Simona Cappelletti

List of Publications by Year
in descending order

24 papers	572 citations	567281 15 h-index	642732 23 g-index
24 all docs	24 docs citations	24 times ranked	905 citing authors

#	Article	IF	Citations
1	Comorbidities in Dravet Syndrome and Lennoxâ€“Gastaut Syndrome. SN Comprehensive Clinical Medicine, 2021, 3, 2167-2179.	0.6	6
2	Purified Cannabidiol for Treatment of Refractory Epilepsies in Pediatric Patients with Developmental and Epileptic Encephalopathy. Paediatric Drugs, 2019, 21, 283-290.	3.1	19
3	Symptoms of anxiety and depression and family's quality of life in children and adolescents with epilepsy. Epilepsy and Behavior, 2018, 79, 146-153.	1.7	28
4	Defining the electroclinical phenotype and outcome of PCDH19â€related epilepsy: A multicenter study. Epilepsia, 2018, 59, 2260-2271.	5.1	39
5	PCDH19-related epilepsy and Dravet Syndrome: Face-off between two early-onset epilepsies with fever sensitivity. Epilepsy Research, 2016, 125, 32-36.	1.6	28
6	Mutation of <scp>CHRNA</scp>2 in a family with benign familial infantile seizures: Potential role of nicotinic acetylcholine receptor in various phenotypes of epilepsy. Epilepsia, 2015, 56, e53-7.	5.1	19
7	Acute intralesional recording in hypothalamic hamartoma: description of 4 cases. Acta Neurologica Belgica, 2015, 115, 233-239.	1.1	6
8	CHD2 mutations are a rare cause of generalized epilepsy with myoclonicâ€“atonic seizures. Epilepsy and Behavior, 2015, 51, 53-56.	1.7	28
9	Structural Focal Temporal Lobe Seizures in a Child With Lipoproteinosis. Pediatric Neurology, 2015, 52, 104-106.	2.1	1
10	Cognitive development in females with PCDH19 gene-related epilepsy. Epilepsy and Behavior, 2015, 42, 36-40.	1.7	32
11	Behavioral disorders as unusual presentation of pediatric extraventricular neurocytoma: report on two cases and review of the literature. BMC Neurology, 2014, 14, 242.	1.8	10
12	Occipital seizures induced by Intermittent Photic Stimulation in Dravet syndrome. Seizure: the Journal of the British Epilepsy Association, 2014, 23, 309-313.	2.0	10
13	Epilepsy in Patients With Duplications of Chromosome 14 Harboring FOXG1. Pediatric Neurology, 2014, 50, 530-535.	2.1	17
14	PRRT2 is mutated in familial and non-familial benign infantile seizures. European Journal of Paediatric Neurology, 2013, 17, 77-81.	1.6	22
15	Cognitive Findings and Behavior in Children and Adolescents with Phenylketonuria. Journal of Developmental and Behavioral Pediatrics, 2013, 34, 392-398.	1.1	22
16	Electroencephalographic Features in Dravet Syndrome. Journal of Child Neurology, 2012, 27, 439-444.	1.4	34
17	Epilepsy in ring 14 chromosome syndrome. Epilepsy and Behavior, 2012, 25, 585-592.	1.7	17
18	Neonatal hemifacial spasm and fourth ventricle mass. Developmental Medicine and Child Neurology, 2012, 54, 697-703.	2.1	9

#	Article	IF	Citations
19	Prolonged episode of dystonia and dyskinesia resembling status epilepticus following acute intrathecal baclofen withdrawal. Epilepsy and Behavior, 2011, 21, 321-323.	1.7	15
20	Ictal yawning in a patient with drug-resistant focal epilepsy: Video/EEG documentation and review of literature reports. Epilepsy and Behavior, 2011, 22, 602-605.	1.7	3
21	Cognitive development in Dravet syndrome: A retrospective, multicenter study of 26 patients. Epilepsia, 2011, 52, 386-392.	5.1	99
22	Myoclonic astatic epilepsy: An age-dependent epileptic syndrome with favorable seizure outcome but variable cognitive evolution. Epilepsy Research, 2011, 97, 133-141.	1.6	33
23	Ring 21 chromosome presenting with epilepsy and intellectual disability: Clinical report and review of the literature. , 2011, 155, 911-914.		13
24	Panayiotopoulos syndrome: A clinical, EEG, and neuropsychological study of 93 consecutive patients. Epilepsia, 2010, 51, 2098-2107.	5.1	62