

Andrea Citterio

List of Publications by Year in descending order

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17
papers

444
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840585

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839398

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1044
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#	ARTICLE	IF	CITATIONS
1	Novel <i>SPTBN2</i> gene mutation and first intragenic deletion in early onset spinocerebellar ataxia type 5. <i>Annals of Clinical and Translational Neurology</i> , 2021, 8, 956-963.	1.7	12
2	Loss of <i>ap4s1</i> in zebrafish leads to neurodevelopmental defects resembling spastic paraplegia 52. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 584-589.	1.7	15
3	ZFYVE26/SPASTIZIN and SPG11/SPATACSIN mutations in hereditary spastic paraplegia types AR-SPG15 and AR-SPG11 have different effects on autophagy and endocytosis. <i>Autophagy</i> , 2019, 15, 34-57.	4.3	41
4	Antipsychotics Promote Metabolic Disorders Disrupting Cellular Lipid Metabolism and Trafficking. <i>Trends in Endocrinology and Metabolism</i> , 2019, 30, 189-210.	3.1	30
5	A Novel CAPN1 Mutation Causes a Pure Hereditary Spastic Paraplegia in an Italian Family. <i>Frontiers in Neurology</i> , 2019, 10, 580.	1.1	14
6	KIF5A and ALS2 Variants in a Family With Hereditary Spastic Paraplegia and Amyotrophic Lateral Sclerosis. <i>Frontiers in Neurology</i> , 2018, 9, 1078.	1.1	12
7	Very preterm birth is associated with <i>PLAGL1</i> gene hypomethylation at birth and discharge. <i>Epigenomics</i> , 2018, 10, 1121-1130.	1.0	7
8	Exome sequencing reveals a novel homozygous mutation in ACP33 gene in the first Italian family with SPG21. <i>Journal of Neurology</i> , 2017, 264, 2021-2023.	1.8	4
9	Tubulin-related cerebellar dysplasia: definition of a distinct pattern of cerebellar malformation. <i>European Radiology</i> , 2017, 27, 5080-5092.	2.3	36
10	Clinical and Paraclinical Indicators of Motor System Impairment in Hereditary Spastic Paraplegia: A Pilot Study. <i>PLoS ONE</i> , 2016, 11, e0153283.	1.1	41
11	Variants in KIF1A gene in dominant and sporadic forms of hereditary spastic paraparesis. <i>Journal of Neurology</i> , 2015, 262, 2684-2690.	1.8	55
12	Mutations in CYP2U1, DDHD2 and GBA2 genes are rare causes of complicated forms of hereditary spastic paraparesis. <i>Journal of Neurology</i> , 2014, 261, 373-381.	1.8	62
13	Impairment of brain and muscle energy metabolism detected by magnetic resonance spectroscopy in hereditary spastic paraparesis type 28 patients with DDHD1 mutations. <i>Journal of Neurology</i> , 2014, 261, 1789-1793.	1.8	25
14	Novel SETX variants in a patient with ataxia, neuropathy, and oculomotor apraxia are associated with normal sensitivity to oxidative DNA damaging agents. <i>Brain and Development</i> , 2014, 36, 682-689.	0.6	8
15	Cerebroretinal Microangiopathy With Calcifications and Cysts Associated With <i>CTC1</i> and <i>NDP</i> Mutations. <i>Journal of Child Neurology</i> , 2013, 28, 1702-1708.	0.7	11
16	Atypical adult onset complicated spastic paraparesis with thin corpus callosum in two patients carrying a novel <i>FA2H</i> mutation. <i>European Journal of Neurology</i> , 2012, 19, e127-9.	1.7	22
17	Processing of Meiotic DNA Double Strand Breaks Requires Cyclin-dependent Kinase and Multiple Nucleases. <i>Journal of Biological Chemistry</i> , 2010, 285, 11628-11637.	1.6	46