Chitra Prasad

List of Publications by Year
in descending order

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