

Tommy Ståldberg

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7775735/publications.pdf>

Version: 2024-02-01

7
papers

411
citations

1478505

6
h-index

1720034

7
g-index

7
all docs

7
docs citations

7
times ranked

975
citing authors

#	ARTICLE	IF	CITATIONS
1	Outcome at age 7 of epilepsy presenting in the first 2% years of life. A population-based study. <i>Epilepsia</i> , 2022, 63, 2096-2107.	5.1	5
2	Integration of whole genome sequencing into a healthcare setting: high diagnostic rates across multiple clinical entities in 3219 rare disease patients. <i>Genome Medicine</i> , 2021, 13, 40.	8.2	116
3	<i>SLC12A2</i> mutations cause NKCC1 deficiency with encephalopathy and impaired secretory epithelia. <i>Neurology: Genetics</i> , 2020, 6, e478.	1.9	20
4	Epilepsy syndromes, etiologies, and the use of next-generation sequencing in epilepsy presenting in the first 2 years of life: A population-based study. <i>Epilepsia</i> , 2020, 61, 2486-2499.	5.1	24
5	Genetic, Phenotypic, and Interferon Biomarker Status in ADAR1-Related Neurological Disease. <i>Neuropediatrics</i> , 2017, 48, 166-184.	0.6	62
6	An LC-MS/MS-Based Method for the Quantification of Pyridox(am)ine 5-Phosphate Oxidase Activity in Dried Blood Spots from Patients with Epilepsy. <i>Analytical Chemistry</i> , 2017, 89, 8892-8900.	6.5	24
7	Mutations in <i>SLC12A5</i> in epilepsy of infancy with migrating focal seizures. <i>Nature Communications</i> , 2015, 6, 8038.	12.8	160