Lineu Cesar Werneck

List of Publications by Year
in descending order

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47 papers	428 citations	840119 11 h-index	794141 19 g-index
49 all docs	49 docs citations	49 times ranked	749 citing authors

#	Article	IF	Citations
1	Seventy years since the invention of the averaging technique in Neurophysiology: Tribute to George Duncan Dawson. Arquivos De Neuro-Psiquiatria, 2022, 80, 208-210.	0.3	0
2	Pain-related nociceptive evoked potential and skin wrinkle test in small fiber neuropathy. Arquivos De Neuro-Psiquiatria, 2022, , .	0.3	3
3	Horner syndrome: tribute to Professor Horner on his 190th birthday. Arquivos De Neuro-Psiquiatria, 2021, 79, 647-649.	0.3	3
4	Myasthenia gravis during pregnancy: what care should be taken?. Arquivos De Neuro-Psiquiatria, 2021, 79, 624-629.	0.3	2
5	Charcot-Marie-Tooth disease type 4C associated with myasthenia gravis: coincidental or a foreseeable association?. Neurological Sciences, 2021, , 1.	0.9	1
6	Denny-Brown and Pennybacker: 80 years after their pioneering article on electromyography, fibrillation and fasciculation. Arquivos De Neuro-Psiquiatria, 2021, 79, 81-83.	0.3	0
7	Somatosensory evoked potentials in clinical practice: a review. Arquivos De Neuro-Psiquiatria, 2021, 79, 824-831.	0.3	3
8	Peripheral polyneuropathy from electrodiagnostic tests: a 10-year etiology and neurophysiology overview. Arquivos De Neuro-Psiquiatria, 2021, , .	0.3	0
9	Congenital myasthenic syndrome in a cohort of patients with â€˜doubleâ€™ seronegative myasthenia gravis. Arquivos De Neuro-Psiquiatria, 2021, , .	0.3	3
10	Single-centre experience on genotypic and phenotypic features of southern Brazilian patients with McArdle disease. Acta Neurologica Belgica, 2020, 120, 303-311.	0.5	3
11	Characterization of the amyotrophic lateral sclerosis-linked P56S mutation of the VAPB gene in Southern Brazil. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 286-290.	1.1	6
12	Myasthenia gravis and azathioprine treatment: Adverse events related to thiopurine S-methyl-transferase (TPMT) polymorphisms. Journal of the Neurological Sciences, 2020, 412, 116734.	0.3	14
13	Localized sporotrichosis during natalizumab treatment in Multiple Sclerosis. Multiple Sclerosis and Related Disorders, 2020, 41, 102029.	0.9	1
14	Congenital myasthenic syndrome due to DOK7 mutation in a cohort of patients with â€˜unexplainedâ€™ limb-girdle muscular weakness. Journal of Clinical Neuroscience, 2020, 75, 195-198.	0.8	2
15	Celebrating the 70 years of pyridostigmine on therapy of Myasthenia Gravis: historical aspects of the preliminary trials. Arquivos De Neuro-Psiquiatria, 2020, 78, 179-181.	0.3	7
16	Somatosensory evoked potentials in Hirayama disease: A Brazilian study. , 2020, 11, 464.		2
17	Reply. Arquivos De Neuro-Psiquiatria, 2020, 78, 315-315.	0.3	0
18	Duchenne muscular dystrophy: an historical treatment review. Arquivos De Neuro-Psiquiatria, 2019, 77, 579-589.	0.3	29

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19	HLA-alleles class I and II associated with genetic susceptibility to neuromyelitis optica in Brazilian patients. Arquivos De Neuro-Psiquiatria, 2019, 77, 239-247.	0.3	11
20	Evaluation of Left-Sided Heart Chambers With Novel Echocardiographic Techniques in Men With Duchenne or Becker Muscular Dystrophy. American Journal of Cardiology, 2019, 123, 972-978.	0.7	5
21	140 Years of the LeÃ§ons sur lâ€™histologie du systÃ¨me nerveux: the pioneering description of the nodes of Ranvier. Arquivos De Neuro-Psiquiatria, 2019, 77, 749-751.	0.3	1
22	Predictors of early left ventricular systolic dysfunction in duchenne muscular dystrophy patients. Muscle and Nerve, 2018, 58, 84-89.	1.0	6
23	How to Spot Congenital Myasthenic Syndromes Resembling the Lambertâ€“Eaton Myasthenic Syndrome? A Brief Review of Clinical, Electrophysiological, and Genetics Features. NeuroMolecular Medicine, 2018, 20, 205-214.	1.8	4
24	Multiple sclerosis: disease modifying therapy and the human leukocyte antigen. Arquivos De Neuro-Psiquiatria, 2018, 76, 697-704.	0.3	9
25	Late-onset Pompe disease: what is the prevalence of limb-girdle muscular weakness presentation?. Arquivos De Neuro-Psiquiatria, 2018, 76, 247-251.	0.3	7
26	Lambert-Eaton myasthenic syndrome: the 60th anniversary of Eaton and Lambert's pioneering article. Arquivos De Neuro-Psiquiatria, 2018, 76, 124-126.	0.3	1
27	Motor neuron disease in patients with HIV infection: Report of two cases and brief review of the literature. Clinical Neurology and Neurosurgery, 2018, 171, 139-142.	0.6	9
28	Immune-mediated rippling muscle disease in a patient with treated hypothyroidism. Journal of the Neurological Sciences, 2017, 383, 53-55.	0.3	3
29	Treatment of epilepsy in patients with myasthenia gravis: Is really harder than it looks?. Journal of Clinical Neuroscience, 2017, 44, 353-356.	0.8	3
30	Is there a relationship between narcolepsy, multiple sclerosis and HLA-DQB1*06:02?. Arquivos De Neuro-Psiquiatria, 2017, 75, 345-348.	0.3	5
31	Hereditary neuropathy with liability to pressure palsies: a single-center experience in southern Brazil. Neurology International, 2016, 8, 6677.	1.3	1
32	Botulinum neurotoxin type-A when utilized in animals with trigeminal sensitization induced a antinociceptive effect. Arquivos De Neuro-Psiquiatria, 2016, 74, 462-469.	0.3	2
33	The immunogenetics of multiple sclerosis. The frequency of HLA-alleles class 1 and 2 is lower in Southern Brazil than in the European population. Arquivos De Neuro-Psiquiatria, 2016, 74, 607-616.	0.3	6
34	Management of Stable Angina with Ivabradine as Safe Alternative to Patients with Myasthenia Gravis. Case Reports in Neurological Medicine, 2016, 2016, 1-3.	0.3	1
35	Necrotizing myopathy: An uncommon initial manifestation of human immunodeficiency virus. Muscle and Nerve, 2016, 54, 334-335.	1.0	2
36	When should MELAS (Mitochondrial myopathy, Encephalopathy, Lactic Acidosis, and Stroke-like) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 6	0.3	49

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37	A importÃ¢ncia de Ã¡cido lÃ¡ctico na enxaqueca e na fibromialgia. Revista Brasileira De Reumatologia, 2015, 55, 471-476.	0.8	3
38	When should MERRF (myoclonus epilepsy associated with ragged-red fibers) be the diagnosis?. Arquivos De Neuro-Psiquiatria, 2014, 72, 803-811.	0.3	36
39	Congenital myasthenic syndrome and minicoreâ€like myopathy with <scp>DOK7</scp> mutation. Muscle and Nerve, 2013, 48, 151-152.	1.0	12
40	Muscle biopsy in Pompe disease. Arquivos De Neuro-Psiquiatria, 2013, 71, 284-289.	0.3	24
41	Congenital Myasthenic Syndrome: A Brief Review. Pediatric Neurology, 2012, 46, 141-148.	1.0	63
42	Influence of treatment in multiple sclerosis dysability: an open, retrospective, non-randomized long-term analysis. Arquivos De Neuro-Psiquiatria, 2010, 68, 511-521.	0.3	2
43	A clinical epidemiological study of 251 cases of amyotrophic lateral sclerosis in the south of Brazil. Arquivos De Neuro-Psiquiatria, 2007, 65, 189-195.	0.3	28
44	Glioma and multiple sclerosis: case report. Arquivos De Neuro-Psiquiatria, 2002, 60, 469-474.	0.3	14
45	Myasthenic crisis: report of 24 cases. Arquivos De Neuro-Psiquiatria, 2002, 60, 519-526.	0.3	16
46	Comparative analysis of PCR-deletion detection and immunohistochemistry in Brazilian Duchenne and Becker muscular dystrophy patients. American Journal of Medical Genetics Part A, 2001, 103, 115-120.	2.4	15
47	Atenolol Prophylaxis in Migraine Secondary to an Arteriovenous Malformation. Headache, 1996, 36, 625-627.	1.8	9