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	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
2	Spinocerebellar Ataxia Type 5 (SCA5) Mimicking Cerebral Palsy: a Very Early Onset Autosomal Dominant Hereditary Ataxia. Cerebellum, 2023, 22, 316-318.	1.4	4
3	An Exploratory Survey on the Care for Ataxic Patients in the American Continents and the Caribbean. Cerebellum, 2023, 22, 708-718.	1.4	1
4	Gluten Ataxia: an Overestimated Condition?. Cerebellum, 2022, 21, 617-619.	1.4	2
5	Diagnostic Yield of Whole Exome Sequencing for Adults with Ataxia: a Brazilian Perspective. Cerebellum, 2022, 21, 49-54.	1.4	6
6	The Intersection Between Cerebellar Ataxia and Neuropathy: a Proposed Classification and a Diagnostic Approach. Cerebellum, 2022, 21, 497-513.	1.4	4
7	A Diagnostic Approach to Spastic ataxia Syndromes. Cerebellum, 2022, 21, 1073-1084.	1.4	5
8	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
9	Rehabilitation in patients with cerebellar ataxias. Arquivos De Neuro-Psiquiatria, 2022, 80, 306-315.	0.3	3
10	A Woman with Progressive Ataxia and Abnormal Eye Movements. Movement Disorders Clinical Practice, 2022, 9, 397-398.	0.8	0
11	Characterization of Retinal Architecture in Spinocerebellar Ataxia Type 3 and Correlation with Disease Severity. Movement Disorders, 2022, 37, 758-766.	2.2	5
12	Clinical and Genetic Characterization of Brazilian Patients with Ataxia and Oculomotor Apraxia. Movement Disorders, 2022, , .	2.2	2
13	Paving the Way Toward Meaningful Trials in Ataxias: An Ataxia Global Initiative Perspective. Movement Disorders, 2022, 37, 1125-1130.	2.2	21
14	Autosomal Recessive Cerebellar Ataxias in South America: A Multicenter Study of 1338 Patients. Movement Disorders, 2022, 37, 1773-1774.	2.2	12
15	Combined assessment by transcranial sonography and Sniffinâ€™ Sticks test has a similar diagnostic accuracy compared to brain SPECT for Parkinson's disease diagnosis.. Clinical Neurology and Neurosurgery, 2022, 220, 107333.	0.6	3
16	A Proposal for Classification of Retinal Degeneration in Spinocerebellar Ataxia Type 7. Cerebellum, 2021, 20, 384-391.	1.4	7
17	Family history as a clue to the diagnosis of orofacial abnormal movements in a 30-year-old man. Parkinsonism and Related Disorders, 2021, 85, 146-148.	1.1	1
18	Clinical and Epidemiological Characterization of Neurological Consults: When a Neurological Evaluation Is Requested. Neurohospitalist, The, 2021, 11, 114-118.	0.3	1

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19	Teaching NeuroImages: Trigeminal Ganglia Hypoplasia as Imaging Clue for the Diagnosis of GÃ³mez-LÃ³pez-HernÃ¡ndez Syndrome. Neurology, 2021, 96, e1593-e1594.	1.5	1
20	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
21	Beyond Typical Ataxia Telangiectasia: How to Identify the Ataxia Telangiectasiaâ€Like Disorders. Movement Disorders Clinical Practice, 2021, 8, 118-125.	0.8	8
22	Teaching Video NeuroImages: Bereitschaftspotential. Neurology, 2021, 96, e1267-e1268.	1.5	0
23	Corticospinal tract involvement in spinocerebellar ataxia type 3: a diffusion tensor imaging study. Neuroradiology, 2021, 63, 217-224.	1.1	7
24	A Proposed Clinical Classification and a Diagnostic Approach for Congenital Ataxias. Neurology: Clinical Practice, 2021, 11, e328-e336.	0.8	4
25	Brain Damage and Gene Expression Across Hereditary Spastic Paraplegia Subtypes. Movement Disorders, 2021, 36, 1644-1653.	2.2	18
26	PRPS1 Gene Mutation Causes Complex X-Linked Adult-Onset Cerebellar Ataxia in Women. Neurology: Genetics, 2021, 7, e563.	0.9	5
27	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
28	Correlation between neurological features, nutritional status, and metabolic changes in patients with Ataxia-telangiectasia. Expert Opinion on Orphan Drugs, 2021, 9, 133-137.	0.5	2
29	X-linked adrenoleukodystrophy presenting as progressive ataxia and pure cerebellar involvement. Arquivos De Neuro-Psiquiatria, 2021, 79, 463-464.	0.3	0
30	Asymptomatic retinal nerve fiber layer thickening in a patient with ataxia. Retinal Cases and Brief Reports, 2021, Publish Ahead of Print, S7-S10.	0.3	0
31	Corpus callosum impingement syndrome. Practical Neurology, 2021, 21, 546-547.	0.5	0
32	Spinocerebellar ataxia type 3 presenting simultaneously with motor neuron disease and cerebellar ataxia. Arquivos De Neuro-Psiquiatria, 2021, 79, 851-852.	0.3	0
33	Brain Structural Signature of <scp>RFC1</scp>â€Related Disorder. Movement Disorders, 2021, 36, 2634-2641.	2.2	19
34	Neuroimaging features in diethylene glycol poisoning. Arquivos De Neuro-Psiquiatria, 2021, 79, 654-655.	0.3	0
35	Clinical and Neuroimaging Features of Encephalocraniocutaneous Lipomatosis. Neurology, 2021, 97, 10.1212/WNL.0000000000012704.	1.5	0
36	A journey through the history of Neurogenetics. Arquivos De Neuro-Psiquiatria, 2021, 79, 929-932.	0.3	1

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37	Professor Ã‚ngelo Machado: career, scientific contributions, and the iconic neuroanatomy book. Arquivos De Neuro-Psiquiatria, 2021, , .	0.3	0
38	Clinical and molecular evaluation of 13 Brazilian patients with Gomezâ€ŁÃ³pezâ€ŁHernÃ¡ndez syndrome. American Journal of Medical Genetics, Part A, 2021, 185, 1047-1058.	0.7	8
39	Nystagmus may be the first neurological sign in early stages of spinocerebellar ataxia type 3. Arquivos De Neuro-Psiquiatria, 2021, 79, 891-894.	0.3	1
40	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.	2.2	1
41	Limb myorhythmia from spinal cord glioma. Practical Neurology, 2021, , practneurol-2021-003147.	0.5	1
42	DRPLA: An unusual disease or an underestimated cause of ataxia in Brazil?. Parkinsonism and Related Disorders, 2021, 92, 67-71.	1.1	2
43	Quadrupedal gait and cerebellar hypoplasia, the Uner Tan syndrome, caused by ITPR1 gene mutation. Parkinsonism and Related Disorders, 2021, 92, 33-35.	1.1	1
44	Small-Expanded Allele Spinocerebellar Ataxia Type 17 Leading to Broad Movement Disorder Phenotype in a Brazilian Patient. Cerebellum, 2021, , 1.	1.4	0
45	Clinical and molecular characterization of a large cohort of childhood onset hereditary spastic paraplegias. Scientific Reports, 2021, 11, 22248.	1.6	8
46	Immunosuppressors and immunomodulators in Neurology - Part I: a guide for management of patients underimmunotherapy. Arquivos De Neuro-Psiquiatria, 2021, 79, 1012-1025.	0.3	1
47	Xâ€Linked Adrenoleukodystrophy Mimicking Hereditary Spastic Paraplegia. Movement Disorders Clinical Practice, 2020, 7, 109-110.	0.8	4
48	Expanding the Phenotype of Dystoniaâ€Deafness Syndrome Caused by ACTB Gene Mutation. Movement Disorders Clinical Practice, 2020, 7, 86-87.	0.8	10
49	Ophthalmological changes in hereditary spastic paraplegia and other genetic diseases with spastic paraplegia. Journal of the Neurological Sciences, 2020, 409, 116620.	0.3	12
50	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
51	Should we investigate mitochondrial disorders in progressive adult-onset undetermined ataxias?. Cerebellum and Ataxias, 2020, 7, 13.	1.9	2
52	Is Ataxia an Underestimated Symptom of Huntington's Disease?. Frontiers in Neurology, 2020, 11, 571843.	1.1	11
53	Postsurgical Myoclonus of the Pectoralis Major. Movement Disorders Clinical Practice, 2020, 7, 716-717.	0.8	0
54	Beyond the Typical Syndrome: Understanding Non-motor Features in Niemann-Pick Type C Disease. Cerebellum, 2020, 19, 722-738.	1.4	6

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55	Facial myokymia in inherited peripheral nerve hyperexcitability syndrome. Practical Neurology, 2020, 20, 253-255.	0.5	3
56	Teaching Video NeuroImages: Disorder of sweat, tonic pupil, and areflexia. Neurology, 2020, 95, e1292-e1293.	1.5	0
57	Huntington's disease as an unexpected cause of deafness with dystonia and chorea. Parkinsonism and Related Disorders, 2020, 76, 10-12.	1.1	0
58	The cerebellar form of acquired hepatocerebral degeneration: The hepatic ataxia. Parkinsonism and Related Disorders, 2020, 72, 72-74.	1.1	0
59	Reconstructing the History of Machado-Joseph Disease. European Neurology, 2020, 83, 99-104.	0.6	2
60	Reversible Acute Parkinsonism and Unusual Neuroimaging Findings in Systemic Lupus Erythematosus. Movement Disorders Clinical Practice, 2020, 7, 459-461.	0.8	3
61	<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
62	Characterisation of ataxia in Sjogrenâ€™s syndrome. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 446-448.	0.9	6
63	Free carnitine and branched chain amino acids are not good biomarkers in Huntingtonâ€™s disease. Arquivos De Neuro-Psiquiatria, 2020, 78, 81-87.	0.3	2
64	Neurological complications in patients with SARS-CoV-2 infection: a systematic review. Arquivos De Neuro-Psiquiatria, 2020, 78, 290-300.	0.3	68
65	Paroxysmal Autonomic Instability with Dystonia after Severe Traumatic Brain Injury. Tremor and Other Hyperkinetic Movements, 2020, 10, 12.	1.1	3
66	Aniridia as a clue for the diagnosis of Gillespie syndrome. Arquivos De Neuro-Psiquiatria, 2020, 78, 383-383.	0.3	1
67	A clinical approach to hypertrophic pachymeningitis. Arquivos De Neuro-Psiquiatria, 2020, 78, 797-804.	0.3	10
68	Malignant cerebral edema: an unusual neurological manifestation of systemic lupus erythematosus. Arquivos De Neuro-Psiquiatria, 2020, 78, 815-815.	0.3	1
69	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
70	Sleep Disorders in Hereditary Ataxias. Current Neurology and Neuroscience Reports, 2019, 19, 59.	2.0	12
71	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
72	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

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73	Teaching Video NeuroImages: Hepatic myelopathy. Neurology, 2019, 93, e320-e321.	1.5	2
74	Late-onset hummingbird sign in a woman with fragile X premutation. Journal of the Neurological Sciences, 2019, 403, 75-77.	0.3	1
75	Deafness and Vestibulopathy in Cerebellar Diseases: a Practical Approach. Cerebellum, 2019, 18, 1011-1016.	1.4	10
76	Cognitive and Psychiatric Evaluation in SYNE1 Ataxia. Cerebellum, 2019, 18, 731-737.	1.4	6
77	Acute cerebellar ataxia: differential diagnosis and clinical approach. Arquivos De Neuro-Psiquiatria, 2019, 77, 184-193.	0.3	35
78	Movement Disorders in Metabolic Disorders. Current Neurology and Neuroscience Reports, 2019, 19, 7.	2.0	9
79	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
80	Selective Forces Related to Spinocerebellar Ataxia Type 2. Cerebellum, 2019, 18, 188-194.	1.4	10
81	Clinical, ophthalmological, imaging and genetic features in Brazilian patients with ARSACS. Parkinsonism and Related Disorders, 2019, 62, 148-155.	1.1	21
82	Facial grimacing and clinical correlates in spinocerebellar ataxia type 3. Journal of the Neurological Sciences, 2019, 397, 138-140.	0.3	3
83	Autonomic dysfunction in hereditary spastic paraplegia type 4. European Journal of Neurology, 2019, 26, 687-693.	1.7	2
84	Septo-optic dysplasia with late-onset seizure: MRI and ophthalmological features. Arquivos De Neuro-Psiquiatria, 2019, 77, 294-295.	0.3	0
85	Rett syndrome: the Brazilian contribution to the gene discovery. Arquivos De Neuro-Psiquiatria, 2019, 77, 896-899.	0.3	1
86	Movement Disorders in Prionopathies: A Systematic Review. Tremor and Other Hyperkinetic Movements, 2019, 9, .	1.1	0
87	Video NeuroImages: Head titubation in anti-mGluR1 autoantibody-associated cerebellitis. Neurology, 2018, 90, 746-747.	1.5	12
88	Cerebellar degeneration and progressive ataxia associated with HIV-virus infection. Parkinsonism and Related Disorders, 2018, 54, 95-98.	1.1	8
89	Tractography study in a patient with hemidystoniaâ€hemiatrophy syndrome. Movement Disorders Clinical Practice, 2018, 5, 200-202.	0.8	0
90	SPG7 with parkinsonism responsive to levodopa and dopaminergic deficit. Parkinsonism and Related Disorders, 2018, 47, 88-90.	1.1	22

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91	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
92	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.	1.8	1
93	Perioral and tongue fasciculations in Kennedyâ€™s disease. Neurological Sciences, 2018, 39, 777-779.	0.9	3
94	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
95	SPG11â€related parkinsonism: Clinical profile, molecular imaging and <scp>l</scp>â€dopa response. Movement Disorders, 2018, 33, 1650-1656.	2.2	22
96	An Unusual Fundus Finding in a Teenage Girl. JAMA Neurology, 2018, 75, 1566.	4.5	1
97	Structural signature in SCA1: clinical correlates, determinants and natural history. Journal of Neurology, 2018, 265, 2949-2959.	1.8	18
98	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
99	SPG11 mutations cause widespread white matter and basal ganglia abnormalities, but restricted cortical damage. NeuroImage: Clinical, 2018, 19, 848-857.	1.4	33
100	Multimodal neuroimaging analysis in patients with SYNE1 Ataxia. Journal of the Neurological Sciences, 2018, 390, 227-230.	0.3	11
101	Neurological manifestations of xeroderma pigmentosum due to XPA gene mutation. Practical Neurology, 2018, 18, 489-491.	0.5	3
102	Autoimmune encephalitis: a review of diagnosis and treatment. Arquivos De Neuro-Psiquiatria, 2018, 76, 41-49.	0.3	84
103	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
104	Structural signature of SCA3: From presymptomatic to late disease stages. Annals of Neurology, 2018, 84, 401-408.	2.8	90
105	The cerebellar histiocytosis. Neurology, 2018, 91, 357-359.	1.5	3
106	Sleep apnea in Machado-Joseph disease: a clinical and polysomnographic evaluation. Sleep Medicine, 2018, 48, 23-26.	0.8	10
107	The progression rate of spinocerebellar ataxia type 2 changes with stage of disease. Orphanet Journal of Rare Diseases, 2018, 13, 20.	1.2	24
108	Neuroimaging in Hereditary Spastic Paraplegias: Current Use and Future Perspectives. Frontiers in Neurology, 2018, 9, 1117.	1.1	53

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109	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
110	Nonneurological Involvement in Late-Onset Friedreich Ataxia (LOFA): Exploring the Phenotypes. Cerebellum, 2017, 16, 253-256.	1.4	22
111	Non-motor and Extracerebellar Features in Spinocerebellar Ataxia Type 2. Cerebellum, 2017, 16, 34-39.	1.4	27
112	Teaching Neuro Images : Clinical and neuroimaging features in Gorlin-Goltz syndrome. Neurology, 2017, 88, e53-e54.	1.5	1
113	Spinal Cord Damage in Spinocerebellar Ataxia Type 1. Cerebellum, 2017, 16, 792-796.	1.4	25
114	Case 241: Hemiparkinsonism- Hemiatrophyâ€"SPECT with ^{99m}Tc TRODAT-1 and Muscle MR Imaging Abnormalities. Radiology, 2017, 283, 613-619.	3.6	0
115	Dentatorubro-Pallidoluysian Atrophy (DRPLA) among 700 Families with Ataxia in Brazil. Cerebellum, 2017, 16, 812-816.	1.4	11
116	NESSCA Validation and Responsiveness of Several Rating Scales in Spinocerebellar Ataxia Type 2. Cerebellum, 2017, 16, 852-858.	1.4	11
117	LMNB1 mutation causes cerebellar involvement and a genome instability defect. Journal of the Neurological Sciences, 2017, 379, 249-252.	0.3	5
118	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
119	Spinocerebellar ataxia type 10: common haplotype and disease progression rate in Peru and Brazil. European Journal of Neurology, 2017, 24, 892.	1.7	12
120	Morvan syndrome as a paraneoplastic disorder of thymoma with anti-CASPR2 antibodies. Lancet, The, 2017, 389, 1367-1368.	6.3	20
121	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.	0.3	2
122	Case 241. Radiology, 2017, 282, 289-292.	3.6	0
123	Teaching Neuro Images : Typical neuroimaging features in high-altitude cerebral edema. Neurology, 2017, 89, e176-e177.	1.5	6
124	Teaching Neuro Images : Spinocerebellar ataxia type 3 presenting with a cock-walk gait phenotype. Neurology, 2017, 89, e192.	1.5	3
125	Teaching Neuro Images : The Charcot shoulder. Neurology, 2017, 89, e38-e39.	1.5	4
126	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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127	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
128	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.	1.4	3
129	Lack of decussation of pyramids in Kallmann syndrome presenting with mirror movements. Journal of the Neurological Sciences, 2017, 372, 220-222.	0.3	3
130	Neurological complications of solid organ transplantation. Arquivos De Neuro-Psiquiatria, 2017, 75, 736-747.	0.3	12
131	Professor Wadiaâ€™s contributions to neurology and spinocerebellar ataxia type 2. Arquivos De Neuro-Psiquiatria, 2017, 75, 255-257.	0.3	1
132	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
133	Dissecting superior cerebellar artery aneurysm: spontaneous resolution in a long-term follow-up. Arquivos De Neuro-Psiquiatria, 2017, 75, 758-759.	0.3	1
134	Central nervous system vasculitis in a patient with HIV infection: a diagnostic challenge. Arquivos De Neuro-Psiquiatria, 2017, 75, 897-898.	0.3	1
135	Frontal lobes white matter abnormalities mimicking cystic leukodystrophy in Wilsonâ€™s disease. Arquivos De Neuro-Psiquiatria, 2017, 75, 260-261.	0.3	1
136	Pontine calcification in late stage cerebellar multiple system atrophy: a marker of synucleinopathy neurodegeneration?. Arquivos De Neuro-Psiquiatria, 2017, 75, 833-834.	0.3	0
137	Clinical and epidemiological profiles of non-traumatic myelopathies. Arquivos De Neuro-Psiquiatria, 2016, 74, 161-165.	0.3	8
138	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
139	Neuropsychiatric Lupus in clinical practice. Arquivos De Neuro-Psiquiatria, 2016, 74, 1021-1030.	0.3	12
140	Current concepts in the treatment of hereditary ataxias. Arquivos De Neuro-Psiquiatria, 2016, 74, 244-252.	0.3	23
141	Teaching Neuro Images : Cytomegalovirus infection mimicking a brain tumor in a kidney transplant recipient. Neurology, 2016, 87, e281-e282.	1.5	2
142	Clonic Perseveration after Acute Ischemic Stroke: An Insight into the Pathophysiological Mechanisms. Journal of Stroke and Cerebrovascular Diseases, 2016, 25, e109-e110.	0.7	1
143	SCA23 and prodynorphin: is it time for gene retraction?. Brain, 2016, 139, e42-e42.	3.7	2
144	Anterior horn degeneration in Machado-Joseph disease. Journal of the Neurological Sciences, 2016, 368, 290-291.	0.3	4

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145	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
146	NREMâ€related parasomnias in Machadoâ€“Joseph disease: clinical and polysomnographic evaluation. Journal of Sleep Research, 2016, 25, 11-15.	1.7	21
147	Polysomnography findings in spinocerebellar ataxia type 6. Journal of Sleep Research, 2016, 25, 720-723.	1.7	11
148	SYNE1 mutations cause autosomalâ€recessive ataxia with retained reflexes in Brazilian patients. Movement Disorders, 2016, 31, 1754-1756.	2.2	11
149	Rapid Eye Movement Sleep Behavior Disorder in Paraneoplastic Cerebellar Degeneration: Improvement with Immunotherapy. Sleep, 2016, 39, 117-120.	0.6	16
150	Sleep disorders in Machadoâ€“Joseph disease. Current Opinion in Psychiatry, 2016, 29, 402-408.	3.1	16
151	Pattern of Peripheral Nerve Involvement in Spinocerebellar Ataxia Type 2: a Neurophysiological Assessment. Cerebellum, 2016, 15, 767-773.	1.4	8
152	ACTH-induced dyskinesia in a child with West syndrome (infantile spasms). Parkinsonism and Related Disorders, 2016, 24, 145-146.	1.1	9
153	ALS5/SPG11/KIAA1840mutations cause autosomal recessive axonal Charcotâ€“Marieâ€“Tooth disease. Brain, 2016, 139, 73-85.	3.7	80
154	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
155	Gene Expression Profile in Peripheral Blood Cells of Friedreich Ataxia Patients. Cerebellum, 2016, 15, 306-313.	1.4	3
156	Late-Onset Psychogenic Chronic Phonic-Tics. Tremor and Other Hyperkinetic Movements, 2016, 6, 387.	1.1	4
157	Are Gait Stereotypies a Marker for Neurodegeneration in Down Syndrome? A Prospective Observation. Tremor and Other Hyperkinetic Movements, 2016, 6, 403.	1.1	0
158	Hepatic Expression Of Sirtuin-1 In Response To Caloric Restriction And Leucine Supplementation In Detrained Rats. Medicine and Science in Sports and Exercise, 2015, 47, 126.	0.2	0
159	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.	0.8	1
160	Non-progressive cerebellar ataxia and previous undetermined acute cerebellar injury: a mysterious clinical condition. Arquivos De Neuro-Psiquiatria, 2015, 73, 823-827.	0.3	2
161	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
162	Mutation in PNKP presenting initially as axonal Charcot-Marie-Tooth disease. Neurology: Genetics, 2015, 1, e30.	0.9	28

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163	Milestones in Friedreich ataxia: more than a century and still learning. Neurogenetics, 2015, 16, 151-160.	0.7	40
164	Mutations in PNPLA6 are linked to photoreceptor degeneration and various forms of childhood blindness. Nature Communications, 2015, 6, 5614.	5.8	77
165	Cognition in SCA21 reflects developmental and adult onset cerebellar cognitive affective syndrome: Table 1. Brain, 2015, 138, e364-e364.	3.7	5
166	Fatty acid 2-hydroxylase deficiency. Neurology, 2015, 84, 960-961.	1.5	10
167	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
168	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
169	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
170	Familial striatal degeneration: New mutation and neuroimaging clues. Neurology, 2015, 85, 1816-1818.	1.5	6
171	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
172	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
173	Hypomelanosis of Ito presenting with adult-onset dementia and marked enlarged Virchow-Robin spaces. Arquivos De Neuro-Psiquiatria, 2015, 73, 366-368.	0.3	3
174	Association of optic atrophy and type 1 diabetes: clinical hallmarks for the diagnosis of Wolfram syndrome. Arquivos De Neuro-Psiquiatria, 2015, 73, 466-468.	0.3	3
175	PET-CT imaging in a patient with progressive supranuclear palsy. Arquivos De Neuro-Psiquiatria, 2015, 73, 364-365.	0.3	2
176	Neuroimaging features of progressive ataxia and palatal tremor. Arquivos De Neuro-Psiquiatria, 2015, 73, 633-633.	0.3	3
177	Low signal intensity of motor cortex in SWI sequence: a radiological marker for motor neuron disease?. Arquivos De Neuro-Psiquiatria, 2015, 73, 362-363.	0.3	1
178	Adult onset sporadic ataxias: a diagnostic challenge. Arquivos De Neuro-Psiquiatria, 2014, 72, 232-240.	0.3	25
179	The relationship between Marcel Proust and Joseph Babinski: the encounter of two geniuses. Arquivos De Neuro-Psiquiatria, 2014, 72, 469-470.	0.3	2
180	Teaching Neuro Images : â€œMini brainâ€•sign. Neurology, 2014, 82, e210-1.	1.5	8

#	Article	IF	Citations
181	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.	0.8	2
182	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
183	Cognitive dysfunction in spinocerebellar ataxia type 3: Variable topographies and patterns. Movement Disorders, 2014, 29, 156-157.	2.2	12
184	Huntington disease and Huntington diseaseâ€like in a case series from Brazil. Clinical Genetics, 2014, 86, 373-377.	1.0	26
185	Spinocerebellar Ataxias in Brazilâ€"Frequencies and Modulating Effects of Related Genes. Cerebellum, 2014, 13, 17-28.	1.4	93
186	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
187	Urbach-Wiethe disease presenting with partial seizures, skin lesions and typical neuroimaging features. Clinical Neurology and Neurosurgery, 2014, 126, 169-170.	0.6	5
188	Neuroimaging Features in Congenital Trichomegaly: The Oliverâ€McFarlane Syndrome. Journal of Neuroimaging, 2014, 24, 418-420.	1.0	4
189	Adult onset Alexander disease presenting with progressive spastic paraplegia. Parkinsonism and Related Disorders, 2014, 20, 241-242.	1.1	4
190	Inherited manganism: The â€œcock-walkâ€•gait and typical neuroimaging features. Journal of the Neurological Sciences, 2014, 341, 150-152.	0.3	29
191	Excessive fragmentary myoclonus in Machadoâ€"Joseph disease. Sleep Medicine, 2014, 15, 355-358.	0.8	15
192	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
193	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
194	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
195	Dengue virus infection presenting with diffuse brain hemorrhage. Arquivos De Neuro-Psiquiatria, 2014, 72, 815-815.	0.3	1
196	Machadoâ€Joseph Disease Progressing to Truncal Dystonia. Movement Disorders Clinical Practice, 2014, 1, 364-365.	0.8	1
197	Vitamin B12 deficiency mimicking neuroimaging features of motor neuron disease. Arquivos De Neuro-Psiquiatria, 2014, 72, 85-85.	0.3	3
198	Broad thumbs and broad hallux: the hallmarks for the Rubinstein-Taybi syndrome. Arquivos De Neuro-Psiquiatria, 2014, 72, 81-82.	0.3	2

#	Article	IF	Citations
199	Acute ischemic stroke resembling Saturday night palsy. Arquivos De Neuro-Psiquiatria, 2014, 72, 391-391.	0.3	2
200	Neurogenic T waves as clues for diagnosing hemorrhagic stroke. Arquivos De Neuro-Psiquiatria, 2014, 72, 472-472.	0.3	1
201	Retinitis pigmentosa in pantothenate kinase-associated neurodegeneration. Arquivos De Neuro-Psiquiatria, 2014, 72, 816-817.	0.3	1
202	Tethered cord syndrome resembling Charcot-Marie-Tooth disease in closed spinal dysraphism. Arquivos De Neuro-Psiquiatria, 2014, 72, 170-171.	0.3	1
203	Eyelid retraction is not a pathognomonic sign of Machado-Joseph disease in the context of spinocerebellar ataxias. Arquivos De Neuro-Psiquiatria, 2014, 72, 326-327.	0.3	0
204	Subacute compressive myelopathy secondary to extramedullary hematopoiesis. Arquivos De Neuro-Psiquiatria, 2014, 72, 740-740.	0.3	0
205	Substantia nigra echogenicity is correlated with nigrostriatal impairment in Machado-Joseph disease. Parkinsonism and Related Disorders, 2013, 19, 742-745.	1.1	7
206	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
207	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
208	Phelan-McDermid syndrome presenting with autistic spectrum: are we underdiagnosing chromosomal diseases in patients with autism?. Journal of Neurology, 2013, 260, 2900-2902.	1.8	1
209	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
210	Nonmotor and extracerebellar features in Machadoâ€Joseph disease: A review. Movement Disorders, 2013, 28, 1200-1208.	2.2	79
211	The Essential Can Be Invisible to the Eyes: The â€œFogging Effectâ€•Phenomenon in the Subacute Stage of Ischemic Stroke. Journal of Stroke and Cerebrovascular Diseases, 2013, 22, e628-e629.	0.7	2
212	Unusual movement disorders in spinocerebellar ataxias. Parkinsonism and Related Disorders, 2013, 19, 834-835.	1.1	8
213	Sleep disorders in Machadoâ€“Joseph disease: A dopamine transporter imaging study. Journal of the Neurological Sciences, 2013, 324, 90-93.	0.3	22
214	The cerebellum in Parkinsonâ€™s disease and parkinsonism in cerebellar disorders. Brain, 2013, 136, e248-e248.	3.7	12
215	Complex movement disorders in fatal familial insomnia: A clinical and genetic discussion. Neurology, 2013, 81, 1098-1099.	1.5	16
216	Teaching Neuro Images : Rosai-Dorfman disease presenting with progressive early-onset cerebellar ataxia. Neurology, 2013, 81, e27-8.	1.5	4

#	Article	IF	Citations
217	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
218	Noninvasive Management of Hemangioma and Vascular Malformation Using Intralesional Bleomycin Injection. Annals of Plastic Surgery, 2013, 70, 70-73.	0.5	49
219	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
220	Babinski's contributions to cerebellar symptomatology: building the basis of the neurological examination. Arquivos De Neuro-Psiquiatria, 2013, 71, 973-975.	0.3	0
221	Neurological contributions from William Osler. Arquivos De Neuro-Psiquiatria, 2013, 71, 258-260.	0.3	2
222	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
223	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
224	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
225	DykeDavidoffMasson syndrome: a combination of clinical and radiological signs not to be missed. Arquivos De Neuro-Psiquiatria, 2013, 71, 911-911.	0.3	3
226	Spinal cord atrophy in spinocerebellar ataxia type 1. Arquivos De Neuro-Psiquiatria, 2013, 71, 977-977.	0.3	8
227	Marfanoid features and X-linked mental retardation associated with craniofacial abnormalities: the Lujan-Fryns syndrome. Arquivos De Neuro-Psiquiatria, 2013, 71, 68-69.	0.3	2
228	Congenital ataxia due to cerebellar malformation presenting with unilateral hypoplasia. Arquivos De Neuro-Psiquiatria, 2013, 71, 198-198.	0.3	0
229	Diagnosis at a first glance? "Bulging eyes" as a clue for a more accurate diagnosis in spinocerebellar ataxias. Arquivos De Neuro-Psiquiatria, 2013, 71, 421-422.	0.3	1
230	Hyperintense signal in pyramidal tract neurons in postoperative brain tumor: wallerian degeneration or neoplastic dissemination?. Arquivos De Neuro-Psiquiatria, 2013, 71, 907-908.	0.3	0
231	Teaching NeuroImages: A prematurely aging patient presenting with severe leukoaraiosis and stroke. Neurology, 2012, 78, e113-e114.	1.5	0
232	Cognitive Deficits in Machadoâ€“Joseph Disease Correlate with Hypoperfusion of Visual System Areas. Cerebellum, 2012, 11, 1037-1044.	1.4	30
233	Teaching Video Neuro Images : Gelastic cataplexy as the first neurologic manifestation of Niemann-Pick disease type C. Neurology, 2012, 79, e189.	1.5	15
234	Cognitive and olfactory deficits in Machadoâ€“Joseph disease: A dopamine transporter study. Parkinsonism and Related Disorders, 2012, 18, 854-858.	1.1	23

#	Article	IF	Citations
235	Earlyâ€onset epilepsy as the main neurological manifestation of cerebrotendinous xanthomatosis. Epilepsy and Behavior, 2012, 24, 380-381.	0.9	12
236	Acute parkinsonism in Cryptococcus gattii meningoencephalitis: Extensive lesions in basal ganglia. Movement Disorders, 2012, 27, 1372-1372.	2.2	3
237	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
238	The relevance of electroencephalogram as a follow-up test in Hashimoto encephalopathy course after corticosteroids therapy. Acta Neurologica Belgica, 2012, 112, 409-411.	0.5	3
239	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
240	Transcranial sonography in Parkinson's disease. Einstein (Sao Paulo, Brazil), 2012, 10, 242-246.	0.3	5
241	Neurosarcoidosis: guidance for the general neurologist. Arquivos De Neuro-Psiquiatria, 2012, 70, 293-299.	0.3	15
242	Cerebellar Cognitive Affective Syndrome in Machado Joseph Disease: Core Clinical Features. Cerebellum, 2012, 11, 549-556.	1.4	68
243	Atypical manifestations in Brazilian patients with neuro-BehÃ§etâ€™s disease. Journal of Neurology, 2012, 259, 1159-1165.	1.8	24
244	Transcranial sonography: Brazilian experience. Arquivos De Neuro-Psiquiatria, 2012, 70, 313-314.	0.3	3
245	Delirium, psychosis, and visual hallucinations induced by pregabalin. Arquivos De Neuro-Psiquiatria, 2012, 70, 960-961.	0.3	14
246	Superior cerebellar hyperintense sign on FLAIR-weighted magnetic resonance imaging in paraneoplastic cerebellar degeneration. Arquivos De Neuro-Psiquiatria, 2012, 70, 967-967.	0.3	1
247	Sneddon's syndrome: case report and review of its relationship with antiphospholipid syndrome. Einstein (Sao Paulo, Brazil), 2012, 10, 230-232.	0.3	14
248	Esteroides Anabolizantes e o Sistema Nervoso. Revista Neurociencias, 2012, 20, 181-182.	0.0	2
249	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
250	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
251	Transcranial sonography findings in spinocerebellar ataxia type 3 (Machadoâ€“Joseph disease): A cross-sectional study. Neuroscience Letters, 2011, 504, 98-101.	1.0	23
252	Clinical correlates of olfactory dysfunction in spinocerebellar ataxia type 3. Parkinsonism and Related Disorders, 2011, 17, 353-356.	1.1	28

#	Article	IF	Citations
253	Akathisia: An unusual movement disorder in Machadoâ€“Joseph disease. Parkinsonism and Related Disorders, 2011, 17, 712-713.	1.1	9
254	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
255	Cognitive impairment in multiple system atrophy: Changing concepts. Dementia E Neuropsychologia, 2011, 5, 303-309.	0.3	3
256	Sleep disorders in cerebellar ataxias. Arquivos De Neuro-Psiquiatria, 2011, 69, 253-257.	0.3	71
257	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
258	Movement disorders in spinocerebellar ataxias. Movement Disorders, 2011, 26, 2302-2302.	2.2	4
259	Gradenigoâ€™s Syndrome: Beyond the Classical Triad of Diplopia, Facial Pain and Otorrhea. Case Reports in Neurology, 2011, 3, 45-47.	0.3	19
260	Ginkgo biloba and Cerebral Bleeding. Neurologist, 2011, 17, 89-90.	0.4	25
261	Is Neuropathy Involved with Restless Legs Syndrome in Machado-Joseph Disease?. European Neurology, 2011, 66, 200-203.	0.6	13
262	Teaching NeuroImages: Brucellosis mimicking demyelinating disease. Neurology, 2011, 76, e51-e51.	1.5	2
263	Hepatitis C virus: A rare manifestation - Remitting relapsing central and peripheral demyelination. Neurology India, 2011, 59, 114.	0.2	10
264	SCA2 presenting as an ataxia-parkinsonism-motor neuron disease syndrome. Arquivos De Neuro-Psiquiatria, 2011, 69, 405-406.	0.3	8
265	Facial grimacing as a clue for the diagnosis of GM1 type 3 gangliosidosis. Arquivos De Neuro-Psiquiatria, 2011, 69, 406-407.	0.3	3
266	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
267	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
268	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
269	Progressive supranuclear palsy: new concepts. Arquivos De Neuro-Psiquiatria, 2010, 68, 938-946.	0.3	28
270	Tic Disorder: An Unusual Presentation of Neurotoxoplasmosis in a Patient with AIDS. Case Reports in Neurology, 2010, 2, 145-149.	0.3	5

#	Article	IF	Citations
271	Restless legs syndrome associated with Guillain-BarrÃ© syndrome: A report of two cases. Parkinsonism and Related Disorders, 2010, 16, 418-419.	1.1	5
272	Alpha-fetoprotein as a biomarker for recessive ataxias. Arquivos De Neuro-Psiquiatria, 2010, 68, 953-955.	0.3	6
273	Ondineâ€²s curse after brainstem infarction. Neurology India, 2009, 57, 206.	0.2	20
274	Spontaneous carotid dissection with hypoglossal nerve palsy as residual deficit: the importance of magnetic resonance evaluation. Arquivos De Neuro-Psiquiatria, 2009, 67, 1109-1110.	0.3	3
275	Venlafaxine induced-myoclonus in a patient with mixed dementia. Arquivos De Neuro-Psiquiatria, 2008, 66, 894-895.	0.3	5
276	Multi-lacunar strokes mimicking atypical parkinsonism with an unusual neuroimaging presentation: Ã©tat criblÃ©. Arquivos De Neuro-Psiquiatria, 2008, 66, 906-907.	0.3	2
277	Les temps changent et avec eux la famille. Les mutations dâ€™accÃ¨s au droit et Ã la justice de la famille et des enfants au Portugal. Revista Critica De Ciencias Sociais, 2008, , 53-83.	0.0	3
278	TO TREAT OR NOT TO TREAT: BIAS AND BARRIERS TO HEPATITIS C CARE AMONG THE HOMELESS OF SKID ROW.. Journal of Investigative Medicine, 2007, 55, S92.	0.7	0
279	SjoÂÌˆgren-Larsson syndrome. Journal of Pediatrics, 2000, 136, 261.	0.9	3