

HÃ©lio van der Linden

List of Publications by Year in descending order

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Version: 2024-02-01

18
papers

818
citations

840776

11
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839539

18
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times ranked

1866
citing authors

#	ARTICLE	IF	CITATIONS
1	Description of 13 Infants Born During October 2015â€“January 2016 With Congenital Zika Virus Infection Without Microcephaly at Birth â€” Brazil. <i>Morbidity and Mortality Weekly Report</i> , 2016, 65, 1343-1348.	15.1	368
2	Spinocerebellar Ataxias in Brazilâ€”Frequencies and Modulating Effects of Related Genes. <i>Cerebellum</i> , 2014, 13, 17-28.	2.5	93
3	Discordant congenital Zika syndrome twins show differential in vitro viral susceptibility of neural progenitor cells. <i>Nature Communications</i> , 2018, 9, 475.	12.8	86
4	Heterozygous Variants in KMT2E Cause a Spectrum of Neurodevelopmental Disorders and Epilepsy. <i>American Journal of Human Genetics</i> , 2019, 104, 1210-1222.	6.2	56
5	Epilepsy Profile in Infants with Congenital Zika Virus Infection. <i>New England Journal of Medicine</i> , 2018, 379, 891-892.	27.0	45
6	Discordant clinical outcomes of congenital Zika virus infection in twin pregnancies. <i>Arquivos De Neuro-Psiquiatria</i> , 2017, 75, 381-386.	0.8	30
7	Association of Severe Hydrocephalus With Congenital Zika Syndrome. <i>JAMA Neurology</i> , 2019, 76, 203.	9.0	28
8	Clinical and biochemical study of 29 Brazilian patients with metachromatic leukodystrophy. <i>Journal of Inherited Metabolic Disease</i> , 2010, 33, 257-262.	3.6	18
9	Clinical and Molecular Characterization of Mcardleâ€™s Disease in Brazilian Patients. <i>NeuroMolecular Medicine</i> , 2013, 15, 470-475.	3.4	15
10	Recessive congenital myasthenic syndrome caused by a homozygous mutation in SYT2 altering a highly conserved Câ€“terminal amino acid sequence. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 1744-1749.	1.2	14
11	Movement disorders in children with congenital Zika virus syndrome. <i>Brain and Development</i> , 2020, 42, 720-729.	1.1	12
12	Neurodevelopment in Children Exposed to Zika in utero: Clinical and Molecular Aspects. <i>Frontiers in Genetics</i> , 2022, 13, 758715.	2.3	12
13	The approach to patients with psychogenic nonepileptic seizures in epilepsy surgery centers regarding diagnosis, treatment, and education. <i>Epilepsy and Behavior</i> , 2017, 68, 78-83.	1.7	11
14	ATXN3, ATXN7, CACNA1A, and RAI1 Genes and Mitochondrial Polymorphism A10398G Did Not Modify Age at Onset in Spinocerebellar Ataxia Type 2 Patients from South America. <i>Cerebellum</i> , 2015, 14, 728-730.	2.5	10
15	Genetic polymorphisms of the 5HT receptors are not related with depression in temporal lobe epilepsy caused by hippocampal sclerosis. <i>Epilepsy and Behavior</i> , 2018, 83, 181-185.	1.7	6
16	Continuous epileptiform discharges during sleep as an evolutionary pattern in patients with congenital Zika virus syndrome. <i>Epilepsia</i> , 2020, 61, e107-e115.	5.1	6
17	Higher transcription alleles of the MAOA-uVNTR polymorphism are associated with higher seizure frequency in temporal lobe epilepsy. <i>Epilepsy Research</i> , 2019, 149, 26-29.	1.6	5
18	Epilepsy and EEG Abnormalities in Congenital Zika Syndrome. <i>Journal of Clinical Neurophysiology</i> , 2022, 39, 248-252.	1.7	3