Sevim Erdem-Ozdamar

List of Publications by Year
in descending order

26	224	1163117 8	1058476 14
papers	citations	h-index	g-index

26	26	26	432
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A novel desmin mutation leading to autosomal recessive limb-girdle muscular dystrophy: distinct histopathological outcomes compared with desminopathies. Journal of Medical Genetics, 2013, 50, 437-443.	3.2	63
2	Transcript levels of plastin 3 and neuritin 1 modifier genes in spinal muscular atrophy siblings. Pediatrics International, 2017, 59, 53-56.	0.5	17
3	The Course of Myasthenia Gravis with Systemic Lupus Erythematosus. European Neurology, 2014, 72, 326-329.	1.4	15
4	The altered expression of perineuronal net elements during neural differentiation. Cellular and Molecular Biology Letters, 2018, 23, 5.	7.0	14
5	The histopathological evaluation of small fiber neuropathy in patients with vitamin B12 deficiency. Acta Neurologica Belgica, 2018, 118, 405-410.	1.1	14
6	Fulminant Central Plus Peripheral Nervous System Demyelination without Antibodies to Neurofascin. Canadian Journal of Neurological Sciences, 2016, 43, 149-156.	0.5	11
7	Myophosphorylase (PYGM) mutations determined by next generation sequencing in a cohort from Turkey with McArdle disease. Neuromuscular Disorders, 2017, 27, 997-1008.	0.6	11
8	Three Turkish families with different transthyretin mutations. Neuromuscular Disorders, 2015, 25, 686-692.	0.6	10
9	New mutations and genotypeâ€“phenotype correlation in late-onset Pompe patients. Acta Neurologica Belgica, 2017, 117, 269-275.	1.1	8
10	The functional and structural evaluation of small fibers in asymptomatic carriers of TTR p.Val50Met (Val30Met) mutation. Neuromuscular Disorders, 2022, 32, 50-56.	0.6	8
11	Remission with fingolimod in a case of demyelinating polyneuropathy. Muscle and Nerve, 2014, 50, 615-617.	2.2	7
12	A database for screening and registering late onset Pompe disease in Turkey. Neuromuscular Disorders, 2018, 28, 262-267.	0.6	6
13	Do Perineuronal Net Elements Contribute to Pathophysiology of Spinal Muscular Atrophy? In Vitro and Transcriptomics Insights. OMICS A Journal of Integrative Biology, 2018, 22, 598-606.	2.0	6
14	Toscana virus associated with Guillainâ€“BarrÃ© syndrome: a caseâ€“control study. Acta Neurologica Belgica, 2021, 121, 661-668.	1.1	6
15	Spinal muscular atrophy type III: Molecular genetic characterization of Turkish patients. European Journal of Medical Genetics, 2015, 58, 654-658.	1.3	5
16	Overexpression of amyloid beta precursor protein enhances expression and secretion of ST6Gal1 in C2C12 myogenic cell line. Cell Biology International, 2011, 35, 9-13.	3.0	4
17	Comprehensive clinical, biochemical, radiological and genetic analysis of 28 Turkish cases with suspected metachromatic leukodystrophy and their relatives. Molecular Genetics and Metabolism Reports, 2020, 25, 100688.	1.1	4
18	Diabetic muscular infarct: an unusual cause of extremity pain and dysfunction. Rheumatology International, 2012, 32, 525-528.	3.0	3

#	Article	IF	Citations
19	Ocular surface alterations and in vivo confocal microscopic characteristics of corneas in patients with myasthenia gravis. European Journal of Ophthalmology, 2018, 28, 541-546.	1.3	3
20	Two distinct skeletal muscle microRNA signatures revealing the complex mechanism of sporadic ALS. Acta Neurologica Belgica, 2022, 122, 1499-1509.	1.1	2
21	One Disease with two Faces: Semidominant Inheritance of a Novel HTRA1 Mutation in a Consanguineous Family. Journal of Stroke and Cerebrovascular Diseases, 2021, 30, 105997.	1.6	2
22	The Activation of RAGE and NF-ÎšB in Nerve Biopsies of Patients with Axonal and Vasculitic Neuropathy. Noropsikiyatri Arsivi, 2015, 52, 279-282.	0.7	2
23	The effects of spinal stabilization exercises in patients with myasthenia gravis: a randomized crossover study. Disability and Rehabilitation, 2022, 44, 8442-8449.	1.8	2
24	Neuropathic Pain Frequency in Neurology Outpatients: A Multicenter Study. Noropsikiyatri Arsivi, 2021, 58, 257-260.	0.3	1
25	Proinflammatory effect of AbetaPP induced ST6GAL1 secretion from C2C12 myogenic cell line. Turkish Journal of Biochemistry, 2015, 40, 31-36.	0.5	0
26	Comprehensive evaluation of velopharyngeal function in myasthenia gravis patients. Acta Neurologica Belgica, 2022, , 1.	1.1	0