

Ersin Tan

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1637820/publications.pdf>

Version: 2024-02-01

18
papers

405
citations

1163117

8
h-index

839539

18
g-index

18
all docs

18
docs citations

18
times ranked

696
citing authors

#	ARTICLE	IF	CITATIONS
1	Two distinct skeletal muscle microRNA signatures revealing the complex mechanism of sporadic ALS. <i>Acta Neurologica Belgica</i> , 2022, 122, 1499-1509.	1.1	2
2	The functional and structural evaluation of small fibers in asymptomatic carriers of TTR p.Val50Met (Val30Met) mutation. <i>Neuromuscular Disorders</i> , 2022, 32, 50-56.	0.6	8
3	A combined clinical and computational approach to understand the SOD1A4T-mediated pathogenesis of rapidly progressive familial amyotrophic lateral sclerosis. <i>Acta Neurologica Belgica</i> , 2021, , 1.	1.1	1
4	Neuropathic Pain Frequency in Neurology Outpatients: A Multicenter Study. <i>Noropsikiyatri Arsivi</i> , 2021, 58, 257-260.	0.3	1
5	The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice. <i>Movement Disorders</i> , 2021, 36, 1676-1688.	3.9	9
6	Cognitive impairment and affective disorder: A rare presentation of cerebellar stroke. <i>Clinical Neurology and Neurosurgery</i> , 2021, 206, 106690.	1.4	3
7	A case with Neurofascin-155 IgG antibody-associated Combined Central and Peripheral Demyelination: Successfully treated with anti-CD20 monoclonal antibody. <i>Clinical Neurology and Neurosurgery</i> , 2021, 210, 106961.	1.4	5
8	Myasthenic Syndrome Caused by Hydroxychloroquine Used for COVID-19 Prophylaxis. <i>Journal of Clinical Neuromuscular Disease</i> , 2020, 22, 60-62.	0.7	19
9	Comprehensive clinical, biochemical, radiological and genetic analysis of 28 Turkish cases with suspected metachromatic leukodystrophy and their relatives. <i>Molecular Genetics and Metabolism Reports</i> , 2020, 25, 100688.	1.1	4
10	The histopathological evaluation of small fiber neuropathy in patients with vitamin B12 deficiency. <i>Acta Neurologica Belgica</i> , 2018, 118, 405-410.	1.1	14
11	Establishment of primary myoblast cell cultures from cryopreserved skeletal muscle biopsies to serve as a tool in related research & development studies. <i>Journal of the Neurological Sciences</i> , 2018, 393, 100-104.	0.6	3
12	Paraneoplastic striatal encephalitis and myelitis associated with anti-CV2/CRMP-5 antibodies in a patient with small cell lung cancer. <i>Clinical Neurology and Neurosurgery</i> , 2018, 170, 117-119.	1.4	12
13	Myophosphorylase (PYGM) mutations determined by next generation sequencing in a cohort from Turkey with McArdle disease. <i>Neuromuscular Disorders</i> , 2017, 27, 997-1008.	0.6	11
14	New mutations and genotypeâ€“phenotype correlation in late-onset Pompe patients. <i>Acta Neurologica Belgica</i> , 2017, 117, 269-275.	1.1	8
15	The distinct genetic pattern of ALS in Turkey and novel mutations. <i>Neurobiology of Aging</i> , 2015, 36, 1764.e9-1764.e18.	3.1	78
16	Three Turkish families with different transthyretin mutations. <i>Neuromuscular Disorders</i> , 2015, 25, 686-692.	0.6	10
17	The spectrum of mutations causing end-plate acetylcholinesterase deficiency. <i>Annals of Neurology</i> , 2000, 47, 162-170.	5.3	123
18	A biochemical, genetic, and clinical survey of autosomal recessive limb girdle muscular dystrophies in Turkey. <i>Annals of Neurology</i> , 1997, 42, 222-229.	5.3	94