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List of Publications by Year
in descending order

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Version: 2024-02-01

46	395	932766	839053
papers	citations	10 h-index	18 g-index

46	46	46	693
all docs	docs citations	times ranked	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	â€œOn the Ophthalmoscopic Signs of Spinal Diseaseâ€•150 Years Later: A Tribute to Professor Sir Thomas Clifford Allbutt. Journal of Neuro-Ophthalmology, 2021, 41, 126-127.	0.4	1
8	Somatosensory evoked potentials in clinical practice: a review. Arquivos De Neuro-Psiquiatria, 2021, 79, 824-831.	0.3	3
9	Peripheral polyneuropathy from electrodiagnostic tests: a 10-year etiology and neurophysiology overview. Arquivos De Neuro-Psiquiatria, 2021, , .	0.3	0
10	Congenital myasthenic syndrome in a cohort of patients with â€˜doubleâ€™ seronegative myasthenia gravis. Arquivos De Neuro-Psiquiatria, 2021, , .	0.3	3
11	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
12	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
13	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
14	Localized sporotrichosis during natalizumab treatment in Multiple Sclerosis. Multiple Sclerosis and Related Disorders, 2020, 41, 102029.	0.9	1
15	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
16	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
17	Somatosensory evoked potentials in Hirayama disease: A Brazilian study. , 2020, 11, 464.		2
18	Reply. Arquivos De Neuro-Psiquiatria, 2020, 78, 315-315.	0.3	0

#	ARTICLE	IF	CITATIONS
19	Duchenne muscular dystrophy: an historical treatment review. Arquivos De Neuro-Psiquiatria, 2019, 77, 579-589.	0.3	29
20	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
21	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
22	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
23	Predictors of early left ventricular systolic dysfunction in duchenne muscular dystrophy patients. Muscle and Nerve, 2018, 58, 84-89.	1.0	6
24	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
25	Multiple sclerosis: disease modifying therapy and the human leukocyte antigen. Arquivos De Neuro-Psiquiatria, 2018, 76, 697-704.	0.3	9
26	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
27	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
28	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
29	Immune-mediated rippling muscle disease in a patient with treated hypothyroidism. Journal of the Neurological Sciences, 2017, 383, 53-55.	0.3	3
30	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
31	Hereditary neuropathy with liability to pressure palsies: a single-center experience in southern Brazil. Neurology International, 2016, 8, 6677.	1.3	1
32	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
33	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
34	Necrotizing myopathy: An uncommon initial manifestation of human immunodeficiency virus. Muscle and Nerve, 2016, 54, 334-335.	1.0	2
35	When should MELAS (Mitochondrial myopathy, Encephalopathy, Lactic Acidosis, and Stroke-like) Tj ETQq1 1 0.784314 rgBT /Overlock	0.3	49
36	When should MERRF (myoclonus epilepsy associated with ragged-red fibers) be the diagnosis?. Arquivos De Neuro-Psiquiatria, 2014, 72, 803-811.	0.3	36

#	ARTICLE	IF	CITATIONS
37	Identification and Functional Characterization of a Novel Mutation in theNKX2-1Gene: Comparison with the Data in the Literature. Thyroid, 2013, 23, 675-682.	2.4	29
38	Myasthenia gravis and thymus: long-term follow-up screening of thymectomized and non-thymectomized patients. Arquivos De Neuro-Psiquiatria, 2013, 71, 462-464.	0.3	3
39	Muscle biopsy in Pompe disease. Arquivos De Neuro-Psiquiatria, 2013, 71, 284-289.	0.3	24
40	Congenital Myasthenic Syndrome: A Brief Review. Pediatric Neurology, 2012, 46, 141-148.	1.0	63
41	Muscle biopsy features in critical ill patients with 2009 influenza A (H1N1) virus infection. Arquivos De Neuro-Psiquiatria, 2012, 70, 325-329.	0.3	6
42	The clinical value of laryngeal electromyography in laryngeal immobility. Journal of Clinical Neuroscience, 2011, 18, 524-527.	0.8	11
43	Idiopathic Inflammatory Myopathies in Childhood: A Brief Review of 27 Cases. Pediatric Neurology, 2011, 45, 17-22.	1.0	16
44	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
45	An electrophysiological study of the intermediate syndrome of organophosphate poisoning. Journal of Clinical Neuroscience, 2010, 17, 1217-1219.	0.8	4
46	Electrophysiological study in synaptic congenital myasthenic syndrome: end-plate acetylcholinesterase deficiency. Arquivos De Neuro-Psiquiatria, 2009, 67, 502-504.	0.3	3