Andrea Citterio

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9318084/publications.pdf

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17	444	840585 11	839398 18
papers	citations	h-index	g-index

18	18	18	1044
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Mutations in CYP2U1, DDHD2 and GBA2 genes are rare causes of complicated forms of hereditary spastic paraparesis. Journal of Neurology, 2014, 261, 373-381.	1.8	62
2	Variants in KIF1A gene in dominant and sporadic forms of hereditary spastic paraparesis. Journal of Neurology, 2015, 262, 2684-2690.	1.8	55
3	Processing of Meiotic DNA Double Strand Breaks Requires Cyclin-dependent Kinase and Multiple Nucleases. Journal of Biological Chemistry, 2010, 285, 11628-11637.	1.6	46
4	Clinical and Paraclinical Indicators of Motor System Impairment in Hereditary Spastic Paraplegia: A Pilot Study. PLoS ONE, 2016, 11, e0153283.	1.1	41
5	ZFYVE26/SPASTIZIN and SPG11/SPATACSIN mutations in hereditary spastic paraplegia types AR-SPG15 and AR-SPG11 have different effects on autophagy and endocytosis. Autophagy, 2019, 15, 34-57.	4.3	41
6	Tubulin-related cerebellar dysplasia: definition of a distinct pattern of cerebellar malformation. European Radiology, 2017, 27, 5080-5092.	2.3	36
7	Antipsychotics Promote Metabolic Disorders Disrupting Cellular Lipid Metabolism and Trafficking. Trends in Endocrinology and Metabolism, 2019, 30, 189-210.	3.1	30
8	Impairment of brain and muscle energy metabolism detected by magnetic resonance spectroscopy in hereditary spastic paraparesis type 28 patients with DDHD1 mutations. Journal of Neurology, 2014, 261, 1789-1793.	1.8	25
9	Atypical adult onset complicated spastic paraparesis with thin corpus callosum in two patients carrying a novel <scp>FA2H</scp> mutation. European Journal of Neurology, 2012, 19, e127-9.	1.7	22
10	Loss of ap4s1 in zebrafish leads to neurodevelopmental defects resembling spastic paraplegia 52. Annals of Clinical and Translational Neurology, 2020, 7, 584-589.	1.7	15
11	A Novel CAPN1 Mutation Causes a Pure Hereditary Spastic Paraplegia in an Italian Family. Frontiers in Neurology, 2019, 10, 580.	1.1	14
12	KIF5A and ALS2 Variants in a Family With Hereditary Spastic Paraplegia and Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 2018, 9, 1078.	1.1	12
13	Novel SPTBN2 gene mutation and first intragenic deletion in early onset spinocerebellar ataxia type 5. Annals of Clinical and Translational Neurology, 2021, 8, 956-963.	1.7	12
14	Cerebroretinal Microangiopathy With Calcifications and Cysts Associated With CTC1 and NDP Mutations. Journal of Child Neurology, 2013, 28, 1702-1708.	0.7	11
15	Novel SETX variants in a patient with ataxia, neuropathy, and oculomotor apraxia are associated with normal sensitivity to oxidative DNA damaging agents. Brain and Development, 2014, 36, 682-689.	0.6	8
16	Very preterm birth is associated with PLAGL1 gene hypomethylation at birth and discharge. Epigenomics, 2018, 10, 1121-1130.	1.0	7
17	Exome sequencing reveals a novel homozygous mutation in ACP33 gene in the first Italian family with SPG21. Journal of Neurology, 2017, 264, 2021-2023.	1.8	4