adult patient presenting with optic atrophy, movement disorders, premature ovarian failure and mitochondrial DNA depletion. European Journal of Medical Genetics, 2020, 63, 103821.	1.3	5
21	Comparison in detection of prodromal Parkinson's disease patients using original and updated MDS research criteria in two independent cohorts. Parkinsonism and Related Disorders, 2021, 87, 48-55.	2.2	5
22	Validation of the Official Slovak Version of the Unified Dyskinesia Rating Scale (UDysRS). Parkinson's Disease, 2015, 2015, 1-7.	1.1	4
23	Alzheimerâ€™s Disease-Associated SNP rs708727 in SLC41A1 May Increase Risk for Parkinsonâ€™s Disease: Report from Enlarged Slovak Study. International Journal of Molecular Sciences, 2022, 23, 1604.	4.1	4
24	Recessive null-allele variants in MAG associated with spastic ataxia, nystagmus, neuropathy, and dystonia. Parkinsonism and Related Disorders, 2020, 77, 70-75.	2.2	3
25	Prevalence of Fabry Disease among Patients with Parkinsonâ€™s Disease. Parkinson's Disease, 2022, 2022, 1-8.	1.1	3
26	Atypical presentations of DYT1 dystonia with acute craniocervical onset. Parkinsonism and Related Disorders, 2021, 83, 54-55.	2.2	1
27	LRRK2 mutations in Parkinson's disease patients from Central Europe: A case control study. Parkinsonism and Related Disorders, 2021, 83, 110-112.	2.2	1
28	Answer to Finsterer about â€œMultisystem presentation of a homozygous POLG2 variantâ€. European Journal of Medical Genetics, 2020, 63, 103900.	1.3	0