Tahseen Mozaffar

List of Publications by Year in descending order

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Version: 2024-02-01

62 papers 4,380 citations

147801 31 h-index 62 g-index

65 all docs

65 does citations

65 times ranked 4675 citing authors

#	Article	IF	Citations
1	Patient reported quality of life in limb girdle muscular dystrophy. Neuromuscular Disorders, 2022, 32, 57-64.	0.6	3
2	Loss of TDP-43 function and rimmed vacuoles persist after T cell depletion in a xenograft model of sporadic inclusion body myositis. Science Translational Medicine, 2022, 14, eabi9196.	12.4	27
3	Development of a standard of care for patients with valosin-containing protein associated multisystem proteinopathy. Orphanet Journal of Rare Diseases, 2022, 17, 23.	2.7	19
4	Multisystem proteinopathy: Where myopathy and motor neuron disease converge. Muscle and Nerve, 2021, 63, 442-454.	2.2	33
5	Clinical utility of anti ytosolic 5'â€nucleotidase 1A antibody in idiopathic inflammatory myopathies. Annals of Clinical and Translational Neurology, 2021, 8, 571-578.	3.7	18
6	Results from a 3-year Non-interventional, Observational Disease Monitoring Program in Adults with GNE Myopathy. Journal of Neuromuscular Diseases, 2021, 8, 225-234.	2.6	9
7	A stromal progenitor and ILC2 niche promotes muscle eosinophilia and fibrosis-associated gene expression. Cell Reports, 2021, 35, 108997.	6.4	28
8	Safety, efficacy, and tolerability of efgartigimod in patients with generalised myasthenia gravis (ADAPT): a multicentre, randomised, placebo-controlled, phase 3 trial. Lancet Neurology, The, 2021, 20, 526-536.	10.2	194
9	IgG regulation through FcRn blocking: A novel mechanism for the treatment of myasthenia gravis. Journal of the Neurological Sciences, 2021, 430, 118074.	0.6	24
10	Safety and efficacy of cipaglucosidase alfa plus miglustat versus alglucosidase alfa plus placebo in late-onset Pompe disease (PROPEL): an international, randomised, double-blind, parallel-group, phase 3 trial. Lancet Neurology, The, 2021, 20, 1027-1037.	10.2	42
11	Safety and efficacy of avalglucosidase alfa versus alglucosidase alfa in patients with late-onset Pompe disease (COMET): a phase 3, randomised, multicentre trial. Lancet Neurology, The, 2021, 20, 1012-1026.	10.2	59
12	Examination of the human motor endplate after brachial plexus injury with twoâ€photon microscopy. Muscle and Nerve, 2020, 61, 390-395.	2.2	6
13	Update on immuneâ€mediated therapies for myasthenia gravis. Muscle and Nerve, 2020, 62, 579-592.	2.2	9
14	Minimal manifestation status and prednisone withdrawal in the MGTX trial. Neurology, 2020, 95, e755-e766.	1.1	17
15	Clinical Effects of the Self-administered Subcutaneous Complement Inhibitor Zilucoplan in Patients With Moderate to Severe Generalized Myasthenia Gravis. JAMA Neurology, 2020, 77, 582.	9.0	126
16	Plasma creatinine and oxidative stress biomarkers in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 263-272.	1.7	20
17	Limbâ€girdle muscular dystrophy: A perspective from adult patients on what matters most. Muscle and Nerve, 2019, 60, 419-424.	2.2	15
18	A phase 3 randomized study evaluating sialic acid extended-release for GNE myopathy. Neurology, 2019, 92, e2109-e2117.	1.1	40

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19	Long-term effect of thymectomy plus prednisone versus prednisone alone in patients with non-thymomatous myasthenia gravis: 2-year extension of the MGTX randomised trial. Lancet Neurology, The, 2019, 18, 259-268.	10.2	139
20	Randomized phase 2 study of FcRn antagonist efgartigimod in generalized myasthenia gravis. Neurology, 2019, 92, e2661-e2673.	1.1	169
21	First-in-human study of advanced and targeted acid $\hat{l}\pm$ -glucosidase (AT-GAA) (ATB200/AT2221) in patients with Pompe disease: preliminary functional assessment results from the ATB200-02 trial. Molecular Genetics and Metabolism, 2019, 126, S86.	1.1	4
22	237th ENMC International Workshop: GNE myopathy \hat{a} €" current and future research Hoofddorp, The Netherlands, 14 \hat{a} €"16 September 2018. Neuromuscular Disorders, 2019, 29, 401-410.	0.6	5
23	New family with <i>HSPB</i> 8-associated autosomal dominant rimmed vacuolar myopathy. Neurology: Genetics, 2019, 5, e349.	1.9	24
24	QuantiMus: A Machine Learning-Based Approach for High Precision Analysis of Skeletal Muscle Morphology. Frontiers in Physiology, 2019, 10, 1416.	2.8	35
25	Rasagiline for amyotrophic lateral sclerosis: A randomized, controlled trial. Muscle and Nerve, 2019, 59, 201-207.	2.2	35
26	Longitudinal Screening Detects Cognitive Stability and Behavioral Deterioration in ALS Patients. Behavioural Neurology, 2018, 2018, 1-7.	2.1	19
27	Review process for IVIg treatment. Neurology: Clinical Practice, 2018, 8, 429-436.	1.6	9
28	Novel Therapeutic Options in Treatment of Idiopathic Inflammatory Myopathies. Current Treatment Options in Neurology, 2018, 20, 37.	1.8	2
29	A cross-sectional analysis of clinical evaluation in 35 individuals with mutations of the valosin-containing protein gene. Neuromuscular Disorders, 2018, 28, 778-786.	0.6	7
30	Mortality and Causes of Death in Patients with Sporadic Inclusion Body Myositis: Survey Study Based on the Clinical Experience of Specialists in Australia, Europe and the USA. Journal of Neuromuscular Diseases, 2016, 3, 67-75.	2.6	44
31	Sporadic inclusion body myositis misdiagnosed as idiopathic granulomatous myositis. Neuromuscular Disorders, 2016, 26, 741-743.	0.6	7
32	Randomized Trial of Thymectomy in Myasthenia Gravis. New England Journal of Medicine, 2016, 375, 511-522.	27.0	695
33	215th ENMC International Workshop VCP-related multi-system proteinopathy (IBMPFD) 13–15 November 2015, Heemskerk, The Netherlands. Neuromuscular Disorders, 2016, 26, 535-547.	0.6	38
34	Prospective exploratory muscle biopsy, imaging, and functional assessment in patients with late-onset Pompe disease treated with alglucosidase alfa: The EMBASSY Study. Molecular Genetics and Metabolism, 2016, 119, 115-123.	1.1	49
35	A randomized controlled trial of methotrexate for patients with generalized myasthenia gravis. Neurology, 2016, 87, 57-64.	1.1	106
36	Cognitive-behavioral screening reveals prevalent impairment in a large multicenter ALS cohort. Neurology, 2016, 86, 813-820.	1.1	70

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37	Pulmonary function tests (maximum inspiratory pressure, maximum expiratory pressure, vital capacity,) Tj ETQq1 2016, 26, 136-145.	1 0.78431 0.6	.4 rgBT /Ov 31
38	An overview of polymyositis and dermatomyositis. Muscle and Nerve, 2015, 51, 638-656.	2.2	176
39	Topical Tranexamic Acid Does Not Affect Electrophysiologic or Neurovascular Sciatic Nerve Markers in an Animal Model. Clinical Orthopaedics and Related Research, 2015, 473, 1074-1082.	1.5	8
40	Desert hedgehog is a mediator of demyelination in compression neuropathies. Experimental Neurology, 2015, 271, 84-94.	4.1	17
41	ALS Multicenter Cohort Study of Oxidative Stress (ALS COSMOS): Study methodology, recruitment, and baseline demographic and disease characteristics. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2014, 15, 192-203.	1.7	35
42	Lipid-enriched diet rescues lethality and slows down progression in a murine model of VCP-associated disease. Human Molecular Genetics, 2014, 23, 1333-1344.	2.9	20
43	Targeting the Wnt/ß-Catenin Signaling Pathway After Traumatic Nerve Injury to Improve Functional Recovery. Journal of Hand Surgery, 2014, 39, e13-e14.	1.6	O
44	Experimental trials in amyotrophic lateral sclerosis: a review of recently completed, ongoing and planned trials using existing and novel drugs. Expert Opinion on Investigational Drugs, 2014, 23, 1541-1551.	4.1	35
45	Early Surgical Decompression Restores Neurovascular Blood Flow and Ischemic Parameters in an in Vivo Animal Model of Nerve Compression Injury. Journal of Bone and Joint Surgery - Series A, 2014, 96, 897-906.	3.0	29
46	A randomized, doubleâ€blind, placeboâ€controlled phase II study of eculizumab in patients with refractory generalized myasthenia gravis. Muscle and Nerve, 2013, 48, 76-84.	2.2	187
47	A progressive translational mouse model of human valosinâ€containing protein disease: The <i>VCP</i> ^{R155H/+} mouse. Muscle and Nerve, 2013, 47, 260-270.	2.2	58
48	Matrix metalloproteinase 3 deletion preserves denervated motor endplates after traumatic nerve injury. Annals of Neurology, 2013, 73, 210-223.	5.3	47
49	Valosin-containing protein mutation and Parkinson's disease. Parkinsonism and Related Disorders, 2012, 18, 107-109.	2.2	53
50	Biophysical stimulation induces demyelination via an integrinâ€dependent mechanism. Annals of Neurology, 2012, 72, 112-123.	5.3	14
51	Chronic nerve compression alters schwann cell myelin architecture in a murine model. Muscle and Nerve, 2012, 45, 231-241.	2.2	50
52	The Homozygote VCPR155H/R155H Mouse Model Exhibits Accelerated Human VCP-Associated Disease Pathology. PLoS ONE, 2012, 7, e46308.	2.5	56
53	Compressive Neuropathies of the Upper Extremity: Update on Pathophysiology, Classification, and Electrodiagnostic Findings. Journal of Hand Surgery, 2010, 35, 668-677.	1.6	76
54	Neuromuscular junction integrity after chronic nerve compression injury. Journal of Orthopaedic Research, 2009, 27, 114-119.	2.3	14

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55	A novel, efficient, randomized selection trial comparing combinations of drug therapy for ALS. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2008, 9, 212-222.	2.1	67
56	The Role of Neurodiagnostic Studies in Nerve Injuries and Other Orthopedic Disorders. Journal of Hand Surgery, 2007, 32, 1280-1290.	1.6	13
57	Efficacy of minocycline in patients with amyotrophic lateral sclerosis: a phase III randomised trial. Lancet Neurology, The, 2007, 6, 1045-1053.	10.2	610
58	Local down-regulation of myelin-associated glycoprotein permits axonal sprouting with chronic nerve compression injury. Experimental Neurology, 2006, 200, 418-429.	4.1	54
59	Schwann cells upregulate vascular endothelial growth factor secondary to chronic nerve compression injury. Muscle and Nerve, 2005, 31, 452-460.	2.2	52
60	Chronic nerve compression induces local demyelination and remyelination in a rat model of carpal tunnel syndrome. Experimental Neurology, 2004, 187, 500-508.	4.1	110
61	Macrophage Recruitment Follows the Pattern of Inducible Nitric Oxide Synthase Expression in a Model for Carpal Tunnel Syndrome. Journal of Neurotrauma, 2003, 20, 671-680.	3.4	26
62	Clinical correlates of granulomas in muscle. Journal of Neurology, 1998, 245, 519-524.	3.6	56