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	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
2	PolifarmÃ¡cia e multimorbidade na SÃndrome PÃ³s-poliomielite: EvidÃªncia de riscos?. Research, Society and Development, 2022, 11, e35111124951.	0.1	0
3	Sobreviventes de Poliomielite em Ã©poca de pandemia por COVID-19 (SARS CoV-2). Research, Society and Development, 2022, 11, e54811730014.	0.1	0
4	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
5	<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
6	Cervical Spondylotic Myelopathy Secondary to Ochronotic Vertebral Arthropathy. Neurology, 2021, 96, 627-628.	1.1	1
7	MR imaging of inherited myopathies: a review and proposal of imaging algorithms. European Radiology, 2021, 31, 8498-8512.	4.5	10
8	Neuromuscular choristoma: a rare cause of congenital non-progressive lower limb amyotrophy. Arquivos De Neuro-Psiquiatria, 2021, 79, 465-466.	0.8	0
9	Spinocerebellar ataxia type 3 presenting simultaneously with motor neuron disease and cerebellar ataxia. Arquivos De Neuro-Psiquiatria, 2021, 79, 851-852.	0.8	0
10	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
11	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
12	Posicionamento sobre DiagnÃ³stico e Tratamento da Amiloidose CardÃaca â€“ 2021. Arquivos Brasileiros De Cardiologia, 2021, 117, 561-598.	0.8	35
13	Acute Hepatic Porphyria: Pathophysiological Basis of Neuromuscular Manifestations. Frontiers in Neuroscience, 2021, 15, 715523.	2.8	15
14	Acute hepatic porphyrias for the neurologist: current concepts and perspectives. Arquivos De Neuro-Psiquiatria, 2021, 79, 68-80.	0.8	8
15	Adult-onset non-5q proximal spinal muscular atrophy: a comprehensive review. Arquivos De Neuro-Psiquiatria, 2021, 79, 912-923.	0.8	3
16	DRPLA: An unusual disease or an underestimated cause of ataxia in Brazil?. Parkinsonism and Related Disorders, 2021, 92, 67-71.	2.2	2
17	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
18	Motor neuron disease with leukodystrophy due to CSF1R mutation. Revue Neurologique, 2020, 176, 219-221.	1.5	1

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19	Rapidly progressive bulbar-onset ALS due to SS18L1 mutation. Revue Neurologique, 2020, 176, 217-219.	1.5	1
20	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
21	Intragenic variants in the SMN1 gene determine the clinical phenotype in 5q spinal muscular atrophy. Neurology: Genetics, 2020, 6, e505.	1.9	24
22	Should we investigate mitochondrial disorders in progressive adult-onset undetermined ataxias?. Cerebellum and Ataxias, 2020, 7, 13.	1.9	2
23	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
24	Teaching NeuroImages: Slowly progressive hypertrophic brachial plexopathy due to SEPT9 mutation. Neurology, 2020, 95, e109-e110.	1.1	0
25	Teaching NeuroImages: Hopkins syndrome. Neurology, 2020, 94, e996-e997.	1.1	0
26	Immune-mediated inflammatory polyneuropathy overlapping Charcot-Marie-Tooth 1B. Journal of Clinical Neuroscience, 2020, 75, 228-231.	1.5	5
27	Functional performance and muscular strength in symptomatic female carriers of Duchenne muscular dystrophy. Arquivos De Neuro-Psiquiatria, 2020, 78, 143-148.	0.8	12
28	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
29	Adult-onset cerebral X-linked adrenoleukodystrophy presenting as obsessive-compulsive disorder. Arquivos De Neuro-Psiquiatria, 2020, 78, 126-127.	0.8	0
30	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
31	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
32	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
33	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
34	Leigh syndrome caused by mitochondrial DNA-maintenance defects revealed by whole exome sequencing. Mitochondrion, 2019, 49, 25-34.	3.4	3
35	SPG76: An extremely rare hereditary spastic paraplegia with a new expanding complicated phenotype. Revue Neurologique, 2019, 175, 572-574.	1.5	5
36	Tonic pupils: an unusual autonomic involvement in chronic inflammatory demyelinating polyneuropathy (CIDP). Neurological Sciences, 2019, 40, 1725-1727.	1.9	4

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37	Finger extension weakness and downbeat nystagmus motor neurone disease (FEWDON-MND). Practical Neurology, 2019, 19, 424-426.	1.1	3
38	Atypical Motor Neuron Disease variants: Still a diagnostic challenge in Neurology. Revue Neurologique, 2019, 175, 221-232.	1.5	21
39	O'Sullivanâ€"McLeod syndrome: Unmasking a rare atypical motor neuron disease. Revue Neurologique, 2019, 175, 81-86.	1.5	4
40	New findings in facial-onset sensory and motor neuronopathy (FOSMN) syndrome. Revue Neurologique, 2019, 175, 238-246.	1.5	18
41	Paraneoplastic motor neuronopathy and malignant acanthosis nigricans. Arquivos De Neuro-Psiquiatria, 2019, 77, 527-527.	0.8	3
42	Estrogens: possible protection against Amyotrophic Lateral Sclerosis?. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2019, 65, 576-577.	0.7	4
43	Pseudoxanthoma elasticum presenting as akinetic-rigid parkinsonism and dementia. Arquivos De Neuro-Psiquiatria, 2019, 77, 525-526.	0.8	0
44	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
45	INCIDENCE OF MUSCULOSKELETAL INJURIES IN SOCCER REFEREES: A THREE-YEAR STUDY. Revista Brasileira De Medicina Do Esporte, 2019, 25, 258-262.	0.2	1
46	Leukodystrophy with disorders of sex development due to WT1 mutations. Journal of the Neurological Sciences, 2018, 390, 94-98.	0.6	3
47	NFU1-Related Disorders as Key Differential Diagnosis of Cavitating Leukoencephalopathy. Journal of Pediatric Genetics, 2018, 07, 040-042.	0.7	9
48	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
49	Distal myopathy due to BICD2 mutations. Clinical Neurology and Neurosurgery, 2018, 165, 47-49.	1.4	5
50	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
51	Early-onset axonal Charcot-Marie-Tooth disease due to SACS mutation. Neuromuscular Disorders, 2018, 28, 169-172.	0.6	11
52	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
53	Hereditary Spastic Paraplegia: Clinical and Genetic Hallmarks. Cerebellum, 2017, 16, 525-551.	2.5	169
54	New genetic causes for complex hereditary spastic paraplegia. Journal of the Neurological Sciences, 2017, 379, 283-292.	0.6	24

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55	Collagen type VI-related myopathy. Practical Neurology, 2017, 17, 406-407.	1.1	3
56	Familial progressive bilateral facial paralysis in Finnish type hereditary amyloidosis. Practical Neurology, 2017, 17, 408-409.	1.1	3
57	Teaching Neuro Images : Typical neuroimaging features in high-altitude cerebral edema. Neurology, 2017, 89, e176-e177.	1.1	6
58	Why averaging multiple MUNIX measures in the longitudinal assessment of patients with ALS?. Clinical Neurophysiology, 2017, 128, 2392-2396.	1.5	12
59	Teaching NeuroImages: MR neurography for the diagnosis of hypertrophic neuropathies. Neurology, 2017, 89, e201-e201.	1.1	0
60	Improving the reproducibility of motor unit number index. Muscle and Nerve, 2017, 55, 635-638.	2.2	8
61	Teaching Neuro Images : Macrocerebellum and optic atrophy in a young boy. Neurology, 2016, 86, e201.	1.1	0
62	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
63	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
64	Teaching Neuro Images : An extremely rare cause of treatable acute encephalopathy. Neurology, 2016, 87, e116.	1.1	3
65	Teaching Neuro Images : Coats disease revealing facioscapulohumeral muscular dystrophy. Neurology, 2016, 87, e39.	1.1	1
66	Infantile-onset ascending spastic paraplegia phenotype associated with SPAST mutation. Journal of the Neurological Sciences, 2016, 371, 34-35.	0.6	8
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 Neuro Images : Longitudinally extensive transverse myelitis in MELAS. Neurology, 2016, 86, e37.	1.1	2
69	Teaching Neuro Images : Leukodystrophy and progressive myoclonic epilepsy disclosing DRPLA. Neurology, 2016, 86, e58-9.	1.1	3
70	Black coated tongue in integrative medicine: An alarm signal. Revista Da AssociaÃ§Ã£o MÃ©dica Brasileira, 2016, 62, 822-824.	0.7	0
71	Anthropometry of arm: nutritional risk indicator in amyotrophic lateral sclerosis. Neurology International, 2015, 7, 5952.	2.8	2
72	Clinical and genetic basis of familial amyotrophic lateral sclerosis. Arquivos De Neuro-Psiquiatria, 2015, 73, 1026-1037.	0.8	23

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73	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
74	Restless leg syndrome exacerbated by amytriptiline in a patient with Duchenne Muscular Dystrophy. Sleep Science, 2014, 7, 178-180.	1.0	6
75	Acupuncture for Treating Persistent Pain in Brazilian Para-Athletes. Medical Acupuncture, 2014, 26, 50-56.	0.6	4
76	Malignant Hyperthermia in Brazil: Analysis of Hotline Activity in 2009. Brazilian Journal of Anesthesiology (Elsevier), 2013, 63, 13-19.	0.4	2
77	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
78	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
79	Amyotrophic lateral sclerosis in Brazil: 1998 national survey. Arquivos De Neuro-Psiquiatria, 2000, 58, 607-615.	0.8	47
80	FasciculaÃ§Ã£o. Revista Neurociencias, 2000, 8, 31-34.	0.0	2
81	The influence of grandparents on grandchildren. Scoping review.. Revista Neurociencias, 0, 30, 1-30.	0.0	0