

Francesca Marchese

List of Publications by Year in descending order

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Version: 2024-02-01

16
papers

198
citations

1040056

9
h-index

1058476

14
g-index

16
all docs

16
docs citations

16
times ranked

345
citing authors

#	ARTICLE	IF	CITATIONS
1	Migraine and cluster headache – the common link. <i>Journal of Headache and Pain</i> , 2018, 19, 89.	6.0	44
2	Genotype-phenotype correlations in patients with de novo <i>KCNQ2</i> pathogenic variants. <i>Neurology: Genetics</i> , 2020, 6, e528.	1.9	24
3	Epilepsy Course and Developmental Trajectories in <i>STXBP1</i> -DEE. <i>Neurology: Genetics</i> , 2022, 8, .	1.9	24
4	A further contribution to the delineation of epileptic phenotype in PACS2-related syndrome. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2020, 79, 53-55.	2.0	15
5	Clinical spectrum and genotype-phenotype correlations in <i>PRRT2</i> Italian patients. <i>European Journal of Paediatric Neurology</i> , 2020, 28, 193-197.	1.6	14
6	Cannabidiol Determination on Peripheral Capillary Blood Using a Microsampling Method and Ultra-High-Performance Liquid Chromatography Tandem Mass Spectrometry with On-Line Sample Preparation. <i>Molecules</i> , 2020, 25, 3608.	3.8	10
7	Complex Neurological Phenotype Associated with a De Novo <i>DHDDS</i> Mutation in a Boy with Intellectual Disability, Refractory Epilepsy, and Movement Disorder. <i>Journal of Pediatric Genetics</i> , 2021, 10, 236-238.	0.7	10
8	Migraine in children under 6 years of age: A long-term follow-up study. <i>European Journal of Paediatric Neurology</i> , 2020, 27, 67-71.	1.6	10
9	<p><p>Adjunctive Rufinamide in Children with Lennox-Gastaut Syndrome: A Literature Review</p></p>. <i>Neuropsychiatric Disease and Treatment</i> , 2020, Volume 16, 369-379.	2.2	9
10	A nationwide study on Sydenham's chorea: Clinical features, treatment and prognostic factors. <i>European Journal of Paediatric Neurology</i> , 2022, 36, 1-6.	1.6	9
11	A Volumetric Absorptive Microsampling Technique to Monitor Cannabidiol Levels in Epilepsy Patients. <i>Frontiers in Pharmacology</i> , 2020, 11, 582286.	3.5	7
12	Electroclinical features of <i>MEF2C</i> haploinsufficiency-related epilepsy: A multicenter European study. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2021, 88, 60-72.	2.0	6
13	Comorbidities in Dravet Syndrome and Lennox-Gastaut Syndrome. <i>SN Comprehensive Clinical Medicine</i> , 2021, 3, 2167-2179.	0.6	6
14	Abnormal circadian rhythm in patients with <i>GRIN1</i> -related developmental epileptic encephalopathy. <i>European Journal of Paediatric Neurology</i> , 2019, 23, 657-661.	1.6	4
15	Atypical Presentation of Aromatic L-Amino Acid Decarboxylase Deficiency with Developmental Epileptic Encephalopathy. <i>Journal of Pediatric Epilepsy</i> , 2021, 10, 124-127.	0.2	4
16	Temporal-parietal-occipital epilepsy in GEFS+ associated with <i>SCN1A</i> mutation. <i>Epileptic Disorders</i> , 2021, 23, 397-401.	1.3	2