Acary Souza Bulle Oliveira

List of Publications by Year
in descending order

81	741	759233 12	610901 24
papers	citations	h-index	g-index

82	82	82	1198
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Hereditary Spastic Paraplegia: Clinical and Genetic Hallmarks. Cerebellum, 2017, 16, 525-551.	2.5	169
2	Amyotrophic lateral sclerosis (ALS): three letters that change the people's life. For ever. Arquivos De Neuro-Psiquiatria, 2009, 67, 750-782.	0.8	61
3	Amyotrophic lateral sclerosis in Brazil: 1998 national survey. Arquivos De Neuro-Psiquiatria, 2000, 58, 607-615.	0.8	47
4	Posicionamento sobre DiagnÃ³stico e Tratamento da Amiloidose CardÃaca â€“ 2021. Arquivos Brasileiros De Cardiologia, 2021, 117, 561-598.	0.8	35
5	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
6	Motor unit number index and neurophysiological index as candidate biomarkers of presymptomatic motor neuron loss in amyotrophic lateral sclerosis. Muscle and Nerve, 2018, 58, 204-212.	2.2	33
7	New genetic causes for complex hereditary spastic paraplegia. Journal of the Neurological Sciences, 2017, 379, 283-292.	0.6	24
8	Intragenic variants in the SMN1 gene determine the clinical phenotype in 5q spinal muscular atrophy. Neurology: Genetics, 2020, 6, e505.	1.9	24
9	Clinical and genetic basis of familial amyotrophic lateral sclerosis. Arquivos De Neuro-Psiquiatria, 2015, 73, 1026-1037.	0.8	23
10	Atypical Motor Neuron Disease variants: Still a diagnostic challenge in Neurology. Revue Neurologique, 2019, 175, 221-232.	1.5	21
11	New findings in facial-onset sensory and motor neuronopathy (FOSMN) syndrome. Revue Neurologique, 2019, 175, 238-246.	1.5	18
12	<scp>GBE1</scp>â€related disorders: Adult polyglucosan body disease and its neuromuscular phenotypes. Journal of Inherited Metabolic Disease, 2021, 44, 534-543.	3.6	15
13	Acute Hepatic Porphyria: Pathophysiological Basis of Neuromuscular Manifestations. Frontiers in Neuroscience, 2021, 15, 715523.	2.8	15
14	Why averaging multiple MUNIX measures in the longitudinal assessment of patients with ALS?. Clinical Neurophysiology, 2017, 128, 2392-2396.	1.5	12
15	Functional performance and muscular strength in symptomatic female carriers of Duchenne muscular dystrophy. Arquivos De Neuro-Psiquiatria, 2020, 78, 143-148.	0.8	12
16	Ultrasound versus electromyography for the detection of fasciculation in amyotrophic lateral sclerosis: systematic review and meta-analysis. Radiologia Brasileira, 2020, 53, 116-121.	0.7	12
17	Early-onset axonal Charcot-Marie-Tooth disease due to SACS mutation. Neuromuscular Disorders, 2018, 28, 169-172.	0.6	11
18	Intermediate-length CAG repeat in ATXN2 is associated with increased risk for amyotrophic lateral sclerosis in Brazilian patients. Neurobiology of Aging, 2018, 69, 292.e15-292.e18.	3.1	10

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19	MR imaging of inherited myopathies: a review and proposal of imaging algorithms. European Radiology, 2021, 31, 8498-8512.	4.5	10
20	NFU1-Related Disorders as Key Differential Diagnosis of Cavitating Leukoencephalopathy. Journal of Pediatric Genetics, 2018, 07, 040-042.	0.7	9
21	Infantile-onset ascending spastic paraplegia phenotype associated with SPAST mutation. Journal of the Neurological Sciences, 2016, 371, 34-35.	0.6	8
22	Improving the reproducibility of motor unit number index. Muscle and Nerve, 2017, 55, 635-638.	2.2	8
23	Acute hepatic porphyrias for the neurologist: current concepts and perspectives. Arquivos De Neuro-Psiquiatria, 2021, 79, 68-80.	0.8	8
24	Restless leg syndrome exacerbated by amytriptiline in a patient with Duchenne Muscular Dystrophy. Sleep Science, 2014, 7, 178-180.	1.0	6
25	Teaching Neuro Images : Typical neuroimaging features in high-altitude cerebral edema. Neurology, 2017, 89, e176-e177.	1.1	6
26	Laparoscopic implantation of electrodes for bilateral neuromodulation of the pudendal nerves and S3 nerve roots for treating pelvic pain and voiding dysfunction. International Urogynecology Journal, 2018, 29, 1061-1064.	1.4	6
27	Whole exome sequencing identifies three recessive FIG4-mutations in an apparently dominant pedigree with Charcot-Marie-Tooth disease. Neuromuscular Disorders, 2015, 25, 359-360.	0.6	5
28	Distal myopathy due to BICD2 mutations. Clinical Neurology and Neurosurgery, 2018, 165, 47-49.	1.4	5
29	SPG76: An extremely rare hereditary spastic paraplegia with a new expanding complicated phenotype. Revue Neurologique, 2019, 175, 572-574.	1.5	5
30	Immune-mediated inflammatory polyneuropathy overlapping Charcot-Marie-Tooth 1B. Journal of Clinical Neuroscience, 2020, 75, 228-231.	1.5	5
31	Progressive spastic tetraplegia and axial hypotonia (STAHP) due to SOD1 deficiency: is it really a new entity?. Orphanet Journal of Rare Diseases, 2021, 16, 360.	2.7	5
32	Acupuncture for Treating Persistent Pain in Brazilian Para-Athletes. Medical Acupuncture, 2014, 26, 50-56.	0.6	4
33	Integrated analysis of the involvement of nitric oxide synthesis in mitochondrial proliferation, mitochondrial deficiency and apoptosis in skeletal muscle fibres. Scientific Reports, 2016, 6, 20780.	3.3	4
34	Tonic pupils: an unusual autonomic involvement in chronic inflammatory demyelinating polyneuropathy (CIDP). Neurological Sciences, 2019, 40, 1725-1727.	1.9	4
35	O'Sullivanâ€“McLeod syndrome: Unmasking a rare atypical motor neuron disease. Revue Neurologique, 2019, 175, 81-86.	1.5	4
36	Cross-cultural adaptation and validation for the Brazilian population of the instrument Amyotrophic Lateral Sclerosis-Specific Quality of Lifeâ€“Short Form (ALSSQOL-SF). Quality of Life Research, 2020, 29, 805-813.	3.1	4

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37	Estrogens: possible protection against Amyotrophic Lateral Sclerosis?. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2019, 65, 576-577.	0.7	4
38	Teaching Neuro Images : An extremely rare cause of treatable acute encephalopathy. Neurology, 2016, 87, e116.	1.1	3
39	Teaching Neuro Images : Leukodystrophy and progressive myoclonic epilepsy disclosing DRPLA. Neurology, 2016, 86, e58-9.	1.1	3
40	Collagen type VI-related myopathy. Practical Neurology, 2017, 17, 406-407.	1.1	3
41	Familial progressive bilateral facial paralysis in Finnish type hereditary amyloidosis. Practical Neurology, 2017, 17, 408-409.	1.1	3
42	Leukodystrophy with disorders of sex development due to WT1 mutations. Journal of the Neurological Sciences, 2018, 390, 94-98.	0.6	3
43	Leigh syndrome caused by mitochondrial DNA-maintenance defects revealed by whole exome sequencing. Mitochondrion, 2019, 49, 25-34.	3.4	3
44	Finger extension weakness and downbeat nystagmus motor neurone disease (FEWDON-MND). Practical Neurology, 2019, 19, 424-426.	1.1	3
45	Paraneoplastic motor neuronopathy and malignant acanthosis nigricans. Arquivos De Neuro-Psiquiatria, 2019, 77, 527-527.	0.8	3
46	Adult-onset non-5q proximal spinal muscular atrophy: a comprehensive review. Arquivos De Neuro-Psiquiatria, 2021, 79, 912-923.	0.8	3
47	Malignant Hyperthermia in Brazil: Analysis of Hotline Activity in 2009. Brazilian Journal of Anesthesiology (Elsevier), 2013, 63, 13-19.	0.4	2
48	Anthropometry of arm: nutritional risk indicator in amyotrophic lateral sclerosis. Neurology International, 2015, 7, 5952.	2.8	2
49	Teaching Neuro Images : Longitudinally extensive transverse myelitis in MELAS. Neurology, 2016, 86, e37.	1.1	2
50	Should we investigate mitochondrial disorders in progressive adult-onset undetermined ataxias?. Cerebellum and Ataxias, 2020, 7, 13.	1.9	2
51	CAG repeatsâ€¯â‰¥â€¯34 in Ataxin-1 gene are associated with amyotrophic lateral sclerosis in a Brazilian cohort. Journal of the Neurological Sciences, 2020, 414, 116842.	0.6	2
52	FasciculaÃ§Ã£o. Revista Neurociencias, 2000, 8, 31-34.	0.0	2
53	DRPLA: An unusual disease or an underestimated cause of ataxia in Brazil?. Parkinsonism and Related Disorders, 2021, 92, 67-71.	2.2	2
54	Brazilian Nursing and Psychology students' visits to patients with amyotrophic lateral sclerosis: prospective analysis. Arquivos De Neuro-Psiquiatria, 2019, 77, 782-791.	0.8	2

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55	Action of hormonal therapy in amyotrophic lateral sclerosis: a systematic review. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2020, 66, 1589-1594.	0.7	2
56	Teaching Neuro Images : Coats disease revealing facioscapulohumeral muscular dystrophy. Neurology, 2016, 87, e39.	1.1	1
57	Motor neuron disease with leukodystrophy due to CSF1R mutation. Revue Neurologique, 2020, 176, 219-221.	1.5	1
58	Rapidly progressive bulbar-onset ALS due to SS18L1 mutation. Revue Neurologique, 2020, 176, 217-219.	1.5	1
59	Cervical Spondylotic Myelopathy Secondary to Ochronotic Vertebral Arthropathy. Neurology, 2021, 96, 627-628.	1.1	1
60	Computed tomography with low-dose radiation versus standard-dose radiation for diagnosing fractures: systematic review and meta-analysis. Sao Paulo Medical Journal, 2021, 139, 388-397.	0.9	1
61	Hardy-Weinberg Equilibrium in different mitochondrial haplogroups of four genes associated with neuroprotection and neurodegeneration. Arquivos De Neuro-Psiquiatria, 2020, 78, 269-276.	0.8	1
62	As Principais Formas de AquisiÃ§Ã£o de Tecnologia Assistiva: Um Estudo Realizado em ClÃnicas de ReabilitaÃ§Ã£o na Cidade de AnÃ¡polis/GO. Fronteiras, 2020, 9, 524-544.	0.1	1
63	INCIDENCE OF MUSCULOSKELETAL INJURIES IN SOCCER REFEREES: A THREE-YEAR STUDY. Revista Brasileira De Medicina Do Esporte, 2019, 25, 258-262.	0.2	1
64	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
65	Predictive factors of the contracture test for diagnosing malignant hyperthermia in a Brazilian population sample: a retrospective observational study. Brazilian Journal of Anesthesiology (Elsevier), 2022, , .	0.4	1
66	Teaching Neuro Images : Macrocerebellum and optic atrophy in a young boy. Neurology, 2016, 86, e201.	1.1	0
67	Teaching Neuro Images : Facial grimacing and sensorineural hearing loss in a woman with cirrhosis of the liver. Neurology, 2016, 87, e239.	1.1	0
68	Teaching NeuroImages: MR neurography for the diagnosis of hypertrophic neuropathies. Neurology, 2017, 89, e201-e201.	1.1	0
69	Teaching NeuroImages: Slowly progressive hypertrophic brachial plexopathy due to SEPT9 mutation. Neurology, 2020, 95, e109-e110.	1.1	0
70	Teaching NeuroImages: Hopkins syndrome. Neurology, 2020, 94, e996-e997.	1.1	0
71	Neuromuscular choristoma: a rare cause of congenital non-progressive lower limb amyotrophy. Arquivos De Neuro-Psiquiatria, 2021, 79, 465-466.	0.8	0
72	Spinocerebellar ataxia type 3 presenting simultaneously with motor neuron disease and cerebellar ataxia. Arquivos De Neuro-Psiquiatria, 2021, 79, 851-852.	0.8	0

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73	Hyperintense signal in pyramidal tract neurons in postoperative brain tumor: wallerian degeneration or neoplastic dissemination?. Arquivos De Neuro-Psiquiatria, 2013, 71, 907-908.	0.8	0
74	Black coated tongue in integrative medicine: An alarm signal. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2016, 62, 822-824.	0.7	0
75	Pseudoxanthoma elasticum presenting as akinetic-rigid parkinsonism and dementia. Arquivos De Neuro-Psiquiatria, 2019, 77, 525-526.	0.8	0
76	Adult-onset cerebral X-linked adrenoleukodystrophy presenting as obsessive-compulsive disorder. Arquivos De Neuro-Psiquiatria, 2020, 78, 126-127.	0.8	0
77	PolifarmÃ¡cia e multimorbidade na SÃndrome Pós-poliomielite: EvidÃªncia de riscos?. Research, Society and Development, 2022, 11, e35111124951.	0.1	0
78	L-carnitine+piracetam for fatigue and muscular strength of patients with post-poliomyelitis. International Physical Medicine & Rehabilitation Journal, 2020, 5, 220-228.	0.1	0
79	Clinical and Genetic Aspects of Childhood-Onset Demyelinating Charcotâ€“Marieâ€“Tooth's Disease in Brazil. Journal of Pediatric Genetics, 2023, 12, 301-307.	0.7	0
80	Sobreviventes de Poliomielite em Ã©poca de pandemia por COVID-19 (SARS CoV-2). Research, Society and Development, 2022, 11, e54811730014.	0.1	0
81	The influence of grandparents on grandchildren. Scoping review.. Revista Neurociencias, 0, 30, 1-30.	0.0	0