Tiffany Grider

List of Publications by Year
in descending order

11 papers	307 citations	1040056 9 h-index	1281871 11 g-index
11 all docs	11 docs citations	11 times ranked	491 citing authors

#	Article	IF	Citations
1	Genotypeâ€“phenotype characteristics and baseline natural history of heritable neuropathies caused by mutations in theMPZgene. Brain, 2015, 138, 3180-3192.	7.6	80
2	Psychometrics evaluation of Charcotâ€Marieâ€Tooth Neuropathy Score (<scp>CMTNSv2</scp>) second version, using Rasch analysis. Journal of the Peripheral Nervous System, 2014, 19, 192-196.	3.1	59
3	Validation of MRC Centre MRI calf muscle fat fraction protocol as an outcome measure in CMT1A. Neurology, 2018, 91, e1125-e1129.	1.1	43
4	A longitudinal study of CMT1A using Rasch analysis based CMT neuropathy and examination scores. Neurology, 2020, 94, e884-e896.	1.1	29
5	Transmembrane protease serine 5: a novel Schwann cell plasma marker for CMT1A. Annals of Clinical and Translational Neurology, 2020, 7, 69-82.	3.7	25
6	Reduced neurofilament expression in cutaneous nerve fibers of patients with CMT2E. Neurology, 2015, 85, 228-234.	1.1	21
7	Rare Manifestation of a c.290 C>T, p.Gly97GluVCPMutation. Case Reports in Genetics, 2015, 2015, 1-5.	0.2	16
8	MicroRNAs as Biomarkers of Charcot-Marie-Tooth Disease Type 1A. Neurology, 2021, 97, e489-e500.	1.1	14
9	The audiologic profile of patients with Charcot-Marie Tooth neuropathy can be characterised by both cochlear and neural deficits. International Journal of Audiology, 2019, 58, 902-912.	1.7	12
10	Coexistence of a T118M PMP22 missense mutation and chromosome 17 (17p11.2â€p12) deletion. Muscle and Nerve, 2015, 52, 905-908.	2.2	4
11	Progressive Lower Extremity Weakness and Axonal Sensorimotor Polyneuropathy from a Mutation inKIF5A(c.611G>A;p.Arg204Gln). Case Reports in Genetics, 2015, 2015, 1-5.	0.2	4