JosÃ© Luiz Pedroso

List of Publications by Year
in descending order

279	3,252	218592	276775
papers	citations	26 h-index	41 g-index

285	285	285	3978
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Spinocerebellar Ataxias in Brazilâ€"Frequencies and Modulating Effects of Related Genes. Cerebellum, 2014, 13, 17-28.	1.4	93
2	Structural signature of SCA3: From presymptomatic to late disease stages. Annals of Neurology, 2018, 84, 401-408.	2.8	90
3	Autoimmune encephalitis: a review of diagnosis and treatment. Arquivos De Neuro-Psiquiatria, 2018, 76, 41-49.	0.3	84
4	ALS5/SPG11/KIAA1840mutations cause autosomal recessive axonal Charcotâ€"Marieâ€"Tooth disease. Brain, 2016, 139, 73-85.	3.7	80
5	The Classification of Autosomal Recessive Cerebellar Ataxias: a Consensus Statement from the Society for Research on the Cerebellum and Ataxias Task Force. Cerebellum, 2019, 18, 1098-1125.	1.4	80
6	Nonmotor and extracerebellar features in Machadoâ€Joseph disease: A review. Movement Disorders, 2013, 28, 1200-1208.	2.2	79
7	Mutations in PNPLA6 are linked to photoreceptor degeneration and various forms of childhood blindness. Nature Communications, 2015, 6, 5614.	5.8	77
8	Sleep disorders in cerebellar ataxias. Arquivos De Neuro-Psiquiatria, 2011, 69, 253-257.	0.3	71
9	Translation and validation into Brazilian version of the Scale of the Assessment and Rating of Ataxia (SARA). Arquivos De Neuro-Psiquiatria, 2010, 68, 228-230.	0.3	70
10	Cerebellar Cognitive Affective Syndrome in Machado Joseph Disease: Core Clinical Features. Cerebellum, 2012, 11, 549-556.	1.4	68
11	Neurological complications in patients with SARS-CoV-2 infection: a systematic review. Arquivos De Neuro-Psiquiatria, 2020, 78, 290-300.	0.3	68
12	Neuroimaging in Hereditary Spastic Paraplegias: Current Use and Future Perspectives. Frontiers in Neurology, 2018, 9, 1117.	1.1	53
13	Sleep Disorders in Machadoâ€"Joseph Disease: Frequency, Discriminative Thresholds, Predictive Values, and Correlation with Ataxia-Related Motor and Non-Motor Features. Cerebellum, 2011, 10, 291-295.	1.4	52
14	Investigation of the RFC1 Repeat Expansion in a Canadian and a Brazilian Ataxia Cohort: Identification of Novel Conformations. Frontiers in Genetics, 2019, 10, 1219.	1.1	51
15	Noninvasive Management of Hemangioma and Vascular Malformation Using Intralesional Bleomycin Injection. Annals of Plastic Surgery, 2013, 70, 70-73.	0.5	49
16	Prolactin-sensitive neurons express estrogen receptor-Î± and depend on sex hormones for normal responsiveness to prolactin. Brain Research, 2014, 1566, 47-59.	1.1	43
17	Milestones in Friedreich ataxia: more than a century and still learning. Neurogenetics, 2015, 16, 151-160.	0.7	40
18	A diagnostic approach for neurodegeneration with brain iron accumulation: clinical features, genetics and brain imaging. Arquivos De Neuro-Psiquiatria, 2016, 74, 587-596.	0.3	39

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19	Acute cerebellar ataxia: differential diagnosis and clinical approach. Arquivos De Neuro-Psiquiatria, 2019, 77, 184-193.	0.3	35
20	One family, one gene and three phenotypes: A novel VCP (valosin-containing protein) mutation associated with myopathy with rimmed vacuoles, amyotrophic lateral sclerosis and frontotemporal dementia. Journal of the Neurological Sciences, 2016, 368, 352-358.	0.3	34
21	SPG11 mutations cause widespread white matter and basal ganglia abnormalities, but restricted cortical damage. NeuroImage: Clinical, 2018, 19, 848-857.	1.4	33
22	Cognitive Deficits in Machadoâ€“Joseph Disease Correlate with Hypoperfusion of Visual System Areas. Cerebellum, 2012, 11, 1037-1044.	1.4	30
23	Inherited manganism: The â€œcock-walkâ€•gait and typical neuroimaging features. Journal of the Neurological Sciences, 2014, 341, 150-152.	0.3	29
24	Progressive supranuclear palsy: new concepts. Arquivos De Neuro-Psiquiatria, 2010, 68, 938-946.	0.3	28
25	Clinical correlates of olfactory dysfunction in spinocerebellar ataxia type 3. Parkinsonism and Related Disorders, 2011, 17, 353-356.	1.1	28
26	Mutation in PNKP presenting initially as axonal Charcot-Marie-Tooth disease. Neurology: Genetics, 2015, 1, e30.	0.9	28
27	Substantia nigra echogenicity and imaging of striatal dopamine transporters in Parkinson's disease: A cross-sectional study. Parkinsonism and Related Disorders, 2014, 20, 477-481.	1.1	27
28	Non-motor and Extracerebellar Features in Spinocerebellar Ataxia Type 2. Cerebellum, 2017, 16, 34-39.	1.4	27
29	Huntington disease and Huntington diseaseâ€like in a case series from Brazil. Clinical Genetics, 2014, 86, 373-377.	1.0	26
30	Ginkgo biloba and Cerebral Bleeding. Neurologist, 2011, 17, 89-90.	0.4	25
31	Adult onset sporadic ataxias: a diagnostic challenge. Arquivos De Neuro-Psiquiatria, 2014, 72, 232-240.	0.3	25
32	Spinal Cord Damage in Spinocerebellar Ataxia Type 1. Cerebellum, 2017, 16, 792-796.	1.4	25
33	Severity of restless legs syndrome is inversely correlated with echogenicity of the substantia nigra in different neurodegenerative movement disorders. A preliminary observation. Journal of the Neurological Sciences, 2012, 319, 59-62.	0.3	24
34	Atypical manifestations in Brazilian patients with neuro-BehÃ§etâ€™s disease. Journal of Neurology, 2012, 259, 1159-1165.	1.8	24
35	A Novel de novo Exon 21 DNMT1 Mutation Causes Cerebellar Ataxia, Deafness, and Narcolepsy in a Brazilian Patient. Sleep, 2013, 36, 1257-1259.	0.6	24
36	MR Imaging Features of Adult-Onset Neuronal Intranuclear Inclusion Disease May Be Indistinguishable from Fragile Xâ€“Associated Tremor/Ataxia Syndrome. American Journal of Neuroradiology, 2018, 39, E100-E101.	1.2	24

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37	The progression rate of spinocerebellar ataxia type 2 changes with stage of disease. Orphanet Journal of Rare Diseases, 2018, 13, 20.	1.2	24
38	Transcranial sonography findings in spinocerebellar ataxia type 3 (Machadoâ€“Joseph disease): A cross-sectional study. Neuroscience Letters, 2011, 504, 98-101.	1.0	23
39	Cognitive and olfactory deficits in Machadoâ€“Joseph disease: A dopamine transporter study. Parkinsonism and Related Disorders, 2012, 18, 854-858.	1.1	23
40	Current concepts in the treatment of hereditary ataxias. Arquivos De Neuro-Psiquiatria, 2016, 74, 244-252.	0.3	23
41	Sleep disorders in Machadoâ€“Joseph disease: A dopamine transporter imaging study. Journal of the Neurological Sciences, 2013, 324, 90-93.	0.3	22
42	Nonneurological Involvement in Late-Onset Friedreich Ataxia (LOFA): Exploring the Phenotypes. Cerebellum, 2017, 16, 253-256.	1.4	22
43	SPG7 with parkinsonism responsive to levodopa and dopaminergic deficit. Parkinsonism and Related Disorders, 2018, 47, 88-90.	1.1	22
44	SPG11â€related parkinsonism: Clinical profile, molecular imaging and <scp>l</scp>â€dopa response. Movement Disorders, 2018, 33, 1650-1656.	2.2	22
45	NREMâ€related parasomnias in Machadoâ€“Joseph disease: clinical and polysomnographic evaluation. Journal of Sleep Research, 2016, 25, 11-15.	1.7	21
46	Clinical, ophthalmological, imaging and genetic features in Brazilian patients with ARSACS. Parkinsonism and Related Disorders, 2019, 62, 148-155.	1.1	21
47	Cerebellar ataxia associated to anti-glutamic acid decarboxylase autoantibody (anti-GAD): partial improvement with intravenous immunoglobulin therapy. Arquivos De Neuro-Psiquiatria, 2011, 69, 993-993.	0.3	21
48	Paving the Way Toward Meaningful Trials in Ataxias: An Ataxia Global Initiative Perspective. Movement Disorders, 2022, 37, 1125-1130.	2.2	21
49	Ondineâ€²s curse after brainstem infarction. Neurology India, 2009, 57, 206.	0.2	20
50	Morvan syndrome as a paraneoplastic disorder of thymoma with anti-CASPR2 antibodies. Lancet, The, 2017, 389, 1367-1368.	6.3	20
51	Gradenigoâ€™s Syndrome: Beyond the Classical Triad of Diplopia, Facial Pain and Otorrhea. Case Reports in Neurology, 2011, 3, 45-47.	0.3	19
52	Brain Structural Signature of <scp>RFC1</scp>â€Related Disorder. Movement Disorders, 2021, 36, 2634-2641.	2.2	19
53	Spontaneous downbeat nystagmus as a clue for the diagnosis of ataxia associated with anti-GAD antibodies. Journal of the Neurological Sciences, 2015, 359, 21-23.	0.3	18
54	Structural signature in SCA1: clinical correlates, determinants and natural history. Journal of Neurology, 2018, 265, 2949-2959.	1.8	18

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55	Brain Damage and Gene Expression Across Hereditary Spastic Paraplegia Subtypes. Movement Disorders, 2021, 36, 1644-1653.	2.2	18
56	Cognitive impairment in Brazilian patients with BehÃ§et's disease occurs independently of neurologic manifestation. Journal of the Neurological Sciences, 2013, 327, 1-5.	0.3	17
57	Patients with autosomal dominant spinocerebellar ataxia have more risk of falls, important balance impairment, and decreased ability to function. Arquivos De Neuro-Psiquiatria, 2013, 71, 508-511.	0.3	17
58	Complex movement disorders in fatal familial insomnia: A clinical and genetic discussion. Neurology, 2013, 81, 1098-1099.	1.5	16
59	Rapid Eye Movement Sleep Behavior Disorder in Paraneoplastic Cerebellar Degeneration: Improvement with Immunotherapy. Sleep, 2016, 39, 117-120.	0.6	16
60	Sleep disorders in Machadoâ€“Joseph disease. Current Opinion in Psychiatry, 2016, 29, 402-408.	3.1	16
61	Teaching Video Neuro Images : Gelastic cataplexy as the first neurologic manifestation of Niemann-Pick disease type C. Neurology, 2012, 79, e189.	1.5	15
62	Neurosarcoidosis: guidance for the general neurologist. Arquivos De Neuro-Psiquiatria, 2012, 70, 293-299.	0.3	15
63	Excessive fragmentary myoclonus in Machadoâ€“Joseph disease. Sleep Medicine, 2014, 15, 355-358.	0.8	15
64	Autosomal recessive spastic ataxia of Charlevoix-Saguenay (ARSACS): typical clinical and neuroimaging features in a Brazilian family. Arquivos De Neuro-Psiquiatria, 2011, 69, 288-291.	0.3	14
65	Machado-Joseph disease in Brazil: from the first descriptions to the emergence as the most common spinocerebellar ataxia. Arquivos De Neuro-Psiquiatria, 2012, 70, 630-632.	0.3	14
66	SCA1 patients may present as hereditary spastic paraplegia and must be included in spastic-ataxias group. Parkinsonism and Related Disorders, 2015, 21, 1243-1246.	1.1	14
67	Neurological phenotypes in spinocerebellar ataxia type 2: Role of mitochondrial polymorphism A10398G and other risk factors. Parkinsonism and Related Disorders, 2017, 42, 54-60.	1.1	14
68	Delirium, psychosis, and visual hallucinations induced by pregabalin. Arquivos De Neuro-Psiquiatria, 2012, 70, 960-961.	0.3	14
69	Sneddon's syndrome: case report and review of its relationship with antiphospholipid syndrome. Einstein (Sao Paulo, Brazil), 2012, 10, 230-232.	0.3	14
70	Is Neuropathy Involved with Restless Legs Syndrome in Machado-Joseph Disease?. European Neurology, 2011, 66, 200-203.	0.6	13
71	Should spinocerebellar ataxias be included in the differential diagnosis for Huntington's diseases-like syndromes?. Journal of the Neurological Sciences, 2014, 347, 356-358.	0.3	13
72	Psychosis in Machadoâ€“Joseph Disease: Clinical Correlates, Pathophysiological Discussion, and Functional Brain Imaging. Expanding the Cerebellar Cognitive Affective Syndrome. Cerebellum, 2016, 15, 483-490.	1.4	13

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73	Structural signature of classical versus lateâ€onset friedreich's ataxia by Multimodality brain M<scp>RI</scp>. Human Brain Mapping, 2017, 38, 4157-4168.	1.9	13
74	Natural history and epidemiology of the spinocerebellar ataxias: Insights from the first description to nowadays. Journal of the Neurological Sciences, 2020, 417, 117082.	0.3	13
75	Heterozygous exon 3 deletion in the Parkin gene in a patient with clinical and radiological MSA-C phenotype. Clinical Neurology and Neurosurgery, 2011, 113, 404-406.	0.6	12
76	Earlyâ€onset epilepsy as the main neurological manifestation of cerebrotendinous xanthomatosis. Epilepsy and Behavior, 2012, 24, 380-381.	0.9	12
77	The cerebellum in Parkinsonâ€™s disease and parkinsonism in cerebellar disorders. Brain, 2013, 136, e248-e248.	3.7	12
78	Cognitive dysfunction in spinocerebellar ataxia type 3: Variable topographies and patterns. Movement Disorders, 2014, 29, 156-157.	2.2	12
79	Neurophysiological Studies and Non-Motor Symptoms Prior to Ataxia in a Patient with Machadoâ€“Joseph Disease: Trying to Understand the Natural History of Brain Degeneration. Cerebellum, 2014, 13, 447-51.	1.4	12
80	Neuropsychiatric Lupus in clinical practice. Arquivos De Neuro-Psiquiatria, 2016, 74, 1021-1030.	0.3	12
81	Spinocerebellar ataxia type 10: common haplotype and disease progression rate in Peru and Brazil. European Journal of Neurology, 2017, 24, 892.	1.7	12
82	Neurological complications of solid organ transplantation. Arquivos De Neuro-Psiquiatria, 2017, 75, 736-747.	0.3	12
83	Video NeuroImages: Head titubation in anti-mGluR1 autoantibody-associated cerebellitis. Neurology, 2018, 90, 746-747.	1.5	12
84	Progressive Myoclonic Epilepsy Type 8 Due to CERS1 Deficiency: A Novel Mutation with Prominent Ataxia. Movement Disorders Clinical Practice, 2018, 5, 330-332.	0.8	12
85	Sleep Disorders in Hereditary Ataxias. Current Neurology and Neuroscience Reports, 2019, 19, 59.	2.0	12
86	Ophthalmological changes in hereditary spastic paraplegia and other genetic diseases with spastic paraplegia. Journal of the Neurological Sciences, 2020, 409, 116620.	0.3	12
87	Autosomal Recessive Cerebellar Ataxias in South America: A Multicenter Study of 1338 Patients. Movement Disorders, 2022, 37, 1773-1774.	2.2	12
88	Epilepsy and BehÃ§et's disease: Cortical and hippocampal involvement in Brazilian patients. Journal of the Neurological Sciences, 2011, 309, 1-4.	0.3	11
89	When should we test patients with familial ataxias for SCA31? A misdiagnosed condition outside Japan?. Journal of the Neurological Sciences, 2015, 355, 206-208.	0.3	11
90	Polysomnography findings in spinocerebellar ataxia type 6. Journal of Sleep Research, 2016, 25, 720-723.	1.7	11

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91	SYNE1 mutations cause autosomalâ€recessive ataxia with retained reflexes in Brazilian patients. Movement Disorders, 2016, 31, 1754-1756.	2.2	11
92	Dentatorubro-Pallidoluysian Atrophy (DRPLA) among 700 Families with Ataxia in Brazil. Cerebellum, 2017, 16, 812-816.	1.4	11
93	NESSCA Validation and Responsiveness of Several Rating Scales in Spinocerebellar Ataxia Type 2. Cerebellum, 2017, 16, 852-858.	1.4	11
94	Multimodal neuroimaging analysis in patients with SYNE1 Ataxia. Journal of the Neurological Sciences, 2018, 390, 227-230.	0.3	11
95	Is Ataxia an Underestimated Symptom of Huntington's Disease?. Frontiers in Neurology, 2020, 11, 571843.	1.1	11
96	Hepatitis C virus: A rare manifestation - Remitting relapsing central and peripheral demyelination. Neurology India, 2011, 59, 114.	0.2	10
97	Phenotype variability and early onset ataxia symptoms in spinocerebellar ataxia type 7: comparison and correlation with other spinocerebellar ataxias. Arquivos De Neuro-Psiquiatria, 2015, 73, 18-21.	0.3	10
98	Fatty acid 2-hydroxylase deficiency. Neurology, 2015, 84, 960-961.	1.5	10
99	ATXN3, ATXN7, CACNA1A, and RAI1 Genes and Mitochondrial Polymorphism A10398G Did Not Modify Age at Onset in Spinocerebellar Ataxia Type 2 Patients from South America. Cerebellum, 2015, 14, 728-730.	1.4	10
100	Cervical and ocular vestibular evoked potentials in Machadoâ€“Joseph disease: Functional involvement of otolith pathways. Journal of the Neurological Sciences, 2015, 358, 294-298.	0.3	10
101	Cross-cultural adaptation and validation of the International Cooperative Ataxia Rating Scale (ICARS) to Brazilian Portuguese. Arquivos De Neuro-Psiquiatria, 2018, 76, 674-684.	0.3	10
102	Sleep apnea in Machado-Joseph disease: a clinical and polysomnographic evaluation. Sleep Medicine, 2018, 48, 23-26.	0.8	10
103	Deafness and Vestibulopathy in Cerebellar Diseases: a Practical Approach. Cerebellum, 2019, 18, 1011-1016.	1.4	10
104	Selective Forces Related to Spinocerebellar Ataxia Type 2. Cerebellum, 2019, 18, 188-194.	1.4	10
105	Expanding the Phenotype of Dystoniaâ€Deafness Syndrome Caused by ACTB Gene Mutation. Movement Disorders Clinical Practice, 2020, 7, 86-87.	0.8	10
106	A clinical approach to hypertrophic pachymeningitis. Arquivos De Neuro-Psiquiatria, 2020, 78, 797-804.	0.3	10
107	Akathisia: An unusual movement disorder in Machadoâ€“Joseph disease. Parkinsonism and Related Disorders, 2011, 17, 712-713.	1.1	9
108	Early-onset familial Alzheimerâ€™s disease related to presenilin 1 mutation resembling autosomal dominant spinocerebellar ataxia. Journal of Neurology, 2013, 260, 1177-1179.	1.8	9

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109	Pattern of Peripheral Nerve Involvement in Machado-Joseph Disease: Neuronopathy or Distal Axonopathy? A Clinical and Neurophysiological Evaluation. European Neurology, 2013, 69, 129-133.	0.6	9
110	ACTH-induced dyskinesia in a child with West syndrome (infantile spasms). Parkinsonism and Related Disorders, 2016, 24, 145-146.	1.1	9
111	Movement Disorders in Metabolic Disorders. Current Neurology and Neuroscience Reports, 2019, 19, 7.	2.0	9
112	Unusual movement disorders in spinocerebellar ataxias. Parkinsonism and Related Disorders, 2013, 19, 834-835.	1.1	8
113	Teaching Neuro Images : â€œMini brainâ€•sign. Neurology, 2014, 82, e210-1.	1.5	8
114	Acute Foot Drop Syndrome Mimicking Peroneal Nerve Injury: An Atypical Presentation of Ischemic Stroke. Journal of Stroke and Cerebrovascular Diseases, 2014, 23, 1229-1231.	0.7	8
115	Clinical and epidemiological profiles of non-traumatic myelopathies. Arquivos De Neuro-Psiquiatria, 2016, 74, 161-165.	0.3	8
116	Pattern of Peripheral Nerve Involvement in Spinocerebellar Ataxia Type 2: a Neurophysiological Assessment. Cerebellum, 2016, 15, 767-773.	1.4	8
117	Cerebellar degeneration and progressive ataxia associated with HIV-virus infection. Parkinsonism and Related Disorders, 2018, 54, 95-98.	1.1	8
118	Beyond Typical Ataxia Telangiectasia: How to Identify the Ataxia Telangiectasiaâ€Like Disorders. Movement Disorders Clinical Practice, 2021, 8, 118-125.	0.8	8
119	Clinical and molecular evaluation of 13 Brazilian patients with Gomezâ€Ã³pezâ€HernÃ¡ndez syndrome. American Journal of Medical Genetics, Part A, 2021, 185, 1047-1058.	0.7	8
120	Spinal cord atrophy in spinocerebellar ataxia type 1. Arquivos De Neuro-Psiquiatria, 2013, 71, 977-977.	0.3	8
121	SCA2 presenting as an ataxia-parkinsonism-motor neuron disease syndrome. Arquivos De Neuro-Psiquiatria, 2011, 69, 405-406.	0.3	8
122	Clinical and molecular characterization of a large cohort of childhood onset hereditary spastic paraplegias. Scientific Reports, 2021, 11, 22248.	1.6	8
123	Substantia nigra echogenicity is correlated with nigrostriatal impairment in Machado-Joseph disease. Parkinsonism and Related Disorders, 2013, 19, 742-745.	1.1	7
124	Minimal prevalence of Huntingtonâ€™s disease in the South of Brazil and instability of the expanded CAG tract during intergenerational transmissions. Genetics and Molecular Biology, 2019, 42, 329-336.	0.6	7
125	A Proposal for Classification of Retinal Degeneration in Spinocerebellar Ataxia Type 7. Cerebellum, 2021, 20, 384-391.	1.4	7
126	<scp>POLR3Aâ€Related</scp> Disorder Presenting with <scp>Lateâ€Onset</scp> Dystonia and Spastic Paraplegia. Movement Disorders Clinical Practice, 2020, 7, 467-469.	0.8	7

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127	Corticospinal tract involvement in spinocerebellar ataxia type 3: a diffusion tensor imaging study. Neuroradiology, 2021, 63, 217-224.	1.1	7
128	Retinal Architecture in Autosomal Recessive Spastic Ataxia of Charlevoixâ€Saguenay <scp>(ARSACS)</scp>: Insights into Disease Pathogenesis and Biomarkers. Movement Disorders, 2021, 36, 2027-2035.	2.2	7
129	Hot cross bun sign resembling multiple system atrophy in a patient with Machado-Joseph disease. Arquivos De Neuro-Psiquiatria, 2013, 71, 824-824.	0.3	7
130	Variable phenotype and severity of sialidosis expressed in two siblings presenting with ataxia and macular cherry-red spots. Journal of Clinical Neuroscience, 2013, 20, 1327-1328.	0.8	6
131	Familial striatal degeneration: New mutation and neuroimaging clues. Neurology, 2015, 85, 1816-1818.	1.5	6
132	Teaching Neuro Images : Typical neuroimaging features in high-altitude cerebral edema. Neurology, 2017, 89, e176-e177.	1.5	6
133	The relationship between the First World War and neurology: 100 years of â€œShell Shockâ€• Arquivos De Neuro-Psiquiatria, 2017, 75, 317-319.	0.3	6
134	Cognitive and Psychiatric Evaluation in SYNE1 Ataxia. Cerebellum, 2019, 18, 731-737.	1.4	6
135	Beyond the Typical Syndrome: Understanding Non-motor Features in Niemann-Pick Type C Disease. Cerebellum, 2020, 19, 722-738.	1.4	6
136	Spinal cord stimulation improves motor function and gait in spastic paraplegia type 4 (SPG4): Clinical and neurophysiological evaluation. Parkinsonism and Related Disorders, 2021, 83, 1-5.	1.1	6
137	Diagnostic Yield of Whole Exome Sequencing for Adults with Ataxia: a Brazilian Perspective. Cerebellum, 2022, 21, 49-54.	1.4	6
138	Characterisation of ataxia in Sjogrenâ€™s syndrome. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 446-448.	0.9	6
139	Alpha-fetoprotein as a biomarker for recessive ataxias. Arquivos De Neuro-Psiquiatria, 2010, 68, 953-955.	0.3	6
140	Subacute cognitive impairment, hyponatremia and mesial temporal lobe lesions: a typical presentation of voltage-gated potassium channel (VGKC) antibody-associated limbic encephalitis. Arquivos De Neuro-Psiquiatria, 2011, 69, 990-991.	0.3	6
141	Translation, Cross-Cultural Adaptation, and Validation to Brazilian Portuguese of the Cerebellar Cognitive Affective/Schmahmann Syndrome Scale. Cerebellum, 2023, 22, 282-294.	1.4	6
142	Tic Disorder: An Unusual Presentation of Neurotoxoplasmosis in a Patient with AIDS. Case Reports in Neurology, 2010, 2, 145-149.	0.3	5
143	Restless legs syndrome associated with Guillain-BarrÃ© syndrome: A report of two cases. Parkinsonism and Related Disorders, 2010, 16, 418-419.	1.1	5
144	Transcranial sonography in Parkinson's disease. Einstein (Sao Paulo, Brazil), 2012, 10, 242-246.	0.3	5

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145	Urbach-Wiethe disease presenting with partial seizures, skin lesions and typical neuroimaging features. Clinical Neurology and Neurosurgery, 2014, 126, 169-170.	0.6	5
146	Cognition in SCA21 reflects developmental and adult onset cerebellar cognitive affective syndrome: Table 1. Brain, 2015, 138, e364-e364.	3.7	5
147	LMNB1 mutation causes cerebellar involvement and a genome instability defect. Journal of the Neurological Sciences, 2017, 379, 249-252.	0.3	5
148	Twenty-five years since the identification of the first SCA gene: history, clinical features and perspectives for SCA1. Arquivos De Neuro-Psiquiatria, 2018, 76, 555-562.	0.3	5
149	PRPS1 Gene Mutation Causes Complex X-Linked Adult-Onset Cerebellar Ataxia in Women. Neurology: Genetics, 2021, 7, e563.	0.9	5
150	Venlafaxine induced-myoclonus in a patient with mixed dementia. Arquivos De Neuro-Psiquiatria, 2008, 66, 894-895.	0.3	5
151	Arm Levitation as Initial Manifestation of Creutzfeldt-Jakob Disease: CaseÂReport and Review of the Literature. Tremor and Other Hyperkinetic Movements, 2018, 8, 572.	1.1	5
152	A Diagnostic Approach to Spastic ataxia Syndromes. Cerebellum, 2022, 21, 1073-1084.	1.4	5
153	Biallelic Lossâ€ofâ€Function NDUFA12 Variants Cause a Wide Phenotypic Spectrum from Leigh/Leighâ€Like Syndrome to Isolated Optic Atrophy. Movement Disorders Clinical Practice, 2022, 9, 218-228.	0.8	5
154	Characterization of Retinal Architecture in Spinocerebellar Ataxia Type 3 and Correlation with Disease Severity. Movement Disorders, 2022, 37, 758-766.	2.2	5
155	Movement disorders in spinocerebellar ataxias. Movement Disorders, 2011, 26, 2302-2302.	2.2	4
156	Teaching Neuro Images : Rosai-Dorfman disease presenting with progressive early-onset cerebellar ataxia. Neurology, 2013, 81, e27-8.	1.5	4
157	Clinical spectrum of early onset cerebellar ataxia with retained tendon reflexes: an autosomal recessive ataxia not to be missed. Arquivos De Neuro-Psiquiatria, 2013, 71, 345-348.	0.3	4
158	Neuroimaging Features in Congenital Trichomegaly: The Oliverâ€McFarlane Syndrome. Journal of Neuroimaging, 2014, 24, 418-420.	1.0	4
159	Adult onset Alexander disease presenting with progressive spastic paraplegia. Parkinsonism and Related Disorders, 2014, 20, 241-242.	1.1	4
160	Anterior horn degeneration in Machado-Joseph disease. Journal of the Neurological Sciences, 2016, 368, 290-291.	0.3	4
161	Teaching Neuro Images : The Charcot shoulder. Neurology, 2017, 89, e38-e39.	1.5	4
162	Lentiform â€œFork Signâ€•and Parkinsonism After Acute Myocardial Infarction and Cardiac Failure. Movement Disorders Clinical Practice, 2017, 4, 646-646.	0.8	4

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163	Xâ€Linked Adrenoleukodystrophy Mimicking Hereditary Spastic Paraplegia. Movement Disorders Clinical Practice, 2020, 7, 109-110.	0.8	4
164	A Proposed Clinical Classification and a Diagnostic Approach for Congenital Ataxias. Neurology: Clinical Practice, 2021, 11, e328-e336.	0.8	4
165	The Intersection Between Cerebellar Ataxia and Neuropathy: a Proposed Classification and a Diagnostic Approach. Cerebellum, 2022, 21, 497-513.	1.4	4
166	Late-Onset Psychogenic Chronic Phonic-Tics. Tremor and Other Hyperkinetic Movements, 2016, 6, 387.	1.1	4
167	Complex Movement Disorders in Ataxia with Oculomotor Apraxia Type 1: Beyond the Cerebellar Syndrome. Tremor and Other Hyperkinetic Movements, 2020, 10, 39.	1.1	4
168	Spinocerebellar Ataxia Type 5 (SCA5) Mimicking Cerebral Palsy: a Very Early Onset Autosomal Dominant Hereditary Ataxia. Cerebellum, 2023, 22, 316-318.	1.4	4
169	SjoÂÌˆgren-Larsson syndrome. Journal of Pediatrics, 2000, 136, 261.	0.9	3
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