Ichraf Kraoua

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/929583/publications.pdf

Version: 2024-02-01

10 papers	29 citations	2258059 3 h-index	2053705 5 g-index

11 all docs	11 docs citations	11 times ranked	79 citing authors

#	Article	IF	Citations
1	Neuronal ceroÃ¯dâ€lipofuscinosis: Clinical, electroencephalographic, imaging, and genetic study of a maghrebian series. Clinical Genetics, 2022, 102, 157-160.	2.0	0
2	Hypomyelination and Congenital Cataract: Clinical, Imaging, and Genetic Findings in Three Tunisian Families and Literature Review. Neuropediatrics, 2021, 52, 302-309.	0.6	0
3	Alpha-mannosidosis in Tunisian consanguineous families: Potential involvement of variants in GHR and SLC19A3 genes in the variable expressivity of cognitive impairment. PLoS ONE, 2021, 16, e0258202.	2.5	2
4	SQSTM1 mutation: Description of the first Tunisian case and literature review. Molecular Genetics & Genomic Medicine, 2020, 8, e1543.	1.2	4
5	Novel POLR1C mutation in RNA polymerase IIIâ€related leukodystrophy with severe myoclonus and dystonia. Molecular Genetics & Genomic Medicine, 2019, 7, e914.	1.2	8
6	Pyridoxine-dependent epilepsy: A novel mutation in a Tunisian child. Archives De Pediatrie, 2017, 24, 241-243.	1.0	6
7	Letter to the editor: â€œA case of Guillain-BarrÃ© syndrome with meningeal irritationâ€. Brain and Development, 2017, 39, 815.	1.1	0
8	Childhood opsoclonusâ€“myoclonus syndrome: A case series from Tunisia. Brain and Development, 2017, 39, 751-755.	1.1	5
9	Elevated aspartate aminotransferase and lactate dehydrogenase levels are a constant finding in <scp>PLA</scp>2G6â€associated neurodegeneration. European Journal of Neurology, 2016, 23, e24-5.	3.3	2
10	Lesch Nyhan syndrome: A novel complex mutation in a Tunisian child. Brain and Development, 2014, 36, 921-923.	1.1	2