Marina A J Tijssen

List of Publications by Year in descending order

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Version: 2024-02-01

198 papers	5,328 citations	87888 38 h-index	144013 57 g-index
201 all docs	201 docs citations	201 times ranked	4518 citing authors

#	Article	IF	Citations
1	WARS2 mutations cause dopa-responsive early-onset parkinsonism and progressive myoclonus ataxia. Parkinsonism and Related Disorders, 2022, 94, 54-61.	2.2	13
2	Electrophysiological testing aids the diagnosis of tremor and myoclonus in clinically challenging patients. Clinical Neurophysiology Practice, 2022, 7, 51-58.	1.4	4
3	A diagnosis of progressive myoclonic ataxia guided by blood biomarkers: Expert commentary. Parkinsonism and Related Disorders, 2022, 94, 127-128.	2.2	0
4	DBS for dystonia: Should we take our patients to the swimming pool?. Parkinsonism and Related Disorders, 2022, 96, 36-37.	2.2	1
5	Serotonergic system in vivo with [11C]DASB PET scans in GTP-cyclohydrolase deficient dopa-responsive dystonia patients. Scientific Reports, 2022, 12, 6292.	3.3	3
6	Nomenclature of Genetic Movement Disorders: Recommendations of the International Parkinson and Movement Disorder Society Task Force â€“ An Update. Movement Disorders, 2022, 37, 905-935.	3.9	49
7	Multi-centre classification of functional neurological disorders based on resting-state functional connectivity. NeuroImage: Clinical, 2022, 35, 103090.	2.7	6
8	A novel diagnostic approach for patients with adult-onset dystonia. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 1039-1048.	1.9	3
9	Diagnostic approach to paediatric movement disorders: a clinical practice guide. Developmental Medicine and Child Neurology, 2021, 63, 252-258.	2.1	11
10	Cross-disease analysis of depression, ataxia and dystonia highlights a role for synaptic plasticity and the cerebellum in the pathophysiology of these comorbid diseases. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2021, 1867, 165976.	3.8	13
11	<scp>Myoclonusâ€Ataxia</scp> Syndromes: A Diagnostic Approach. Movement Disorders Clinical Practice, 2021, 8, 9-24.	1.5	11
12	A Gainâ€ofâ€Function Variant in Dopamine <scp>D2</scp> Receptor and Progressive Chorea and Dystonia Phenotype. Movement Disorders, 2021, 36, 729-739.	3.9	20
13	Bilateral Pallidotomy for Dystonia: A Systematic Review. Movement Disorders, 2021, 36, 547-557.	3.9	19
14	Functional or not functional; thatâ€™s the question. European Journal of Neurology, 2021, 28, 33-39.	3.3	15
15	Rare functional missense variants in CACNA1H: What can we learn from Writerâ€™s cramp?. Molecular Brain, 2021, 14, 18.	2.6	3
16	Sleep disturbance in movement disorders: insights, treatments and challenges. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 723-736.	1.9	12
17	Challenges in Clinicogenetic Correlations: One Phenotype â€“ Many Genes. Movement Disorders Clinical Practice, 2021, 8, 311-321.	1.5	12
18	WDR45, one gene associated with multiple neurodevelopmental disorders. Autophagy, 2021, 17, 3908-3923.	9.1	20

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19	Signaling-Biased and Constitutively Active Dopamine D2 Receptor Variant. ACS Chemical Neuroscience, 2021, 12, 1873-1884.	3.5	9
20	Dystonia Management: What to Expect From the Future? The Perspectives of Patients and Clinicians Within DystoniaNet Europe. Frontiers in Neurology, 2021, 12, 646841.	2.4	10
21	Reply to: â€œChildhood Onset Chorea Caused by a Recurrent De Novo <scp>DRD2</scp> Variantâ€. Movement Disorders, 2021, 36, 1473-1474.	3.9	2
22	The auditory startle response in relation to outcome in functional movement disorders. Parkinsonism and Related Disorders, 2021, 89, 113-117.	2.2	0
23	Cognition in children and young adults with myoclonus dystonia â€“ A case control study. Parkinsonism and Related Disorders, 2021, 89, 162-166.	2.2	4
24	The diagnostic value of clinical neurophysiology in hyperkinetic movement disorders: A systematic review. Parkinsonism and Related Disorders, 2021, 89, 176-185.	2.2	19
25	Altered Posterior Midline Activity in Patients with Jerky and Tremulous Functional Movement Disorders. Brain Connectivity, 2021, 11, 584-593.	1.7	3
26	Three Days of Measurement Provide Reliable Estimates of Daily Tremor Characteristics: A Pilot Study in Organic and Functional Tremor Patients. Tremor and Other Hyperkinetic Movements, 2021, 11, 13.	2.0	0
27	Neuroimaging in Functional Neurological Disorder: State of the Field and Research Agenda. NeuroImage: Clinical, 2021, 30, 102623.	2.7	79
28	Next move in movement disorders (NEMO): developing a computer-aided classification tool for hyperkinetic movement disorders. BMJ Open, 2021, 11, e055068.	1.9	1
29	Long-term follow-up of acute functional stroke mimic in comparison to mild acute ischaemic stroke. Journal of Neurology, Neurosurgery and Psychiatry, 2021, , jnnp-2021-327379.	1.9	0
30	Clinical Practice Patterns in Tic Disorders Among Movement Disorder Society Members. Tremor and Other Hyperkinetic Movements, 2021, 11, 43.	2.0	8
31	Shared demographics and comorbidities in different functional motor disorders. Parkinsonism and Related Disorders, 2020, 70, 1-6.	2.2	26
32	Outcome Measures for Functional Neurological Disorder: A Review of the Theoretical Complexities. Journal of Neuropsychiatry and Clinical Neurosciences, 2020, 32, 33-42.	1.8	65
33	Inborn Errors of Metabolism in Adults: Two Patients with Movement Disorders Caused by Glutaric Aciduria Type 1. Movement Disorders Clinical Practice, 2020, 7, S85-S88.	1.5	2
34	Pentameric repeat expansions: cortical myoclonus or cortical tremor?. Brain, 2020, 143, e86-e86.	7.6	9
35	Botulinum neurotoxin (BoNT) treatment in functional movement disorders: long-term follow-up. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1120-1121.	1.9	1
36	Internet-based self-help randomized trial for motor functional neurologic disorder (SHIFT). Neurology, 2020, 95, e1883-e1896.	1.1	22

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37	The chronnectome as a model for Charcotâ€™s â€˜dynamic lesionâ€™ in functional movement disorders. NeuroImage: Clinical, 2020, 28, 102381.	2.7	5
38	Low-frequency oscillation suppression in dystonia: Implications for adaptive deep brain stimulation. Parkinsonism and Related Disorders, 2020, 79, 105-109.	2.2	14
39	Lossâ€ofâ€Function Variants in <scp>HOPS</scp> Complex Genes <scp>VPS16</scp> and <scp>VPS41</scp> Cause Early Onset Dystonia Associated with Lysosomal Abnormalities. Annals of Neurology, 2020, 88, 867-877.	5.3	70
40	Early Onset Ataxia with Comorbid Dystonia: Clinical, Anatomical and Biological Pathway Analysis Expose Shared Pathophysiology. Diagnostics, 2020, 10, 997.	2.6	7
41	The use of nextâ€generation sequencing to unravel new genes: overcoming challenges posed by rare neurological disorders such as myoclonusâ€dystonia. European Journal of Neurology, 2020, 27, 1459-1460.	3.3	1
42	Management of rare movement disorders in Europe: outcome of surveys of the European Reference Network for Rare Neurological Diseases. European Journal of Neurology, 2020, 27, 1493-1500.	3.3	15
43	Systematic clinical approach for diagnosing upper limb tremor. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 822-830.	1.9	35
44	The Effect of Escitalopram on Central Serotonergic and Dopaminergic Systems in Patients with Cervical Dystonia, and Its Relationship with Clinical Treatment Effects: A Double-Blind Placebo-Controlled Trial. Biomolecules, 2020, 10, 880.	4.0	8
45	De novo variants in CAMTA1 cause a syndrome variably associated with spasticity, ataxia, and intellectual disability. European Journal of Human Genetics, 2020, 28, 763-769.	2.8	7
46	Natural Course of <scp>Myoclonusâ€Dystonia</scp> in Adulthood: Stable Motor Signs But Increased Psychiatry. Movement Disorders, 2020, 35, 1077-1078.	3.9	6
47	Outcome measurement in functional neurological disorder: a systematic review and recommendations. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 638-649.	1.9	77
48	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
49	A detailed description of the phenotypic spectrum of North Sea Progressive Myoclonus Epilepsy in a large cohort of seventeen patients. Parkinsonism and Related Disorders, 2020, 72, 44-48.	2.2	9
50	Not every excessive startle is hyperekplexia, the curious case of SOD1. Brain, 2020, 143, e11-e11.	7.6	4
51	Driving Performance in Patients With Idiopathic Cervical Dystonia; A Driving Simulator Pilot Study. Frontiers in Neurology, 2020, 11, 229.	2.4	4
52	The European Reference Network for Rare Neurological Diseases. Frontiers in Neurology, 2020, 11, 616569.	2.4	26
53	The Effectiveness of Deep Brain Stimulation in Dystonia: A Patient-Centered Approach. Tremor and Other Hyperkinetic Movements, 2020, 10, 2.	2.0	5
54	TheÂMovement disorder associated with NMDAR antibody-encephalitis is complex and characteristic: an expert video-rating study. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 724-726.	1.9	71

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55	A clinical diagnostic algorithm for early onset cerebellar ataxia. European Journal of Paediatric Neurology, 2019, 23, 692-706.	1.6	37
56	Tremor and myoclonus. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2019, 161, 149-165.	1.8	10
57	Botulinum neurotoxin treatment in jerky and tremulous functional movement disorders: a double-blind, randomised placebo-controlled trial with an open-label extension. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 1244-1250.	1.9	23
58	Cortical pencil lining on SWI MRI in NBIA and healthy aging. BMC Neurology, 2019, 19, 233.	1.8	3
59	Skater's Cramp: A Possible Taskâ€Specific Dystonia in Dutch Ice Skaters. Movement Disorders Clinical Practice, 2019, 6, 559-566.	1.5	7
60	Nomenclature of Genetically Determined Myoclonus Syndromes: Recommendations of the International Parkinson and Movement Disorder Society Task Force. Movement Disorders, 2019, 34, 1602-1613.	3.9	23
61	Intronic ATTTC repeat expansions in STARD7 in familial adult myoclonic epilepsy linked to chromosome 2. Nature Communications, 2019, 10, 4920.	12.8	99
62	Myoclonus-dystonia: Distinctive motor and non-motor phenotype from other dystonia syndromes. Parkinsonism and Related Disorders, 2019, 69, 85-90.	2.2	24
63	Which disease features run in essential tremor families? A systematic review. Parkinsonism and Related Disorders, 2019, 69, 71-78.	2.2	4
64	The spectrum of involuntary vocalizations in humans: A video atlas. Movement Disorders, 2019, 34, 1774-1791.	3.9	24
65	Intermuscular coherence as biomarker for pallidal deep brain stimulation efficacy in dystonia. Clinical Neurophysiology, 2019, 130, 1351-1357.	1.5	15
66	Similar association between objective and subjective symptoms in functional and organic tremor. Parkinsonism and Related Disorders, 2019, 64, 2-7.	2.2	11
67	The prognosis of functional limb weakness: a 14-year case-control study. Brain, 2019, 142, 2137-2148.	7.6	60
68	Hiding in Plain Sight: Functional Neurological Disorders in the News. Journal of Neuropsychiatry and Clinical Neurosciences, 2019, 31, 361-367.	1.8	13
69	Direct comparison of oscillatory activity in the motor system of Parkinsonâ€™s disease and dystonia: A review of the literature and meta-analysis. Clinical Neurophysiology, 2019, 130, 917-924.	1.5	24
70	Long-Term Specialized Physical Therapy in Cervical Dystonia: Outcomes of a Randomized Controlled Trial. Archives of Physical Medicine and Rehabilitation, 2019, 100, 1417-1425.	0.9	24
71	Variable Interpretation of the Dystonia Consensus Classification Items Compromises Its Solidity. Movement Disorders, 2019, 34, 317-320.	3.9	12
72	Movement disorders and nonmotor neuropsychological symptoms in children and adults with classical galactosemia. Journal of Inherited Metabolic Disease, 2019, 42, 451-458.	3.6	27

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73	CoAâ€dependent activation of mitochondrial acyl carrier protein links four neurodegenerative diseases. EMBO Molecular Medicine, 2019, 11, e10488.	6.9	46
74	Unstable TTTTA/TTTCA expansions in MARCH6 are associated with Familial Adult Myoclonic Epilepsy type 3. Nature Communications, 2019, 10, 4919.	12.8	111
75	The characteristics of pallidal low-frequency and beta bursts could help implementing adaptive brain stimulation in the parkinsonian and dystonic internal globus pallidus. Neurobiology of Disease, 2019, 121, 47-57.	4.4	49
76	Motor and non-motor determinants of health-related quality of life in young dystonia patients. Parkinsonism and Related Disorders, 2019, 58, 50-55.	2.2	20
77	Involuntary Thumb Flexion on Neurological Examination: An Unusual Form of Upper Limb Dystonia in the Faroe Islands. Tremor and Other Hyperkinetic Movements, 2019, 9, .	2.0	2
78	Improving neurophysiological biomarkers for functional myoclonic movements. Parkinsonism and Related Disorders, 2018, 51, 3-8.	2.2	16
79	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
80	Electrophysiologic testing aids diagnosis and subtyping of myoclonus. Neurology, 2018, 90, e647-e657.	1.1	31
81	Fever-Induced Paroxysmal Weakness and Encephalopathy (FIPWE)â€"Part of a Phenotypic Continuum in Patients With ATP1A3 Mutations?. Pediatric Neurology, 2018, 81, 57-58.	2.1	7
82	Cognition in childhood dystonia: a systematic review. Developmental Medicine and Child Neurology, 2018, 60, 244-255.	2.1	7
83	The interrelation between clinical presentation and neurophysiology of posthypoxic myoclonus. Annals of Clinical and Translational Neurology, 2018, 5, 386-396.	3.7	7
84	Physiological movement disorder-like features during typical motor development. European Journal of Paediatric Neurology, 2018, 22, 595-601.	1.6	12
85	Reproducibility of standardized fine motor control tasks and age effects in healthy adults. Measurement: Journal of the International Measurement Confederation, 2018, 114, 177-184.	5.0	4
86	Quality and reporting of guidelines on the diagnosis and management of dystonia. European Journal of Neurology, 2018, 25, 275-283.	3.3	9
87	Wavelet coherence analysis: A new approach to distinguish organic and functional tremor types. Clinical Neurophysiology, 2018, 129, 13-20.	1.5	20
88	Response to â€˜Classification of cerebral palsy and potential role of video recordingâ€™. European Journal of Paediatric Neurology, 2018, 22, 211-212.	1.6	0
89	The premotor syndrome of cervical dystonia: Disordered processing of salient environmental stimuli. Movement Disorders, 2018, 33, 232-237.	3.9	13
90	Eye movement disorders and neurological symptoms in lateâ€onset inborn errors of metabolism. Movement Disorders, 2018, 33, 1844-1856.	3.9	12

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91	Distinctive tics suppression network in Gilles de la Tourette syndrome distinguished from suppression of natural urges using multimodal imaging. NeuroImage: Clinical, 2018, 20, 783-792.	2.7	29
92	Expanding the ADCY5 phenotype toward spastic paraparesis. Neurology: Genetics, 2018, 4, e214.	1.9	11
93	Non-motor effects of deep brain stimulation in dystonia: A systematic review. Parkinsonism and Related Disorders, 2018, 55, 26-44.	2.2	22
94	Toward adaptive deep brain stimulation for dystonia. Neurosurgical Focus, 2018, 45, E3.	2.3	38
95	Relationships between Serotonin Transporter Binding in the Raphe Nuclei, Basal Ganglia, and Hippocampus with Clinical Symptoms in Cervical Dystonia: A [11C]DASB Positron Emission Tomography Study. Frontiers in Neurology, 2018, 9, 88.	2.4	18
96	Crossing barriers: a multidisciplinary approach to children and adults with young-onset movement disorders. Journal of Clinical Movement Disorders, 2018, 5, 3.	2.2	10
97	Progressive myoclonus ataxia: Time for a new definition?. Movement Disorders, 2018, 33, 1281-1286.	3.9	18
98	Fatigue, not self-rated motor symptom severity, affects quality of life in functional motor disorders. Journal of Neurology, 2018, 265, 1803-1809.	3.6	48
99	Reversal of Status Dystonicus after Relocation of Pallidal Electrodes in DYT6 Generalized Dystonia. Tremor and Other Hyperkinetic Movements, 2018, 8, 530.	2.0	9
100	Dynamic head-neck stabilization in cervical dystonia. Clinical Biomechanics, 2017, 42, 120-127.	1.2	1
101	Using the shared genetics of dystonia and ataxia to unravel their pathogenesis. Neuroscience and Biobehavioral Reviews, 2017, 75, 22-39.	6.1	41
102	Clinician and patient perceptions of free will in movement disorders: mind the gap. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 532-533.	1.9	5
103	A post hoc study on gene panel analysis for the diagnosis of dystonia. Movement Disorders, 2017, 32, 569-575.	3.9	59
104	Spasticity, dyskinesia and ataxia in cerebral palsy: Are we sure we can differentiate them?. European Journal of Paediatric Neurology, 2017, 21, 703-706.	1.6	34
105	Clinical decision-making in functional and hyperkinetic movement disorders. Neurology, 2017, 88, 118-123.	1.1	9
106	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
107	A review of psychiatric co-morbidity described in genetic and immune mediated movement disorders. Neuroscience and Biobehavioral Reviews, 2017, 80, 23-35.	6.1	11
108	Adaptive DBS in a Parkinson's patient with chronically implanted DBS: A proof of principle. Movement Disorders, 2017, 32, 1253-1254.	3.9	73

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109	Startle responses in functional jerky movement disorders are increased but have a normal pattern. Parkinsonism and Related Disorders, 2017, 40, 27-32.	2.2	17
110	The efficacy of the modified Atkins diet in North Sea Progressive Myoclonus Epilepsy: an observational prospective open-label study. Orphanet Journal of Rare Diseases, 2017, 12, 45.	2.7	11
111	The relationship between the dopaminergic system and depressive symptoms in cervical dystonia. European Journal of Nuclear Medicine and Molecular Imaging, 2017, 44, 1375-1382.	6.4	26
112	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
113	Non-motor symptoms and quality of life in dopa-responsive dystonia patients. Parkinsonism and Related Disorders, 2017, 45, 57-62.	2.2	19
114	The presence of depression and anxiety do not distinguish between functional jerks and cortical myoclonus. Parkinsonism and Related Disorders, 2017, 45, 90-93.	2.2	8
115	Dystonic neck muscles show a shift in relative autospectral power during isometric contractions. Clinical Neurophysiology, 2017, 128, 1937-1945.	1.5	5
116	Î´-Catenin (CTNND2) missense mutation in familial cortical myoclonic tremor and epilepsy. Neurology, 2017, 89, 2341-2350.	1.1	22
117	Clinical characterization of dystonia in adult patients with Huntington's disease. European Journal of Neurology, 2017, 24, 1140-1147.	3.3	33
118	Myoclonus subtypes in tertiary referral center. Cortical myoclonus and functional jerks are common. Clinical Neurophysiology, 2017, 128, 253-259.	1.5	28
119	Dystoniaâ€deafness syndrome caused by a Î²â€actin gene mutation and response to deep brain stimulation. Movement Disorders, 2017, 32, 162-165.	3.9	13
120	Spectral EMG Changes in Cervical Dystonia Patients and the Influence of Botulinum Toxin Treatment. Toxins, 2017, 9, 256.	3.4	5
121	Clinical Practice: Evidence-Based Recommendations for the Treatment of Cervical Dystonia with Botulinum Toxin. Frontiers in Neurology, 2017, 8, 35.	2.4	63
122	Blink rate is associated with drug-induced parkinsonism in patients with severe mental illness, but does not meet requirements to serve as a clinical test: the Curacao extrapyramidal syndromes study XIII. Journal of Negative Results in BioMedicine, 2017, 16, 15.	1.4	1
123	Lower serotonin transporter binding in patients with cervical dystonia is associated with psychiatric symptoms. EJNMMI Research, 2017, 7, 87.	2.5	18
124	Effect of Antipsychotic Type and Dose Changes on Tardive Dyskinesia and Parkinsonism Severity in Patients With a Serious Mental Illness. Journal of Clinical Psychiatry, 2017, 78, e279-e285.	2.2	33
125	The Inter-rater Variability of Clinical Assessment in Post-anoxic Myoclonus. Tremor and Other Hyperkinetic Movements, 2017, 7, 470.	2.0	5
126	Risk Factors for Tremor in a Population of Patients with Severe Mental Illness: An 18-year Prospective Study in a Geographically Representative Sample (The Curacao Extrapyramidal Syndromes Study XI). Tremor and Other Hyperkinetic Movements, 2017, 7, 468.	2.0	0

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127	Distribution and Coexistence of Myoclonus and Dystonia as Clinical Predictors of SGCE Mutation Status: A Pilot Study. Frontiers in Neurology, 2016, 7, 72.	2.4	9
128	Unmet Needs in the Management of Cervical Dystonia. Frontiers in Neurology, 2016, 7, 165.	2.4	20
129	The Burkeâ€Fahnâ€Marsden Dystonia Rating Scale is Ageâ€Dependent in Healthy Children. Movement Disorders Clinical Practice, 2016, 3, 580-586.	1.5	19
130	Validation of â€œlaboratoryâ€supportedâ€•criteria for functional (psychogenic) tremor. Movement Disorders, 2016, 31, 555-562.	3.9	86
131	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
132	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
133	Serotonergic perturbations in dystonia disordersâ€”a systematic review. Neuroscience and Biobehavioral Reviews, 2016, 65, 264-275.	6.1	24
134	Ataxia, dystonia and myoclonus in adult patients with Niemann-Pick type C. Orphanet Journal of Rare Diseases, 2016, 11, 121.	2.7	29
135	Patience is the key: Contraceptive induced chorea in a girl with Down Syndrome. European Journal of Paediatric Neurology, 2016, 20, 671-673.	1.6	1
136	Event related desynchronisation predicts functional propriospinal myoclonus. Parkinsonism and Related Disorders, 2016, 31, 116-118.	2.2	13
137	The Symptomatic Treatment of Acquired Dystonia: A Systematic Review. Movement Disorders Clinical Practice, 2016, 3, 548-558.	1.5	10
138	Psychiatric disorders, myoclonus dystonia and <scp>SGCE</scp> : an international study. Annals of Clinical and Translational Neurology, 2016, 3, 4-11.	3.7	43
139	Neurometabolic disorders are treatable causes of dystonia. Revue Neurologique, 2016, 172, 455-464.	1.5	8
140	Clinical Pearls - how my patients taught me: The fainting lark symptom. Journal of Clinical Movement Disorders, 2016, 3, 16.	2.2	1
141	Determinants of disability in cervical dystonia. Parkinsonism and Related Disorders, 2016, 32, 48-53.	2.2	58
142	How typical are â€˜typicalâ€™ tremor characteristics? Sensitivity and specificity of five tremor phenomena. Parkinsonism and Related Disorders, 2016, 30, 23-28.	2.2	48
143	Cerebellar Atrophy in Cortical Myoclonic Tremor and Not in Hereditary Essential Tremorâ€”a Voxel-Based Morphometry Study. Cerebellum, 2016, 15, 696-704.	2.5	34
144	Reliability of phenotypic earlyâ€onset ataxia assessment: a pilot study. Developmental Medicine and Child Neurology, 2016, 58, 70-76.	2.1	13

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145	Electroencephalographic Findings in Posthypoxic Myoclonus. Journal of Intensive Care Medicine, 2016, 31, 270-275.	2.8	31
146	Cortical Myoclonus in a Young Boy with <scp>GOSR</scp>2 Mutation Mimics Chorea. Movement Disorders Clinical Practice, 2015, 2, 61-63.	1.5	7
147	RELN rare variants in myoclonusâ€dystonia. Movement Disorders, 2015, 30, 415-419.	3.9	27
148	Movement Disorders and Psychosis, a Complex Marriage. Frontiers in Psychiatry, 2015, 5, 190.	2.6	10
149	Tic related local field potentials in the thalamus and the effect of deep brain stimulation in Tourette syndrome: Report of three cases. Clinical Neurophysiology, 2015, 126, 1578-1588.	1.5	36
150	Genetic advances spark a revolution in dystonia phenotyping. Nature Reviews Neurology, 2015, 11, 78-79.	10.1	5
151	Usefulness of intermuscular coherence and cumulant analysis in the diagnosis of postural tremor. Clinical Neurophysiology, 2015, 126, 1564-1569.	1.5	24
152	Non-motor symptoms in genetically defined dystonia: Homogenous groups require systematic assessment. Parkinsonism and Related Disorders, 2015, 21, 1031-1040.	2.2	31
153	Myoclonus in childhood-onset neurogenetic disorders: The importance of early identification and treatment. European Journal of Paediatric Neurology, 2015, 19, 726-729.	1.6	20
154	A novel diagnostic approach to patients with myoclonus. Nature Reviews Neurology, 2015, 11, 687-697.	10.1	67
155	Dystonia in children and adolescents: a systematic review and a new diagnostic algorithm. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 774-781.	1.9	124
156	CACNA1B mutation is linked to unique myoclonus-dystonia syndrome. Human Molecular Genetics, 2015, 24, 987-993.	2.9	70
157	Reticular Myoclonus: It Really Comes From the Brainstem!. Movement Disorders Clinical Practice, 2014, 1, 258-260.	1.5	10
158	Propriospinal myoclonus. Neurology, 2014, 83, 1862-1870.	1.1	162
159	Assessment of speech in early-onset ataxia: a pilot study. Developmental Medicine and Child Neurology, 2014, 56, 1202-1206.	2.1	7
160	DRD1 rare variants associated with tardive-like dystonia: A pilot pathway sequencing study in dystonia. Parkinsonism and Related Disorders, 2014, 20, 782-785.	2.2	7
161	Ramsay hunt syndrome: Clinical characterization of progressive myoclonus ataxia caused by GOSR2 mutation. Movement Disorders, 2014, 29, 139-143.	3.9	113
162	Latah: An indonesian startle syndrome. Movement Disorders, 2013, 28, 370-379.	3.9	25

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163	The eye of the beholder: inter-rater agreement among experts on psychogenic jerky movement disorders. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 742-747.	1.9	56
164	Is TOR1A a risk factor in adultâ€onset primary torsion dystonia?. Movement Disorders, 2013, 28, 827-831.	3.9	14
165	Phenotypes and genetic architecture of focal primary torsion dystonia. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 1006-1011.	1.9	22
166	White matter abnormalities in geneâ€positive myoclonusâ€dystonia. Movement Disorders, 2012, 27, 1666-1672.	3.9	34
167	Cognition and psychopathology in myoclonus-dystonia. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 814-820.	1.9	58
168	The startle syndromes: Physiology and treatment. Epilepsia, 2012, 53, 3-11.	5.1	76
169	A new hyperekplexia family with a recessive frameshift mutation in the GLRA1 gene. Movement Disorders, 2012, 27, 795-796.	3.9	5
170	Association of BDNF Met66Met polymorphism with arm tremor in cervical dystonia. Movement Disorders, 2012, 27, 796-797.	3.9	13
171	THAP1 mutations are infrequent in spasmodic dysphonia. Movement Disorders, 2011, 26, 1952-1954.	3.9	12
172	Management of patients with myoclonus: available therapies and the need for an evidence-based approach. Lancet Neurology, The, 2010, 9, 1028-1036.	10.2	76
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174	Myoclonus-dystonia and spinocerebellar ataxia type 14 presenting with similar phenotypes: Trunk tremor, myoclonus, and dystonia. Parkinsonism and Related Disorders, 2010, 16, 288-289.	2.2	21
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