# Kailash Phatechand Bhatia

## List of Publications by Citations



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|---|---|---|---|
| 495 papers | 23,888 citations | 76 h-index | 141 g-index |
| 520 ext. papers | 28,578 ext. citations | 6.1 avg, IF | 6.99 L-index |

| # | Paper | IF | Citations |
|---|---|---|---|
| 495 | Theta burst stimulation of the human motor cortex. *Neuron*, **2005**, 45, 201-6 | 13.9 | 2414 |
| 494 | Phenomenology and classification of dystonia: a consensus update. *Movement Disorders*, **2013**, 28, 863-73 | 7 | 1217 |
| 493 | Criteria for the diagnosis of corticobasal degeneration. *Neurology*, **2013**, 80, 496-503 | 6.5 | 1004 |
| 492 | Clinical diagnosis of progressive supranuclear palsy: The movement disorder society criteria. *Movement Disorders*, **2017**, 32, 853-864 | 7 | 840 |
| 491 | Consensus Statement on the classification of tremors. from the task force on tremor of the International Parkinson and Movement Disorder Society. *Movement Disorders*, **2018**, 33, 75-87 | 7 | 504 |
| 490 | Characterization of PLA2G6 as a locus for dystonia-parkinsonism. *Annals of Neurology*, **2009**, 65, 19-23 | 9.4 | 320 |
| 489 | The syndrome of fixed dystonia: an evaluation of 103 patients. *Brain*, **2004**, 127, 2360-72 | 11.2 | 286 |
| 488 | Exome sequencing reveals de novo WDR45 mutations causing a phenotypically distinct, X-linked dominant form of NBIA. *American Journal of Human Genetics*, **2012**, 91, 1144-9 | 11 | 268 |
| 487 | GLUT1 mutations are a cause of paroxysmal exertion-induced dyskinesias and induce hemolytic anemia by a cation leak. *Journal of Clinical Investigation*, **2008**, 118, 2157-68 | 15.9 | 263 |
| 486 | Clinical approach to Parkinson's disease: features, diagnosis, and principles of management. *Cold Spring Harbor Perspectives in Medicine*, **2012**, 2, a008870 | 5.4 | 225 |
| 485 | Slater revisited: 6 year follow up study of patients with medically unexplained motor symptoms. *BMJ: British Medical Journal*, **1998**, 316, 582-6 | | 212 |
| 484 | Pallidal neurostimulation in patients with medication-refractory cervical dystonia: a randomised, sham-controlled trial. *Lancet Neurology, The*, **2014**, 13, 875-84 | 24.1 | 210 |
| 483 | Long-term clinical outcome of fetal cell transplantation for Parkinson disease: two case reports. *JAMA Neurology*, **2014**, 71, 83-7 | 17.2 | 205 |
| 482 | Mutations in the gene PRRT2 cause paroxysmal kinesigenic dyskinesia with infantile convulsions. *Cell Reports*, **2012**, 1, 2-12 | 10.6 | 205 |
| 481 | Functional (psychogenic) movement disorders: merging mind and brain. *Lancet Neurology, The*, **2012**, 11, 250-60 | 24.1 | 202 |
| 480 | The non-motor syndrome of primary dystonia: clinical and pathophysiological implications. *Brain*, **2012**, 135, 1668-81 | 11.2 | 191 |
| 479 | Reward Pays the Cost of Noise Reduction in Motor and Cognitive Control. *Current Biology*, **2015**, 25, 1707-16 | 7.5 | 180 |

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| 478 | Dissection of the genetics of Parkinson's disease identifies an additional association 5' of SNCA and multiple associated haplotypes at 17q21. *Human Molecular Genetics*, **2011**, 20, 345-53 | 5.6 | 178 |
| 477 | Distinguishing SWEDDs patients with asymmetric resting tremor from Parkinson's disease: a clinical and electrophysiological study. *Movement Disorders*, **2010**, 25, 560-9 | 7 | 176 |
| 476 | Mutations in ANO3 cause dominant craniocervical dystonia: ion channel implicated in pathogenesis. *American Journal of Human Genetics*, **2012**, 91, 1041-50 | 11 | 172 |
| 475 | Early-onset L-dopa-responsive parkinsonism with pyramidal signs due to ATP13A2, PLA2G6, FBXO7 and spatacsin mutations. *Movement Disorders*, **2010**, 25, 1791-800 | 7 | 171 |
| 474 | Propeller protein-associated neurodegeneration: a new X-linked dominant disorder with brain iron accumulation. *Brain*, **2013**, 136, 1708-17 | 11.2 | 167 |
| 473 | Widespread Lewy body and tau accumulation in childhood and adult onset dystonia-parkinsonism cases with PLA2G6 mutations. *Neurobiology of Aging*, **2012**, 33, 814-23 | 5.6 | 151 |
| 472 | ATP13A2 mutations (PARK9) cause neurodegeneration with brain iron accumulation. *Movement Disorders*, **2010**, 25, 979-84 | 7 | 147 |
| 471 | PRRT2 gene mutations: from paroxysmal dyskinesia to episodic ataxia and hemiplegic migraine. *Neurology*, **2012**, 79, 2115-21 | 6.5 | 132 |
| 470 | Paroxysmal dyskinesias. *Movement Disorders*, **2011**, 26, 1157-65 | 7 | 130 |
| 469 | Natural history and syndromic associations of orthostatic tremor: a review of 41 patients. *Movement Disorders*, **2004**, 19, 788-795 | 7 | 130 |
| 468 | Complex movement disorders at disease onset in childhood narcolepsy with cataplexy. *Brain*, **2011**, 134, 3477-89 | 11.2 | 129 |
| 467 | Myoclonic disorders: a practical approach for diagnosis and treatment. *Therapeutic Advances in Neurological Disorders*, **2011**, 4, 47-62 | 6.6 | 128 |
| 466 | Parkinson's disease in GTP cyclohydrolase 1 mutation carriers. *Brain*, **2014**, 137, 2480-92 | 11.2 | 127 |
| 465 | 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 | 7 | 126 |
| 464 | Current Opinions and Areas of Consensus on the Role of the Cerebellum in Dystonia. *Cerebellum*, **2017**, 16, 577-594 | 4.3 | 125 |
| 463 | Genetic and phenotypic characterization of complex hereditary spastic paraplegia. *Brain*, **2016**, 139, 1904-18 | 11.2 | 123 |
| 462 | Paroxysmal dyskinesias revisited: a review of 500 genetically proven cases and a new classification. *Movement Disorders*, **2014**, 29, 1108-16 | 7 | 123 |
| 461 | Lysine 27 ubiquitination of the mitochondrial transport protein Miro is dependent on serine 65 of the Parkin ubiquitin ligase. *Journal of Biological Chemistry*, **2014**, 289, 14569-82 | 5.4 | 120 |

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| 460 | Different patterns of electrophysiological deficits in manifesting and non-manifesting carriers of the DYT1 gene mutation. *Brain*, **2003**, 126, 2074-80 | 11.2 | 118 |
| 459 | The expanding universe of disorders of the basal ganglia. *Lancet, The*, **2014**, 384, 523-31 | 40 | 117 |
| 458 | Dystonia. *Nature Reviews Disease Primers*, **2018**, 4, 25 | 51.1 | 117 |
| 457 | Mutations in the histone methyltransferase gene KMT2B cause complex early-onset dystonia. *Nature Genetics*, **2017**, 49, 223-237 | 36.3 | 116 |
| 456 | Abnormalities in motor cortical plasticity differentiate manifesting and nonmanifesting DYT1 carriers. *Movement Disorders*, **2006**, 21, 2181-6 | 7 | 116 |
| 455 | Dopamine and performance in a reinforcement learning task: evidence from Parkinson's disease. *Brain*, **2012**, 135, 1871-83 | 11.2 | 115 |
| 454 | Mutations in the autoregulatory domain of Ⅱubulin 4a cause hereditary dystonia. *Annals of Neurology*, **2013**, 73, 546-53 | 9.4 | 114 |
| 453 | Psychogenic movement disorders in children: a report of 15 cases and a review of the literature. *Movement Disorders*, **2008**, 23, 1882-8 | 7 | 114 |
| 452 | Myoclonus-dystonia syndrome: epsilon-sarcoglycan mutations and phenotype. *Annals of Neurology*, **2002**, 52, 489-92 | 9.4 | 114 |
| 451 | The paroxysmal dyskinesias. *Journal of Neurology*, **1999**, 246, 149-55 | 5.5 | 114 |
| 450 | Physical precipitating factors in functional movement disorders. *Journal of the Neurological Sciences*, **2014**, 338, 174-7 | 3.2 | 113 |
| 449 | Moving toward "laboratory-supported" criteria for psychogenic tremor. *Movement Disorders*, **2011**, 26, 2509-15 | 7 | 110 |
| 448 | Believing is perceiving: mismatch between self-report and actigraphy in psychogenic tremor. *Brain*, **2012**, 135, 117-23 | 11.2 | 102 |
| 447 | Defective temporal processing of sensory stimuli in DYT1 mutation carriers: a new endophenotype of dystonia?. *Brain*, **2007**, 130, 134-42 | 11.2 | 102 |
| 446 | Movement disorders with neuronal antibodies: syndromic approach, genetic parallels and pathophysiology. *Brain*, **2018**, 141, 13-36 | 11.2 | 100 |
| 445 | Loss of PLA2G6 leads to elevated mitochondrial lipid peroxidation and mitochondrial dysfunction. *Brain*, **2015**, 138, 1801-16 | 11.2 | 100 |
| 444 | Mutations in C2orf37, encoding a nucleolar protein, cause hypogonadism, alopecia, diabetes mellitus, mental retardation, and extrapyramidal syndrome. *American Journal of Human Genetics*, **2008**, 83, 684-91 | 11 | 100 |
| 443 | Paroxysmal kinesigenic choreoathetosis: a report of 26 patients. *Journal of Neurology*, **1999**, 246, 120-6 | 5.5 | 98 |

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| 442 | The clinical and genetic heterogeneity of paroxysmal dyskinesias. *Brain*, **2015**, 138, 3567-80 | 11.2 | 96 |
| 441 | Premonitory urge to tic in Tourette's is associated with interoceptive awareness. *Movement Disorders*, **2015**, 30, 1198-202 | 7 | 96 |
| 440 | A pathway-based analysis provides additional support for an immune-related genetic susceptibility to Parkinson's disease. *Human Molecular Genetics*, **2013**, 22, 1039-49 | 5.6 | 96 |
| 439 | Paroxysmal exercise-induced dystonia: eight new sporadic cases and a review of the literature. *Movement Disorders*, **1997**, 12, 1007-12 | 7 | 96 |
| 438 | Clinical and polysomnographic course of childhood narcolepsy with cataplexy. *Brain*, **2013**, 136, 3787-95 | 11.2 | 95 |
| 437 | Deep brain stimulation effects in dystonia: time course of electrophysiological changes in early treatment. *Movement Disorders*, **2011**, 26, 1913-21 | 7 | 95 |
| 436 | GLUT1 gene mutations cause sporadic paroxysmal exercise-induced dyskinesias. *Movement Disorders*, **2009**, 24, 1684-8 | 7 | 94 |
| 435 | Sensory functions in dystonia: insights from behavioral studies. *Movement Disorders*, **2009**, 24, 1427-36 | 7 | 93 |
| 434 | Huntington's disease phenocopies are clinically and genetically heterogeneous. *Movement Disorders*, **2008**, 23, 716-20 | 7 | 93 |
| 433 | Cortical excitability is abnormal in patients with the "fixed dystonia" syndrome. *Movement Disorders* , **2008**, 23, 646-52 | 7 | 91 |
| 432 | Which ante mortem clinical features predict progressive supranuclear palsy pathology?. *Movement Disorders*, **2017**, 32, 995-1005 | 7 | 88 |
| 431 | The glucocerobrosidase E326K variant predisposes to Parkinson's disease, but does not cause Gaucher's disease. *Movement Disorders*, **2013**, 28, 232-236 | 7 | 86 |
| 430 | Tardive dyskinesia is caused by maladaptive synaptic plasticity: a hypothesis. *Movement Disorders*, **2012**, 27, 1205-15 | 7 | 86 |
| 429 | Atypical and typical cranial dystonia following dental procedures. *Movement Disorders*, **1999**, 14, 492-6 | 7 | 86 |
| 428 | What do patients with scans without evidence of dopaminergic deficit (SWEDD) have? New evidence and continuing controversies. *Journal of Neurology, Neurosurgery and Psychiatry*, **2016**, 87, 319-23 | 5.5 | 85 |
| 427 | Dystonia with brain manganese accumulation resulting from SLC30A10 mutations: a new treatable disorder. *Movement Disorders*, **2012**, 27, 1317-22 | 7 | 85 |
| 426 | Genetics and Pathophysiology of Neurodegeneration with Brain Iron Accumulation (NBIA). *Current Neuropharmacology*, **2013**, 11, 59-79 | 7.6 | 85 |
| 425 | Secondary and primary dystonia: pathophysiological differences. *Brain*, **2013**, 136, 2038-49 | 11.2 | 84 |

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| 424 | Clinical relevance of serum antibodies to extracellular N-methyl-D-aspartate receptor epitopes. *Journal of Neurology, Neurosurgery and Psychiatry*, **2015**, 86, 708-13 | 5.5 | 81 |
| 423 | The genetics of dystonia: new twists in an old tale. *Brain*, **2013**, 136, 2017-37 | 11.2 | 80 |
| 422 | A missense mutation in KCTD17 causes autosomal dominant myoclonus-dystonia. *American Journal of Human Genetics*, **2015**, 96, 938-47 | 11 | 77 |
| 421 | "Atypical" atypical parkinsonism: new genetic conditions presenting with features of progressive supranuclear palsy, corticobasal degeneration, or multiple system atrophy-a diagnostic guide. *Movement Disorders*, **2013**, 28, 1184-99 | 7 | 76 |
| 420 | Stereotypies: a critical appraisal and suggestion of a clinically useful definition. *Movement Disorders* , **2012**, 27, 179-85 | 7 | 74 |
| 419 | De Novo Mutations in PDE10A Cause Childhood-Onset Chorea with Bilateral Striatal Lesions. *American Journal of Human Genetics*, **2016**, 98, 763-71 | 11 | 74 |
| 418 | Network localization of cervical dystonia based on causal brain lesions. *Brain*, **2019**, 142, 1660-1674 | 11.2 | 73 |
| 417 | Psychogenic facial movement disorders: clinical features and associated conditions. *Movement Disorders*, **2012**, 27, 1544-51 | 7 | 73 |
| 416 | The functional neuroimaging correlates of psychogenic versus organic dystonia. *Brain*, **2013**, 136, 770-81 | 11.2 | 73 |
| 415 | Excessive daytime sleepiness in multiple system atrophy (SLEEMSA study). *Archives of Neurology*, **2011**, 68, 223-30 | | 73 |
| 414 | Genetic and clinical heterogeneity in paroxysmal kinesigenic dyskinesia: evidence for a third EKD gene. *Movement Disorders*, **2002**, 17, 717-25 | 7 | 72 |
| 413 | Benign hereditary chorea--entity or syndrome?. *Movement Disorders*, **2000**, 15, 280-8 | 7 | 72 |
| 412 | Abnormal motor cortex plasticity in premanifest and very early manifest Huntington disease. *Journal of Neurology, Neurosurgery and Psychiatry*, **2010**, 81, 267-70 | 5.5 | 71 |
| 411 | Propriospinal myoclonus: clinical reappraisal and review of literature. *Neurology*, **2014**, 83, 1862-70 | 6.5 | 69 |
| 410 | Loss of sensory attenuation in patients with functional (psychogenic) movement disorders. *Brain*, **2014**, 137, 2916-21 | 11.2 | 67 |
| 409 | The phenotypic spectrum of DYT24 due to ANO3 mutations. *Movement Disorders*, **2014**, 29, 928-34 | 7 | 67 |
| 408 | Repetitive transcranial magnetic stimulation for levodopa-induced dyskinesias in Parkinson's disease. *Movement Disorders*, **2009**, 24, 246-53 | 7 | 67 |
| 407 | Differentiating drug-induced parkinsonism from Parkinson's disease: an update on non-motor symptoms and investigations. *Parkinsonism and Related Disorders*, **2014**, 20, 808-14 | 3.6 | 66 |

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| 406 | Psychogenic paroxysmal movement disorders--clinical features and diagnostic clues. *Parkinsonism and Related Disorders*, **2014**, 20, 41-6 | 3.6 | 66 |
| 405 | Abnormal sense of intention preceding voluntary movement in patients with psychogenic tremor. *Neuropsychologia*, **2011**, 49, 2791-3 | 3.2 | 66 |
| 404 | Focal task-specific tremors. *Movement Disorders*, **1996**, 11, 665-70 | 7 | 65 |
| 403 | ADCY5 mutations are another cause of benign hereditary chorea. *Neurology*, **2015**, 85, 80-8 | 6.5 | 63 |
| 402 | RAD51 haploinsufficiency causes congenital mirror movements in humans. *American Journal of Human Genetics*, **2012**, 90, 301-7 | 11 | 63 |
| 401 | Assessment of patients with isolated or combined dystonia: an update on dystonia syndromes. *Movement Disorders*, **2013**, 28, 889-98 | 7 | 62 |
| 400 | Paroxysmal exercise-induced dystonia as a presenting feature of young-onset Parkinson's disease. *Movement Disorders*, **2003**, 18, 1545-7 | 7 | 61 |
| 399 | Rest and other types of tremor in adult-onset primary dystonia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2014**, 85, 965-8 | 5.5 | 60 |
| 398 | Mutations in HPCA cause autosomal-recessive primary isolated dystonia. *American Journal of Human Genetics*, **2015**, 96, 657-65 | 11 | 59 |
| 397 | Movement disorders on YouTube--caveat spectator. *New England Journal of Medicine*, **2011**, 365, 1160-1 | 59.2 | 59 |
| 396 | Validation of "laboratory-supported" criteria for functional (psychogenic) tremor. *Movement Disorders*, **2016**, 31, 555-62 | 7 | 59 |
| 395 | Tremor stability index: a new tool for differential diagnosis in tremor syndromes. *Brain*, **2017**, 140, 1977-1986 | 11.2 | 58 |
| 394 | The differential diagnosis of Huntington's disease-like syndromes: 'red flags' for the clinician. *Journal of Neurology, Neurosurgery and Psychiatry*, **2013**, 84, 650-6 | 5.5 | 58 |
| 393 | Treatable inherited rare movement disorders. *Movement Disorders*, **2018**, 33, 21-35 | 7 | 57 |
| 392 | Dystonia: an update on phenomenology, classification, pathogenesis and treatment. *Current Opinion in Neurology*, **2014**, 27, 468-76 | 7.1 | 57 |
| 391 | Dystonia in the Woodhouse Sakati syndrome: A new family and literature review. *Movement Disorders*, **2008**, 23, 592-6 | 7 | 57 |
| 390 | How to apply the movement disorder society criteria for diagnosis of progressive supranuclear palsy. *Movement Disorders*, **2019**, 34, 1228-1232 | 7 | 56 |
| 389 | ADCY5-related movement disorders: Frequency, disease course and phenotypic variability in a cohort of paediatric patients. *Parkinsonism and Related Disorders*, **2017**, 41, 37-43 | 3.6 | 54 |

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| 388 | Hereditary geniospasm: two new families. *Movement Disorders*, **1996**, 11, 744-6 | 7 | 53 |
| 387 | Essential pitfalls in "essential" tremor. *Movement Disorders*, **2017**, 32, 325-331 | 7 | 52 |
| 386 | Ataxia telangiectasia presenting as dopa-responsive cervical dystonia. *Neurology*, **2013**, 81, 1148-51 | 6.5 | 52 |
| 385 | 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. *Movement Disorders*, **2006**, 21, 1737-41 | 7 | 52 |
| 384 | Pathogenesis of dystonia: is it of cerebellar or basal ganglia origin?. *Journal of Neurology, Neurosurgery and Psychiatry*, **2018**, 89, 488-492 | 5.5 | 51 |
| 383 | A 6.4 Mb duplication of the □synuclein locus causing frontotemporal dementia and Parkinsonism: phenotype-genotype correlations. *JAMA Neurology*, **2014**, 71, 1162-71 | 17.2 | 51 |
| 382 | Unusual phenotypes in DYT1 dystonia: a report of five cases and a review of the literature. *Movement Disorders*, **2003**, 18, 706-11 | 7 | 51 |
| 381 | Neurophysiological correlates of abnormal somatosensory temporal discrimination in dystonia. *Movement Disorders*, **2017**, 32, 141-148 | 7 | 50 |
| 380 | The epileptic and nonepileptic spectrum of paroxysmal dyskinesias: Channelopathies, synaptopathies, and transportopathies. *Movement Disorders*, **2017**, 32, 310-318 | 7 | 49 |
| 379 | Functional neurological disorders in Parkinson disease. *Journal of Neurology, Neurosurgery and Psychiatry*, **2018**, 89, 566-571 | 5.5 | 49 |
| 378 | Psychogenic palatal tremor may be underrecognized: reappraisal of a large series of cases. *Movement Disorders*, **2012**, 27, 1164-1168 | 7 | 49 |
| 377 | Deconstructing Fahr's disease/syndrome of brain calcification in the era of new genes. *Parkinsonism and Related Disorders*, **2017**, 37, 1-10 | 3.6 | 48 |
| 376 | The□Movement 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 | 5.5 | 48 |
| 375 | Syndromes of neurodegeneration with brain iron accumulation. *Seminars in Pediatric Neurology*, **2012**, 19, 57-66 | 2.9 | 48 |
| 374 | Oculogyric crises: Etiology, pathophysiology and therapeutic approaches. *Parkinsonism and Related Disorders*, **2017**, 36, 3-9 | 3.6 | 48 |
| 373 | THAP1 mutations and dystonia phenotypes: genotype phenotype correlations. *Movement Disorders*, **2012**, 27, 1290-4 | 7 | 46 |
| 372 | Dystonia in corticobasal degeneration: a review of the literature on 404 pathologically proven cases. *Movement Disorders*, **2012**, 27, 696-702 | 7 | 46 |
| 371 | Dopaminergic neuronal imaging in genetic Parkinson's disease: insights into pathogenesis. *PLoS ONE*, **2013**, 8, e69190 | 3.7 | 46 |

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| 370 | One-Hz repetitive transcranial magnetic stimulation of the premotor cortex alters reciprocal inhibition in DYT1 dystonia. *Movement Disorders*, **2004**, 19, 54-9 | 7 | 46 |
| 369 | Choreic syndrome and coeliac disease: a hitherto unrecognised association. *Movement Disorders*, **2004**, 19, 478-82 | 7 | 46 |
| 368 | Tics and functional tic-like movements: Can we tell them apart?. *Neurology*, **2019**, 93, 750-758 | 6.5 | 45 |
| 367 | Screening for VPS35 mutations in Parkinson's disease. *Neurobiology of Aging*, **2012**, 33, 838.e1-5 | 5.6 | 45 |
| 366 | Functional (psychogenic) symptoms in Parkinson's disease. *Movement Disorders*, **2013**, 28, 1622-7 | 7 | 44 |
| 365 | Pallidal stimulation for primary generalised dystonia: effect on cognition, mood and quality of life. *Journal of Neurology*, **2014**, 261, 164-73 | 5.5 | 43 |
| 364 | Clinical Practice: Evidence-Based Recommendations for the Treatment of Cervical Dystonia with Botulinum Toxin. *Frontiers in Neurology*, **2017**, 8, 35 | 4.1 | 43 |
| 363 | Slow (1 Hz) repetitive transcranial magnetic stimulation (rTMS) induces a sustained change in cortical excitability in patients with Parkinson's disease. *Clinical Neurophysiology*, **2010**, 121, 1129-37 | 4.3 | 43 |
| 362 | Atypical movement disorders in antiphospholipid syndrome. *Movement Disorders*, **2006**, 21, 944-9 | 7 | 43 |
| 361 | Transducer-based evaluation of tremor. *Movement Disorders*, **2016**, 31, 1327-36 | 7 | 43 |
| 360 | Clinical diagnosis of propriospinal myoclonus is unreliable: an electrophysiologic study. *Movement Disorders*, **2013**, 28, 1868-73 | 7 | 42 |
| 359 | Movement disorders in adult patients with classical galactosemia. *Movement Disorders*, **2013**, 28, 804-107 | | 42 |
| 358 | Botulinum toxin injections reduce associative plasticity in patients with primary dystonia. *Movement Disorders*, **2011**, 26, 1282-9 | 7 | 42 |
| 357 | Attention to self in psychogenic tremor. *Movement Disorders*, **2011**, 26, 2575-6 | 7 | 42 |
| 356 | Characteristic head drops and axial extension in advanced chorea-acanthocytosis. *Movement Disorders*, **2010**, 25, 1487-91 | 7 | 42 |
| 355 | Botulinum toxin A may be efficacious as treatment for jaw tremor in Parkinson's disease. *Movement Disorders*, **2006**, 21, 1722-4 | 7 | 42 |
| 354 | Unravelling of the paroxysmal dyskinesias. *Journal of Neurology, Neurosurgery and Psychiatry*, **2019**, 90, 227-234 | 5.5 | 42 |
| 353 | Tremor in inflammatory neuropathies. *Journal of Neurology, Neurosurgery and Psychiatry*, **2013**, 84, 1282-75 | 5.5 | 41 |

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| 352 | Early and marked benefit with GPi DBS for Lubag syndrome presenting with rapidly progressive life-threatening dystonia. *Movement Disorders*, **2009**, 24, 1710-2 | 7 | 41 |
| 351 | Cerebellar stimulation fails to modulate motor cortex plasticity in writing dystonia. *Movement Disorders*, **2014**, 29, 1304-7 | 7 | 40 |
| 350 | The distinct movement disorder in anti-NMDA receptor encephalitis may be related to Status Dissociatus: a hypothesis. *Movement Disorders*, **2012**, 27, 1360-3 | 7 | 40 |
| 349 | Transcranial magnetic stimulation follow-up study in early Parkinson's disease: A decline in compensation with disease progression?. *Movement Disorders*, **2015**, 30, 1098-106 | 7 | 39 |
| 348 | From state dissociation to status dissociatus. *Sleep Medicine Reviews*, **2016**, 28, 5-17 | 10.2 | 39 |
| 347 | Excess iron harms the brain: the syndromes of neurodegeneration with brain iron accumulation (NBIA). *Journal of Neural Transmission*, **2013**, 120, 695-703 | 4.3 | 38 |
| 346 | Motivation and movement: the effect of monetary incentive on performance speed. *Experimental Brain Research*, **2011**, 209, 551-9 | 2.3 | 38 |
| 345 | Dystonia and Parkinson's disease: What is the relationship?. *Neurobiology of Disease*, **2019**, 132, 104462 | 7.5 | 37 |
| 344 | SLC25A46 mutations underlie progressive myoclonic ataxia with optic atrophy and neuropathy. *Movement Disorders*, **2016**, 31, 1249-51 | 7 | 37 |
| 343 | Failure of explicit movement control in patients with functional motor symptoms. *Movement Disorders*, **2013**, 28, 517-23 | 7 | 37 |
| 342 | 'Jumping to conclusions' bias in functional movement disorders. *Journal of Neurology, Neurosurgery and Psychiatry*, **2012**, 83, 460-3 | 5.5 | 37 |
| 341 | Parkinsonism following neuroleptic exposure: A double-hit hypothesis?. *Movement Disorders*, **2015**, 30, 780-5 | 7 | 36 |
| 340 | H-ABC syndrome and DYT4: Variable expressivity or pleiotropy of TUBB4 mutations?. *Movement Disorders*, **2015**, 30, 828-33 | 7 | 36 |
| 339 | A Yorkshire family with adult-onset cranio-cervical primary torsion dystonia. *Movement Disorders*, **2000**, 15, 954-9 | 7 | 36 |
| 338 | Immediate response to botulinum toxin injections in patients with fixed dystonia. *Movement Disorders*, **2011**, 26, 917-8 | 7 | 35 |
| 337 | Limb amputations in fixed dystonia: a form of body integrity identity disorder?. *Movement Disorders* , **2011**, 26, 1410-4 | 7 | 34 |
| 336 | Prominent oromandibular dystonia and pharyngeal telangiectasia in atypical ataxia telangiectasia. *Cerebellum*, **2009**, 8, 22-7 | 4.3 | 34 |
| 335 | Unilateral cerebellothalamic tract ablation in essential tremor by MRI-guided focused ultrasound. *Neurology*, **2017**, 88, 1329-1333 | 6.5 | 33 |

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| 334 | Functional movement disorders are not uncommon in the elderly. *Movement Disorders*, **2013**, 28, 540-3 | 7 | 33 |
| 333 | Knowledge gaps and research recommendations for essential tremor. *Parkinsonism and Related Disorders*, **2016**, 33, 27-35 | 3.6 | 33 |
| 332 | High frequency somatosensory stimulation increases sensori-motor inhibition and leads to perceptual improvement in healthy subjects. *Clinical Neurophysiology*, **2017**, 128, 1015-1025 | 4.3 | 32 |
| 331 | The role of the cerebellum in the pathogenesis of cortical myoclonus. *Movement Disorders*, **2014**, 29, 437-43 | 7 | 32 |
| 330 | Causes of failure of pallidal deep brain stimulation in cases with pre-operative diagnosis of isolated dystonia. *Parkinsonism and Related Disorders*, **2017**, 43, 38-48 | 3.6 | 32 |
| 329 | Analysis of C9orf72 repeat expansions in a large series of clinically and pathologically diagnosed cases with atypical parkinsonism. *Neurobiology of Aging*, **2015**, 36, 1221.e1-6 | 5.6 | 32 |
| 328 | Myoclonus-dystonia syndrome due to tyrosine hydroxylase deficiency. *Neurology*, **2012**, 79, 435-41 | 6.5 | 32 |
| 327 | Short and long term outcome of bilateral pallidal stimulation in chorea-acanthocytosis. *PLoS ONE*, **2013**, 8, e79241 | 3.7 | 32 |
| 326 | Valproate-Associated Parkinsonism: A Critical Review of the Literature. *CNS Drugs*, **2016**, 30, 527-40 | 6.7 | 31 |
| 325 | Dystonic opisthotonus: a "red flag" for neurodegeneration with brain iron accumulation syndromes?. *Movement Disorders*, **2013**, 28, 1325-9 | 7 | 31 |
| 324 | Cerebellar brain inhibition is decreased in active and surround muscles at the onset of voluntary movement. *Experimental Brain Research*, **2011**, 209, 437-42 | 2.3 | 31 |
| 323 | Rare causes of dystonia parkinsonism. *Current Neurology and Neuroscience Reports*, **2010**, 10, 431-9 | 6.6 | 31 |
| 322 | Dystonia as the major manifestation of Leigh's syndrome. *Movement Disorders*, **1994**, 9, 642-9 | 7 | 31 |
| 321 | High frequency somatosensory stimulation in dystonia: Evidence fordefective inhibitory plasticity. *Movement Disorders*, **2018**, 33, 1902-1909 | 7 | 31 |
| 320 | A reflection on plasticity research in writing dystonia. *Movement Disorders*, **2014**, 29, 980-7 | 7 | 30 |
| 319 | The interindividual variability of transcranial magnetic stimulation effects: Implications for diagnostic use in movement disorders. *Movement Disorders*, **2019**, 34, 936-949 | 7 | 29 |
| 318 | The long-term outcome of orthostatic tremor. *Journal of Neurology, Neurosurgery and Psychiatry*, **2016**, 87, 167-72 | 5.5 | 29 |
| 317 | Atypical parkinsonism: diagnosis and treatment. *Neurologic Clinics*, **2015**, 33, 39-56 | 4.5 | 29 |

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