

# Shwetha Chiplunkar

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/11271668/publications.pdf>

Version: 2024-02-01

13  
papers

160  
citations

1163117

8  
h-index

1281871

11  
g-index

13  
all docs

13  
docs citations

13  
times ranked

265  
citing authors

#	ARTICLE	IF	CITATIONS
1	Infantile Onset Encephalomyopathy, Heart Block, and Sensorineural Hearing Loss: RMND1-Associated Mitochondrial Disease. <i>Journal of Pediatric Neurology</i> , 2021, 19, 183-188.	0.2	0
2	Serum fibroblast growth factor 21 and growth differentiation factor 15: Two sensitive biomarkers in the diagnosis of mitochondrial disorders. <i>Mitochondrion</i> , 2021, 60, 170-177.	3.4	9
3	Clinico-pathological and Molecular Spectrum of Mitochondrial Polymerase $\hat{I}^3$ Mutations in a Cohort from India. <i>Journal of Molecular Neuroscience</i> , 2021, 71, 2219-2228.	2.3	6
4	Leukodystrophies and Genetic Leukoencephalopathies in Children Specified by Exome Sequencing in an Expanded Gene Panel. <i>Journal of Child Neurology</i> , 2020, 35, 433-441.	1.4	11
5	Mitochondrial leukoencephalopathies: A border zone between acquired and inherited white matter disorders in children?. <i>Multiple Sclerosis and Related Disorders</i> , 2018, 20, 84-92.	2.0	27
6	Outcome of epilepsy in patients with mitochondrial disorders: Phenotype genotype and magnetic resonance imaging correlations. <i>Clinical Neurology and Neurosurgery</i> , 2018, 164, 182-189.	1.4	17
7	Clinical Reasoning: West syndrome, pontocerebellar hypoplasia, and hypomyelination in a 6-month-old boy. <i>Neurology</i> , 2018, 91, e1652-e1656.	1.1	1
8	Child Neurology: Sjögren-Larsson syndrome. <i>Neurology</i> , 2017, 88, e1-e4.	1.1	16
9	Mitochondrial oxidative phosphorylation disorders in children: Phenotypic, genotypic and biochemical correlations in 85 patients from South India. <i>Mitochondrion</i> , 2017, 32, 42-49.	3.4	17
10	Reply to Letter to the Editor: Hearing impairment in m.3243A>G carriers requires comprehensive work- and follow-up. <i>Clinical Neurology and Neurosurgery</i> , 2016, 150, 198-199.	1.4	0
11	Audiological manifestations in mitochondrial encephalomyopathy lactic acidosis and stroke like episodes (MELAS) syndrome. <i>Clinical Neurology and Neurosurgery</i> , 2016, 148, 17-21.	1.4	18
12	Peripheral neuropathy in genetically characterized patients with mitochondrial disorders: A study from south India. <i>Mitochondrion</i> , 2016, 27, 1-5.	3.4	10
13	Magnetic resonance imaging correlates of genetically characterized patients with mitochondrial disorders: A study from south India. <i>Mitochondrion</i> , 2015, 25, 6-16.	3.4	28