

# Susana Teijeira

## List of Publications by Citations

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33  
papers

680  
citations

16  
h-index

25  
g-index

40  
ext. papers

773  
ext. citations

3.7  
avg, IF

2.86  
L-index

#	Paper	IF	Citations
33	Genotypic and phenotypic features of McArdle disease: insights from the Spanish national registry. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2012</b> , 83, 322-8	5.5	90
32	Oxidation of methionine residues in the prion protein by hydrogen peroxide. <i>Archives of Biochemistry and Biophysics</i> , <b>2004</b> , 432, 188-95	4.1	76
31	Molecular heterogeneity of myophosphorylase deficiency (McArdle's disease): A genotype-phenotype correlation study. <i>Annals of Neurology</i> , <b>2001</b> , 50, 574-581	9.4	58
30	Permanent muscle weakness in McArdle disease. <i>Muscle and Nerve</i> , <b>2009</b> , 40, 350-7	3.4	45
29	Chorionic villi ultrastructure in the prenatal diagnosis of glycogenosis type II. <i>Journal of Inherited Metabolic Disease</i> , <b>2010</b> , 33 Suppl 3, S105-11	5.4	43
28	A novel MYH7 mutation links congenital fiber type disproportion and myosin storage myopathy. <i>Neuromuscular Disorders</i> , <b>2011</b> , 21, 254-62	2.9	42
27	Evaluation of heart involvement in gamma-sarcoglycanopathy (LGMD2C). A study of ten patients. <i>Neuromuscular Disorders</i> , <b>2000</b> , 10, 560-6	2.9	32
26	Molecular and clinical study of McArdle's disease in a cohort of 123 European patients. Identification of 20 novel mutations. <i>Neuromuscular Disorders</i> , <b>2011</b> , 21, 817-23	2.9	26
25	Chimeric Peptide Species Contribute to Divergent Dipeptide Repeat Pathology in c9ALS/FTD and SCA36. <i>Neuron</i> , <b>2020</b> , 107, 292-305.e6	13.9	25
24	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). <i>Neuromuscular Disorders</i> , <b>2002</b> , 12, 159-66	2.9	24
23	Fabry disease: an ultrastructural comparative study of skin in hemizygous and heterozygous patients. <i>Acta Neuropathologica</i> , <b>2006</b> , 111, 178-85	14.3	22
22	Miliary brain metastases presenting as rapidly progressive dementia. <i>Neuropathology</i> , <b>2005</b> , 25, 153-8	2	19
21	Adult glycogenosis II with paracrystalline mitochondrial inclusions and Hirano bodies in skeletal muscle. <i>Neuromuscular Disorders</i> , <b>1999</b> , 9, 136-43	2.9	19
20	A novel missense mutation (W797R) in the myophosphorylase gene in Spanish patients with McArdle disease. <i>Archives of Neurology</i> , <b>2000</b> , 57, 217-9		17
19	Myoadenylate deaminase deficiency: clinico-pathological and molecular study of a series of 27 Spanish cases <b>2009</b> , 28, 136-42		17
18	Acute necrotizing encephalopathy of childhood: report of a Spanish case. <i>Pediatric Neurology</i> , <b>2007</b> , 37, 438-41	2.9	16
17	Phenotypic variability in a Spanish family with a Caveolin-3 mutation. <i>Journal of the Neurological Sciences</i> , <b>2009</b> , 276, 95-8	3.2	15

16	Autosomal dominant congenital fibre type disproportion: a clinicopathological and imaging study of a large family. <i>Brain</i> , <b>2005</b> , 128, 1716-27	11.2	15
15	A new rare mutation (691delCC/insAAA) in exon 17 of the PYGM gene causing McArdle disease. <i>Archives of Neurology</i> , <b>2004</b> , 61, 1108-10		11
14	Fabry disease in the Spanish population: observational study with detection of 77 patients. <i>Orphanet Journal of Rare Diseases</i> , <b>2018</b> , 13, 52	4.2	10
13	Characterisation of Lafora-like bodies and other polyglucosan bodies in two aged dogs with neurological disease. <i>Veterinary Journal</i> , <b>2010</b> , 183, 222-5	2.5	10
12	Shrinking lung syndrome caused by lupus myopathy. <i>QJM - Monthly Journal of the Association of Physicians</i> , <b>2011</b> , 104, 259-62	2.7	9
11	Chloroquine-induced myopathy. <i>Lupus</i> , <b>2011</b> , 20, 773-4	2.6	8
10	Subsarcolemmal expression of utrophin in neuromuscular disorders: an immunohistochemical study of 80 cases. <i>Acta Neuropathologica</i> , <b>1998</b> , 96, 481-6	14.3	6
9	Unusual clinical findings and Complex III deficiency in a family with myotonic dystrophy. <i>Journal of the Neurological Sciences</i> , <b>2003</b> , 208, 87-91	3.2	6
8	Spinal muscular atrophy with respiratory distress type 1 (SMARD1) Report of a Spanish case with extended clinicopathological follow-up <b>2016</b> , 35, 58-65		6
7	Muscle weakness: Understanding the principles of myopathy and neuropathy in the critically ill patient and the management options. <i>Clinical Nutrition</i> , <b>2020</b> , 39, 1331-1344	5.9	3
6	Beta-sarcoglycanopathy (LGMD 2E) in a Spanish family. <i>Acta Myologica</i> , <b>2004</b> , 23, 159-62	1.6	3
5	Pathology and diagnosis of muscular dystrophies. <i>Advances in Experimental Medicine and Biology</i> , <b>2009</b> , 652, 1-11	3.6	2
4	Histopathology of Skin in Fabry Disease <b>2010</b> , 275-292		1
3	Clear cell clusters in the kidney: a rare finding that should not be misdiagnosed as renal cell carcinoma. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , <b>2021</b> , 479, 57-67	5.1	1
2	Description of the first Spanish case of Gerstmann-Strüssler-Scheinker disease with A117V variant: clinical, histopathological and biochemical characterization.. <i>Journal of Neurology</i> , <b>2022</b> ,	5.5	1
1	Molecular diagnosis of muscular dystrophies, focused on limb girdle muscular dystrophies. <i>Expert Opinion on Medical Diagnostics</i> , <b>2009</b> , 3, 631-47		