Matej Skorvanek

List of Publications by Year in descending order

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66 1,632 22 36 papers citations h-index g-index

66 66 66 2465

times ranked

citing authors

docs citations

all docs

#	Article	IF	CITATIONS
1	Monogenic variants in dystonia: an exome-wide sequencing study. Lancet Neurology, The, 2020, 19, 908-918.	10.2	139
2	Global scales for cognitive screening in Parkinson's disease: Critique and recommendations. Movement Disorders, 2018, 33, 208-218.	3.9	138
3	The Clinical Syndrome of Paroxysmal Exercise-Induced Dystonia: Diagnostic Outcomes and an Algorithm. Movement Disorders Clinical Practice, 2014, 1, 57-61.	1.5	100
4	The associations between fatigue, apathy, and depression in Parkinson's disease. Acta Neurologica Scandinavica, 2015, 131, 80-87.	2.1	93
5	Lossâ€ofâ€Function Variants in <scp>HOPS</scp> Complex Genes <scp><i>VPS16</i></scp> and <scp><i>VPS41</i></scp> Cause Early Onset Dystonia Associated with Lysosomal Abnormalities. Annals of Neurology, 2020, 88, 867-877.	5.3	70
6	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
7	<i>De novo</i> variants in neurodevelopmental disordersâ€"experiences from a tertiary care center. Clinical Genetics, 2021, 100, 14-28.	2.0	64
8	Clinical Practice: Evidence-Based Recommendations for the Treatment of Cervical Dystonia with Botulinum Toxin. Frontiers in Neurology, 2017, 8, 35.	2.4	63
9	Relationship between the non-motor items of the MDS–UPDRS and Quality of Life in patients with Parkinson's disease. Journal of the Neurological Sciences, 2015, 353, 87-91.	0.6	58
10	Paroxysmal exercise-induced dystonia within the phenotypic spectrum of <i>ECHS1</i> deficiency. Movement Disorders, 2016, 31, 1041-1048.	3.9	58
11	Assessment of Ataxia Rating Scales and Cerebellar Functional Tests: Critique and Recommendations. Movement Disorders, 2021, 36, 283-297.	3.9	52
12	Scales to assess impulsive and compulsive behaviors in Parkinson's disease: Critique and recommendations. Movement Disorders, 2019, 34, 791-798.	3.9	49
13	Rating Scales for Pain in Parkinson's Disease: Critique and Recommendations. Movement Disorders Clinical Practice, 2016, 3, 527-537.	1.5	46
14	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
15	Accuracy of Rating Scales and Clinical Measures for Screening of Rapid Eye Movement Sleep Behavior Disorder and for Predicting Conversion to Parkinson's Disease and Other Synucleinopathies. Frontiers in Neurology, 2018, 9, 376.	2.4	39
16	The Skin and Parkinson's Disease: Review of Clinical, Diagnostic, and Therapeutic Issues. Movement Disorders Clinical Practice, 2017, 4, 21-31.	1.5	37
17	Adherence to Pharmacotherapy in Patients With Parkinson's Disease Taking Three and More Daily Doses of Medication. Frontiers in Neurology, 2019, 10, 799.	2.4	34
18	Update on the Management of Parkinson's Disease for General Neurologists. Parkinson's Disease, 2020, 2020, 1-13.	1.1	33

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19	Validation study of the hoehn and yahr scale included in the MDSâ€UPDRS. Movement Disorders, 2018, 33, 651-652.	3.9	32
20	Apathy in Elderly Nondemented Patients With Parkinson's Disease. Journal of Geriatric Psychiatry and Neurology, 2013, 26, 237-243.	2.3	30
21	Clinical determinants of primary and secondary fatigue in patients with Parkinson's disease. Journal of Neurology, 2013, 260, 1554-1561.	3.6	29
22	The role of social support in anxiety and depression among Parkinson's disease patients. Disability and Rehabilitation, 2014, 36, 2044-2049.	1.8	24
23	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
24	Neurodevelopmental disorder associated with IRF2BPL gene mutation: Expanding the phenotype?. Parkinsonism and Related Disorders, 2019, 62, 239-241.	2.2	20
25	Measurement of Nonmotor Symptoms in Clinical Practice. International Review of Neurobiology, 2017, 133, 291-345.	2.0	18
26	Blood DNA methylation provides an accurate biomarker of <i>KMT2B</i> -related dystonia and predicts onset. Brain, 2022, 145, 644-654.	7.6	18
27	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
28	Randomised controlled trial of escitalopram for cervical dystonia with dystonic jerks/tremor. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 579-585.	1.9	15
29	Clinically relevant copy-number variants in exome sequencing data of patients with dystonia. Parkinsonism and Related Disorders, 2021, 84, 129-134.	2.2	15
30	Influence of Disease Severity on Fatigue in Patients with Parkinson's Disease Is Mainly Mediated by Symptoms of Depression. European Neurology, 2013, 70, 201-209.	1.4	13
31	WARS2 mutations cause dopa-responsive early-onset parkinsonism and progressive myoclonus ataxia. Parkinsonism and Related Disorders, 2022, 94, 54-61.	2.2	13
32	Management of dystonia in Europe: a survey of the European network for the study of the dystonia syndromes. European Journal of Neurology, 2016, 23, 772-779.	3.3	12
33	Prevalence of Prodromal Parkinson'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
34	α‧ynuclein antibody 5G4 identifies manifest and prodromal Parkinson's disease in colonic mucosa. Movement Disorders, 2018, 33, 1366-1368.	3.9	12
35	Does the 5–2-1 criteria identify patients with advanced Parkinson's disease? Real-world screening accuracy and burden of 5–2-1-positive patients in 7 countries. BMC Neurology, 2022, 22, 35.	1.8	12
36	Ataxia Telangiectasia Gene Mutation in Isolated Segmental Dystonia Without Ataxia and Telangiectasia. Movement Disorders Clinical Practice, 2018, 5, 89-91.	1.5	11

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37	The neurological and neuropsychiatric spectrum of adults with late-treated phenylketonuria. Parkinsonism and Related Disorders, 2021, 89, 167-175.	2.2	11
38	Exploding head syndrome – a rare parasomnia or a dissociative episode?. Sleep Medicine, 2014, 15, 728-730.	1.6	10
39	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
40	Extensive validation study of the Parkinson's Disease Composite Scale. European Journal of Neurology, 2019, 26, 1281-1288.	3.3	8
41	Genetic overlap between dystonia and other neurologic disorders: A study of 1,100 exomes. Parkinsonism and Related Disorders, 2022, 102, 1-6.	2.2	8
42	Opsoclonus-myoclonus ataxia syndrome secondary to venlafaxine intoxication. Journal of the Neurological Sciences, 2017, 372, 19-20.	0.6	7
43	A Recurrent <scp><i>VPS16</i></scp> p.Arg187* Nonsense Variant in Earlyâ€Onset Generalized Dystonia. Movement Disorders, 2021, 36, 1984-1985.	3.9	7
44	Scoring Algorithmâ€Based Genomic Testing in Dystonia: A Prospective Validation Study. Movement Disorders, 2021, 36, 1959-1964.	3.9	7
45	Variant recurrence confirms the existence of a <i>FBXO31</i> i>â€related spasticâ€dystonic cerebral palsy syndrome. Annals of Clinical and Translational Neurology, 2021, 8, 951-955.	3.7	6
46	Selfâ€rated health mediates the association between functional status and healthâ€related quality of life in <scp>P</scp> arkinson's disease. Journal of Clinical Nursing, 2014, 23, 1970-1977.	3.0	5
47	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
48	Predictors of outcome events and 6-year mortality after carotid endarterectomy and carotid stenting in patients with carotid artery stenosis. Neurologia I Neurochirurgia Polska, 2021, 55, 67-73.	1.2	5
49	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
50	Validation of the Official Slovak Version of the Unified Dyskinesia Rating Scale (UDysRS). Parkinson's Disease, 2015, 2015, 1-7.	1.1	4
51	Alzheimer's Disease-Associated SNP rs708727 in SLC41A1 May Increase Risk for Parkinson's Disease: Report from Enlarged Slovak Study. International Journal of Molecular Sciences, 2022, 23, 1604.	4.1	4
52	Lack of Accredited Clinical Training in Movement Disorders in Europe, Egypt, and Tunisia. Journal of Parkinson's Disease, 2020, 10, 1833-1843.	2.8	3
53	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
54	Prevalence of Fabry Disease among Patients with Parkinson's Disease. Parkinson's Disease, 2022, 2022, 1-8.	1.1	3

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55	Skin Conditions and Movement Disorders: Hiding in Plain Sight. Movement Disorders Clinical Practice, 2022, 9, 566-583.	1.5	3
56	Reply: MoCA for cognitive screening in Parkinson's disease: Beware of floor effect. Movement Disorders, 2018, 33, 499-500.	3.9	2
57	Neuromodulation Options and Patient Selection for Parkinson's Disease. Neurology India, 2020, 68, 170.	0.4	2
58	Effect of Pillbox Organizers with Alarms on Adherence to Pharmacotherapy in Parkinson Disease Patients Taking Three and More Daily Doses of Dopaminergic Medications. Journal of Personalized Medicine, 2022, 12, 179.	2.5	2
59	Validation of the Arabic Version of the Movement Disorder Societyâ€Unified Parkinson's Disease Rating Scale. Movement Disorders, 2022, 37, 826-841.	3.9	2
60	Atypical presentations of DYT1 dystonia with acute craniocervical onset. Parkinsonism and Related Disorders, 2021, 83, 54-55.	2.2	1
61	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
62	Reply: Hoehn and Yahr stage 3 and Postural stability item in the Movement Disorder Societyâ€Unified Parkinson's Disease Rating Scale. Movement Disorders, 2018, 33, 1189-1190.	3.9	0
63	Answer to Finsterer about "Multisystem presentation of a homozygous POLG2 variant― European Journal of Medical Genetics, 2020, 63, 103900.	1.3	O
64	for , , , and in Parkinson's. Neuromethods, 2021, , 227-269.	0.3	0
65	Impact of the COVID-19 outbreak on neurological consultation in an emergency department. Acta Neurologica Belgica, 2022, , 1.	1.1	0
66	Progressive choreodystonia in Xâ€linked <scp>hyperâ€lgM</scp> immunodeficiency: a rare but recurrent presentation. Annals of Clinical and Translational Neurology, 2022, , .	3.7	0