Ersin Tan

List of Publications by Year
in descending order

18	405	1163117 8	839539 18
papers	citations	h-index	g-index

18	18	18	696
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The spectrum of mutations causing end-plate acetylcholinesterase deficiency. Annals of Neurology, 2000, 47, 162-170.	5.3	123
2	A biochemical, genetic, and clinical survey of autosomal recessive limb girdle muscular dystrophies in Turkey. Annals of Neurology, 1997, 42, 222-229.	5.3	94
3	The distinct genetic pattern of ALS in Turkey and novel mutations. Neurobiology of Aging, 2015, 36, 1764.e9-1764.e18.	3.1	78
4	Myasthenic Syndrome Caused by Hydroxychloroquine Used for COVID-19 Prophylaxis. Journal of Clinical Neuromuscular Disease, 2020, 22, 60-62.	0.7	19
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	Paraneoplastic striatal encephalitis and myelitis associated with anti-CV2/CRMP-5 antibodies in a patient with small cell lung cancer. Clinical Neurology and Neurosurgery, 2018, 170, 117-119.	1.4	12
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	The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice. Movement Disorders, 2021, 36, 1676-1688.	3.9	9
10	New mutations and genotypeâ€“phenotype correlation in late-onset Pompe patients. Acta Neurologica Belgica, 2017, 117, 269-275.	1.1	8
11	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
12	A case with Neurofascin-155 IgG antibody-associated Combined Central and Peripheral Demyelination: Succesfully treated with anti-CD20 monoclonal antibody. Clinical Neurology and Neurosurgery, 2021, 210, 106961.	1.4	5
13	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
14	Establishment of primary myoblast cell cultures from cryopreserved skeletal muscle biopsies to serve as a tool in related research & development studies. Journal of the Neurological Sciences, 2018, 393, 100-104.	0.6	3
15	Cognitive impairment and affective disorder: A rare presentation of cerebellar stroke. Clinical Neurology and Neurosurgery, 2021, 206, 106690.	1.4	3
16	Two distinct skeletal muscle microRNA signatures revealing the complex mechanism of sporadic ALS. Acta Neurologica Belgica, 2022, 122, 1499-1509.	1.1	2
17	A combined clinical and computational approach to understand the SOD1A4T-mediated pathogenesis of rapidly progressive familial amyotrophic lateral sclerosis. Acta Neurologica Belgica, 2021, , 1.	1.1	1
18	Neuropathic Pain Frequency in Neurology Outpatients: A Multicenter Study. Noropsikiyatri Arsivi, 2021, 58, 257-260.	0.3	1