Cláudia Suemi Kamoi Kay

List of Publications by Year in descending order

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Version: 2024-02-01

46 papers

395 citations

932766 10 h-index 18 g-index

46 all docs 46 docs citations

46 times ranked

693 citing authors

#	Article	IF	CITATIONS
1	Congenital Myasthenic Syndrome: A Brief Review. Pediatric Neurology, 2012, 46, 141-148.	1.0	63
2	When should MELAS (Mitochondrial myopathy, Encephalopathy, Lactic Acidosis, and Stroke-like) Tj ETQq0 0 0 rg	gBT/Qverl	ock_10 Tf 50 7
3	When should MERRF (myoclonus epilepsy associated with ragged-red fibers) be the diagnosis?. Arquivos De Neuro-Psiquiatria, 2014, 72, 803-811.	0.3	36
4	Identification and Functional Characterization of a Novel Mutation in the <i>NKX2-1 </i> Comparison with the Data in the Literature. Thyroid, 2013, 23, 675-682.	2.4	29
5	Duchenne muscular dystrophy: an historical treatment review. Arquivos De Neuro-Psiquiatria, 2019, 77, 579-589.	0.3	29
6	Muscle biopsy in Pompe disease. Arquivos De Neuro-Psiquiatria, 2013, 71, 284-289.	0.3	24
7	Idiopathic Inflammatory Myopathies in Childhood: A Brief Review of 27 Cases. Pediatric Neurology, 2011, 45, 17-22.	1.0	16
8	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
9	The clinical value of laryngeal electromyography in laryngeal immobility. Journal of Clinical Neuroscience, 2011, 18, 524-527.	0.8	11
10	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
11	Multiple sclerosis: disease modifying therapy and the human leukocyte antigen. Arquivos De Neuro-Psiquiatria, 2018, 76, 697-704.	0.3	9
12	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
13	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
14	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
15	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
16	Predictors of early left ventricular systolic dysfunction in duchenne muscular dystrophy patients. Muscle and Nerve, 2018, 58, 84-89.	1.0	6
17	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
18	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

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19	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
20	An electrophysiological study of the intermediate syndrome of organophosphate poisoning. Journal of Clinical Neuroscience, 2010, 17, 1217-1219.	0.8	4
21	How to Spot Congenital Myasthenic Syndromes Resembling the Lambert–Eaton Myasthenic Syndrome? A Brief Review of Clinical, Electrophysiological, and Genetics Features. NeuroMolecular Medicine, 2018, 20, 205-214.	1.8	4
22	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
23	Immune-mediated rippling muscle disease in a patient with treated hypothyroidism. Journal of the Neurological Sciences, 2017, 383, 53-55.	0.3	3
24	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
25	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
26	Horner syndrome: tribute to Professor Horner on his 190th birthday. Arquivos De Neuro-Psiquiatria, 2021, 79, 647-649.	0.3	3
27	Electrophysiological study in synaptic congenital myasthenic syndrome: end-plate acetylcholinesterase deficiency. Arquivos De Neuro-Psiquiatria, 2009, 67, 502-504.	0.3	3
28	Somatosensory evoked potentials in clinical practice: a review. Arquivos De Neuro-Psiquiatria, 2021, 79, 824-831.	0.3	3
29	Congenital myasthenic syndrome in a cohort of patients with †double†seronegative myasthenia gravis. Arquivos De Neuro-Psiquiatria, 2021, , .	0.3	3
30	Pain-related nociceptive evoked potential and skin wrinkle test in small fiber neuropathy. Arquivos De Neuro-Psiquiatria, 2022, , .	0.3	3
31	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
32	Necrotizing myopathy: An uncommon initial manifestation of human immunodeficiency virus. Muscle and Nerve, 2016, 54, 334-335.	1.0	2
33	Congenital myasthenic syndrome due to DOK7 mutation in a cohort of patients with †unexplained†limb-girdle muscular weakness. Journal of Clinical Neuroscience, 2020, 75, 195-198.	0.8	2
34	Myasthenia gravis during pregnancy: what care should be taken?. Arquivos De Neuro-Psiquiatria, 2021, 79, 624-629.	0.3	2
35	Somatosensory evoked potentials in Hirayama disease: A Brazilian study. , 2020, 11, 464.		2
36	Hereditary neuropathy with liability to pressure palsies: a single-center experience in southern Brazil. Neurology International, 2016, 8, 6677.	1.3	1

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37	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
38	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
39	Localized sporotrichosis during natalizumab treatment in Multiple Sclerosis. Multiple Sclerosis and Related Disorders, 2020, 41, 102029.	0.9	1
40	Charcot-Marie-Tooth disease type 4C associated with myasthenia gravis: coincidental or a foreseeable association?. Neurological Sciences, 2021, , 1.	0.9	1
41	140 Years of the Leçons sur l'histologie du système nerveux: the pioneering description of the nodes of Ranvier. Arquivos De Neuro-Psiquiatria, 2019, 77, 749-751.	0.3	1
42	"On the Ophthalmoscopic Signs of Spinal Disease―150 Years Later: A Tribute to Professor Sir Thomas Clifford Allbutt. Journal of Neuro-Ophthalmology, 2021, 41, 126-127.	0.4	1
43	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	O
44	Reply. Arquivos De Neuro-Psiquiatria, 2020, 78, 315-315.	0.3	0
45	Peripheral polyneuropathy from electrodiagnostic tests: a 10-year etiology and neurophysiology overview. Arquivos De Neuro-Psiquiatria, 2021, , .	0.3	O
46	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	O