

Noriko Sangu

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1693554/publications.pdf>

Version: 2024-02-01

8
papers

100
citations

1684188
5
h-index

1588992
8
g-index

8
all docs

8
docs citations

8
times ranked

359
citing authors

#	ARTICLE	IF	CITATIONS
1	Genial Tubercle Fracture. Journal of Craniofacial Surgery, 2019, 30, 161-162.	0.7	5
2	Single Nucleotide Variations in CLCN6 Identified in Patients with Benign Partial Epilepsies in Infancy and/or Febrile Seizures. PLoS ONE, 2015, 10, e0118946.	2.5	13
3	Microarray analysis of 50 patients reveals the critical chromosomal regions responsible for 1p36 deletion syndrome-related complications. Brain and Development, 2015, 37, 515-526.	1.1	43
4	Mutations in the genes encoding eukaryotic translation initiation factor 2B in Japanese patients with vanishing white matter disease. Brain and Development, 2015, 37, 960-966.	1.1	10
5	Characteristics of patients with benign partial epilepsy in infancy without PRRT2 mutations. Epilepsy Research, 2015, 118, 10-13.	1.6	8
6	Growth patterns of patients with 1p36 deletion syndrome. Congenital Anomalies (discontinued), 2014, 54, 82-86.	0.6	4
7	Overlapping microdeletions involving 15q22.2 narrow the critical region for intellectual disability to NARG2 and RORA. European Journal of Medical Genetics, 2014, 57, 163-168.	1.3	11
8	Novel nucleotide mutation leading to a recurrent amino acid alteration in <i>SH3BP2</i> in a patient with cherubism. Congenital Anomalies (discontinued), 2013, 53, 166-169.	0.6	6