Mĥica Santos

List of Publications by Year in Descending Order

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Version: 2024-04-23

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

27 papers	563 citations	13 h-index	23 g-index
29 ext. papers	642 ext. citations	4.4 avg, IF	2.74 L-index

#	Paper	IF	Citations
27	MeCP2 haplodeficiency and early-life stress interaction on anxiety-like behavior in adolescent female mice.. *Journal of Neurodevelopmental Disorders*, **2021**, 13, 59	4.6	2
26	Male-specific features are reduced in Mecp2-null mice: analyses of vasopressinergic innervation, pheromone production and social behaviour. *Brain Structure and Function*, **2020**, 225, 2219-2238	4	4
25	Lack of MeCP2 leads to region-specific increase of doublecortin in the olfactory system. *Brain Structure and Function*, **2019**, 224, 1647-1658	4	4
24	Selective rescue of heightened anxiety but not gait ataxia in a premutation 90CGG mouse model of Fragile X-associated tremor/ataxia syndrome. *Human Molecular Genetics*, **2017**, 26, 2133-2145	5.6	10
23	Infralimbic Neurotrophin-3 Infusion Rescues Fear Extinction Impairment in a Mouse Model of Pathological Fear. *Neuropsychopharmacology*, **2017**, 42, 462-472	8.7	6
22	Aberrant neuronal activity-induced signaling and gene expression in a mouse model of RASopathy. *PLoS Genetics*, **2017**, 13, e1006684	6	11
21	Fragile X-Associated Tremor/Ataxia Syndrome (FXTAS) Motor Dysfunction Modeled in Mice. *Cerebellum*, **2016**, 15, 611-22	4.3	7
20	From neural to genetic substrates of panic disorder: Insights from human and mouse studies. *European Journal of Pharmacology*, **2015**, 759, 127-41	5.3	9
19	Genome-wide miR-155 and miR-802 target gene identification in the hippocampus of Ts65Dn Down syndrome mouse model by miRNA sponges. *BMC Genomics*, **2015**, 16, 907	4.5	24
18	Identificación de genes clave implicados en el síndrome de Down mediante terapia génica. *Revista Médica Internacional Sobre El Síndrome De Down*, **2014**, 18, 21-28		
17	Hippocampal hyperexcitability underlies enhanced fear memories in TgNTRK3, a panic disorder mouse model. *Journal of Neuroscience*, **2013**, 33, 15259-71	6.6	25
16	Rett syndrome with and without detected MECP2 mutations: an attempt to redefine phenotypes. *Brain and Development*, **2011**, 33, 69-76	2.2	18
15	Monoamine deficits in the brain of methyl-CpG binding protein 2 null mice suggest the involvement of the cerebral cortex in early stages of Rett syndrome. *Neuroscience*, **2010**, 170, 453-67	3.9	37
14	Mutations in the MECP2 gene are not a major cause of Rett syndrome-like or related neurodevelopmental phenotype in male patients. *Journal of Child Neurology*, **2009**, 24, 49-55	2.5	10
13	T-1237C polymorphism of TLR9 gene is not associated with multiple sclerosis in the Portuguese population. *Multiple Sclerosis Journal*, **2008**, 14, 550-2	5	3
12	The C677T polymorphism in MTHFR is not associated with migraine in Portugal. *Disease Markers*, **2008**, 25, 107-13	3.2	19
11	Analysis of highly conserved regions of the 3'UTR of MECP2 gene in patients with clinical diagnosis of Rett syndrome and other disorders associated with mental retardation. *Disease Markers*, **2008**, 24, 319-24	3.2	7

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10	MECP2 coding sequence and 3ՍTR variation in 172 unrelated autistic patients. *American Journal of Medical Genetics Part B: Neuropsychiatric Genetics*, **2007**, 144B, 475-83	3.5	49
9	An explanation for another familial case of Rett syndrome: maternal germline mosaicism. *European Journal of Human Genetics*, **2007**, 15, 902-4	5.3	25
8	Evidence for abnormal early development in a mouse model of Rett syndrome. *Genes, Brain and Behavior*, **2007**, 6, 277-86	3.6	71
7	Stereotypies in Rett syndrome: analysis of 83 patients with and without detected MECP2 mutations. *Neurology*, **2007**, 68, 1183-7	6.5	65
6	Chromatin remodeling and neuronal function: exciting links. *Genes, Brain and Behavior*, **2006**, 5 Suppl 2, 80-91	3.6	7
5	Linkage disequilibrium screening for multiple sclerosis implicates JAG1 and POU2AF1 as susceptibility genes in Europeans. *Journal of Neuroimmunology*, **2006**, 179, 108-16	3.5	22
4	APOE epsilon variation in multiple sclerosis susceptibility and disease severity: some answers. *Neurology*, **2006**, 66, 1373-83	6.5	72
3	Detection of heterozygous deletions and duplications in the MECP2 gene in Rett syndrome by Robust Dosage PCR (RD-PCR). *Human Mutation*, **2005**, 25, 505	4.7	22
2	Genotypes at the APOE and SCA2 loci do not predict the course of multiple sclerosis in patients of Portuguese origin. *Multiple Sclerosis Journal*, **2004**, 10, 153-7	5	22
1	A whole genome screen for association with multiple sclerosis in Portuguese patients. *Journal of Neuroimmunology*, **2003**, 143, 112-5	3.5	10