Micaela Milani

List of Publications by Year
in descending order

18	369	933447 10	940533 16
papers	citations	h-index	g-index

18	18	18	647
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Deciphering the Labyrinthine System of the Immune Microenvironment in Recurrent Glioblastoma: Recent Original Advances and Lessons from Clinical Immunotherapeutic Approaches. Cancers, 2021, 13, 6156.	3.7	8
2	Expansion of effector and memory T cells is associated with increased survival in recurrent glioblastomas treated with dendritic cell immunotherapy. Neuro-Oncology Advances, 2019, 1, vdz022.	0.7	16
3	A novel <scp>NDRG1</scp> mutation in a nonâ€Romani patient with <scp>CMT4D</scp>/<scp>HMSN</scp>â€Lom. Journal of the Peripheral Nervous System, 2017, 22, 47-50.	3.1	6
4	Mutational mechanisms in <scp>MFN2</scp>â€related neuropathy: compound heterozygosity for recessive and semidominant mutations. Journal of the Peripheral Nervous System, 2015, 20, 380-386.	3.1	23
5	Xâ€linked Charcotâ€Marieâ€Tooth type 1: strokeâ€like presentation of a novel <scp>GJB1</scp> mutation. Journal of the Peripheral Nervous System, 2014, 19, 183-186.	3.1	20
6	Doubleâ€trouble in pediatric neurology: Myotonia congenita combined with charcotâ€“marieâ€“tooth disease type 1a. Muscle and Nerve, 2014, 50, 145-147.	2.2	6
7	Clinical, electrophysiological and pathological findings in a patient with Charcotâ€“Marieâ€“Tooth disease 4D caused by the NDRG1 Lom mutation. Journal of the Neurological Sciences, 2014, 345, 271-273.	0.6	6
8	Myelin protein zero <scp>Arg36Gly</scp> mutation with very late onset and rapidly progressive painful neuropathy. Journal of the Peripheral Nervous System, 2012, 17, 422-425.	3.1	9
9	Co-occurrence of amyotrophic lateral sclerosis and Charcot-Marie-Tooth disease type 2A in a patient with a novel mutation in the mitofusin-2 gene. Neuromuscular Disorders, 2011, 21, 129-131.	0.6	19
10	Recessive axonal Charcot-Marie-Tooth disease due to compound heterozygous mitofusin 2 mutations. Neurology, 2011, 77, 168-173.	1.1	72
11	Four novel cases of periaxin-related neuropathy and review of the literature. Neurology, 2010, 75, 1830-1838.	1.1	47
12	P60 Variable severity of early onset CMT2 with compound heterozygous MFN2 mutations. Neuromuscular Disorders, 2010, 20, S21.	0.6	0
13	Novel mutations in the GDAP1 gene in patients affected with early-onset axonal Charcot-Marie-Tooth type 4A. Neuromuscular Disorders, 2009, 19, 476-480.	0.6	23
14	A novel founder mutation in the MFN2 gene associated with variable Charcot-Marie-Tooth type 2 phenotype in two families from Southern Italy. BMJ Case Reports, 2009, 2009, bcr0820080652-bcr0820080652.	0.5	0
15	A novel founder mutation in the MFN2 gene associated with variable Charcot Marie Tooth type 2 phenotype in two families from Southern Italy. Journal of Neurology, Neurosurgery and Psychiatry, 2007, 78, 1286-1287.	1.9	10
16	Rapid progression of late onset axonal Charcot Marie Tooth disease associated with a novel MPZ mutation in the extracellular domain. Journal of Neurology, Neurosurgery and Psychiatry, 2007, 78, 1263-1266.	1.9	29
17	Does CMT1A homozygosity cause more severe disease with root hypertrophy and higher CSF proteins?. Neurology, 2003, 60, 1721-1722.	1.1	21
18	Charcot-Marie-Tooth disease type I and related demyelinating neuropathies: Mutation analysis in a large cohort of Italian families. Human Mutation, 2001, 18, 32-41.	2.5	54