

Mônica Santos

List of Publications by Citations

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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	APOE epsilon variation in multiple sclerosis susceptibility and disease severity: some answers. <i>Neurology</i> , 2006 , 66, 1373-83	6.5	72
26	Evidence for abnormal early development in a mouse model of Rett syndrome. <i>Genes, Brain and Behavior</i> , 2007 , 6, 277-86	3.6	71
25	Stereotypies in Rett syndrome: analysis of 83 patients with and without detected MECP2 mutations. <i>Neurology</i> , 2007 , 68, 1183-7	6.5	65
24	MECP2 coding sequence and 3ΨTR variation in 172 unrelated autistic patients. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2007 , 144B, 475-83	3.5	49
23	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. <i>Neuroscience</i> , 2010 , 170, 453-67	3.9	37
22	Hippocampal hyperexcitability underlies enhanced fear memories in TgNTRK3, a panic disorder mouse model. <i>Journal of Neuroscience</i> , 2013 , 33, 15259-71	6.6	25
21	An explanation for another familial case of Rett syndrome: maternal germline mosaicism. <i>European Journal of Human Genetics</i> , 2007 , 15, 902-4	5.3	25
20	Genome-wide miR-155 and miR-802 target gene identification in the hippocampus of Ts65Dn Down syndrome mouse model by miRNA sponges. <i>BMC Genomics</i> , 2015 , 16, 907	4.5	24
19	Linkage disequilibrium screening for multiple sclerosis implicates JAG1 and POU2AF1 as susceptibility genes in Europeans. <i>Journal of Neuroimmunology</i> , 2006 , 179, 108-16	3.5	22
18	Genotypes at the APOE and SCA2 loci do not predict the course of multiple sclerosis in patients of Portuguese origin. <i>Multiple Sclerosis Journal</i> , 2004 , 10, 153-7	5	22
17	Detection of heterozygous deletions and duplications in the MECP2 gene in Rett syndrome by Robust Dosage PCR (RD-PCR). <i>Human Mutation</i> , 2005 , 25, 505	4.7	22
16	The C677T polymorphism in MTHFR is not associated with migraine in Portugal. <i>Disease Markers</i> , 2008 , 25, 107-13	3.2	19
15	Rett syndrome with and without detected MECP2 mutations: an attempt to redefine phenotypes. <i>Brain and Development</i> , 2011 , 33, 69-76	2.2	18
14	Aberrant neuronal activity-induced signaling and gene expression in a mouse model of RASopathy. <i>PLoS Genetics</i> , 2017 , 13, e1006684	6	11
13	Selective rescue of heightened anxiety but not gait ataxia in a premutation 90CGG mouse model of Fragile X-associated tremor/ataxia syndrome. <i>Human Molecular Genetics</i> , 2017 , 26, 2133-2145	5.6	10
12	Mutations in the MECP2 gene are not a major cause of Rett syndrome-like or related neurodevelopmental phenotype in male patients. <i>Journal of Child Neurology</i> , 2009 , 24, 49-55	2.5	10
11	A whole genome screen for association with multiple sclerosis in Portuguese patients. <i>Journal of Neuroimmunology</i> , 2003 , 143, 112-5	3.5	10

10	From neural to genetic substrates of panic disorder: Insights from human and mouse studies. <i>European Journal of Pharmacology</i> , 2015 , 759, 127-41	5.3	9
9	Fragile X-Associated Tremor/Ataxia Syndrome (FXTAS) Motor Dysfunction Modeled in Mice. <i>Cerebellum</i> , 2016 , 15, 611-22	4.3	7
8	Analysis of highly conserved regions of the 3ΨTR of MECP2 gene in patients with clinical diagnosis of Rett syndrome and other disorders associated with mental retardation. <i>Disease Markers</i> , 2008 , 24, 319-24	3.2	7
7	Chromatin remodeling and neuronal function: exciting links. <i>Genes, Brain and Behavior</i> , 2006 , 5 Suppl 2, 80-91	3.6	7
6	Infralimbic Neurotrophin-3 Infusion Rescues Fear Extinction Impairment in a Mouse Model of Pathological Fear. <i>Neuropsychopharmacology</i> , 2017 , 42, 462-472	8.7	6
5	Lack of MeCP2 leads to region-specific increase of doublecortin in the olfactory system. <i>Brain Structure and Function</i> , 2019 , 224, 1647-1658	4	4
4	Male-specific features are reduced in Mecp2-null mice: analyses of vasopressinergic innervation, pheromone production and social behaviour. <i>Brain Structure and Function</i> , 2020 , 225, 2219-2238	4	4
3	T-1237C polymorphism of TLR9 gene is not associated with multiple sclerosis in the Portuguese population. <i>Multiple Sclerosis Journal</i> , 2008 , 14, 550-2	5	3
2	MeCP2 haplodeficiency and early-life stress interaction on anxiety-like behavior in adolescent female mice.. <i>Journal of Neurodevelopmental Disorders</i> , 2021 , 13, 59	4.6	2
1	Identificaci3n de genes clave implicados en el s3ndrome de Down mediante terapia g3nica. <i>Revista M3dica Internacional Sobre El S3ndrome De Down</i> , 2014 , 18, 21-28		