Susana Teijeira

List of Publications by Year
in descending order

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40	874	471061	476904
papers	citations	17	29
		h-index	g-index

40	40	40	1161
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genotypic and phenotypic features of McArdle disease: insights from the Spanish national registry. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 322-328.	0.9	114
2	Molecular heterogeneity of myophosphorylase deficiency (Mcardle's disease): A genotype-phenotype correlation study. Annals of Neurology, 2001, 50, 574-581.	2.8	86
3	Oxidation of methionine residues in the prion protein by hydrogen peroxide. Archives of Biochemistry and Biophysics, 2004, 432, 188-195.	1.4	82
4	Permanent muscle weakness in M<scp>C</scp>Ardle disease. Muscle and Nerve, 2009, 40, 350-357.	1.0	59
5	Chimeric Peptide Species Contribute to Divergent Dipeptide Repeat Pathology in c9ALS/FTD and SCA36. Neuron, 2020, 107, 292-305.e6.	3.8	51
6	A novel MYH7 mutation links congenital fiber type disproportion and myosin storage myopathy. Neuromuscular Disorders, 2011, 21, 254-262.	0.3	47
7	Chorionic villi ultrastructure in the prenatal diagnosis of glycogenosis type II. Journal of Inherited Metabolic Disease, 2010, 33, 105-111.	1.7	46
8	Evaluation of heart involvement in gamma-sarcoglycanopathy (LGMD2C). A study of ten patients. Neuromuscular Disorders, 2000, 10, 560-566.	0.3	40
9	Molecular and clinical study of McArdleâ€™s disease in a cohort of 123 European patients. Identification of 20 novel mutations. Neuromuscular Disorders, 2011, 21, 817-823.	0.3	34
10	Identification of six novel mutations in the acid alpha-glucosidase gene in three Spanish patients with infantile onset glycogen storage disease type II (Pompe disease). Neuromuscular Disorders, 2002, 12, 159-166.	0.3	27
11	Fabry disease: an ultrastructural comparative study of skin in hemizygous and heterozygous patients. Acta Neuropathologica, 2006, 111, 178-185.	3.9	27
12	Myoadenylate deaminase deficiency: clinico-pathological and molecular study of a series of 27 Spanish cases. , 2009, 28, 136-142.		26
13	Adult glycogenosis II with paracrystalline mitochondrial inclusions and Hirano bodies in skeletal muscle. Neuromuscular Disorders, 1999, 9, 136-143.	0.3	24
14	Acute Necrotizing Encephalopathy of Childhood: Report of a Spanish Case. Pediatric Neurology, 2007, 37, 438-441.	1.0	23
15	Miliary brain metastases presenting as rapidly progressive dementia. Neuropathology, 2005, 25, 153-158.	0.7	21
16	Phenotypic variability in a Spanish family with a Caveolin-3 mutation. Journal of the Neurological Sciences, 2009, 276, 95-98.	0.3	20
17	Fabry disease in the Spanish population: observational study with detection of 77 patients. Orphanet Journal of Rare Diseases, 2018, 13, 52.	1.2	20
18	A Novel Missense Mutation (W797R) in the Myophosphorylase Gene in Spanish Patients With McArdle Disease. Archives of Neurology, 2000, 57, 217.	4.9	19

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19	Autosomal dominant congenital fibre type disproportion: a clinicopathological and imaging study of a large family. Brain, 2005, 128, 1716-1727.	3.7	18
20	Characterisation of Lafora-like bodies and other polyglucosan bodies in two aged dogs with neurological disease. Veterinary Journal, 2010, 183, 222-225.	0.6	16
21	A New Rare Mutation (691delCC/insAAA) in Exon 17 of the PYGM Gene Causing McArdle Disease. Archives of Neurology, 2004, 61, 1108-10.	4.9	11
22	Chloroquine-induced myopathy. Lupus, 2011, 20, 773-774.	0.8	11
23	Shrinking lung syndrome caused by lupus myopathy. QJM - Monthly Journal of the Association of Physicians, 2011, 104, 259-262.	0.2	10
24	Muscle weakness: Understanding the principles of myopathy and neuropathy in the critically ill patient and the management options. Clinical Nutrition, 2020, 39, 1331-1344.	2.3	9
25	Subsarcolemmal expression of utrophin in neuromuscular disorders: an immunohistochemical study of 80 cases. Acta Neuropathologica, 1998, 96, 481-486.	3.9	7
26	Unusual clinical findings and Complex III deficiency in a family with myotonic dystrophy. Journal of the Neurological Sciences, 2003, 208, 87-91.	0.3	7
27	Spinal muscular atrophy with respiratory distress type 1 (SMARD1) Report of a Spanish case with extended clinicopathological follow-up. , 2016, 35, 58-65.		7
28	Beta-sarcoglycanopathy (LGMD 2E) in a Spanish family. Acta Myologica, 2004, 23, 159-62.	1.5	3
29	Description of the first Spanish case of Gerstmannâ€“StrÃ¤usslerâ€“Scheinker disease with A117V variant: clinical, histopathological and biochemical characterization. Journal of Neurology, 2022, , .	1.8	3
30	Pathology and Diagnosis of Muscular Dystrophies. Advances in Experimental Medicine and Biology, 2009, 652, 1-11.	0.8	2
31	M.P.4.02 Permanent muscle weakness in McArdle disease. Neuromuscular Disorders, 2007, 17, 859-860.	0.3	1
32	Variable skeletal muscle involvement in VARS2 mitochondrial encephalomyopathy. Neuromuscular Disorders, 2016, 26, S178.	0.3	1
33	Clear cell clusters in the kidney: a rare finding that should not be misdiagnosed as renal cell carcinoma. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2021, 479, 57-67.	1.4	1
34	Histopathology of Skin in Fabry Disease. , 2010, , 275-292.		1
35	M.P.2.11 Morphological studies in an eleven week-gestation foetus with glycogenosis II (Pompe) Tj ETQq1 1 0.784314 rgBT /Overlock	0.3	0
36	G.P.13.16 Vacuolar myopathies. A morphological approach. Neuromuscular Disorders, 2007, 17, 853.	0.3	0

#	Article	IF	Citations
37	G.P.16.05. McArdle disease. Molecular study of the myophosphorylase gene (PYGM) in a series of 120 patients. Neuromuscular Disorders, 2008, 18, 828.	0.3	0
38	Molecular diagnosis of muscular dystrophies, focused on limb girdle muscular dystrophies. Expert Opinion on Medical Diagnostics, 2009, 3, 631-647.	1.6	0
39	P3.4 Spinal muscular atrophy with respiratory distress type 1 (SMARD-1). A clinico-pathological follow-up. Neuromuscular Disorders, 2011, 21, 683.	0.3	0
40	LGMD1F: A morphological study. Neuromuscular Disorders, 2015, 25, S236.	0.3	0