Susana Teijeira

List of Publications by Year in Descending Order

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	Description of the first Spanish case of Gerstmann-Str□ssler-Scheinker disease with A117V variant: clinical, histopathological and biochemical characterization.. *Journal of Neurology*, **2022**,	5.5	1
32	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	5.1	1
31	Chimeric Peptide Species Contribute to Divergent Dipeptide Repeat Pathology in c9ALS/FTD and SCA36. *Neuron*, **2020**, 107, 292-305.e6	13.9	25
30	Muscle weakness: Understanding the principles of myopathy and neuropathy in the critically ill patient and the management options. *Clinical Nutrition*, **2020**, 39, 1331-1344	5.9	3
29	Fabry disease in the Spanish population: observational study with detection of 77 patients. *Orphanet Journal of Rare Diseases*, **2018**, 13, 52	4.2	10
28	Spinal muscular atrophy with respiratory distress type 1 (SMARD1) Report of a Spanish case with extended clinicopathological follow-up **2016**, 35, 58-65		6
27	Genotypic and phenotypic features of McArdle disease: insights from the Spanish national registry. *Journal of Neurology, Neurosurgery and Psychiatry*, **2012**, 83, 322-8	5.5	90
26	A novel MYH7 mutation links congenital fiber type disproportion and myosin storage myopathy. *Neuromuscular Disorders*, **2011**, 21, 254-62	2.9	42
25	Molecular and clinical study of McArdle¥s disease in a cohort of 123 European patients. Identification of 20 novel mutations. *Neuromuscular Disorders*, **2011**, 21, 817-23	2.9	26
24	Chloroquine-induced myopathy. *Lupus*, **2011**, 20, 773-4	2.6	8
23	Shrinking lung syndrome caused by lupus myopathy. *QJM - Monthly Journal of the Association of Physicians*, **2011**, 104, 259-62	2.7	9
22	Chorionic villi ultrastructure in the prenatal diagnosis of glycogenosis type II. *Journal of Inherited Metabolic Disease*, **2010**, 33 Suppl 3, S105-11	5.4	43
21	Characterisation of Lafora-like bodies and other polyglucosan bodies in two aged dogs with neurological disease. *Veterinary Journal*, **2010**, 183, 222-5	2.5	10
20	Histopathology of Skin in Fabry Disease **2010**, 275-292		1
19	Molecular diagnosis of muscular dystrophies, focused on limb girdle muscular dystrophies. *Expert Opinion on Medical Diagnostics*, **2009**, 3, 631-47		
18	Permanent muscle weakness in McArdle disease. *Muscle and Nerve*, **2009**, 40, 350-7	3.4	45
17	Phenotypic variability in a Spanish family with a Caveolin-3 mutation. *Journal of the Neurological Sciences*, **2009**, 276, 95-8	3.2	15

16	Pathology and diagnosis of muscular dystrophies. *Advances in Experimental Medicine and Biology*, **2009**, 652, 1-11	3.6	2
15	Myoadenylate deaminase deficiency: clinico-pathological and molecular study of a series of 27 Spanish cases **2009**, 28, 136-42		17
14	Acute necrotizing encephalopathy of childhood: report of a Spanish case. *Pediatric Neurology*, **2007**, 37, 438-41	2.9	16
13	Fabry disease: an ultrastructural comparative study of skin in hemizygous and heterozygous patients. *Acta Neuropathologica*, **2006**, 111, 178-85	14.3	22
12	Autosomal dominant congenital fibre type disproportion: a clinicopathological and imaging study of a large family. *Brain*, **2005**, 128, 1716-27	11.2	15
11	Miliary brain metastases presenting as rapidly progressive dementia. *Neuropathology*, **2005**, 25, 153-8	2	19
10	A new rare mutation (691delCC/insAAA) in exon 17 of the PYGM gene causing McArdle disease. *Archives of Neurology*, **2004**, 61, 1108-10		11
9	Oxidation of methionine residues in the prion protein by hydrogen peroxide. *Archives of Biochemistry and Biophysics*, **2004**, 432, 188-95	4.1	76
8	Beta-sarcoglycanopathy (LGMD 2E) in a Spanish family. *Acta Myologica*, **2004**, 23, 159-62	1.6	3
7	Unusual clinical findings and Complex III deficiency in a family with myotonic dystrophy. *Journal of the Neurological Sciences*, **2003**, 208, 87-91	3.2	6
6	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, 159-66	2.9	24
5	Molecular heterogeneity of myophosphorylase deficiency (Mcardle¥s disease): A genotype-phenotype correlation study. *Annals of Neurology*, **2001**, 50, 574-581	9.4	58
4	A novel missense mutation (W797R) in the myophosphorylase gene in Spanish patients with McArdle disease. *Archives of Neurology*, **2000**, 57, 217-9		17
3	Evaluation of heart involvement in gamma-sarcoglycanopathy (LGMD2C). A study of ten patients. *Neuromuscular Disorders*, **2000**, 10, 560-6	2.9	32
2	Adult glycogenosis II with paracrystalline mitochondrial inclusions and Hirano bodies in skeletal muscle. *Neuromuscular Disorders*, **1999**, 9, 136-43	2.9	19
1	Subsarcolemmal expression of utrophin in neuromuscular disorders: an immunohistochemical study of 80 cases. *Acta Neuropathologica*, **1998**, 96, 481-6	14.3	6