Marta Arpone

List of Publications by Year
in descending order

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Version: 2024-02-01

15 papers	258 citations	1039880 9 h-index	996849 15 g-index
17 all docs	17 docs citations	17 times ranked	433 citing authors

#	Article	IF	Citations
1	Microstructural white matter correlates of emotion recognition impairment in Amyotrophic Lateral Sclerosis. Cortex, 2014, 53, 1-8.	1.1	71
2	Intellectual functioning and behavioural features associated with mosaicism in fragile X syndrome. Journal of Neurodevelopmental Disorders, 2019, 11, 41.	1.5	26
3	Dysarthria and broader motor speech deficits in Dravet syndrome. Neurology, 2017, 88, 743-749.	1.5	22
4	Cognitive, behavioral, and social functioning in children and adults with Dravet syndrome. Epilepsy and Behavior, 2020, 112, 107319.	0.9	21
5	Partially methylated alleles, microdeletion, and tissue mosaicism in a fragile X male with tremor and ataxia at 30 years of age: A case report. American Journal of Medical Genetics, Part A, 2016, 170, 3327-3332.	0.7	20
6	Incomplete silencing of full mutation alleles in males with fragile X syndrome is associated with autistic features. Molecular Autism, 2019, 10, 21.	2.6	20
7	Intragenic DNA methylation in buccal epithelial cells and intellectual functioning in a paediatric cohort of males with fragile X. Scientific Reports, 2018, 8, 3644.	1.6	17
8	Significantly Elevated FMR1 mRNA and Mosaicism for Methylated Premutation and Full Mutation Alleles in Two Brothers with Autism Features Referred for Fragile X Testing. International Journal of Molecular Sciences, 2019, 20, 3907.	1.8	12
9	Development and Usability of a Novel Interactive Tablet App (PediAppRREST) to Support the Management of Pediatric Cardiac Arrest: Pilot High-Fidelity Simulation-Based Study. JMIR MHealth and UHealth, 2020, 8, e19070.	1.8	12
10	FMR1 mRNA from full mutation alleles is associated with ABC-CFX scores in males with fragile X syndrome. Scientific Reports, 2020, 10, 11701.	1.6	11
11	DNA Methylation at Birth Predicts Intellectual Functioning and Autism Features in Children with Fragile X Syndrome. International Journal of Molecular Sciences, 2020, 21, 7735.	1.8	10
12	Molecular Inconsistencies in a Fragile X Male with Early Onset Ataxia. Genes, 2016, 7, 68.	1.0	8
13	Clinical and Molecular Differences between 4-Year-Old Monozygous Male Twins Mosaic for Normal, Premutation and Fragile X Full Mutation Alleles. Genes, 2019, 10, 279.	1.0	4
14	Missed intracranial injuries are rare in emergency departments using the PECARN head injury decision rules. Child's Nervous System, 2021, 37, 55-62.	0.6	3
15	PediAppRREST: effectiveness of an interactive cognitive support tablet app in reducing deviations from guidelines in the management of paediatric cardiac arrest: protocol for a simulation-based randomised controlled trial. BMJ Open, 2021, 11, e047208.	0.8	1