Bettina Balint

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9157669/publications.pdf

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81 papers	2,696 citations	236925 25 h-index	206112 48 g-index
83 all docs	83 docs citations	83 times ranked	3255 citing authors

#	Article	IF	Citations
1	Glycine receptor antibodies in PERM and related syndromes: characteristics, clinical features and outcomes. Brain, 2014, 137, 2178-2192.	7.6	430
2	Dystonia. Nature Reviews Disease Primers, 2018, 4, 25.	30.5	223
3	Progressive encephalomyelitis with rigidity and myoclonus. Neurology, 2014, 82, 1521-1528.	1.1	168
4	Movement disorders with neuronal antibodies: syndromic approach, genetic parallels and pathophysiology. Brain, 2018, 141, 13-36.	7.6	145
5	ADCY5 mutations are another cause of benign hereditary chorea. Neurology, 2015, 85, 80-88.	1.1	140
6	De Novo Mutations in PDE10A Cause Childhood-Onset Chorea with Bilateral Striatal Lesions. American Journal of Human Genetics, 2016, 98, 763-771.	6.2	96
7	Cortical pencil lining in neuroferritinopathy: A diagnostic clue. Neurology, 2015, 84, 1816-1818.	1.1	93
8	Isolated and combined dystonia syndromes â€“ an update on new genes and their phenotypes. European Journal of Neurology, 2015, 22, 610-617.	3.3	82
9	Dystonia. Current Opinion in Neurology, 2014, 27, 468-476.	3.6	81
10	Functional neurological disorders in Parkinson disease. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 566-571.	1.9	76
11	TheÂMovement disorder associated with NMDAR antibody-encephalitis is complex and characteristic: an expert video-rating study. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 724-726.	1.9	71
12	Deconstructing Fahr's disease/syndrome of brain calcification in the era of new genes. Parkinsonism and Related Disorders, 2017, 37, 1-10.	2.2	63
13	T-cell homeostasis in pediatric multiple sclerosis. Neurology, 2013, 81, 784-792.	1.1	62
14	The role of cerebellum in patients with late onset cervical/segmental dystonia?â€“Evidence from the clinic. Parkinsonism and Related Disorders, 2015, 21, 1317-1322.	2.2	57
15	KMT2B-related disorders: expansion of the phenotypic spectrum and long-term efficacy of deep brain stimulation. Brain, 2020, 143, 3242-3261.	7.6	57
16	Why is there motor deterioration in Parkinsonâ€™s disease during systemic infections-a hypothetical view. Npj Parkinson's Disease, 2015, 1, 15014.	5.3	54
17	Stiff person syndrome and other immune-mediated movement disorders â€“ new insights. Current Opinion in Neurology, 2016, 29, 496-506.	3.6	47
18	The clinical features, underlying immunology, and treatment of autoantibodyâ€mediated movement disorders. Movement Disorders, 2018, 33, 1376-1389.	3.9	44

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19	PDE10A and ADCY5 mutations linked to molecular and microstructural basal ganglia pathology. Movement Disorders, 2018, 33, 1961-1965.	3.9	38
20	B-cell populations discriminate between pediatric- and adult-onset multiple sclerosis. Neurology: Neuroimmunology and NeuroInflammation, 2017, 4, e309.	6.0	35
21	Caspr2 antibodies in limbic encephalitis with cerebellar ataxia, dyskinesias and myoclonus. Journal of the Neurological Sciences, 2013, 327, 73-74.	0.6	34
22	Partial loss-of-function of sodium channel SCN8A in familial isolated myoclonus. Human Mutation, 2018, 39, 965-969.	2.5	34
23	Dissecting the Phenotype and Genotype of <scp>PLA2G6</scp>â€Related Parkinsonism. Movement Disorders, 2022, 37, 148-161.	3.9	32
24	Pragmatic Treatment of Stiff Person Spectrum Disorders. Movement Disorders Clinical Practice, 2018, 5, 394-401.	1.5	31
25	Twenty years on: Myoclonusâ€dystonia and Îµâ€sarcoglycan â€" neurodevelopment, channel, and signaling dysfunction. Movement Disorders, 2019, 34, 1588-1601.	3.9	31
26	The spectrum of involuntary vocalizations in humans: A video atlas. Movement Disorders, 2019, 34, 1774-1791.	3.9	24
27	KCNN2 mutation in autosomalâ€dominant tremulous myoclonusâ€dystonia. European Journal of Neurology, 2020, 27, 1471-1477.	3.3	21
28	Antibody-related movement disorders â€" a comprehensive review of phenotype-autoantibody correlations and a guide to testing. Neurological Research and Practice, 2020, 2, 6.	2.0	21
29	Worldwide barriers to genetic testing for movement disorders. European Journal of Neurology, 2021, 28, 1901-1909.	3.3	21
30	Validation of a selfâ€completed Dystonia Nonâ€Motor Symptoms Questionnaire. Annals of Clinical and Translational Neurology, 2019, 6, 2054-2065.	3.7	20
31	Biallelic variants in TSPOAP1, encoding the active-zone protein RIMBP1, cause autosomal recessive dystonia. Journal of Clinical Investigation, 2021, 131, .	8.2	18
32	Temporal Discrimination is Altered in Patients With Isolated Asymmetric and Jerky Upper Limb Tremor. Movement Disorders, 2020, 35, 306-315.	3.9	17
33	Early Ataxia and Subsequent Parkinsonism: PLA2G6 Mutations Cause a Continuum Rather Than Three Discrete Phenotypes. Movement Disorders Clinical Practice, 2017, 4, 125-128.	1.5	16
34	Paroxysmal Asymmetric Dystonic Arm Posturingâ€"A Less Recognized but Characteristic Manifestation of ATP1A3â€related disease. Movement Disorders Clinical Practice, 2019, 6, 312-315.	1.5	15
35	Friend or Foe? IgLON5 antibodies in a novel tauopathy with prominent sleep movement disorder, ataxia, and chorea. Movement Disorders, 2014, 29, 989-989.	3.9	13
36	MYORG -related disease is associated with central pontine calcifications and atypical parkinsonism. Neurology: Genetics, 2020, 6, e399.	1.9	13

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37	Isolated and combined genetic tremor syndromes: a critical appraisal based on the 2018 MDS criteria. Parkinsonism and Related Disorders, 2020, 77, 121-140.	2.2	13
38	â€œAtypicalâ€•atypical parkinsonism: Critical appraisal of a cohort. Parkinsonism and Related Disorders, 2017, 37, 36-42.	2.2	12
39	Intermittent head drops: the differential spectrum. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 414-419.	1.9	11
40	â€œAntibody of Unknown Significanceâ€•(AUS): The Issue of Interpreting Antibody Test Results. Movement Disorders, 2021, 36, 1543-1547.	3.9	11
41	Facial tremor in dystonia. Parkinsonism and Related Disorders, 2014, 20, 924-925.	2.2	10
42	The clinical syndrome of dystonia with anarthria/aphonia. Parkinsonism and Related Disorders, 2016, 24, 20-27.	2.2	10
43	Oculomotor apraxia and disrupted sleep with nocturnal ballistic bouts in ADCY5-related disease. Parkinsonism and Related Disorders, 2018, 54, 103-106.	2.2	10
44	Delineating the phenotype of autosomalâ€•recessive HPCA mutations: Not only isolated dystonia!. Movement Disorders, 2019, 34, 589-592.	3.9	10
45	Sensory trick efficacy in cervical dystonia is linked to processing of neck proprioception. Parkinsonism and Related Disorders, 2019, 61, 50-56.	2.2	10
46	Tardive syndromes. Practical Neurology, 2020, 20, 368-376.	1.1	10
47	Movement disorders in systemic autoimmune diseases: Clinical spectrum, ancillary investigations, pathophysiological considerations. Parkinsonism and Related Disorders, 2021, 88, 116-128.	2.2	10
48	Development of parkinsonism after long-standing cervical dystonia â€“ A cohort. Journal of the Neurological Sciences, 2021, 427, 117477.	0.6	10
49	Homerâ€3 Antibody Disease: A Potentially Treatable MSAâ€C Mimic. Movement Disorders Clinical Practice, 2022, 9, 178-182.	1.5	10
50	Movement disorders associated with neuronal antibodies: a data-driven approach. Journal of Neurology, 2022, 269, 3511-3521.	3.6	10
51	Stiff Limb Syndrome Mimicking Corticobasal Syndrome. Movement Disorders Clinical Practice, 2014, 1, 354-356.	1.5	9
52	Mitochondrial complex I NUBPL mutations cause combined dystonia with bilateral striatal necrosis and cerebellar atrophy. European Journal of Neurology, 2019, 26, 1240-1243.	3.3	9
53	Autoimmune movement disorders with neuronal antibodies â€“ an update. Current Opinion in Neurology, 2021, 34, 565-571.	3.6	9
54	Video-tutorial for the Movement Disorder Society criteria for progressive supranuclear palsy. Parkinsonism and Related Disorders, 2020, 78, 200-203.	2.2	8

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55	Parkinsonism and dystonia: Clinical spectrum and diagnostic clues. Journal of the Neurological Sciences, 2022, 433, 120016.	0.6	8
56	Axial Dystonia Mimicking Stiff Person Syndrome. Movement Disorders Clinical Practice, 2016, 3, 176-179.	1.5	7
57	Quick Flicks: Association of Paroxysmal Kinesigenic Dyskinesia and Tics. Movement Disorders Clinical Practice, 2018, 5, 317-320.	1.5	6
58	Stimulus Sensitive Foot Myoclonus: A Clue to Coeliac Disease. Movement Disorders Clinical Practice, 2019, 6, 320-323.	1.5	6
59	Voluntary Inhibitory Control of Chorea: A Case Series. Movement Disorders Clinical Practice, 2020, 7, 308-312.	1.5	6
60	Parkinsonism and Other Movement Disorders Associated with Chediakâ€Higashi Syndrome: Case Report and Systematic Literature Review. Movement Disorders Clinical Practice, 2015, 2, 93-98.	1.5	5
61	Tremor in motor neuron disease may be central rather than peripheral in origin. European Journal of Neurology, 2019, 26, 394.	3.3	5
62	Benign tremulous parkinsonism of the young-consider Parkin. Parkinsonism and Related Disorders, 2019, 65, 270-271.	2.2	5
63	Brachial Neuritis After Botulinum Toxin Injections for Cervical Dystonia: A Need for a Reappraisal?. Movement Disorders Clinical Practice, 2019, 6, 160-165.	1.5	5
64	Teaching Video NeuroImages: Cold-induced laryngeal pseudomyotonia in Isaacs syndrome. Neurology, 2019, 92, e2734.	1.1	4
65	Reopening the case for antiâ€“basal ganglia antibodies (ABGAs): Identification of dopamineâ€2 receptor antibodies associated with movement disorders. Movement Disorders, 2013, 28, 733-733.	3.9	3
66	PKD or Not PKD: That is the question. Annals of Neurology, 2016, 80, 167-168.	5.3	3
67	Development and clinimetric assessment of a nurse-administered screening tool for movement disorders in psychosis. BJPsych Open, 2018, 4, 404-410.	0.7	3
68	Are Antibody Panels <scp>Underâ€Utilized</scp> in Movement Disorders Diagnosis? Yes. Movement Disorders Clinical Practice, 2021, 8, 341-346.	1.5	3
69	KMT2B: A new twist in dystonia genetics. Movement Disorders, 2017, 32, 529-529.	3.9	2
70	Expert comment to: Novel Xp21.1 deletion associated with unusual features in large McLeod syndrome kindred. Parkinsonism and Related Disorders, 2020, 79, 133-134.	2.2	2
71	Self-concocted, curious and creative coping strategies in movement disorders. Parkinsonism and Related Disorders, 2021, 83, 140-143.	2.2	2
72	Hot topic: Recessive mutations in the a3(VI) collagen gene COL6A3 cause early-onset isolated dystonia. Movement Disorders, 2015, 30, 1622-1622.	3.9	1

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73	T1-weighted basal ganglia hyperintensities due to gadolinium deposition â€“ a cautionary note. Parkinsonism and Related Disorders, 2016, 32, 135-136.	2.2	1
74	The Discovery of Central Nervous System Lymphatic Vessels: The Missing Link That Closes the Circle of Brain Immunosurveillance. Movement Disorders Clinical Practice, 2016, 3, 29-30.	1.5	1
75	Conjugal Parkinson's disease â€“ Real or chance?. Parkinsonism and Related Disorders, 2016, 33, 146-148.	2.2	1
76	Reply to: Youngâ€•onset multiple system atrophy. Movement Disorders, 2018, 33, 1975-1976.	3.9	1
77	Editorial: Genetics of Paroxysmal Movement Disorders. Frontiers in Neurology, 2021, 12, 752000.	2.4	1
78	Hot topic: PNKP mutations cause ataxia with oculomotor apraxia type 4. Movement Disorders, 2016, 31, 500-500.	3.9	0
79	Commentary: Progressive Encephalomyelitis with Rigidity and Myoclonus and Myasthenia Gravis Comorbid Status with Thymoma. Movement Disorders Clinical Practice, 2021, 8, S14-S15.	1.5	0
80	Reply to: Comparing <scp>VUS</scp> and <scp>AUS</scp>: Parallels and Differences in Neurogenetics and Neuroimmunology. Movement Disorders, 2021, 36, 2454-2456.	3.9	0
81	Disentangling the phenotypes of <scp>antiâ€IgLON5</scp> disease: when clinical suspicion matters. Movement Disorders Clinical Practice, 2022, 9, 456-457.	1.5	0