Sevim Erdem-Ozdamar

List of Publications by Year
in descending order

26	224	1162367 8	1058022 14
papers	citations	h-index	g-index

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

#	Article	IF	Citations
1	Two distinct skeletal muscle microRNA signatures revealing the complex mechanism of sporadic ALS. Acta Neurologica Belgica, 2022, 122, 1499-1509.	0.5	2
2	The functional and structural evaluation of small fibers in asymptomatic carriers of TTR p.Val50Met (Val30Met) mutation. Neuromuscular Disorders, 2022, 32, 50-56.	0.3	8
3	The effects of spinal stabilization exercises in patients with myasthenia gravis: a randomized crossover study. Disability and Rehabilitation, 2022, 44, 8442-8449.	0.9	2
4	Comprehensive evaluation of velopharyngeal function in myasthenia gravis patients. Acta Neurologica Belgica, 2022, , 1.	0.5	0
5	Toscana virus associated with Guillainâ€“BarrÃ© syndrome: a caseâ€“control study. Acta Neurologica Belgica, 2021, 121, 661-668.	0.5	6
6	Neuropathic Pain Frequency in Neurology Outpatients: A Multicenter Study. Noropsikiyatri Arsivi, 2021, 58, 257-260.	0.2	1
7	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.	0.7	2
8	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.	0.4	4
9	A database for screening and registering late onset Pompe disease in Turkey. Neuromuscular Disorders, 2018, 28, 262-267.	0.3	6
10	The altered expression of perineuronal net elements during neural differentiation. Cellular and Molecular Biology Letters, 2018, 23, 5.	2.7	14
11	Ocular surface alterations and in vivo confocal microscopic characteristics of corneas in patients with myasthenia gravis. European Journal of Ophthalmology, 2018, 28, 541-546.	0.7	3
12	The histopathological evaluation of small fiber neuropathy in patients with vitamin B12 deficiency. Acta Neurologica Belgica, 2018, 118, 405-410.	0.5	14
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.	1.0	6
14	Transcript levels of plastin 3 and neuritin 1 modifier genes in spinal muscular atrophy siblings. Pediatrics International, 2017, 59, 53-56.	0.2	17
15	Myophosphorylase (PYGM) mutations determined by next generation sequencing in a cohort from Turkey with McArdle disease. Neuromuscular Disorders, 2017, 27, 997-1008.	0.3	11
16	New mutations and genotypeâ€“phenotype correlation in late-onset Pompe patients. Acta Neurologica Belgica, 2017, 117, 269-275.	0.5	8
17	Fulminant Central Plus Peripheral Nervous System Demyelination without Antibodies to Neurofascin. Canadian Journal of Neurological Sciences, 2016, 43, 149-156.	0.3	11
18	Proinflammatory effect of AbetaPP induced ST6GAL1 secretion from C2C12 myogenic cell line. Turkish Journal of Biochemistry, 2015, 40, 31-36.	0.3	0

#	Article	IF	Citations
19	Three Turkish families with different transthyretin mutations. Neuromuscular Disorders, 2015, 25, 686-692.	0.3	10
20	Spinal muscular atrophy type III: Molecular genetic characterization of Turkish patients. European Journal of Medical Genetics, 2015, 58, 654-658.	0.7	5
21	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
22	The Course of Myasthenia Gravis with Systemic Lupus Erythematosus. European Neurology, 2014, 72, 326-329.	0.6	15
23	Remission with fingolimod in a case of demyelinating polyneuropathy. Muscle and Nerve, 2014, 50, 615-617.	1.0	7
24	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.	1.5	63
25	Diabetic muscular infarct: an unusual cause of extremity pain and dysfunction. Rheumatology International, 2012, 32, 525-528.	1.5	3
26	Overexpression of amyloid beta precursor protein enhances expression and secretion of ST6Gal1 in C2C12 myogenic cell line. Cell Biology International, 2011, 35, 9-13.	1.4	4