Shwetha Chiplunkar

List of Publications by Year
in descending order

13	160	1163117 8	1281871 11
papers	citations	h-index	g-index

13	13	13	265
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Magnetic resonance imaging correlates of genetically characterized patients with mitochondrial disorders: A study from south India. Mitochondrion, 2015, 25, 6-16.	3.4	28
2	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
3	Audiological manifestations in mitochondrial encephalomyopathy lactic acidosis and stroke like episodes (MELAS) syndrome. Clinical Neurology and Neurosurgery, 2016, 148, 17-21.	1.4	18
4	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
5	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
6	Child Neurology: SjÃ¶gren-Larsson syndrome. Neurology, 2017, 88, e1-e4.	1.1	16
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	Peripheral neuropathy in genetically characterized patients with mitochondrial disorders: A study from south India. Mitochondrion, 2016, 27, 1-5.	3.4	10
9	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
10	Clinico-pathological and Molecular Spectrum of Mitochondrial Polymerase Î³ Mutations in a Cohort from India. Journal of Molecular Neuroscience, 2021, 71, 2219-2228.	2.3	6
11	Clinical Reasoning: West syndrome, pontocerebellar hypoplasia, and hypomyelination in a 6-month-old boy. Neurology, 2018, 91, e1652-e1656.	1.1	1
12	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
13	Infantile Onset Encephalomyopathy, Heart Block, and Sensorineural Hearing Loss: RMND1-Associated Mitochondrial Disease. Journal of Pediatric Neurology, 2021, 19, 183-188.	0.2	0