

# Chitra Prasad

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/482137/publications.pdf>

Version: 2024-02-01

8  
papers

124  
citations

1937685

4  
h-index

1872680

6  
g-index

8  
all docs

8  
docs citations

8  
times ranked

485  
citing authors

#	ARTICLE	IF	CITATIONS
1	Expanding the phenotypic and molecular spectrum of <i>NFS1</i> -related disorders that cause functional deficiencies in mitochondrial and cytosolic iron-sulfur cluster containing enzymes. <i>Human Mutation</i> , 2022, 43, 305-315.	2.5	1
2	Bulging anterior fontanelle and dense bones in an infant. <i>Paediatrics and Child Health</i> , 2020, 25, 69-71.	0.6	0
3	Further delineation of TBCK - Infantile hypotonia with psychomotor retardation and characteristic facies type 3. <i>European Journal of Medical Genetics</i> , 2019, 62, 273-277.	1.3	11
4	Magnetic resonance imaging in the diagnosis of white matter signal abnormalities. <i>Neuroradiology Journal</i> , 2018, 31, 362-371.	1.2	4
5	Nephrological and urological complications of homozygous c.974G>A (p.Arg325Gln) OSGEP mutations. <i>Pediatric Nephrology</i> , 2018, 33, 2201-2204.	1.7	9
6	Molecular characterization of NRXN1 deletions from 19,263 clinical microarray cases identifies exons important for neurodevelopmental disease expression. <i>Genetics in Medicine</i> , 2017, 19, 53-61.	2.4	70
7	Seizures Related to Hypomagnesemia. <i>Child Neurology Open</i> , 2016, 3, 2329048X1667483.	1.1	29
8	MG-123...Exonic and intronic NRXN1 deletions: Novel genotype-phenotype correlations. <i>Journal of Medical Genetics</i> , 2015, 52, A9.1-A9.	3.2	0