

# Kailash Phatechand Bhatia

## List of Publications by Year in descending order

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507  
papers

34,314  
citations

5126

86  
h-index

7043

159  
g-index

520  
all docs

520  
docs citations

520  
times ranked

28155  
citing authors

#	ARTICLE	IF	CITATIONS
1	Parkinsonism and dystonia: Clinical spectrum and diagnostic clues. Journal of the Neurological Sciences, 2022, 433, 120016.	0.3	8
2	Dissecting the Phenotype and Genotype of <i>PLA2G6</i> -Related Parkinsonism. Movement Disorders, 2022, 37, 148-161.	2.2	32
3	Heterozygous <i>EIF2AK2</i> Variant Causes Adolescence-Onset Generalized Dystonia Partially Responsive to DBS. Movement Disorders Clinical Practice, 2022, 9, 268-271.	0.8	7
4	A Note of Caution on Distorted Visual Feedback as a Treatment for Functional Movement Disorders. Movement Disorders Clinical Practice, 2022, 9, 275-277.	0.8	1
5	Homer's Antibody Disease: A Potentially Treatable MSA Mimic. Movement Disorders Clinical Practice, 2022, 9, 178-182.	0.8	10
6	Biallelic Loss of Function NDUFA12 Variants Cause a Wide Phenotypic Spectrum from Leigh/Leigh-Like Syndrome to Isolated Optic Atrophy. Movement Disorders Clinical Practice, 2022, 9, 218-228.	0.8	5
7	Cerebellar and Midbrain Lysosomal Enzyme Deficiency in Isolated Dystonia. Movement Disorders, 2022, 37, 875-877.	2.2	1
8	Altered pituitary morphology as a sign of benign hereditary chorea caused by TITF1/NKX2.1 mutations. Neurogenetics, 2022, 23, 91.	0.7	2
9	Restless Legs Syndrome: Known Knowns and Known Unknowns. Brain Sciences, 2022, 12, 118.	1.1	13
10	HOPS-Associated Neurological Disorders: Lysosomal Dysfunction as an Emerging Concept Underlying Dystonia. Movement Disorders Clinical Practice, 2022, 9, 452-453.	0.8	0
11	Motor Cortical Network Excitability in Parkinson's Disease. Movement Disorders, 2022, 37, 734-744.	2.2	19
12	Milestones in Tremor Research: 10 Years Later. Movement Disorders Clinical Practice, 2022, 9, 429-435.	0.8	19
13	Reply to: Juvenile <i>PLA2G6</i> -parkinsonism due to Indian "Asian" p.R741Q mutation, and response to STN DBS. Movement Disorders, 2022, 37, 658-662.	2.2	5
14	A Critical Investigation of Cerebellar Associative Learning in Isolated Dystonia. Movement Disorders, 2022, 37, 1187-1192.	2.2	8
15	The MDS consensus tremor classification: The best way to classify patients with tremor at present. Journal of the Neurological Sciences, 2022, 435, 120191.	0.3	10
16	Patients' Postjudice of Tele-Neurology for Movement Disorders. Movement Disorders Clinical Practice, 2022, 9, 446-451.	0.8	3
17	Ethnic Differences in Dystonia Prevalence and Phenotype. Movement Disorders, 2022, 37, 1323-1325.	2.2	4
18	Biallelic variants in <i>ZNF142</i> lead to a syndromic neurodevelopmental disorder. Clinical Genetics, 2022, 102, 98-109.	1.0	6

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19	029â€¦ Postural instability in DYT-TOR1A dystonia dynamically dependent on sensory feedback. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, A110.1-A110.	0.9	0
20	Diagnosing Premotor Multiple System Atrophy. Neurology, 2022, 99, .	1.5	4
21	Lateâ€œOnset Chorea in JAK2 â€œAssociated Essential Thrombocythemia. Movement Disorders Clinical Practice, 2021, 8, 145-148.	0.8	5
22	Reversal of Temporal Discrimination in Cervical Dystonia after Lowâ€œFrequency Sensory Stimulation. Movement Disorders, 2021, 36, 761-766.	2.2	11
23	Defective Somatosensory Inhibition and Plasticity Are Not Required to Develop Dystonia. Movement Disorders, 2021, 36, 1015-1021.	2.2	17
24	Symptomâ€œTriggered Attention to Self as a Possible Trigger of Functional Comorbidity. Movement Disorders Clinical Practice, 2021, 8, 159-161.	0.8	5
25	Variability of Movement Disorders: The Influence of Sensation, Action, Cognition, and Emotions. Movement Disorders, 2021, 36, 581-593.	2.2	14
26	Self-concocted, curious and creative coping strategies in movement disorders. Parkinsonism and Related Disorders, 2021, 83, 140-143.	1.1	2
27	Non-invasive suppression of essential tremor via phase-locked disruption of its temporal coherence. Nature Communications, 2021, 12, 363.	5.8	50
28	Huntington disease-like phenotype in a patient with ANO3 mutation. Parkinsonism and Related Disorders, 2021, 90, 120-122.	1.1	5
29	Exploring Interrater Disagreement on Essential Tremor Using a Standardized Tremor Elements Assessment. Movement Disorders Clinical Practice, 2021, 8, 371-376.	0.8	15
30	Throatâ€œClearing Vocalizations in Primary Brain Calcification Syndromes. Movement Disorders Clinical Practice, 2021, 8, 627-630.	0.8	1
31	The Signature of Primary Writing Tremor Is Dystonic. Movement Disorders, 2021, 36, 1715-1720.	2.2	16
32	Exploratory pilot study of exogenous sustainedâ€œrelease melatonin on nocturia in Parkinson's disease. European Journal of Neurology, 2021, 28, 1884-1892.	1.7	19
33	The Phenomenon of Exquisite Motor Control in Tic Disorders and its Pathophysiological Implications. Movement Disorders, 2021, 36, 1308-1315.	2.2	7
34	A geroscience approach for Parkinsonâ€™s disease: Conceptual framework and design of PROPAG-AGEING project. Mechanisms of Ageing and Development, 2021, 194, 111426.	2.2	14
35	No increased suggestibility to placebo in functional neurological disorder. European Journal of Neurology, 2021, 28, 2367-2371.	1.7	4
36	Expanding the Spectrum of Movement Disorders Associated With <i>C9orf72</i> Hexanucleotide Expansions. Neurology: Genetics, 2021, 7, e575.	0.9	20

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37	Challenges in Clinicogenetic Correlations: One Gene “ Many Phenotypes. Movement Disorders Clinical Practice, 2021, 8, 299-310.	0.8	34
38	Management of Secondary Poor Response to Botulinum Toxin in Cervical Dystonia: A Multicenter Audit. Movement Disorders Clinical Practice, 2021, 8, 541-545.	0.8	3
39	Reply to Comment on: Voluntary Inhibitory Control of Chorea. Movement Disorders Clinical Practice, 2021, 8, 636-636.	0.8	0
40	Paroxysmal, exercise-induced, diurnally fluctuating dystonia: Expanding the phenotype of SPG8. Parkinsonism and Related Disorders, 2021, 85, 26-28.	1.1	3
41	Worldwide barriers to genetic testing for movement disorders. European Journal of Neurology, 2021, 28, 1901-1909.	1.7	21
42	Biallelic variants in TSPOAP1, encoding the active-zone protein RIMBP1, cause autosomal recessive dystonia. Journal of Clinical Investigation, 2021, 131, .	3.9	18
43	Autoimmune movement disorders with neuronal antibodies “ an update. Current Opinion in Neurology, 2021, 34, 565-571.	1.8	9
44	Dysphagia in multiple system atrophy consensus statement on diagnosis, prognosis and treatment. Parkinsonism and Related Disorders, 2021, 86, 124-132.	1.1	22
45	Dystonia in a Female Fragile X Premutation Carrier. Movement Disorders Clinical Practice, 2021, 8, 797-799.	0.8	1
46	Movement disorders in systemic autoimmune diseases: Clinical spectrum, ancillary investigations, pathophysiological considerations. Parkinsonism and Related Disorders, 2021, 88, 116-128.	1.1	10
47	Movement Disorders and Liver Disease. Movement Disorders Clinical Practice, 2021, 8, 828-842.	0.8	7
48	Xâ€Linked Parkinsonism: Phenotypic and Genetic Heterogeneity. Movement Disorders, 2021, 36, 1511-1525.	2.2	10
49	â€Antibody of Unknown Significanceâ€(AUS): The Issue of Interpreting Antibody Test Results. Movement Disorders, 2021, 36, 1543-1547.	2.2	11
50	Misdirected attentional focus in functional tremor. Brain, 2021, 144, 3436-3450.	3.7	15
51	A practical guide to troubleshooting pallidal deep brain stimulation issues in patients with dystonia. Parkinsonism and Related Disorders, 2021, 87, 142-154.	1.1	1
52	The Emerging Role of Phosphodiesterases in Movement Disorders. Movement Disorders, 2021, 36, 2225-2243.	2.2	21
53	Development of parkinsonism after long-standing cervical dystonia “ A cohort. Journal of the Neurological Sciences, 2021, 427, 117477.	0.3	10
54	From Collar to Coccyx: Truncal Movement Disorders: A Clinical Review. Movement Disorders Clinical Practice, 2021, 8, 1027-1033.	0.8	2

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55	Reply to: "A Primary Writing Tremor Is a Form of Dystonic Tremor: Is the Debate Settled?" Movement Disorders, 2021, 36, 1996-1997.	2.2	0
56	Commentary: <scp>Andersonâ€Fabry</scp> Disease: A Rare Cause of Levodopaâ€Responsive Early Onset Parkinsonism. Movement Disorders Clinical Practice, 2021, 8, S35-S36.	0.8	0
57	Standing on the Shoulders of Giants: The Most Relevant Papers in Movement Disorders Field from the Second Half of the 20th Century. Movement Disorders Clinical Practice, 2021, 8, 992-992.	0.8	0
58	Two forms of short-interval intracortical inhibition in human motor cortex. Brain Stimulation, 2021, 14, 1340-1352.	0.7	16
59	Heterogeneity of prodromal Parkinson symptoms in siblings of Parkinson disease patients. Npj Parkinson's Disease, 2021, 7, 78.	2.5	2
60	Reply to: Comparing <scp>VUS</scp> and <scp>AUS</scp>: Parallels and Differences in Neurogenetics and Neuroimmunology. Movement Disorders, 2021, 36, 2454-2456.	2.2	0
61	Paroxysmal Dyskinesia: Definitions and Clinical Approach. , 2021, , 1-5.		0
62	Other Paroxysmal Movement Disorders. , 2021, , 119-124.		0
63	Validation of the Movement Disorder Society Criteria for the Diagnosis of "Repeat Tauopathies. Movement Disorders, 2020, 35, 171-176.	2.2	37
64	Risk of Developing Parkinson Disease in Bipolar Disorder. JAMA Neurology, 2020, 77, 192.	4.5	42
65	Temporal Discrimination is Altered in Patients With Isolated Asymmetric and Jerky Upper Limb Tremor. Movement Disorders, 2020, 35, 306-315.	2.2	17
66	Ciliary Dysfunction: The Hairy Explanation of Normal Pressure Hydrocephalus?. Movement Disorders Clinical Practice, 2020, 7, 30-31.	0.8	1
67	The Flip Side of Distractibilityâ€Executive Dysfunction in Functional Movement Disorders. Frontiers in Neurology, 2020, 11, 969.	1.1	9
68	Video-tutorial for the Movement Disorder Society criteria for progressive supranuclear palsy. Parkinsonism and Related Disorders, 2020, 78, 200-203.	1.1	8
69	Modulation of Reaction Times and Sense of Agency via Subliminal Priming in Functional Movement Disorders. Frontiers in Neurology, 2020, 11, 989.	1.1	3
70	Reply: Pentameric repeat expansions: cortical myoclonus or cortical tremor? and Cortical tremor: a tantalizing conundrum between cortex and cerebellum. Brain, 2020, 143, e88-e88.	3.7	1
71	Botulinum Neurotoxin-A Injection in Adult Cervical Dystonia and Spastic Paresis: Results From the INPUT (INjection Practice, Usage and Training) Survey. Frontiers in Neurology, 2020, 11, 570671.	1.1	4
72	The Need to Tic. Movement Disorders Clinical Practice, 2020, 7, 863-864.	0.8	1

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73	A new family with GLRB-related hyperekplexia showing chorea in homo- and heterozygous variant carriers. <i>Parkinsonism and Related Disorders</i> , 2020, 79, 97-99.	1.1	4
74	Psychiatric Manifestations of <sc><i>ATP13A2</i></sc> Mutations. <i>Movement Disorders Clinical Practice</i> , 2020, 7, 838-841.	0.8	6
75	Criss-cross gait. <i>Neurology</i> , 2020, 95, 500-501.	1.5	5
76	Delineating the electrophysiological signature of dystonia. <i>Experimental Brain Research</i> , 2020, 238, 1685-1692.	0.7	25
77	Toward an Early Real-time Quaking-induced Conversion-based Diagnostic Biomarker for Lewy Body-related Synucleinopathies. <i>Movement Disorders Clinical Practice</i> , 2020, 7, 780-781.	0.8	0
78	Telemedicine in Movement Disorders: Leçons du COVID-19. <i>Movement Disorders</i> , 2020, 35, 1893-1896.	2.2	24
79	Dystonia genes functionally converge in specific neurons and share neurobiology with psychiatric disorders. <i>Brain</i> , 2020, 143, 2771-2787.	3.7	50
80	Bilateral polymicrogyria associated with dystonia: A new neurogenetic syndrome?. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 2207-2213.	0.7	0
81	Reply to: "A New Day: The Role of Telemedicine in Reshaping Care for Persons With Movement Disorders". <i>Movement Disorders</i> , 2020, 35, 1903-1904.	2.2	9
82	The CloudUPDRS smartphone software in Parkinson's study: cross-validation against blinded human raters. <i>Npj Parkinson's Disease</i> , 2020, 6, 36.	2.5	18
83	Opicapone Efficacy and Tolerability in Parkinson's Disease Patients Reporting Insufficient Benefit/Failure of Entacapone. <i>Movement Disorders Clinical Practice</i> , 2020, 7, 955-960.	0.8	6
84	<i>KMT2B</i>-related disorders: expansion of the phenotypic spectrum and long-term efficacy of deep brain stimulation. <i>Brain</i> , 2020, 143, 3242-3261.	3.7	57
85	Unravelling the enigma of cortical tremor and other forms of cortical myoclonus. <i>Brain</i> , 2020, 143, 2653-2663.	3.7	38
86	Self-injurious behaviour in movement disorders: systematic review. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020, 91, 712-719.	0.9	19
87	Tardive syndromes. <i>Practical Neurology</i> , 2020, 20, 368-376.	0.5	10
88	<i>MYORG</i>-related disease is associated with central pontine calcifications and atypical parkinsonism. <i>Neurology: Genetics</i> , 2020, 6, e399.	0.9	13
89	Isolated and combined genetic tremor syndromes: a critical appraisal based on the 2018 MDS criteria. <i>Parkinsonism and Related Disorders</i> , 2020, 77, 121-140.	1.1	13
90	Some New and Unexpected Tauopathies in Movement Disorders. <i>Movement Disorders Clinical Practice</i> , 2020, 7, 616-626.	0.8	13

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91	Movement Disorders in the World of <sc>COVID</sc>â€19. Movement Disorders, 2020, 35, 709-710.	2.2	27
92	Voluntary Inhibitory Control of Chorea: A Case Series. Movement Disorders Clinical Practice, 2020, 7, 308-312.	0.8	6
93	Antibody-related movement disorders â€ a comprehensive review of phenotype-autoantibody correlations and a guide to testing. Neurological Research and Practice, 2020, 2, 6.	1.0	21
94	Impaired automatic but intact volitional inhibition in primary tic disorders. Brain, 2020, 143, 906-919.	3.7	35
95	Huntington disease like 2 (HDL-2) with parkinsonism and abnormal DAT-SPECT â€ A novel observation. Parkinsonism and Related Disorders, 2020, 71, 46-48.	1.1	4
96	Reply to J. Dulski and J. Slawek's â€Fibrodysplasia ossificans progressiva as a form of pseudodystoniaâ€. Parkinsonism and Related Disorders, 2020, 71, 49-50.	1.1	0
97	Movement Disorders in the World of <sc>COVID</sc>â€19. Movement Disorders Clinical Practice, 2020, 7, 355-356.	0.8	18
98	Treatment of Paroxysmal Dyskinesia. Neurologic Clinics, 2020, 38, 433-447.	0.8	16
99	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.	0.9	71
100	Bringing order to higher order motor disorders. Journal of Neurology, 2019, 266, 797-805.	1.8	15
101	Reply to: Tics in Paroxysmal Kinesigenic Dyskinesia. Movement Disorders Clinical Practice, 2019, 6, 504-505.	0.8	0
102	ADCY5â€Related Dyskinesia: Improving Clinical Detection of an Evolving Disorder. Movement Disorders Clinical Practice, 2019, 6, 512-520.	0.8	31
103	A case of congenital hypoplasia of the left cerebellar hemisphere and ipsilateral cortical myoclonus. Movement Disorders, 2019, 34, 1745-1747.	2.2	12
104	IRF2BPL mutations cause autosomal dominant dystonia with anarthria, slow saccades and seizures. Parkinsonism and Related Disorders, 2019, 68, 57-59.	1.1	15
105	Management of Spastic Paresis and Cervical Dystonia: Access to Therapeutic Innovations Through an International Program of Practical Courses. Clinical Therapeutics, 2019, 41, 2321-2330.e4.	1.1	5
106	The Gut Microbiome: A Therapeutically Targetable Site of Peripheral Levodopa Metabolism. Movement Disorders Clinical Practice, 2019, 6, 547-548.	0.8	8
107	Validation of a selfâ€completed Dystonia Nonâ€Motor Symptoms Questionnaire. Annals of Clinical and Translational Neurology, 2019, 6, 2054-2065.	1.7	20
108	Twenty years on: Myoclonusâ€dystonia and Î¼â€sarcoglycan â€ neurodevelopment, channel, and signaling dysfunction. Movement Disorders, 2019, 34, 1588-1601.	2.2	31

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109	Sex differences in Parkinson's disease: A transcranial magnetic stimulation study. <i>Movement Disorders</i> , 2019, 34, 1873-1881.	2.2	14
110	The spectrum of involuntary vocalizations in humans: A video atlas. <i>Movement Disorders</i> , 2019, 34, 1774-1791.	2.2	24
111	Tics and functional tic-like movements. <i>Neurology</i> , 2019, 93, 750-758.	1.5	89
112	The long-term outcome of impulsive compulsive behaviours in Parkinson's disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 1288-1289.	0.9	3
113	The interindividual variability of transcranial magnetic stimulation effects: Implications for diagnostic use in movement disorders. <i>Movement Disorders</i> , 2019, 34, 936-949.	2.2	44
114	Pain in cervical dystonia: Evidence of abnormal inhibitory control. <i>Parkinsonism and Related Disorders</i> , 2019, 65, 252-255.	1.1	35
115	A Novel SGCE Nonsense Variant Associated With Marked Intrafamilial Variability in a Turkish Family With Myoclonus-Dystonia. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 479-482.	0.8	3
116	Benign tremulous parkinsonism of the young-consider Parkin. <i>Parkinsonism and Related Disorders</i> , 2019, 65, 270-271.	1.1	5
117	Dystonia and Parkinson's disease: What is the relationship?. <i>Neurobiology of Disease</i> , 2019, 132, 104462.	2.1	71
118	Network localization of cervical dystonia based on causal brain lesions. <i>Brain</i> , 2019, 142, 1660-1674.	3.7	160
119	Paroxysmal Asymmetric Dystonic Arm Posturing: A Less Recognized but Characteristic Manifestation of ATP1A3-related disease. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 312-315.	0.8	15
120	Syringomyelia-Associated Dystonia: Case Series, Literature Review, and Novel Insights. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 387-392.	0.8	6
121	The use of transcranial magnetic stimulation as a treatment for movement disorders: A critical review. <i>Movement Disorders</i> , 2019, 34, 769-782.	2.2	48
122	Delineating the phenotype of autosomal recessive HPCA mutations: Not only isolated dystonia!. <i>Movement Disorders</i> , 2019, 34, 589-592.	2.2	10
123	Dystonia in Handcuffs: A Picture Typical of Lesch-Nyhan Syndrome. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 612-613.	0.8	0
124	How to apply the movement disorder society criteria for diagnosis of progressive supranuclear palsy. <i>Movement Disorders</i> , 2019, 34, 1228-1232.	2.2	93
125	A New Year and a New Era for MDCP. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 95-95.	0.8	0
126	Stimulus Sensitive Foot Myoclonus: A Clue to Coeliac Disease. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 320-323.	0.8	6



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127	Animals in the Brain. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 189-198.	0.8	8
128	Remission in dystonia â€“ Systematic review of the literature and meta-analysis. <i>Parkinsonism and Related Disorders</i> , 2019, 66, 9-15.	1.1	26
129	Abnormal DaTSCAN and Atypical Parkinsonism in SCA12. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 400-402.	0.8	5
130	Response from the Editors RE: DUOPAÂ® is an Excellent Alternative Treatment but with Some Caveats. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 338-338.	0.8	0
131	Pseudodystonia: A new perspective on an old phenomenon. <i>Parkinsonism and Related Disorders</i> , 2019, 62, 44-50.	1.1	21
132	Parkinsonism in essential tremor cases: A clinicopathological studyâ€”were they really essential tremor?. <i>Movement Disorders</i> , 2019, 34, 1749-1749.	2.2	0
133	How to approach a patient with parkinsonism â€“ red flags for atypical parkinsonism. <i>International Review of Neurobiology</i> , 2019, 149, 1-34.	0.9	4
134	Genetic mimics of the non-genetic atypical parkinsonian disorders â€“ the â€˜atypicalâ€™ atypical. <i>International Review of Neurobiology</i> , 2019, 149, 327-351.	0.9	8
135	Brachial Neuritis After Botulinum Toxin Injections for Cervical Dystonia: A Need for a Reappraisal?. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 160-165.	0.8	5
136	Ataxia with Oculomotor Apraxia Type 1â€”New Mutation, Characteristic Phenotype. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 265-266.	0.8	0
137	Combined Dystonia With Selfâ€“Mutilation in 6â€“Pyruvoylâ€“Tetrahydropterin Synthase (PTPS) Deficiency: A Case Report. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 81-82.	0.8	6
138	Sensory trick efficacy in cervical dystonia is linked to processing of neck proprioception. <i>Parkinsonism and Related Disorders</i> , 2019, 61, 50-56.	1.1	10
139	SPG7 : The Great Imitator of MSAâ€“ Within the ILOCAs. <i>Movement Disorders Clinical Practice</i> , 2019, 6, 174-175.	0.8	5
140	Unravelling of the paroxysmal dyskinesias. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 227-234.	0.9	57
141	Self-Injurious Behaviour in SCA17: A New Clinical Observation. <i>Tremor and Other Hyperkinetic Movements</i> , 2019, 9, .	1.1	3
142	Solving Mendelian Mysteries: The Non-coding Genome May Hold the Key. <i>Cell</i> , 2018, 172, 889-891.	13.5	9
143	High motor variability in DYT1 dystonia is associated with impaired visuomotor adaptation. <i>Scientific Reports</i> , 2018, 8, 3653.	1.6	26
144	Reappraising the role of motor surround inhibition in dystonia. <i>Journal of the Neurological Sciences</i> , 2018, 390, 178-183.	0.3	14

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145	Understanding the new tremor classification. <i>Movement Disorders</i> , 2018, 33, 1267-1268.	2.2	6
146	Cerebellar and brainstem functional abnormalities in patients with primary orthostatic tremor. <i>Movement Disorders</i> , 2018, 33, 1024-1025.	2.2	10
147	Oculomotor apraxia and disrupted sleep with nocturnal ballistic bouts in ADCY5-related disease. <i>Parkinsonism and Related Disorders</i> , 2018, 54, 103-106.	1.1	10
148	The distinguishing motor features of cataplexy: a study from video-recorded attacks. <i>Sleep</i> , 2018, 41, .	0.6	26
149	Functional lesional neurosurgery for tremor: back to the future?. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018, 89, 727-735.	0.9	15
150	Functional lesional neurosurgery for tremor: a systematic review and meta-analysis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018, 89, 717-726.	0.9	24
151	Genetic Dystoniaâ€tataxia Syndromes: Clinical Spectrum, Diagnostic Approach, and Treatment Options. <i>Movement Disorders Clinical Practice</i> , 2018, 5, 373-382.	0.8	21
152	Quick Flicks: Association of Paroxysmal Kinesigenic Dyskinesia and Tics. <i>Movement Disorders Clinical Practice</i> , 2018, 5, 317-320.	0.8	6
153	Functional neurological disorders in Parkinson disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018, 89, 566-571.	0.9	76
154	Movement disorders with neuronal antibodies: syndromic approach, genetic parallels and pathophysiology. <i>Brain</i> , 2018, 141, 13-36.	3.7	145
155	Pathogenesis of dystonia: is it of cerebellar or basal ganglia origin?. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018, 89, 488-492.	0.9	85
156	Treatable inherited rare movement disorders. <i>Movement Disorders</i> , 2018, 33, 21-35.	2.2	79
157	Movement disorders in genetically confirmed mitochondrial disease and the putative role of the cerebellum. <i>Movement Disorders</i> , 2018, 33, 146-155.	2.2	21
158	Reappraisal of cortical myoclonus: A retrospective study of clinical neurophysiology. <i>Movement Disorders</i> , 2018, 33, 339-341.	2.2	17
159	Consensus Statement on the classification of tremors. from the task force on tremor of the International Parkinson and Movement Disorder Society. <i>Movement Disorders</i> , 2018, 33, 75-87.	2.2	918
160	Cortical inhibitory function in cervical dystonia. <i>Clinical Neurophysiology</i> , 2018, 129, 466-472.	0.7	23
161	Development and clinimetric assessment of a nurse-administered screening tool for movement disorders in psychosis. <i>BJPsych Open</i> , 2018, 4, 404-410.	0.3	3
162	Complexity of the Genetics and Clinical Presentation of Spinocerebellar Ataxia 17. <i>Frontiers in Cellular Neuroscience</i> , 2018, 12, 429.	1.8	21

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163	Reply to: Young-onset multiple system atrophy. <i>Movement Disorders</i> , 2018, 33, 1975-1976.	2.2	1
164	PDE10A and ADCY5 mutations linked to molecular and microstructural basal ganglia pathology. <i>Movement Disorders</i> , 2018, 33, 1961-1965.	2.2	38
165	High frequency somatosensory stimulation in dystonia: Evidence for defective inhibitory plasticity. <i>Movement Disorders</i> , 2018, 33, 1902-1909.	2.2	43
166	Motor cortical excitability during voluntary inhibition of involuntary tic movements. <i>Movement Disorders</i> , 2018, 33, 1804-1809.	2.2	25
167	Delineating cerebellar mechanisms in DYT11 myoclonus-dystonia. <i>Movement Disorders</i> , 2018, 33, 1956-1961.	2.2	7
168	Dystonia. <i>Nature Reviews Disease Primers</i> , 2018, 4, 25.	18.1	223
169	Reply: "Reappraisal of cortical myoclonus: Electrophysiology is the gold standard" <i>Movement Disorders</i> , 2018, 33, 1191-1191.	2.2	2
170	Parkinsonian signs in patients with cervical dystonia treated with pallidal deep brain stimulation. <i>Brain</i> , 2018, 141, 3023-3034.	3.7	33
171	Young-onset multiple system atrophy: Clinical and pathological features. <i>Movement Disorders</i> , 2018, 33, 1099-1107.	2.2	30
172	Disease-related patterns of in vivo pathology in Corticobasal syndrome. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2018, 45, 2413-2425.	3.3	26
173	Sensitivity and Specificity of the ECAS in Parkinson's Disease and Progressive Supranuclear Palsy. <i>Parkinson's Disease</i> , 2018, 2018, 1-8.	0.6	11
174	Letter to the Editor. Errors in the meta-analysis of outcomes and complications of MRgFUS. <i>Neurosurgical Focus</i> , 2018, 45, E15.	1.0	2
175	Adult Periodic Alternating Nystagmus Masked by Involuntary Head Movements. <i>Frontiers in Neurology</i> , 2018, 9, 326.	1.1	2
176	Cervical dystonia: Normal auditory mismatch negativity and abnormal somatosensory mismatch negativity. <i>Clinical Neurophysiology</i> , 2018, 129, 1947-1954.	0.7	4
177	Partial loss-of-function of sodium channel SCN8A in familial isolated myoclonus. <i>Human Mutation</i> , 2018, 39, 965-969.	1.1	34
178	Neuroimaging advances in Parkinson's disease. <i>Current Opinion in Neurology</i> , 2018, 31, 415-424.	1.8	25
179	Exome Sequencing Identifies a Novel Homozygous Missense <i>ATP13A2</i> Mutation. <i>Movement Disorders Clinical Practice</i> , 2017, 4, 132-135.	0.8	7
180	Early Ataxia and Subsequent Parkinsonism: PLA2G6 Mutations Cause a Continuum Rather Than Three Discrete Phenotypes. <i>Movement Disorders Clinical Practice</i> , 2017, 4, 125-128.	0.8	16

#	ARTICLE	IF	CITATIONS
181	The epileptic and nonepileptic spectrum of paroxysmal dyskinesias: Channelopathies, synaptopathies, and transportopathies. <i>Movement Disorders</i> , 2017, 32, 310-318.	2.2	63
182	Essential pitfalls in "essential" tremor. <i>Movement Disorders</i> , 2017, 32, 325-331.	2.2	74
183	Unilateral cerebellothalamic tract ablation in essential tremor by MRI-guided focused ultrasound. <i>Neurology</i> , 2017, 88, 1329-1333.	1.5	51
184	Advances in the Clinical Differential Diagnosis of Parkinson's Disease. <i>International Review of Neurobiology</i> , 2017, 132, 79-127.	0.9	4
185	Which ante mortem clinical features predict progressive supranuclear palsy pathology?. <i>Movement Disorders</i> , 2017, 32, 995-1005.	2.2	121
186	Functional lesional neurosurgery for tremor—a protocol for a systematic review and meta-analysis. <i>BMJ Open</i> , 2017, 7, e015409.	0.8	7
187	Tremor stability index: a new tool for differential diagnosis in tremor syndromes. <i>Brain</i> , 2017, 140, 1977-1986.	3.7	103
188	Clinical diagnosis of progressive supranuclear palsy: The movement disorder society criteria. <i>Movement Disorders</i> , 2017, 32, 853-864.	2.2	1,402
189	Endophenotyping in idiopathic adult onset cervical dystonia. <i>Clinical Neurophysiology</i> , 2017, 128, 1142-1147.	0.7	12
190	High frequency somatosensory stimulation increases sensori-motor inhibition and leads to perceptual improvement in healthy subjects. <i>Clinical Neurophysiology</i> , 2017, 128, 1015-1025.	0.7	45
191	ADCY5-related movement disorders: Frequency, disease course and phenotypic variability in a cohort of paediatric patients. <i>Parkinsonism and Related Disorders</i> , 2017, 41, 37-43.	1.1	67
192	Deconstructing Fahr's disease/syndrome of brain calcification in the era of new genes. <i>Parkinsonism and Related Disorders</i> , 2017, 37, 1-10.	1.1	63
193	Mutations in the histone methyltransferase gene KMT2B cause complex early-onset dystonia. <i>Nature Genetics</i> , 2017, 49, 223-237.	9.4	186
194	"Atypical" atypical parkinsonism: Critical appraisal of a cohort. <i>Parkinsonism and Related Disorders</i> , 2017, 37, 36-42.	1.1	12
195	Pathophysiological heterogeneity in Parkinson's disease: Neurophysiological insights from LRRK2 mutations. <i>Movement Disorders</i> , 2017, 32, 1333-1335.	2.2	9
196	Causes of failure of pallidal deep brain stimulation in cases with pre-operative diagnosis of isolated dystonia. <i>Parkinsonism and Related Disorders</i> , 2017, 43, 38-48.	1.1	51
197	Nonmotor Features in Atypical Parkinsonism. <i>International Review of Neurobiology</i> , 2017, 134, 1285-1301.	0.9	21
198	Neurophysiological correlates of abnormal somatosensory temporal discrimination in dystonia. <i>Movement Disorders</i> , 2017, 32, 141-148.	2.2	67

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199	Current Opinions and Areas of Consensus on the Role of the Cerebellum in Dystonia. <i>Cerebellum</i> , 2017, 16, 577-594.	1.4	184
200	Neurodegeneration With Brain Iron Accumulation (NBIA) Syndromes Presenting With Late-Onset Craniocervical Dystonia: An Illustrative Case Series. <i>Movement Disorders Clinical Practice</i> , 2017, 4, 254-257.	0.8	6
201	Oculogyric crises: Etiology, pathophysiology and therapeutic approaches. <i>Parkinsonism and Related Disorders</i> , 2017, 36, 3-9.	1.1	67
202	PO081. Myoclonus dystonia and russell-silver syndrome in a patient with a microdeletion of chromosome 7q. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017, 88, A32.4-A33.	0.9	0
203	Clinical Practice: Evidence-Based Recommendations for the Treatment of Cervical Dystonia with Botulinum Toxin. <i>Frontiers in Neurology</i> , 2017, 8, 35.	1.1	63
204	iPSC-derived neuronal models of PANK2-associated neurodegeneration reveal mitochondrial dysfunction contributing to early disease. <i>PLoS ONE</i> , 2017, 12, e0184104.	1.1	39
205	Ixcellence Network: an international educational network to improve current practice in the management of cervical dystonia or spastic paresis by botulinum toxin injection. <i>Functional Neurology</i> , 2017, 32, 103.	1.3	7
206	The long-term outcome of orthostatic tremor. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, jnnp-2014-309942.	0.9	100
207	alpha synuclein and crystallin expression in human lens in parkinson's disease. <i>Movement Disorders</i> , 2016, 31, 600-601.	2.2	18
208	Transducer-based evaluation of tremor. <i>Movement Disorders</i> , 2016, 31, 1327-1336.	2.2	64
209	Validation of laboratory-supported criteria for functional (psychogenic) tremor. <i>Movement Disorders</i> , 2016, 31, 555-562.	2.2	86
210	PKD or Not PKD: That is the question. <i>Annals of Neurology</i> , 2016, 80, 167-168.	2.8	3
211	De Novo Mutations in PDE10A Cause Childhood-Onset Chorea with Bilateral Striatal Lesions. <i>American Journal of Human Genetics</i> , 2016, 98, 763-771.	2.6	96
212	Complex dystonia is not a category in the new 2013 consensus classification. <i>Movement Disorders</i> , 2016, 31, 1758-1759.	2.2	5
213	Mental rotation and working memory in musicians' dystonia. <i>Brain and Cognition</i> , 2016, 109, 124-129.	0.8	3
214	T1-weighted basal ganglia hyperintensities due to gadolinium deposition – a cautionary note. <i>Parkinsonism and Related Disorders</i> , 2016, 32, 135-136.	1.1	1
215	Knowledge gaps and research recommendations for essential tremor. <i>Parkinsonism and Related Disorders</i> , 2016, 33, 27-35.	1.1	46
216	SLC25A46 mutations underlie progressive myoclonic ataxia with optic atrophy and neuropathy. <i>Movement Disorders</i> , 2016, 31, 1249-1251.	2.2	49

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217	Genetic and phenotypic characterization of complex hereditary spastic paraplegia. <i>Brain</i> , 2016, 139, 1904-1918.	3.7	170
218	Conjugal Parkinson's disease – Real or chance?. <i>Parkinsonism and Related Disorders</i> , 2016, 33, 146-148.	1.1	1
219	Reply: Glial mitochondriopathy in infantile neuroaxonal dystrophy: pathophysiological and therapeutic implications. <i>Brain</i> , 2016, 139, e68-e68.	3.7	0
220	Hot topic: PNKP mutations cause ataxia with oculomotor apraxia type 4. <i>Movement Disorders</i> , 2016, 31, 500-500.	2.2	0
221	The role of polymyography in the treatment of cervical dystonia. <i>Journal of Neurology</i> , 2016, 263, 1663-1664.	1.8	6
222	Valproate-Associated Parkinsonism: A Critical Review of the Literature. <i>CNS Drugs</i> , 2016, 30, 527-540.	2.7	41
223	Axial Dystonia Mimicking Stiff Person Syndrome. <i>Movement Disorders Clinical Practice</i> , 2016, 3, 176-179.	0.8	7
224	Novel movement disorder society-Parkinson's disease criteria: What about SWEDD and genetic forms?. <i>Movement Disorders</i> , 2016, 31, 431-431.	2.2	3
225	What do patients with scans without evidence of dopaminergic deficit (SWEDD) have? New evidence and continuing controversies. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, 319-323.	0.9	186
226	The clinical syndrome of dystonia with anarthria/aphonia. <i>Parkinsonism and Related Disorders</i> , 2016, 24, 20-27.	1.1	10
227	Know thyself: Exploring interoceptive sensitivity in Parkinson's disease. <i>Journal of the Neurological Sciences</i> , 2016, 364, 110-115.	0.3	28
228	High frequency repetitive sensory stimulation improves temporal discrimination in healthy subjects. <i>Clinical Neurophysiology</i> , 2016, 127, 817-820.	0.7	21
229	Lower urinary tract dysfunction in patients with functional movement disorders. <i>Journal of the Neurological Sciences</i> , 2016, 361, 192-194.	0.3	12
230	From state dissociation to status dissociatus. <i>Sleep Medicine Reviews</i> , 2016, 28, 5-17.	3.8	56
231	Intermittent head drops: the differential spectrum. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, 414-419.	0.9	11
232	Why is there motor deterioration in Parkinson's disease during systemic infections-a hypothetical view. <i>Npj Parkinson's Disease</i> , 2015, 1, 15014.	2.5	54
233	Nonmotor Symptoms in Dopa-Responsive Dystonia. <i>Movement Disorders Clinical Practice</i> , 2015, 2, 347-356.	0.8	10
234	Premonitory urge to tic in tourette's is associated with interoceptive awareness. <i>Movement Disorders</i> , 2015, 30, 1198-1202.	2.2	118

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235	Reply to letter: Transcranial magnetic stimulation for Parkinson's disease. <i>Movement Disorders</i> , 2015, 30, 1973-1974.	2.2	2
236	Myoclonus, Epilepsy, and Ataxia Resulting From Potassium Channel Gene Mutation: Expanding the Spectrum Underlying Ramsay Hunt Syndrome. <i>Movement Disorders Clinical Practice</i> , 2015, 2, 230-231.	0.8	2
237	Hot topic: Recessive mutations in the $\alpha 3$ (VI) collagen gene COL6A3 cause early-onset isolated dystonia. <i>Movement Disorders</i> , 2015, 30, 1622-1622.	2.2	1
238	Parkinsonism following neuroleptic exposure: A double-hit hypothesis?. <i>Movement Disorders</i> , 2015, 30, 780-785.	2.2	41
239	A Missense Mutation in KCTD17 Causes Autosomal Dominant Myoclonus-Dystonia. <i>American Journal of Human Genetics</i> , 2015, 96, 938-947.	2.6	109
240	Loss of <i>PLA2G6</i> leads to elevated mitochondrial lipid peroxidation and mitochondrial dysfunction. <i>Brain</i> , 2015, 138, 1801-1816.	3.7	143
241	Concomitant fragile X-associated tremor ataxia syndrome and Parkinson's disease: a clinicopathological report of two cases: Table 1. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 934-936.	0.9	20
242	Syndromic associations and RNF216 mutations. <i>Parkinsonism and Related Disorders</i> , 2015, 21, 1389-1390.	1.1	4
243	Episodic akinesia of jaw movements akin to freezing in multiple system atrophy. <i>Parkinsonism and Related Disorders</i> , 2015, 21, 1483-1484.	1.1	2
244	H-ABC syndrome and DYT4: Variable expressivity or pleiotropy of TUBB4 mutations?. <i>Movement Disorders</i> , 2015, 30, 828-833.	2.2	117
245	The analysis of C9orf72 repeat expansions in a large series of clinically and pathologically diagnosed cases with atypical parkinsonism. <i>Neurobiology of Aging</i> , 2015, 36, 1221.e1-1221.e6.	1.5	39
246	Tremor in Charcot-Marie-Tooth disease: No evidence of cerebellar dysfunction. <i>Clinical Neurophysiology</i> , 2015, 126, 1817-1824.	0.7	22
247	A novel TUBB4 mutation suggests that genotype-phenotype correlation of H-ABC syndrome needs to be revisited: Figure 1. <i>Brain</i> , 2015, 138, e370-e370.	3.7	3
248	Clinical relevance of serum antibodies to extracellular <i>N</i> -methyl-d-aspartate receptor epitopes. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 708-713.	0.9	97
249	Feedforward somatosensory inhibition is normal in cervical dystonia. <i>Parkinsonism and Related Disorders</i> , 2015, 21, 266-270.	1.1	1
250	Reward Pays the Cost of Noise Reduction in Motor and Cognitive Control. <i>Current Biology</i> , 2015, 25, 1707-1716.	1.8	272
251	The 4H syndrome due to RNF216 mutation. <i>Parkinsonism and Related Disorders</i> , 2015, 21, 1122-1123.	1.1	15
252	Reply: Parkinson's disease in GTP cyclohydrolase 1 mutation carriers. <i>Brain</i> , 2015, 138, e352-e352.	3.7	4

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253	<i>ADCY5</i> mutations are another cause of benign hereditary chorea. <i>Neurology</i> , 2015, 85, 80-88.	1.5	140
254	Transcranial magnetic stimulation follow-up study in early Parkinson's disease: A decline in compensation with disease progression?. <i>Movement Disorders</i> , 2015, 30, 1098-1106.	2.2	55
255	Mutations in HPCA Cause Autosomal-Recessive Primary Isolated Dystonia. <i>American Journal of Human Genetics</i> , 2015, 96, 657-665.	2.6	151
256	The <i>CACNA1B</i> R1389H variant is not associated with myoclonus-dystonia in a large European multicentric cohort. <i>Human Molecular Genetics</i> , 2015, 24, 5326-5329.	1.4	28
257	Cortical pencil lining in neuroferritinopathy: A diagnostic clue. <i>Neurology</i> , 2015, 84, 1816-1818.	1.5	93
258	A systematic screening to identify <i>de novo</i> mutations causing sporadic early-onset Parkinson's disease. <i>Human Molecular Genetics</i> , 2015, 24, 6711-6720.	1.4	59
259	Primary writing tremor is a dystonic trait: Evidence from an instructive family. <i>Journal of the Neurological Sciences</i> , 2015, 356, 210-211.	0.3	9
260	Classic PD-like rest tremor associated with the tau p.R406W mutation. <i>Parkinsonism and Related Disorders</i> , 2015, 21, 1002-1004.	1.1	3
261	On the long-term outcome of orthostatic tremor. <i>Parkinsonism and Related Disorders</i> , 2015, 21, 1290-1291.	1.1	2
262	The clinical and genetic heterogeneity of paroxysmal dyskinesias. <i>Brain</i> , 2015, 138, 3567-3580.	3.7	129
263	Atypical Parkinsonism. <i>Neurologic Clinics</i> , 2015, 33, 39-56.	0.8	34
264	Rest and other types of tremor in adult-onset primary dystonia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014, 85, 965-968.	0.9	150
265	<i>ALS2</i> mutations. <i>Neurology</i> , 2014, 82, 1065-1067.	1.5	29
266	The phenotypic spectrum of DYT24 due to ANO3 mutations. <i>Movement Disorders</i> , 2014, 29, 928-934.	2.2	161
267	Functional tics and echophenomena. <i>Parkinsonism and Related Disorders</i> , 2014, 20, 1440-1441.	1.1	14
268	Cerebellar stimulation fails to modulate motor cortex plasticity in writing dystonia. <i>Movement Disorders</i> , 2014, 29, 1304-1307.	2.2	50
269	Fixed Dystonia of the Tongue. <i>Movement Disorders Clinical Practice</i> , 2014, 1, 134-135.	0.8	2
270	Reply: Dystonia after severe head injuries. <i>Movement Disorders</i> , 2014, 29, 578-579.	2.2	1



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271	Shaking on Standing: A Critical Review. <i>Movement Disorders Clinical Practice</i> , 2014, 1, 173-179.	0.8	28
272	Genomewide association study in cervical dystonia demonstrates possible association with sodium leak channel. <i>Movement Disorders</i> , 2014, 29, 245-251.	2.2	43
273	A reflection on plasticity research in writing dystonia. <i>Movement Disorders</i> , 2014, 29, 980-987.	2.2	33
274	The Clinical Syndrome of Paroxysmal Exercise-Induced Dystonia: Diagnostic Outcomes and an Algorithm. <i>Movement Disorders Clinical Practice</i> , 2014, 1, 57-61.	0.8	100
275	Neuropathological features of genetically confirmed DYT1 dystonia: investigating disease-specific inclusions. <i>Acta Neuropathologica Communications</i> , 2014, 2, 159.	2.4	21
276	Lysine 27 Ubiquitination of the Mitochondrial Transport Protein Miro Is Dependent on Serine 65 of the Parkin Ubiquitin Ligase. <i>Journal of Biological Chemistry</i> , 2014, 289, 14569-14582.	1.6	152
277	Dystonia. <i>Current Opinion in Neurology</i> , 2014, 27, 468-476.	1.8	81
278	<i>GNAL</i> Mutations and Dystonia. <i>JAMA Neurology</i> , 2014, 71, 1052.	4.5	5
279	Long-term Clinical Outcome of Fetal Cell Transplantation for Parkinson Disease. <i>JAMA Neurology</i> , 2014, 71, 83.	4.5	257
280	A 6.4 Mb Duplication of the $\alpha$ -Synuclein Locus Causing Frontotemporal Dementia and Parkinsonism. <i>JAMA Neurology</i> , 2014, 71, 1162.	4.5	60
281	Propriospinal myoclonus. <i>Neurology</i> , 2014, 83, 1862-1870.	1.5	162
282	Psychogenic paroxysmal movement disorders – Clinical features and diagnostic clues. <i>Parkinsonism and Related Disorders</i> , 2014, 20, 41-46.	1.1	77
283	Physical precipitating factors in functional movement disorders. <i>Journal of the Neurological Sciences</i> , 2014, 338, 174-177.	0.3	136
284	Paroxysmal exercise-induced dystonia due to GLUT1 mutation can be responsive to levodopa: a case report. <i>Journal of Neurology</i> , 2014, 261, 615-616.	1.8	20
285	A new gene for Fahr's syndrome – <i>PDGF<math>\beta</math></i> . <i>Movement Disorders</i> , 2014, 29, 307-307.	2.2	5
286	That DAT gene that causes dystonia-parkinsonism: broadening the phenotype. <i>Brain</i> , 2014, 137, 976-977.	3.7	2
287	The Phenomenology of Functional (Psychogenic) Dystonia. <i>Movement Disorders Clinical Practice</i> , 2014, 1, 36-44.	0.8	40
288	Reply to R. Erro and M. Tinazzi's – Functional (psychogenic) paroxysms: the diagnosis is in the eye of the beholder. <i>Parkinsonism and Related Disorders</i> , 2014, 20, 345.	1.1	0

#	ARTICLE	IF	CITATIONS
289	Pallidal stimulation for primary generalised dystonia: effect on cognition, mood and quality of life. <i>Journal of Neurology</i> , 2014, 261, 164-173.	1.8	51
290	Motor sequence learning and motor adaptation in primary cervical dystonia. <i>Journal of Clinical Neuroscience</i> , 2014, 21, 934-938.	0.8	19
291	Paroxysmal Kinesigenic Dyskinesia May Be Misdiagnosed in Co-occurring Gilles de la Tourette Syndrome. <i>Movement Disorders Clinical Practice</i> , 2014, 1, 84-86.	0.8	11
292	The entity of parkinsonism and associated lipomatosis. <i>Neurology</i> , 2014, 83, 1673-1674.	1.5	2
293	Loss of sensory attenuation in patients with functional (psychogenic) movement disorders. <i>Brain</i> , 2014, 137, 2916-2921.	3.7	104
294	Pallidal neurostimulation in patients with medication-refractory cervical dystonia: a randomised, sham-controlled trial. <i>Lancet Neurology</i> , The, 2014, 13, 875-884.	4.9	281
295	Parkinson's disease in GTP cyclohydrolase 1 mutation carriers. <i>Brain</i> , 2014, 137, 2480-2492.	3.7	169
296	Screening of mutations in NOL3 in a myoclonic syndromes series. <i>Journal of Neurology</i> , 2014, 261, 1830-1831.	1.8	1
297	Motivational modulation of bradykinesia in Parkinson's disease off and on dopaminergic medication. <i>Journal of Neurology</i> , 2014, 261, 1080-1089.	1.8	32
298	Dystonia with aphonia, slow horizontal saccades, epilepsy and photic myoclonus: A novel syndrome?. <i>Parkinsonism and Related Disorders</i> , 2014, 20, 328-331.	1.1	4
299	The role of the cerebellum in the pathogenesis of cortical myoclonus. <i>Movement Disorders</i> , 2014, 29, 437-443.	2.2	110
300	The expanding universe of disorders of the basal ganglia. <i>Lancet</i> , The, 2014, 384, 523-531.	6.3	155
301	Normal Motor Adaptation in Cervical Dystonia: A Fundamental Cerebellar Computation is Intact. <i>Cerebellum</i> , 2014, 13, 558-567.	1.4	34
302	A Novel De Novo Mutation of the TITF1/NKX2-1 Gene Causing Ataxia, Benign Hereditary Chorea, Hypothyroidism and a Pituitary Mass in a UK Family and Review of the Literature. <i>Cerebellum</i> , 2014, 13, 588-595.	1.4	93
303	Paroxysmal dyskinesias revisited: A review of 500 genetically proven cases and a new classification. <i>Movement Disorders</i> , 2014, 29, 1108-1116.	2.2	224
304	Psychogenic axial myoclonus: Clinical features and long-term outcome. <i>Parkinsonism and Related Disorders</i> , 2014, 20, 596-599.	1.1	98
305	Differentiating drug-induced parkinsonism from Parkinson's disease: An update on non-motor symptoms and investigations. <i>Parkinsonism and Related Disorders</i> , 2014, 20, 808-814.	1.1	81
306	Facial tremor in dystonia. <i>Parkinsonism and Related Disorders</i> , 2014, 20, 924-925.	1.1	10

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307	No pathogenic <i>GNAL</i> mutations in 192 sporadic and familial cases of cervical dystonia. <i>Movement Disorders</i> , 2014, 29, 154-155.	2.2	8
308	Assessment of patients with isolated or combined dystonia: An update on dystonia syndromes. <i>Movement Disorders</i> , 2013, 28, 889-898.	2.2	88
309	1â€Hz repetitive transcranial magnetic stimulation and diphasic dyskinesia in Parkinson's disease. <i>Movement Disorders</i> , 2013, 28, 245-245.	2.2	5
310	Extragenetic factors and clinical penetrance of DYT1 dystonia: an exploratory study. <i>Journal of Neurology</i> , 2013, 260, 1081-1086.	1.8	30
311	Patients with rest-tremor and scans with ipsilateral dopaminergic deficit. <i>Journal of Neurology</i> , 2013, 260, 1132-1135.	1.8	10
312	Migraine with aura as the predominant phenotype in a family with a PRRT2 mutation. <i>Journal of Neurology</i> , 2013, 260, 656-660.	1.8	17
313	Increased glucocerebrosidase (GBA) 2 activity in GBA1 deficient mice brains and in Gaucher leucocytes. <i>Journal of Inherited Metabolic Disease</i> , 2013, 36, 869-872.	1.7	28
314	Progressive ataxia and palatal tremor associated with dense pontine calcification: A unique case. <i>Movement Disorders</i> , 2013, 28, 1155-1157.	2.2	5
315	Ropinirole monotherapy induced severe reversible dyskinesias in Parkinson's disease. <i>Movement Disorders</i> , 2013, 28, 1159-1160.	2.2	6
316	The glucocerebrosidase E326K variant predisposes to Parkinson's disease, but does not cause Gaucher's disease. <i>Movement Disorders</i> , 2013, 28, 232-236.	2.2	121
317	Reopening the case for antiâ€basal ganglia antibodies (ABCAs): Identification of dopamineâ€2 receptor antibodies associated with movement disorders. <i>Movement Disorders</i> , 2013, 28, 733-733.	2.2	3
318	Functional (psychogenic) symptoms in Parkinson's disease. <i>Movement Disorders</i> , 2013, 28, 1622-1627.	2.2	52
319	<i>FUS</i> gene mutations cause essential tremor: A surprise but also confirms genetic heterogeneity of essential tremor. <i>Movement Disorders</i> , 2013, 28, 290-290.	2.2	2
320	Dystonic opisthotonus: A â€red flagâ€ for neurodegeneration with brain iron accumulation syndromes?. <i>Movement Disorders</i> , 2013, 28, 1325-1329.	2.2	39
321	Markedly asymmetric presentation in multiple system atrophy. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 901-905.	1.1	21
322	Deep brain stimulation as a treatment for chorea-acanthocytosis. <i>Journal of Neurology</i> , 2013, 260, 303-305.	1.8	19
323	When the levator scapulae becomes a â€rotator capitisâ€ Implications for cervical dystonia. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 705-706.	1.1	10
324	Coactivation sign in fixed dystonia. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 474-476.	1.1	8

#	ARTICLE	IF	CITATIONS
325	Functional movement disorders are not uncommon in the elderly. <i>Movement Disorders</i> , 2013, 28, 540-543.	2.2	40
326	Criteria for the diagnosis of corticobasal degeneration. <i>Neurology</i> , 2013, 80, 496-503.	1.5	1,445
327	Excess iron harms the brain: the syndromes of neurodegeneration with brain iron accumulation (NBIA). <i>Journal of Neural Transmission</i> , 2013, 120, 695-703.	1.4	52
328	Commentary for "Progressive ataxia associated with scarring skin lesions and vertical gaze palsy". <i>Movement Disorders</i> , 2013, 28, 445-446.	2.2	1
329	Familial psychogenic movement disorders. <i>Movement Disorders</i> , 2013, 28, 1295-1298.	2.2	31
330	Phenomenology and classification of dystonia: A consensus update. <i>Movement Disorders</i> , 2013, 28, 863-873.	2.2	1,754
331	"Atypical" atypical parkinsonism: New genetic conditions presenting with features of progressive supranuclear palsy, corticobasal degeneration, or multiple system atrophy" A diagnostic guide. <i>Movement Disorders</i> , 2013, 28, 1184-1199.	2.2	167
332	Late onset rest-tremor in DYT1 dystonia. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 136-137.	1.1	11
333	Tremor in inflammatory neuropathies. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2013, 84, 1282-1287.	0.9	129
334	Genetics and Pathophysiology of Neurodegeneration with Brain Iron Accumulation (NBIA). <i>Current Neuropharmacology</i> , 2013, 11, 59-79.	1.4	100
335	Cerebellar learning distinguishes inflammatory neuropathy with and without tremor. <i>Neurology</i> , 2013, 80, 1867-1873.	1.5	30
336	Secondary and primary dystonia: pathophysiological differences. <i>Brain</i> , 2013, 136, 2038-2049.	3.7	104
337	Clinical and polysomnographic course of childhood narcolepsy with cataplexy. <i>Brain</i> , 2013, 136, 3787-3795.	3.7	113
338	Neuropsychiatric Symptoms as a Putative Crescendo Warning of a Striatocapsular Infarct?. <i>Journal of Neuropsychiatry and Clinical Neurosciences</i> , 2013, 25, E39-E43.	0.9	0
339	The functional neuroimaging correlates of psychogenic versus organic dystonia. <i>Brain</i> , 2013, 136, 770-781.	3.7	83
340	Beta-propeller protein-associated neurodegeneration: a new X-linked dominant disorder with brain iron accumulation. <i>Brain</i> , 2013, 136, 1708-1717.	3.7	203
341	The genetics of dystonia: new twists in an old tale. <i>Brain</i> , 2013, 136, 2017-2037.	3.7	102
342	Ataxia telangiectasia presenting as dopa-responsive cervical dystonia. <i>Neurology</i> , 2013, 81, 1148-1151.	1.5	65

#	ARTICLE	IF	CITATIONS
343	Preface. <i>International Review of Neurobiology</i> , 2013, 110, xv-xviii.	0.9	1
344	A pathway-based analysis provides additional support for an immune-related genetic susceptibility to Parkinson's disease. <i>Human Molecular Genetics</i> , 2013, 22, 1039-1049.	1.4	122
345	Primary and secondary dystonic syndromes. <i>Current Opinion in Neurology</i> , 2013, 26, 406-412.	1.8	13
346	Does rest tremor exclude the diagnosis of adult-onset primary dystonia?. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2013, 84, 708-708.	0.9	5
347	Failure of explicit movement control in patients with functional motor symptoms. <i>Movement Disorders</i> , 2013, 28, 517-523.	2.2	43
348	Mutations in the autoregulatory domain of $\alpha$ -tubulin 4a cause hereditary dystonia. <i>Annals of Neurology</i> , 2013, 73, 546-553.	2.8	148
349	Defining the Epsilon-Sarcoglycan (SGCE) Gene Phenotypic Signature in Myoclonus-Dystonia: A Reappraisal of Genetic Testing Criteria. <i>Movement Disorders</i> , 2013, 28, 787-794.	2.2	31
350	Reply to Drs. Kurlan, Fasano, and Evans: A clinically useful definition of stereotypies. <i>Movement Disorders</i> , 2013, 28, 405-406.	2.2	0
351	The differential diagnosis of Huntington's disease-like syndromes: 'red flags' for the clinician. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2013, 84, 650-656.	0.9	141
352	Clinical diagnosis of propriospinal myoclonus is unreliable: An electrophysiologic study. <i>Movement Disorders</i> , 2013, 28, 1868-1873.	2.2	124
353	The syndrome of deafness-dystonia: Clinical and genetic heterogeneity. <i>Movement Disorders</i> , 2013, 28, 795-803.	2.2	25
354	Movement Disorders in Adult Patients With Classical Galactosemia. <i>Movement Disorders</i> , 2013, 28, 804-810.	2.2	57
355	Pallidal stimulation for cervical dystonia does not correct abnormal temporal discrimination. <i>Movement Disorders</i> , 2013, 28, 1874-1877.	2.2	30
356	Dopaminergic Neuronal Imaging in Genetic Parkinson's Disease: Insights into Pathogenesis. <i>PLoS ONE</i> , 2013, 8, e69190.	1.1	55
357	Short and Long Term Outcome of Bilateral Pallidal Stimulation in Chorea-Acanthocytosis. <i>PLoS ONE</i> , 2013, 8, e79241.	1.1	44
358	Myoclonus-dystonia syndrome due to tyrosine hydroxylase deficiency. <i>Neurology</i> , 2012, 79, 435-441.	1.5	45
359	Clinical Approach to Parkinson's Disease: Features, Diagnosis, and Principles of Management. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2012, 2, a008870-a008870.	2.9	288
360	Dopamine and performance in a reinforcement learning task: evidence from Parkinson's disease. <i>Brain</i> , 2012, 135, 1871-1883.	3.7	137

#	ARTICLE	IF	CITATIONS
361	<i>PRRT2</i> gene mutations. <i>Neurology</i> , 2012, 79, 2115-2121.	1.5	159
362	The non-motor syndrome of primary dystonia: clinical and pathophysiological implications. <i>Brain</i> , 2012, 135, 1668-1681.	3.7	246
363	"Jumping to conclusions" bias in functional movement disorders. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2012, 83, 460-463.	0.9	42
364	Believing is perceiving: mismatch between self-report and actigraphy in psychogenic tremor. <i>Brain</i> , 2012, 135, 117-123.	3.7	123
365	The Brighter Side of Music in Dystonia. <i>Archives of Neurology</i> , 2012, 69, 917-9.	4.9	10
366	Psychogenic facial movement disorders: Clinical features and associated conditions. <i>Movement Disorders</i> , 2012, 27, 1544-1551.	2.2	93
367	Young-onset parkinsonism due to homozygous duplication of $\alpha$ -synuclein in a consanguineous family. <i>Movement Disorders</i> , 2012, 27, 1829-1830.	2.2	27
368	Atypical parkinsonism and cerebrotendinous xanthomatosis: Report of a family with corticobasal syndrome and a literature review. <i>Movement Disorders</i> , 2012, 27, 1769-1774.	2.2	31
369	Camptocormia: don't forget muscle disease in the movement disorder clinic. <i>Journal of Neurology</i> , 2012, 259, 1752-1754.	1.8	8
370	Widespread Lewy body and tau accumulation in childhood and adult onset dystonia-parkinsonism cases with PLA2G6 mutations. <i>Neurobiology of Aging</i> , 2012, 33, 814-823.	1.5	184
371	Screening for VPS35 mutations in Parkinson's disease. <i>Neurobiology of Aging</i> , 2012, 33, 838.e1-838.e5.	1.5	53
372	Analysis of ATP13A2 in large neurodegeneration with brain iron accumulation (NBIA) and dystonia-parkinsonism cohorts. <i>Neuroscience Letters</i> , 2012, 523, 35-38.	1.0	15
373	Mutations in the Gene PRRT2 Cause Paroxysmal Kinesigenic Dyskinesia with Infantile Convulsions. <i>Cell Reports</i> , 2012, 1, 2-12.	2.9	250
374	Secondary cervical dystonia caused by cerebellar cystic lesion " A case study with transcranial magnetic stimulation. <i>Clinical Neurophysiology</i> , 2012, 123, 418-419.	0.7	4
375	Dystonia with brain manganese accumulation resulting from <i>SLC30A10</i> mutations: A new treatable disorder. <i>Movement Disorders</i> , 2012, 27, 1317-1322.	2.2	104
376	THAP1 mutations and dystonia phenotypes: Genotype phenotype correlations. <i>Movement Disorders</i> , 2012, 27, 1290-1294.	2.2	126
377	Exome Sequencing Reveals De Novo WDR45 Mutations Causing a Phenotypically Distinct, X-Linked Dominant Form of NBIA. <i>American Journal of Human Genetics</i> , 2012, 91, 1144-1149.	2.6	309
378	Mutations in ANO3 Cause Dominant Craniocervical Dystonia: Ion Channel Implicated in Pathogenesis. <i>American Journal of Human Genetics</i> , 2012, 91, 1041-1050.	2.6	224

#	ARTICLE	IF	CITATIONS
379	The Effect of Motivation on Movement: A Study of Bradykinesia in Parkinson's Disease. PLoS ONE, 2012, 7, e47138.	1.1	28
380	Treatment of Focal Dystonia. Current Treatment Options in Neurology, 2012, 14, 213-229.	0.7	28
381	Psychogenic palatal tremor may be underrecognized: Reappraisal of a large series of cases. Movement Disorders, 2012, 27, 1164-1168.	2.2	126
382	Dystonia in corticobasal degeneration: A review of the literature on 404 pathologically proven cases. Movement Disorders, 2012, 27, 696-702.	2.2	119
383	Joint hypermobility syndrome: A risk factor for fixed dystonia?. Movement Disorders, 2012, 27, 1070-1070.	2.2	12
384	A new treatable genetic disorder of manganese metabolism causing dystonia, parkinsonism and cirrhosis: The 'New' Wilson's disease?. Movement Disorders, 2012, 27, 962-962.	2.2	9
385	Identification of PRRT2 as the causative gene of paroxysmal kinesigenic dyskinesia. Movement Disorders, 2012, 27, 707-707.	2.2	10
386	The distinct movement disorder in anti-NMDA receptor encephalitis may be related to status dissociatus: A hypothesis. Movement Disorders, 2012, 27, 1360-1363.	2.2	46
387	Is transcranial sonography useful to distinguish scans without evidence of dopaminergic deficit patients from Parkinson's disease?. Movement Disorders, 2012, 27, 1182-1185.	2.2	32
388	Tardive dyskinesia is caused by maladaptive synaptic plasticity: A hypothesis. Movement Disorders, 2012, 27, 1205-1215.	2.2	172
389	RAD51 Haploinsufficiency Causes Congenital Mirror Movements in Humans. American Journal of Human Genetics, 2012, 90, 301-307.	2.6	81
390	Functional (psychogenic) movement disorders: merging mind and brain. Lancet Neurology, The, 2012, 11, 250-260.	4.9	252
391	Syndromes of Neurodegeneration With Brain Iron Accumulation. Seminars in Pediatric Neurology, 2012, 19, 57-66.	1.0	55
392	Syndromes of neurodegeneration with brain iron accumulation (NBIA): An update on clinical presentations, histological and genetic underpinnings, and treatment considerations. Movement Disorders, 2012, 27, 42-53.	2.2	219
393	Stereotypies: A critical appraisal and suggestion of a clinically useful definition. Movement Disorders, 2012, 27, 179-185.	2.2	93
394	Cervical dystonia and joint hypermobility syndrome: A dangerous combination. Movement Disorders, 2012, 27, 203-204.	2.2	7
395	Myoclonic disorders: a practical approach for diagnosis and treatment. Therapeutic Advances in Neurological Disorders, 2011, 4, 47-62.	1.5	159
396	Movement disorders and mitochondrial disease. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2011, 100, 173-192.	1.0	8

#	ARTICLE	IF	CITATIONS
397	A distinctive pattern of cortical excitability in patients with the syndrome of dystonia and cerebellar ataxia. <i>Clinical Neurophysiology</i> , 2011, 122, 1816-1819.	0.7	10
398	Characteristic constant groaning in late stage progressive supranuclear palsy: A case report. <i>Parkinsonism and Related Disorders</i> , 2011, 17, 575-576.	1.1	11
399	Excessive Daytime Sleepiness in Multiple System Atrophy (SLEEMSA Study). <i>Archives of Neurology</i> , 2011, 68, 223-30.	4.9	83
400	Abnormal sense of intention preceding voluntary movement in patients with psychogenic tremor. <i>Neuropsychologia</i> , 2011, 49, 2791-2793.	0.7	81
401	Cerebellar brain inhibition is decreased in active and surround muscles at the onset of voluntary movement. <i>Experimental Brain Research</i> , 2011, 209, 437-442.	0.7	39
402	Motivation and movement: the effect of monetary incentive on performance speed. <i>Experimental Brain Research</i> , 2011, 209, 551-559.	0.7	55
403	Tremor—some controversial aspects. <i>Movement Disorders</i> , 2011, 26, 18-23.	2.2	169
404	The clinical syndrome of primary tic disorder associated with dystonia: A large clinical series and a review of the literature. <i>Movement Disorders</i> , 2011, 26, 679-684.	2.2	83
405	Cervical dystonia associated with facioscapulohumeral dystrophy: Expanding the clinical spectrum?. <i>Movement Disorders</i> , 2011, 26, 765-766.	2.2	3
406	Immediate response to botulinum toxin injections in patients with fixed dystonia. <i>Movement Disorders</i> , 2011, 26, 917-918.	2.2	42
407	Movement disorders in adult surviving patients with maple syrup urine disease. <i>Movement Disorders</i> , 2011, 26, 1324-1328.	2.2	46
408	Limb amputations in fixed dystonia: A form of body integrity identity disorder?. <i>Movement Disorders</i> , 2011, 26, 1410-1414.	2.2	39
409	Late-onset asymmetric myoclonus: An emerging syndrome. <i>Movement Disorders</i> , 2011, 26, 1744-1747.	2.2	71
410	Botulinum toxin injections reduce associative plasticity in patients with primary dystonia. <i>Movement Disorders</i> , 2011, 26, 1282-1289.	2.2	67
411	Gait in SWEDDs patients: Comparison with Parkinson's disease patients and healthy controls. <i>Movement Disorders</i> , 2011, 26, 1266-1273.	2.2	28
412	Deep brain stimulation effects in dystonia: Time course of electrophysiological changes in early treatment. <i>Movement Disorders</i> , 2011, 26, 1913-1921.	2.2	111
413	Paroxysmal dyskinesias. <i>Movement Disorders</i> , 2011, 26, 1157-1165.	2.2	153
414	Ability to cycle despite severe freezing of gait in atypical parkinsonism in Fahr's syndrome. <i>Movement Disorders</i> , 2011, 26, 2141-2142.	2.2	6



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415	Paroxysmal limb dyskinesia induced by weight: An unusual case of cortical reflex seizures. <i>Movement Disorders</i> , 2011, 26, 2438-2439.	2.2	2
416	Attention to self in psychogenic tremor. <i>Movement Disorders</i> , 2011, 26, 2575-2576.	2.2	50
417	Moving toward "laboratory"-supported criteria for psychogenic tremor. <i>Movement Disorders</i> , 2011, 26, 2509-2515.	2.2	132
418	Paroxysmal craniocervical dyskinesia as manifestation of frontal lobe epilepsy. <i>Movement Disorders</i> , 2011, 26, 2580-2582.	2.2	2
419	Recognition of faciobrachial dystonic seizures allowing early intervention with prevention of development of full-blown limbic encephalitis. <i>Movement Disorders</i> , 2011, 26, 2176-2176.	2.2	4
420	The Wilson films "Wilson's disease. <i>Movement Disorders</i> , 2011, 26, 2473-2474.	2.2	0
421	<i>Movement Disorders on YouTube</i> "Caveat Spectator. <i>New England Journal of Medicine</i> , 2011, 365, 1160-1161.	13.9	77
422	Complex movement disorders at disease onset in childhood narcolepsy with cataplexy. <i>Brain</i> , 2011, 134, 3480-3492.	3.7	159
423	Patients with primary cervical dystonia have evidence of discrete deficits in praxis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2011, 82, 615-619.	0.9	7
424	Dissection of the genetics of Parkinson's disease identifies an additional association 5' of SNCA and multiple associated haplotypes at 17q21. <i>Human Molecular Genetics</i> , 2011, 20, 345-353.	1.4	202
425	Huntington's disease look-alikes. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2011, 100, 101-112.	1.0	20
426	Brain structure in movement disorders: a neuroimaging perspective. <i>Current Opinion in Neurology</i> , 2010, 23, 413-419.	1.8	18
427	Rare Causes of Dystonia Parkinsonism. <i>Current Neurology and Neuroscience Reports</i> , 2010, 10, 431-439.	2.0	47
428	Three faces of the same gene: <i>FA2H</i> links neurodegeneration with brain iron accumulation, leukodystrophies, and hereditary spastic paraplegias. <i>Annals of Neurology</i> , 2010, 68, 575-577.	2.8	36
429	Abnormal explicit but normal implicit sequence learning in premanifest and early Huntington's disease. <i>Movement Disorders</i> , 2010, 25, 1343-1349.	2.2	23
430	Nonmotor symptoms in <i>Parkin</i> gene-related parkinsonism. <i>Movement Disorders</i> , 2010, 25, 1279-1284.	2.2	31
431	<i>ATP13A2</i> mutations ( <i>PARK9</i> ) cause neurodegeneration with brain iron accumulation. <i>Movement Disorders</i> , 2010, 25, 979-984.	2.2	163
432	Distinguishing SWEDDs patients with asymmetric resting tremor from Parkinson's disease: A clinical and electrophysiological study. <i>Movement Disorders</i> , 2010, 25, 560-569.	2.2	223

#	ARTICLE	IF	CITATIONS
433	Characteristic head drops and axial extension in advanced choreaâ€acanthocytosis. <i>Movement Disorders</i> , 2010, 25, 1487-1491.	2.2	54
434	Atypical parkinsonism with apraxia and supranuclear gaze abnormalities in type 1 Gaucher disease. Expanding the spectrum: Case report and literature review. <i>Movement Disorders</i> , 2010, 25, 1506-1509.	2.2	21
435	â€œProgressive delayedâ€onset postanoxic dystoniaâ€diagnosed with PANK2 mutations 26 years after onsetâ€ An update. <i>Movement Disorders</i> , 2010, 25, 2889-2891.	2.2	2
436	Earlyâ€onset Lâ€dopaâ€responsive parkinsonism with pyramidal signs due to <i>ATP13A2, PLA2G6, FBOX7</i> and <i>spatacsin</i> mutations. <i>Movement Disorders</i> , 2010, 25, 1791-1800.	2.2	287
437	Levodopaâ€induced belly dancer's dyskinesias in Parkinson's disease: Report of one case. <i>Movement Disorders</i> , 2010, 25, 1760-1762.	2.2	18
438	Endothelial, Sympathetic, and Cardiac Function in Inherited (6 <i>R</i>)- <scp>l</scp>-Erythro-5,6,7,8-Tetrahydro- <scp>l</scp>-Biopterin Deficiency. <i>Circulation: Cardiovascular Genetics</i> , 2010, 3, 513-522.	5.1	15
439	Abnormal motor cortex plasticity in premanifest and very early manifest Huntington disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2010, 81, 267-270.	0.9	92
440	Low-frequency repetitive transcranial magnetic stimulation and off-phase motor symptoms in Parkinson's disease. <i>Journal of the Neurological Sciences</i> , 2010, 291, 1-4.	0.3	36
441	Slow (1 Hz) repetitive transcranial magnetic stimulation (rTMS) induces a sustained change in cortical excitability in patients with Parkinsonâ€™s disease. <i>Clinical Neurophysiology</i> , 2010, 121, 1129-1137.	0.7	51
442	Validation of a dystonia screening questionnaire: Testing in a cohort of mixed neurological disorders. <i>Parkinsonism and Related Disorders</i> , 2010, 16, 620-622.	1.1	2
443	Characterization of PLA2G6 as a locus for dystoniaâ€parkinsonism. <i>Annals of Neurology</i> , 2009, 65, 19-23.	2.8	399
444	Reply: Puschmann â€œUnverrichtâ€Lundborg diseaseâ€”A misnomerâ€• <i>Movement Disorders</i> , 2009, 24, 630-630.2.2		0
445	Repetitive transcranial magnetic stimulation for levodopaâ€induced dyskinesias in Parkinson's disease. <i>Movement Disorders</i> , 2009, 24, 246-253.	2.2	75
446	Sensory functions in dystonia: Insights from behavioral studies. <i>Movement Disorders</i> , 2009, 24, 1427-1436.	2.2	103
447	GLUT1 gene mutations cause sporadic paroxysmal exerciseâ€induced dyskinesias. <i>Movement Disorders</i> , 2009, 24, 1684-1688.	2.2	110
448	Early and marked benefit with GPi DBS for Lubag syndrome presenting with rapidly progressive lifeâ€threatening dystonia. <i>Movement Disorders</i> , 2009, 24, 1710-1712.	2.2	48
449	Tremor on smiling. <i>Movement Disorders</i> , 2009, 24, 1542-1545.	2.2	9
450	Prominent Oromandibular Dystonia and Pharyngeal Telangiectasia in Atypical Ataxia Telangiectasia. <i>Cerebellum</i> , 2009, 8, 22-27.	1.4	38

#	ARTICLE	IF	CITATIONS
451	Secondary Dystonia-Clinical Clues and Syndromic Associations. <i>Journal of Movement Disorders</i> , 2009, 2, 58-63.	0.7	11
452	Cortical excitability is abnormal in patients with the "fixed dystonia" syndrome. <i>Movement Disorders</i> , 2008, 23, 646-652.	2.2	111
453	Dystonia in the Woodhouse Sakati syndrome: A new family and literature review. <i>Movement Disorders</i> , 2008, 23, 592-596.	2.2	70
454	Huntington's disease phenocopies are clinically and genetically heterogeneous. <i>Movement Disorders</i> , 2008, 23, 716-720.	2.2	108
455	Antineuronal antibodies in Parkinson's disease. <i>Movement Disorders</i> , 2008, 23, 958-963.	2.2	17
456	Unusual familial presentation of epsilon-sarcoglycan gene mutation with falls and writer's cramp. <i>Movement Disorders</i> , 2008, 23, 1913-1915.	2.2	27
457	Fragile X syndrome associated with tic disorders. <i>Movement Disorders</i> , 2008, 23, 1108-1112.	2.2	18
458	Motor cortical physiology in patients and asymptomatic carriers of parkin gene mutations. <i>Movement Disorders</i> , 2008, 23, 1812-1819.	2.2	29
459	Psychogenic movement disorders in children: A report of 15 cases and a review of the literature. <i>Movement Disorders</i> , 2008, 23, 1882-1888.	2.2	129
460	Mutations in C2orf37, Encoding a Nucleolar Protein, Cause Hypogonadism, Alopecia, Diabetes Mellitus, Mental Retardation, and Extrapyrmidal Syndrome. <i>American Journal of Human Genetics</i> , 2008, 83, 684-691.	2.6	121
461	GLUT1 mutations are a cause of paroxysmal exertion-induced dyskinesias and induce hemolytic anemia by a cation leak. <i>Journal of Clinical Investigation</i> , 2008, 118, 2157-2168.	3.9	321
462	Blepharospasm and limb dystonia caused by Mohr-Tranebjaerg syndrome with a novel splice-site mutation in the deafness/dystonia peptide gene. <i>Movement Disorders</i> , 2007, 22, 1328-1331.	2.2	27
463	The entity of jaw tremor and dystonia. <i>Movement Disorders</i> , 2007, 22, 1491-1495.	2.2	32
464	Is the anticonvulsant topiramate beneficial in essential tremor?. <i>Nature Clinical Practice Neurology</i> , 2006, 2, 478-479.	2.7	1
465	Atypical movement disorders in antiphospholipid syndrome. <i>Movement Disorders</i> , 2006, 21, 944-949.	2.2	50
466	Botulinum toxin may be efficacious as treatment for jaw tremor in Parkinson's disease. <i>Movement Disorders</i> , 2006, 21, 1722-1724.	2.2	47
467	Secondary nonresponsiveness to botulinum toxin A in cervical dystonia: The role of electromyogram-guided injections, botulinum toxin A antibody assay, and the extensor digitorum brevis test. <i>Movement Disorders</i> , 2006, 21, 1737-1741.	2.2	61
468	Abnormalities in motor cortical plasticity differentiate manifesting and nonmanifesting DYT1 carriers. <i>Movement Disorders</i> , 2006, 21, 2181-2186.	2.2	137

#	ARTICLE	IF	CITATIONS
469	Defective temporal processing of sensory stimuli in DYT1 mutation carriers: a new endophenotype of dystonia?. <i>Brain</i> , 2006, 130, 134-142.	3.7	122
470	Corticobasal degeneration. , 2005, , 682-696.		0
471	Theta Burst Stimulation of the Human Motor Cortex. <i>Neuron</i> , 2005, 45, 201-206.	3.8	3,223
472	One-Hz repetitive transcranial magnetic stimulation of the premotor cortex alters reciprocal inhibition in DYT1 dystonia. <i>Movement Disorders</i> , 2004, 19, 54-59.	2.2	58
473	Choreic syndrome and coeliac disease: A hitherto unrecognised association. <i>Movement Disorders</i> , 2004, 19, 478-482.	2.2	53
474	Natural history and syndromic associations of orthostatic tremor: A review of 41 patients. <i>Movement Disorders</i> , 2004, 19, 788-795.	2.2	224
475	The syndrome of fixed dystonia: an evaluation of 103 patients. <i>Brain</i> , 2004, 127, 2360-2372.	3.7	338
476	Slowly progressive cerebellar ataxia and cervical dystonia: Clinical presentation of a new form of spinocerebellar ataxia?. <i>Movement Disorders</i> , 2003, 18, 200-206.	2.2	35
477	Unusual phenotypes in DYT1 dystonia: A report of five cases and a review of the literature. <i>Movement Disorders</i> , 2003, 18, 706-711.	2.2	137
478	Episodic focal lingual dystonic spasms. <i>Movement Disorders</i> , 2003, 18, 836-837.	2.2	16
479	Paroxysmal exercise-induced dystonia as a presenting feature of young-onset Parkinson's disease. <i>Movement Disorders</i> , 2003, 18, 1545-1547.	2.2	71
480	Different patterns of electrophysiological deficits in manifesting and non-manifesting carriers of the DYT1 gene mutation. <i>Brain</i> , 2003, 126, 2074-2080.	3.7	141
481	Other extrapyramidal syndromes: parkinsonism-plus and other forms of secondary parkinsonism. , 2002, , 490-512.		1
482	The Man who Walks Backwards. <i>Journal of the Royal Society of Medicine</i> , 2002, 95, 273-273.	1.1	3
483	Myoclonus-dystonia syndrome: $\hat{\mu}$ -sarcoglycan mutations and phenotype. <i>Annals of Neurology</i> , 2002, 52, 489-492.	2.8	143
484	Genetic and clinical heterogeneity in paroxysmal kinesigenic dyskinesia: Evidence for a third EKD gene. <i>Movement Disorders</i> , 2002, 17, 717-725.	2.2	85
485	Further case of paroxysmal exercise-induced dystonia and some insights into pathogenesis. <i>Movement Disorders</i> , 2002, 17, 1386-1387.	2.2	13
486	Updated guidelines for the management of Parkinson's disease. <i>British Journal of Hospital Medicine</i> , 2001, 62, 456-470.	0.3	25

#	ARTICLE	IF	CITATIONS
487	International Medical Workshop covering progressive supranuclear palsy, multiple system atrophy and cortico basal degeneration. Movement Disorders, 2001, 16, 382-395.	2.2	6
488	Benign hereditary chorea?Entity or syndrome?. Movement Disorders, 2000, 15, 280-288.	2.2	80
489	Episodic movement disorders as channelopathies. Movement Disorders, 2000, 15, 429-433.	2.2	47
490	A Yorkshire family with adult-onset cranio-cervical primary torsion dystonia. Movement Disorders, 2000, 15, 954-959.	2.2	38
491	Paroxysmal kinesigenic choreoathetosis: a report of 26 patients. Journal of Neurology, 1999, 246, 120-126.	1.8	108
492	The paroxysmal dyskinesias. Journal of Neurology, 1999, 246, 149-155.	1.8	144
493	Machado-Joseph disease presenting as severe generalised dystonia in a German patient. Journal of Neurology, 1999, 246, 840-842.	1.8	23
494	Atypical and typical cranial dystonia following dental procedures. Movement Disorders, 1999, 14, 492-496.	2.2	103
495	An unusual jaw tremor with characteristics of primary orthostatic tremor. Movement Disorders, 1999, 14, 528-530.	2.2	23
496	SSRI-induced reversal of levodopa benefit in two patients with dopa-responsive dystonia. Movement Disorders, 1999, 14, 874-876.	2.2	19
497	The dancing larynx?a variant of palatal tremor?. Movement Disorders, 1999, 14, 882-883.	2.2	6
498	Slater revisited: 6Âyear follow up study of patients with medically unexplained motor symptoms. BMJ: British Medical Journal, 1998, 316, 582-586.	2.4	280
499	Paroxysmal exercise-induced dystonia: Eight new sporadic cases and a review of the literature. Movement Disorders, 1997, 12, 1007-1012.	2.2	106
500	Focal task-specific tremors. Movement Disorders, 1996, 11, 665-670.	2.2	79
501	Hereditary geniospasm: Two new families. Movement Disorders, 1996, 11, 744-746.	2.2	57
502	Dystonia as the major manifestation of Leigh's syndrome. Movement Disorders, 1994, 9, 642-649.	2.2	36
503	Movement disorders in metabolic diseases in adulthood. , 0, , 99-114.		0
504	Treatment of Paroxysmal Dystonia. , 0, , 292-296.		0

#	ARTICLE	IF	CITATIONS
505	Periodic Limb Movements While Awake (PLMA) as a manifestation of Wearingâ€œOff in Parkinson's Disease: A Case Series and Review of the Literature. Movement Disorders Clinical Practice, 0, , .	0.8	0
506	Abnormal <scp>DaTscan</scp> in <scp>GM1</scp> â€œgangliosidosis type <scp>III</scp> manifesting with dystoniaâ€œparkinsonism. Movement Disorders Clinical Practice, 0, , .	0.8	0
507	Breakthrough News in Adenoviral Vectorâ€œMediated <scp>AADC</scp> Gene Therapy: Lessons from the Success in <scp>AADC</scp> Deficiency and Possible Future Applications. Movement Disorders Clinical Practice, 0, , .	0.8	1