

Seyda Besen

List of Publications by Year in descending order

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Version: 2024-02-01

20
papers

97
citations

1684188

5
h-index

1588992

8
g-index

20
all docs

20
docs citations

20
times ranked

115
citing authors

#	ARTICLE	IF	CITATIONS
1	First pediatric case with primary familial brain calcification due to a novel variant on the MYORG gene and review of the literature. <i>Brain and Development</i> , 2021, 43, 789-797.	1.1	6
2	Effect of levetiracetam usage on serum creatine phosphokinase concentration in patients with epilepsy. <i>Journal of Pediatric Neurosciences</i> , 2020, 15, 81.	0.3	5
3	Dynamic thiol/disulphide homeostasis in children with neurofibromatosis type 1 and tuberous sclerosis. <i>Acta Neurologica Belgica</i> , 2019, 119, 419-422.	1.1	3
4	Çocuklarda psikiyatrik serebri: etyoloji, klinik bulgular, prognoz. <i>Cukurova Medical Journal</i> , 2019, 44, 1-1.	0.2	0
5	Mucopolysaccharidosis type VI, 9 sibling pairs and 1 set of three siblings: single center experience from Turkey. <i>Molecular Genetics and Metabolism</i> , 2018, 123, S101.	1.1	0
6	Autoimmune encephalitis associated with glutamic acid decarboxylase antibodies: a case series. <i>Acta Neurologica Belgica</i> , 2018, 118, 411-414.	1.1	12
7	Late-onset Leigh syndrome due to NDUFV1 mutation in a 10-year-old boy initially presenting with ataxia. <i>Journal of Pediatric Neurosciences</i> , 2018, 13, 205.	0.3	11
8	Hereditary Spastic Paraplegia Type 35 with a Novel Mutation in Fatty Acid 2-Hydroxylase Gene and Literature Review of the Clinical Features. <i>Annals of Indian Academy of Neurology</i> , 2018, 21, 335-339.	0.5	3
9	Hereditary spastic paraplegia type 35 with a novel mutation in fatty acid 2-hydroxylase gene and literature review of the clinical features. <i>Annals of Indian Academy of Neurology</i> , 2018, 21, 335.	0.5	6
10	Finger drop sign in a child with acute motor and sensory axonal neuropathy form of Guillain-Barré syndrome. <i>Acta Neurologica Belgica</i> , 2017, 117, 393-394.	1.1	2
11	Risk factors for epilepsy after ischemic stroke in children. <i>Journal of Dr Behcet Uz Children S Hospital</i> , 2017, , .	0.1	2
12	Acute rhabdomyolysis associated with levetiracetam therapy in a child. <i>Acta Neurologica Belgica</i> , 2016, 116, 369-370.	1.1	13
13	Limbic encephalitis associated with anti-leucine-rich glioma-inactivated-1 protein antibodies in a child. <i>Neurology India</i> , 2016, 64, 1321.	0.4	5
14	Intravenous levetiracetam in critically ill children. <i>Annals of Indian Academy of Neurology</i> , 2016, 19, 79.	0.5	5
15	Guillain-Barré syndrome with hyperreflexia and bilateral papillitis in a child. <i>Journal of Pediatric Neurosciences</i> , 2016, 11, 71.	0.3	5
16	Mitochondrial membrane protein-associated neurodegeneration in a Turkish patient. <i>Journal of Pediatric Neurosciences</i> , 2016, 11, 288.	0.3	2
17	P125 – 2716: A presentation of Lyme disease: Pseudotumor cerebri. <i>European Journal of Paediatric Neurology</i> , 2015, 19, S129.	1.6	0
18	Limbic encephalitis with antibodies to glutamic acid decarboxylase presenting with brainstem symptoms. <i>Annals of Indian Academy of Neurology</i> , 2015, 18, 243.	0.5	11

#	ARTICLE	IF	CITATIONS
19	Urinary and fecal incontinence during levetiracetam therapy. <i>Annals of Indian Academy of Neurology</i> , 2015, 18, 479-80.	0.5	3
20	Insulin sensitivity obtained from the oral glucose tolerance test and its relationship with birthweight. <i>Annals of Saudi Medicine</i> , 2007, 27, 13.	1.1	3