Orlando Barsottini

List of Publications by Year
in descending order

279 papers	3,937 citations	186265 28 h-index	214800 47 g-index
286 all docs	286 docs citations	286 times ranked	4924 citing authors

#	Article	IF	Citations
1	Gluten Ataxia: an Overestimated Condition?. Cerebellum, 2022, 21, 617-619.	2.5	2
2	Diagnostic Yield of Whole Exome Sequencing for Adults with Ataxia: a Brazilian Perspective. Cerebellum, 2022, 21, 49-54.	2.5	6
3	Serum BDNF and cognitive dysfunction in SLE: findings from a cohort of 111 patients. Clinical Rheumatology, 2022, 41, 421-428.	2.2	7
4	Guidelines for Parkinsonâ€™s disease treatment: consensus from the Movement Disorders Scientific Department of the Brazilian Academy of Neurology - motor symptoms. Arquivos De Neuro-Psiquiatria, 2022, 80, 316-329.	0.8	4
5	Rehabilitation in patients with cerebellar ataxias. Arquivos De Neuro-Psiquiatria, 2022, 80, 306-315.	0.8	3
6	Characterization of Retinal Architecture in Spinocerebellar Ataxia Type 3 and Correlation with Disease Severity. Movement Disorders, 2022, 37, 758-766.	3.9	5
7	Clinical and Genetic Characterization of Brazilian Patients with Ataxia and Oculomotor Apraxia. Movement Disorders, 2022, , .	3.9	2
8	Autosomal Recessive Cerebellar Ataxias in South America: A Multicenter Study of 1338 Patients. Movement Disorders, 2022, 37, 1773-1774.	3.9	12
9	A Proposal for Classification of Retinal Degeneration in Spinocerebellar Ataxia Type 7. Cerebellum, 2021, 20, 384-391.	2.5	7
10	Clinical and Epidemiological Characterization of Neurological Consults: When a Neurological Evaluation Is Requested. Neurohospitalist, The, 2021, 11, 114-118.	0.8	1
11	Teaching NeuroImages: Trigeminal Ganglia Hypoplasia as Imaging Clue for the Diagnosis of GÃ³mez-LÃ³pez-HernÃ¡ndez Syndrome. Neurology, 2021, 96, e1593-e1594.	1.1	1
12	Beyond Typical Ataxia Telangiectasia: How to Identify the Ataxia Telangiectasiaâ€Like Disorders. Movement Disorders Clinical Practice, 2021, 8, 118-125.	1.5	8
13	Teaching Video NeuroImages: Bereitschaftspotential. Neurology, 2021, 96, e1267-e1268.	1.1	0
14	Corticospinal tract involvement in spinocerebellar ataxia type 3: a diffusion tensor imaging study. Neuroradiology, 2021, 63, 217-224.	2.2	7
15	A Proposed Clinical Classification and a Diagnostic Approach for Congenital Ataxias. Neurology: Clinical Practice, 2021, 11, e328-e336.	1.6	4
16	PRPS1 Gene Mutation Causes Complex X-Linked Adult-Onset Cerebellar Ataxia in Women. Neurology: Genetics, 2021, 7, e563.	1.9	5
17	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.	3.9	7
18	X-linked adrenoleukodystrophy presenting as progressive ataxia and pure cerebellar involvement. Arquivos De Neuro-Psiquiatria, 2021, 79, 463-464.	0.8	0

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19	Spinocerebellar ataxia type 3 presenting simultaneously with motor neuron disease and cerebellar ataxia. Arquivos De Neuro-Psiquiatria, 2021, 79, 851-852.	0.8	0
20	Brain Structural Signature of <scp>RFC1</scp>â€Related Disorder. Movement Disorders, 2021, 36, 2634-2641.	3.9	19
21	Commentary: Juvenile Dystoniaâ€Parkinsonism due to <scp>DNAJC6</scp> Mutation. Movement Disorders Clinical Practice, 2021, 8, S29-S31.	1.5	0
22	Clinical and Neuroimaging Features of Encephalocraniocutaneous Lipomatosis. Neurology, 2021, 97, 10.1212/WNL.0000000000012704.	1.1	0
23	A journey through the history of Neurogenetics. Arquivos De Neuro-Psiquiatria, 2021, 79, 929-932.	0.8	1
24	Professor Ã‚ngelo Machado: career, scientific contributions, and the iconic neuroanatomy book. Arquivos De Neuro-Psiquiatria, 2021, , .	0.8	0
25	Nystagmus may be the first neurological sign in early stages of spinocerebellar ataxia type 3. Arquivos De Neuro-Psiquiatria, 2021, 79, 891-894.	0.8	1
26	From <scp>VUS</scp> to <scp>AUS</scp>: The Connection and the Differences between Genetics and Immuneâ€Mediated Disorders. Movement Disorders, 2021, 36, 2453-2454.	3.9	1
27	Limb myorhythmia from spinal cord glioma. Practical Neurology, 2021, , practneurol-2021-003147.	1.1	1
28	DRPLA: An unusual disease or an underestimated cause of ataxia in Brazil?. Parkinsonism and Related Disorders, 2021, 92, 67-71.	2.2	2
29	Lighthouse in the open sea of spastic ataxia. Parkinsonism and Related Disorders, 2021, 91, 184-185.	2.2	0
30	Small-Expanded Allele Spinocerebellar Ataxia Type 17 Leading to Broad Movement Disorder Phenotype in a Brazilian Patient. Cerebellum, 2021, , 1.	2.5	0
31	Immunosuppressors and immunomodulators in Neurology - Part I: a guide for management of patients underimmunotherapy. Arquivos De Neuro-Psiquiatria, 2021, 79, 1012-1025.	0.8	1
32	Xâ€Linked Adrenoleukodystrophy Mimicking Hereditary Spastic Paraplegia. Movement Disorders Clinical Practice, 2020, 7, 109-110.	1.5	4
33	Expanding the Phenotype of Dystoniaâ€Deafness Syndrome Caused by ACTB Gene Mutation. Movement Disorders Clinical Practice, 2020, 7, 86-87.	1.5	10
34	Ophthalmological changes in hereditary spastic paraplegia and other genetic diseases with spastic paraplegia. Journal of the Neurological Sciences, 2020, 409, 116620.	0.6	12
35	Natural history and epidemiology of the spinocerebellar ataxias: Insights from the first description to nowadays. Journal of the Neurological Sciences, 2020, 417, 117082.	0.6	13
36	Should we investigate mitochondrial disorders in progressive adult-onset undetermined ataxias?. Cerebellum and Ataxias, 2020, 7, 13.	1.9	2

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37	Is Ataxia an Underestimated Symptom of Huntington's Disease?. Frontiers in Neurology, 2020, 11, 571843.	2.4	11
38	Postsurgical Myoclonus of the Pectoralis Major. Movement Disorders Clinical Practice, 2020, 7, 716-717.	1.5	0
39	Beyond the Typical Syndrome: Understanding Non-motor Features in Niemann-Pick Type C Disease. Cerebellum, 2020, 19, 722-738.	2.5	6
40	Teaching Video NeuroImages: Disorder of sweat, tonic pupil, and areflexia. Neurology, 2020, 95, e1292-e1293.	1.1	0
41	Huntington's disease as an unexpected cause of deafness with dystonia and chorea. Parkinsonism and Related Disorders, 2020, 76, 10-12.	2.2	0
42	The cerebellar form of acquired hepatocerebral degeneration: The hepatic ataxia. Parkinsonism and Related Disorders, 2020, 72, 72-74.	2.2	0
43	Reconstructing the History of Machado-Joseph Disease. European Neurology, 2020, 83, 99-104.	1.4	2
44	Reversible Acute Parkinsonism and Unusual Neuroimaging Findings in Systemic Lupus Erythematosus. Movement Disorders Clinical Practice, 2020, 7, 459-461.	1.5	3
45	<scp>POLR3Aâ€Related</scp> Disorder Presenting with <scp>Lateâ€Onset</scp> Dystonia and Spastic Paraplegia. Movement Disorders Clinical Practice, 2020, 7, 467-469.	1.5	7
46	Characterisation of ataxia in Sjogrenâ€™s syndrome. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 446-448.	1.9	6
47	Free carnitine and branched chain amino acids are not good biomarkers in Huntingtonâ€™s disease. Arquivos De Neuro-Psiquiatria, 2020, 78, 81-87.	0.8	2
48	Neurological complications in patients with SARS-CoV-2 infection: a systematic review. Arquivos De Neuro-Psiquiatria, 2020, 78, 290-300.	0.8	68
49	Paroxysmal Autonomic Instability with Dystonia after Severe Traumatic Brain Injury. Tremor and Other Hyperkinetic Movements, 2020, 10, 12.	2.0	3
50	Aniridia as a clue for the diagnosis of Gillespie syndrome. Arquivos De Neuro-Psiquiatria, 2020, 78, 383-383.	0.8	1
51	A clinical approach to hypertrophic pachymeningitis. Arquivos De Neuro-Psiquiatria, 2020, 78, 797-804.	0.8	10
52	Malignant cerebral edema: an unusual neurological manifestation of systemic lupus erythematosus. Arquivos De Neuro-Psiquiatria, 2020, 78, 815-815.	0.8	1
53	Complex Movement Disorders in Ataxia with Oculomotor Apraxia Type 1: Beyond the Cerebellar Syndrome. Tremor and Other Hyperkinetic Movements, 2020, 10, 39.	2.0	4
54	Sleep Disorders in Hereditary Ataxias. Current Neurology and Neuroscience Reports, 2019, 19, 59.	4.2	12

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55	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.	1.3	7
56	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.	2.5	80
57	Teaching Video NeuroImages: Hepatic myelopathy. Neurology, 2019, 93, e320-e321.	1.1	2
58	Late-onset hummingbird sign in a woman with fragile X premutation. Journal of the Neurological Sciences, 2019, 403, 75-77.	0.6	1
59	Deafness and Vestibulopathy in Cerebellar Diseases: a Practical Approach. Cerebellum, 2019, 18, 1011-1016.	2.5	10
60	Cognitive and Psychiatric Evaluation in SYNE1 Ataxia. Cerebellum, 2019, 18, 731-737.	2.5	6
61	Acute cerebellar ataxia: differential diagnosis and clinical approach. Arquivos De Neuro-Psiquiatria, 2019, 77, 184-193.	0.8	35
62	Transcranial Doppler findings in antiphospholipid syndrome. Lupus, 2019, 28, 483-491.	1.6	3
63	Movement Disorders in Metabolic Disorders. Current Neurology and Neuroscience Reports, 2019, 19, 7.	4.2	9
64	Investigation of the RFC1 Repeat Expansion in a Canadian and a Brazilian Ataxia Cohort: Identification of Novel Conformations. Frontiers in Genetics, 2019, 10, 1219.	2.3	51
65	Selective Forces Related to Spinocerebellar Ataxia Type 2. Cerebellum, 2019, 18, 188-194.	2.5	10
66	Clinical, ophthalmological, imaging and genetic features in Brazilian patients with ARSACS. Parkinsonism and Related Disorders, 2019, 62, 148-155.	2.2	21
67	Facial grimacing and clinical correlates in spinocerebellar ataxia type 3. Journal of the Neurological Sciences, 2019, 397, 138-140.	0.6	3
68	Autonomic dysfunction in hereditary spastic paraplegia type 4. European Journal of Neurology, 2019, 26, 687-693.	3.3	2
69	Septo-optic dysplasia with late-onset seizure: MRI and ophthalmological features. Arquivos De Neuro-Psiquiatria, 2019, 77, 294-295.	0.8	0
70	Rett syndrome: the Brazilian contribution to the gene discovery. Arquivos De Neuro-Psiquiatria, 2019, 77, 896-899.	0.8	1
71	Movement Disorders in Prionopathies: A Systematic Review. Tremor and Other Hyperkinetic Movements, 2019, 9, .	2.0	0
72	Video NeuroImages: Head titubation in anti-mGluR1 autoantibody-associated cerebellitis. Neurology, 2018, 90, 746-747.	1.1	12

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73	Cerebellar degeneration and progressive ataxia associated with HIV-virus infection. Parkinsonism and Related Disorders, 2018, 54, 95-98.	2.2	8
74	Tractography study in a patient with hemidystoniaâ€hemiatrophy syndrome. Movement Disorders Clinical Practice, 2018, 5, 200-202.	1.5	0
75	SPG7 with parkinsonism responsive to levodopa and dopaminergic deficit. Parkinsonism and Related Disorders, 2018, 47, 88-90.	2.2	22
76	Progressive Myoclonic Epilepsy Type 8 Due to CERS1 Deficiency: A Novel Mutation with Prominent Ataxia. Movement Disorders Clinical Practice, 2018, 5, 330-332.	1.5	12
77	Metabolic studies of a patient harbouring a novel S487L mutation in the catalytic subunit of AMPK. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2018, 1864, 1896-1903.	3.8	1
78	Perioral and tongue fasciculations in Kennedyâ€™s disease. Neurological Sciences, 2018, 39, 777-779.	1.9	3
79	Cross-cultural adaptation and validation of the International Cooperative Ataxia Rating Scale (ICARS) to Brazilian Portuguese. Arquivos De Neuro-Psiquiatria, 2018, 76, 674-684.	0.8	10
80	SPG11â€related parkinsonism: Clinical profile, molecular imaging and <scp>l</scp>â€dopa response. Movement Disorders, 2018, 33, 1650-1656.	3.9	22
81	An Unusual Fundus Finding in a Teenage Girl. JAMA Neurology, 2018, 75, 1566.	9.0	1
82	Structural signature in SCA1: clinical correlates, determinants and natural history. Journal of Neurology, 2018, 265, 2949-2959.	3.6	18
83	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.	0.8	5
84	Neurologic manifestations of antiphospholipid syndrome. Lupus, 2018, 27, 1404-1414.	1.6	49
85	SPG11 mutations cause widespread white matter and basal ganglia abnormalities, but restricted cortical damage. NeuroImage: Clinical, 2018, 19, 848-857.	2.7	33
86	Multimodal neuroimaging analysis in patients with SYNE1 Ataxia. Journal of the Neurological Sciences, 2018, 390, 227-230.	0.6	11
87	Neurological manifestations of xeroderma pigmentosum due to XPA gene mutation. Practical Neurology, 2018, 18, 489-491.	1.1	3
88	Autoimmune encephalitis: a review of diagnosis and treatment. Arquivos De Neuro-Psiquiatria, 2018, 76, 41-49.	0.8	84
89	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.	2.4	24
90	Structural signature of SCA3: From presymptomatic to late disease stages. Annals of Neurology, 2018, 84, 401-408.	5.3	90

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91	The cerebellar histiocytosis. Neurology, 2018, 91, 357-359.	1.1	3
92	Sleep apnea in Machado-Joseph disease: a clinical and polysomnographic evaluation. Sleep Medicine, 2018, 48, 23-26.	1.6	10
93	The progression rate of spinocerebellar ataxia type 2 changes with stage of disease. Orphanet Journal of Rare Diseases, 2018, 13, 20.	2.7	24
94	Neuroimaging in Hereditary Spastic Paraplegias: Current Use and Future Perspectives. Frontiers in Neurology, 2018, 9, 1117.	2.4	53
95	AB1417-HPRâ€...Predictors of cognitive dysfunction in patients with lupus. , 2018, , .		0
96	Arm Levitation as Initial Manifestation of Creutzfeldt-Jakob Disease: CaseÂReport and Review of the Literature. Tremor and Other Hyperkinetic Movements, 2018, 8, 572.	2.0	5
97	Nonneurological Involvement in Late-Onset Friedreich Ataxia (LOFA): Exploring the Phenotypes. Cerebellum, 2017, 16, 253-256.	2.5	22
98	Non-motor and Extracerebellar Features in Spinocerebellar Ataxia Type 2. Cerebellum, 2017, 16, 34-39.	2.5	27
99	Central nervous system vasculitis in adults: An update. Autoimmunity Reviews, 2017, 16, 123-131.	5.8	45
100	Teaching Neuro Images : Clinical and neuroimaging features in Gorlin-Goltz syndrome. Neurology, 2017, 88, e53-e54.	1.1	1
101	Spinal Cord Damage in Spinocerebellar Ataxia Type 1. Cerebellum, 2017, 16, 792-796.	2.5	25
102	Case 241: Hemiparkinsonism- Hemiatrophyâ€"SPECT with ^{99m}Tc TRODAT-1 and Muscle MR Imaging Abnormalities. Radiology, 2017, 283, 613-619.	7.3	0
103	Dentatorubro-Pallidoluysian Atrophy (DRPLA) among 700 Families with Ataxia in Brazil. Cerebellum, 2017, 16, 812-816.	2.5	11
104	NESSCA Validation and Responsiveness of Several Rating Scales in Spinocerebellar Ataxia Type 2. Cerebellum, 2017, 16, 852-858.	2.5	11
105	LMNB1 mutation causes cerebellar involvement and a genome instability defect. Journal of the Neurological Sciences, 2017, 379, 249-252.	0.6	5
106	Structural signature of classical versus lateâ€onset friedreich's ataxia by Multimodality brain M<scp>RI</scp>. Human Brain Mapping, 2017, 38, 4157-4168.	3.6	13
107	Spinocerebellar ataxia type 10: common haplotype and disease progression rate in Peru and Brazil. European Journal of Neurology, 2017, 24, 892.	3.3	12
108	Morvan syndrome as a paraneoplastic disorder of thymoma with anti-CASPR2 antibodies. Lancet, The, 2017, 389, 1367-1368.	13.7	20

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109	Spinocerebellar ataxia type 6 presenting with parkinsonism, pre-synaptic dopaminergic dysfunction and hyperechogenicity of the substantia nigra. Journal of the Neurological Sciences, 2017, 376, 60-62.	0.6	2
110	Case 241. Radiology, 2017, 282, 289-292.	7.3	0
111	Teaching Neuro Images : Typical neuroimaging features in high-altitude cerebral edema. Neurology, 2017, 89, e176-e177.	1.1	6
112	Teaching Neuro Images : Spinocerebellar ataxia type 3 presenting with a cock-walk gait phenotype. Neurology, 2017, 89, e192.	1.1	3
113	Teaching Neuro Images : The Charcot shoulder. Neurology, 2017, 89, e38-e39.	1.1	4
114	Antiâ€Nâ€methylâ€<scp>D</scp>â€aspartate receptor encephalitis and Epsteinâ€Barr virus: another tale on autoimmunity?. European Journal of Neurology, 2017, 24, e46-e47.	3.3	11
115	Lentiform â€œFork Signâ€•and Parkinsonism After Acute Myocardial Infarction and Cardiac Failure. Movement Disorders Clinical Practice, 2017, 4, 646-646.	1.5	4
116	Neurological phenotypes in spinocerebellar ataxia type 2: Role of mitochondrial polymorphism A10398G and other risk factors. Parkinsonism and Related Disorders, 2017, 42, 54-60.	2.2	14
117	Late-Onset Friedreichâ€™s Ataxia (LOFA) Mimicking Charcotâ€“Marieâ€“Tooth Disease Type 2: What Is Similar and What Is Different?. Cerebellum, 2017, 16, 599-601.	2.5	3
118	Lack of decussation of pyramids in Kallmann syndrome presenting with mirror movements. Journal of the Neurological Sciences, 2017, 372, 220-222.	0.6	3
119	Neurological complications of solid organ transplantation. Arquivos De Neuro-Psiquiatria, 2017, 75, 736-747.	0.8	12
120	Professor Wadiaâ€™s contributions to neurology and spinocerebellar ataxia type 2. Arquivos De Neuro-Psiquiatria, 2017, 75, 255-257.	0.8	1
121	The relationship between the First World War and neurology: 100 years of â€œShell Shockâ€• Arquivos De Neuro-Psiquiatria, 2017, 75, 317-319.	0.8	6
122	Dissecting superior cerebellar artery aneurysm: spontaneous resolution in a long-term follow-up. Arquivos De Neuro-Psiquiatria, 2017, 75, 758-759.	0.8	1
123	Central nervous system vasculitis in a patient with HIV infection: a diagnostic challenge. Arquivos De Neuro-Psiquiatria, 2017, 75, 897-898.	0.8	1
124	Frontal lobes white matter abnormalities mimicking cystic leukodystrophy in Wilsonâ€™s disease. Arquivos De Neuro-Psiquiatria, 2017, 75, 260-261.	0.8	1
125	Pontine calcification in late stage cerebellar multiple system atrophy: a marker of synucleinopathy neurodegeneration?. Arquivos De Neuro-Psiquiatria, 2017, 75, 833-834.	0.8	0
126	Clinical and epidemiological profiles of non-traumatic myelopathies. Arquivos De Neuro-Psiquiatria, 2016, 74, 161-165.	0.8	8

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127	A diagnostic approach for neurodegeneration with brain iron accumulation: clinical features, genetics and brain imaging. Arquivos De Neuro-Psiquiatria, 2016, 74, 587-596.	0.8	39
128	Neuropsychiatric Lupus in clinical practice. Arquivos De Neuro-Psiquiatria, 2016, 74, 1021-1030.	0.8	12
129	Current concepts in the treatment of hereditary ataxias. Arquivos De Neuro-Psiquiatria, 2016, 74, 244-252.	0.8	23
130	Teaching Neuro Images : Cytomegalovirus infection mimicking a brain tumor in a kidney transplant recipient. Neurology, 2016, 87, e281-e282.	1.1	2
131	SCA23 and prodynorphin: is it time for gene retraction?. Brain, 2016, 139, e42-e42.	7.6	2
132	Anterior horn degeneration in Machado-Joseph disease. Journal of the Neurological Sciences, 2016, 368, 290-291.	0.6	4
133	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.	0.6	34
134	NREMâ€related parasomnias in Machadoâ€“Joseph disease: clinical and polysomnographic evaluation. Journal of Sleep Research, 2016, 25, 11-15.	3.2	21
135	Polysomnography findings in spinocerebellar ataxia type 6. Journal of Sleep Research, 2016, 25, 720-723.	3.2	11
136	SYNE1 mutations cause autosomalâ€recessive ataxia with retained reflexes in Brazilian patients. Movement Disorders, 2016, 31, 1754-1756.	3.9	11
137	Rapid Eye Movement Sleep Behavior Disorder in Paraneoplastic Cerebellar Degeneration: Improvement with Immunotherapy. Sleep, 2016, 39, 117-120.	1.1	16
138	Sleep disorders in Machadoâ€“Joseph disease. Current Opinion in Psychiatry, 2016, 29, 402-408.	6.3	16
139	Pattern of Peripheral Nerve Involvement in Spinocerebellar Ataxia Type 2: a Neurophysiological Assessment. Cerebellum, 2016, 15, 767-773.	2.5	8
140	ACTH-induced dyskinesia in a child with West syndrome (infantile spasms). Parkinsonism and Related Disorders, 2016, 24, 145-146.	2.2	9
141	ALS5/SPG11/KIAA1840mutations cause autosomal recessive axonal Charcotâ€“Marieâ€“Tooth disease. Brain, 2016, 139, 73-85.	7.6	80
142	Psychosis in Machadoâ€“Joseph Disease: Clinical Correlates, Pathophysiological Discussion, and Functional Brain Imaging. Expanding the Cerebellar Cognitive Affective Syndrome. Cerebellum, 2016, 15, 483-490.	2.5	13
143	Gene Expression Profile in Peripheral Blood Cells of Friedreich Ataxia Patients. Cerebellum, 2016, 15, 306-313.	2.5	3
144	Brain atrophy after cortical hyperintensities in systemic lupus erythematosus. Arquivos De Neuro-Psiquiatria, 2016, 74, 82-82.	0.8	0

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145	A Brazilian Football Player Still on the Pitch After 10ÂYears of Parkinson's Disease with Severe Freezing of Gait. Movement Disorders Clinical Practice, 2015, 2, 43-44.	1.5	1
146	Non-progressive cerebellar ataxia and previous undetermined acute cerebellar injury: a mysterious clinical condition. Arquivos De Neuro-Psiquiatria, 2015, 73, 823-827.	0.8	2
147	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.	0.8	10
148	Mutation in PNKP presenting initially as axonal Charcot-Marie-Tooth disease. Neurology: Genetics, 2015, 1, e30.	1.9	28
149	Milestones in Friedreich ataxia: more than a century and still learning. Neurogenetics, 2015, 16, 151-160.	1.4	40
150	Mutations in PNPLA6 are linked to photoreceptor degeneration and various forms of childhood blindness. Nature Communications, 2015, 6, 5614.	12.8	77
151	Cognition in SCA21 reflects developmental and adult onset cerebellar cognitive affective syndrome: Table 1. Brain, 2015, 138, e364-e364.	7.6	5
152	Fatty acid 2-hydroxylase deficiency. Neurology, 2015, 84, 960-961.	1.1	10
153	SCA1 patients may present as hereditary spastic paraplegia and must be included in spastic-ataxias group. Parkinsonism and Related Disorders, 2015, 21, 1243-1246.	2.2	14
154	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.	2.5	10
155	Cervical and ocular vestibular evoked potentials in Machadoâ€"Joseph disease: Functional involvement of otolith pathways. Journal of the Neurological Sciences, 2015, 358, 294-298.	0.6	10
156	Familial striatal degeneration: New mutation and neuroimaging clues. Neurology, 2015, 85, 1816-1818.	1.1	6
157	When should we test patients with familial ataxias for SCA31? A misdiagnosed condition outside Japan?. Journal of the Neurological Sciences, 2015, 355, 206-208.	0.6	11
158	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.	0.6	18
159	Hypomelanosis of Ito presenting with adult-onset dementia and marked enlarged Virchow-Robin spaces. Arquivos De Neuro-Psiquiatria, 2015, 73, 366-368.	0.8	3
160	Association of optic atrophy and type 1 diabetes: clinical hallmarks for the diagnosis of Wolfram syndrome. Arquivos De Neuro-Psiquiatria, 2015, 73, 466-468.	0.8	3
161	PET-CT imaging in a patient with progressive supranuclear palsy. Arquivos De Neuro-Psiquiatria, 2015, 73, 364-365.	0.8	2
162	Neuroimaging features of progressive ataxia and palatal tremor. Arquivos De Neuro-Psiquiatria, 2015, 73, 633-633.	0.8	3

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163	Low signal intensity of motor cortex in SWI sequence: a radiological marker for motor neuron disease?. Arquivos De Neuro-Psiquiatria, 2015, 73, 362-363.	0.8	1
164	Adult onset sporadic ataxias: a diagnostic challenge. Arquivos De Neuro-Psiquiatria, 2014, 72, 232-240.	0.8	25
165	The relationship between Marcel Proust and Joseph Babinski: the encounter of two geniuses. Arquivos De Neuro-Psiquiatria, 2014, 72, 469-470.	0.8	2
166	Teaching Neuro Images : â€œMini brainâ€•sign. Neurology, 2014, 82, e210-1.	1.1	8
167	Neonatal <scp>SCA</scp>2 Presenting With Choreic Movements and Dystonia With Dystonic Jerks, Retinitis, Seizures, and Hypotonia. Movement Disorders Clinical Practice, 2014, 1, 252-254.	1.5	2
168	Should spinocerebellar ataxias be included in the differential diagnosis for Huntington's diseases-like syndromes?. Journal of the Neurological Sciences, 2014, 347, 356-358.	0.6	13
169	Cognitive dysfunction in spinocerebellar ataxia type 3: Variable topographies and patterns. Movement Disorders, 2014, 29, 156-157.	3.9	12
170	Huntington disease and Huntington diseaseâ€like in a case series from Brazil. Clinical Genetics, 2014, 86, 373-377.	2.0	26
171	Spinocerebellar Ataxias in Brazilâ€”Frequencies and Modulating Effects of Related Genes. Cerebellum, 2014, 13, 17-28.	2.5	93
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