Guida LandourÃ©

List of Publications by Year
in descending order

10 papers	154 citations	1478505 6 h-index	1474206 9 g-index
10 all docs	10 docs citations	10 times ranked	269 citing authors

#	Article	IF	Citations
1	<scp> GJB1 </scp> variants in <scp>Charcotâ€Marieâ€Tooth</scp> disease Xâ€linked type 1 in Mali. Journal of the Peripheral Nervous System, 2022, , .	3.1	2
2	Increasing African genomic data generation and sharing to resolve rare and undiagnosed diseases in Africa: a call-to-action by the H3Africa rare diseases working group. Orphanet Journal of Rare Diseases, 2022, 17, .	2.7	20
3	Friedreich ataxia in a family from Mali, West Africa/Friedreich ataxia in a Malian family. Clinical Case Reports (discontinued), 2021, 9, e04065.	0.5	0
4	A novel variant in the spatacsin gene causing SPG11 in a Malian family. Journal of the Neurological Sciences, 2020, 411, 116675.	0.6	4
5	Fahr's syndrome with hyperparathyroidism revealed by seizures and proximal weakness. ENeurologicalSci, 2019, 15, 100192.	1.3	4
6	A novel mutation in the GARS gene in a Malian family with Charcotâ€Marieâ€Tooth disease. Molecular Genetics & Genomic Medicine, 2019, 7, e00782.	1.2	12
7	Hereditary spastic paraplegia type 35 in a family from Mali. American Journal of Medical Genetics, Part A, 2019, 179, 1122-1125.	1.2	9
8	A novel mutation in KIF5A in a Malian family with spastic paraplegia and sensory loss. Annals of Clinical and Translational Neurology, 2017, 4, 272-275.	3.7	16
9	Genetics and genomic medicine in Mali: challenges and future perspectives. Molecular Genetics & Genomic Medicine, 2016, 4, 126-134.	1.2	8
10	Hereditary Spastic Paraplegia Type 43 (SPG43) is Caused by Mutation inC19orf12. Human Mutation, 2013, 34, 1357-1360.	2.5	79