# Noriko Sangu

List of Publications by Year
in descending order

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

Version: 2024-02-01

| | | | |
|---|---|---|---|
| 8<br>papers | 100<br>citations | 1684188<br>5<br>h-index | 1588992<br>8<br>g-index |
| 8<br>all docs | 8<br>docs citations | 8<br>times ranked | 359<br>citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | 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 |
| 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 | 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 |
| 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 | Novel nucleotide mutation leading to a recurrent amino acid alteration in <scp>SH3BP2</scp> in a patient with cherubism. Congenital Anomalies (discontinued), 2013, 53, 166-169. | 0.6 | 6 |
| 7 | Genial Tubercle Fracture. Journal of Craniofacial Surgery, 2019, 30, 161-162. | 0.7 | 5 |
| 8 | Growth patterns of patients with 1p36 deletion syndrome. Congenital Anomalies (discontinued), 2014, 54, 82-86. | 0.6 | 4 |