Periyasamy Govindaraj

List of Publications by Year
in descending order

16	142	1307594 7	1281871 11
papers	citations	h-index	g-index

16	16	16	208
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Infantile Onset Encephalomyopathy, Heart Block, and Sensorineural Hearing Loss: RMND1-Associated Mitochondrial Disease. Journal of Pediatric Neurology, 2021, 19, 183-188.	0.2	0
2	Child Neurology: Hereditary Folate Malabsorption. Neurology, 2021, 97, 40-43.	1.1	2
3	Ethylmalonic encephalopathy ETHE1 p. D165H mutation alters the mitochondrial function in human skeletal muscle proteome. Mitochondrion, 2021, 58, 64-71.	3.4	4
4	Serum fibroblast growth factor 21 and growth differentiation factor 15: Two sensitive biomarkers in the diagnosis of mitochondrial disorders. Mitochondrion, 2021, 60, 170-177.	3.4	9
5	Leukodystrophy due to eIF2B mutations in adults. Canadian Journal of Neurological Sciences, 2021, , 1-11.	0.5	1
6	Exposure to the neurotoxin 3-nitropropionic acid in neuronal cells induces unique histone acetylation pattern: Implications for neurodegeneration. Neurochemistry International, 2020, 140, 104846.	3.8	1
7	Leukodystrophies and Genetic Leukoencephalopathies in Children Specified by Exome Sequencing in an Expanded Gene Panel. Journal of Child Neurology, 2020, 35, 433-441.	1.4	11
8	Child Neurology: Ethylmalonic encephalopathy. Neurology, 2020, 94, e1336-e1339.	1.1	13
9	Mitochondrial leukoencephalopathies: A border zone between acquired and inherited white matter disorders in children?. Multiple Sclerosis and Related Disorders, 2018, 20, 84-92.	2.0	27
10	Outcome of epilepsy in patients with mitochondrial disorders: Phenotype genotype and magnetic resonance imaging correlations. Clinical Neurology and Neurosurgery, 2018, 164, 182-189.	1.4	17
11	Clinical Reasoning: West syndrome, pontocerebellar hypoplasia, and hypomyelination in a 6-month-old boy. Neurology, 2018, 91, e1652-e1656.	1.1	1
12	Mitochondrial oxidative phosphorylation disorders in children: Phenotypic, genotypic and biochemical correlations in 85 patients from South India. Mitochondrion, 2017, 32, 42-49.	3.4	17
13	Leber's Hereditary Optic Neuropathyâ€“Specific Mutation m.11778G>A Exists on Diverse Mitochondrial Haplogroups in India. , 2017, 58, 3923.		19
14	Reply to Letter to the Editor: Hearing impairment in m.3243A>G carriers requires comprehensive work- and follow-up. Clinical Neurology and Neurosurgery, 2016, 150, 198-199.	1.4	0
15	Audiological manifestations in mitochondrial encephalomyopathy lactic acidosis and stroke like episodes (MELAS) syndrome. Clinical Neurology and Neurosurgery, 2016, 148, 17-21.	1.4	18
16	Mutational analysis of telomere complex genes in Indian population with acquired aplastic anemia. Leukemia Research, 2015, 39, 1263-1269.	0.8	2