Chiara Terracciano

List of Publications by Year
in descending order

52	1,655	304368	301761
papers	citations	22 h-index	39 g-index

53	53	53	5756
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Lactate dehydrogenase and C-reactive protein as predictors of respiratory failure in CoVID-19 patients. Clinica Chimica Acta, 2020, 509, 135-138.	0.5	168
2	The Baffling Case of Ischemic Stroke Disappearance from the Casualty Department in the COVID-19 Era. European Neurology, 2020, 83, 213-215.	0.6	168
3	p62/SQSTM1 is overexpressed and prominently accumulated in inclusions of sporadic inclusion-body myositis muscle fibers, and can help differentiating it from polymyositis and dermatomyositis. Acta Neuropathologica, 2009, 118, 407-413.	3.9	133
4	Potential of Curcumin in Skin Disorders. Nutrients, 2019, 11, 2169.	1.7	106
5	Impaired Autophagy in Sporadic Inclusion-Body Myositis and in Endoplasmic Reticulum Stress-Provoked Cultured Human Muscle Fibers. American Journal of Pathology, 2010, 177, 1377-1387.	1.9	94
6	ALS5/SPG11/KIAA1840mutations cause autosomal recessive axonal Charcotâ€“Marieâ€“Tooth disease. Brain, 2016, 139, 73-85.	3.7	80
7	Screening of ARHSP-TCC patients expands the spectrum ofSPG11mutations and includes a large scale gene deletion. Human Mutation, 2009, 30, E500-E519.	1.1	53
8	Differential features of muscle fiber atrophy in osteoporosis and osteoarthritis. Osteoporosis International, 2013, 24, 1095-1100.	1.3	53
9	LOPED study: looking for an early diagnosis in a late-onset Pompe disease high-risk population. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, jnnp-2014-310164.	0.9	50
10	Preferential central nucleation of type 2 myofibers is an invariable feature of myotonic dystrophy type 2. Muscle and Nerve, 2008, 38, 1405-1411.	1.0	48
11	Auto-reactions, autoimmunity and psoriatic arthritis. Autoimmunity Reviews, 2015, 14, 1142-1146.	2.5	47
12	Complete loss of the DNAJB6 G/F domain and novel missense mutations cause distal-onset DNAJB6 myopathy. Acta Neuropathologica Communications, 2015, 3, 44.	2.4	45
13	Sleep disorders in myotonic dystrophy type 2: a controlled polysomnographic study and selfâ€reported questionnaires. European Journal of Neurology, 2014, 21, 929-934.	1.7	37
14	An Age-Standardized Prevalence Estimate and a Sex and Age Distribution of Myotonic Dystrophy Types 1 and 2 in the Rome Province, Italy. Neuroepidemiology, 2016, 46, 191-197.	1.1	37
15	In AÎ²PPâ€overexpressing cultured human muscle fibers proteasome inhibition enhances phosphorylation of AÎ²PP751 and GSK3Î² activation: effects mitigated by lithium and apparently relevant to sporadic inclusionâ€body myositis. Journal of Neurochemistry, 2010, 112, 389-396.	2.1	35
16	The Autophagy Signaling Pathway: A Potential Multifunctional Therapeutic Target of Curcumin in Neurological and Neuromuscular Diseases. Nutrients, 2019, 11, 1881.	1.7	35
17	Late-onset MNGIE without peripheral neuropathy due to incomplete loss of thymidine phosphorylase activity. Neuromuscular Disorders, 2009, 19, 837-840.	0.3	34
18	Increased risk of tumor in DM1 is not related to exposure to common lifestyle risk factors. Journal of Neurology, 2016, 263, 492-498.	1.8	32

#	Article	IF	Citations
19	Association of HLA-DQB1*05:02 and DRB1*16 Alleles with Late-Onset, Nonthymomatous, AChR-Ab-Positive Myasthenia Gravis. Autoimmune Diseases, 2012, 2012, 1-3.	2.7	30
20	Recurrent hyperCKemia with normal muscle biopsy in a pediatric patient with neuromyelitis optica. Neurology, 2012, 79, 1182-1184.	1.5	27
21	In inclusion-body myositis muscle fibers Parkinson-associated DJ-1 is increased and oxidized. Free Radical Biology and Medicine, 2008, 45, 773-779.	1.3	24
22	Vitamin D deficiency in myotonic dystrophy type 1. Journal of Neurology, 2013, 260, 2330-2334.	1.8	24
23	Sleep disorders in spinal and bulbar muscular atrophy (Kennedyâ€™s disease): a controlled polysomnographic and self-reported questionnaires study. Journal of Neurology, 2014, 261, 889-893.	1.8	21
24	Stroke in COVID-19 patientsâ€”A case series from Italy. International Journal of Stroke, 2020, 15, 701-702.	2.9	21
25	Overexpression of ErbB2 and ErbB3 receptors in Schwann cells of patients with Charcotâ€“Marieâ€“Tooth disease type 1A. Muscle and Nerve, 2006, 33, 342-349.	1.0	20
26	Aberrant splicing and expression of the non muscle myosin heavy-chain gene MYH14 in DM1 muscle tissues. Neurobiology of Disease, 2012, 45, 264-271.	2.1	20
27	Upper motor neuron involvement in X-linked recessive bulbospinal muscular atrophy. Clinical Neurophysiology, 2007, 118, 262-268.	0.7	19
28	Inverse correlation between VEGF and soluble VEGF receptor 2 in POEMS with AIDP responsive to intravenous immunoglobulin. Muscle and Nerve, 2010, 42, 445-448.	1.0	18
29	Cutaneous features of myotonic dystrophy types 1 and 2: Implication of premature aging and vitamin D homeostasis. Neuromuscular Disorders, 2017, 27, 163-169.	0.3	18
30	Impaired healing of fragility fractures in type 2 diabetes: clinical and radiographic assessments and serum cytokine levels. Aging Clinical and Experimental Research, 2015, 27, 37-44.	1.4	17
31	The myotonic dystrophy type 2 (DM2) gene product zinc finger protein 9 (ZNF9) is associated with sarcomeres and normally localized in DM2 patients' muscles. Neuropathology and Applied Neurobiology, 2010, 36, 275-284.	1.8	15
32	Vacuolated PAS-positive lymphocytes as an hallmark of Pompe disease and other myopathies related to impaired autophagy. Journal of Cellular Physiology, 2018, 233, 5829-5837.	2.0	15
33	Thymomatous myasthenia gravis: novel association with HLA DQB1*05:01 and strengthened evidence of high clinical and serological severity. Journal of Neurology, 2019, 266, 982-989.	1.8	14
34	Subacute demyelinating polyneuropathy in B-cell lymphoma with IgM antibodies against glycolipid GD1b. Neurological Sciences, 2005, 26, 355-357.	0.9	13
35	Early subclinical cochlear dysfunction in myotonic dystrophy type 1. European Journal of Neurology, 2011, 18, 1412-1416.	1.7	12
36	Neutral lipidâ€storage disease with myopathy and extended phenotype with novel PNPLA2 mutation. Muscle and Nerve, 2016, 53, 644-648.	1.0	11

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37	Successful long-term therapy with flecainide in a family with paramyotonia congenita. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 1232-1234.	0.9	7
38	Neuromuscular transmission abnormalities in myotonic dystrophy type 1: A neurophysiological study. Clinical Neurology and Neurosurgery, 2016, 150, 84-88.	0.6	6
39	Diagnostic Accuracy of a Bedside Screening Tool for Dysphagia (BSTD) in Acute Stroke Patients. Journal of Stroke and Cerebrovascular Diseases, 2021, 30, 105470.	0.7	6
40	Quantitative Evaluation of Upright Posture by x-Ray and 3D Stereophotogrammetry with a New Marker Set Protocol in Late Onset Pompe Disease. Journal of Neuromuscular Diseases, 2021, 8, 979-988.	1.1	6
41	Validation of the Nine Hole Peg Test as a measure of dexterity in myotonic dystrophy type 1. Neuromuscular Disorders, 2018, 28, 947-951.	0.3	5
42	COVID-19-Related Stroke: Barking up the Wrong Tree?. European Neurology, 2020, 83, 218-219.	0.6	5
43	Multiple Sclerosis Treatment in the COVID-19 Era: A Risk-Benefit Approach. Neurology International, 2022, 14, 368-377.	1.3	5
44	Aquaporin 4 expression in human skeletal muscle fiber types. Muscle and Nerve, 2018, 57, 856-858.	1.0	4
45	Novel autophagic vacuolar myopathies: Phenotype and genotype features. Neuropathology and Applied Neurobiology, 2021, 47, 664-678.	1.8	4
46	Neurofibromatous neuropathy: An ultrastructural study. Ultrastructural Pathology, 2018, 42, 312-316.	0.4	3
47	Periodic acid-Schiff staining on resin muscle sections: Improvement in the histological diagnosis of late-onset Pompe disease. Muscle and Nerve, 2012, 45, 611-612.	1.0	2
48	Late adult-onset adrenomyeloneuropathy evolving with atypical severe frontal lobe syndrome: Importance of neuroimaging. Radiology Case Reports, 2019, 14, 309-314.	0.2	2
49	Intrafamilial â€œDOAâ€plusâ€•phenotype variability related to different OMI/HTRA2 expression. American Journal of Medical Genetics, Part A, 2020, 182, 176-182.	0.7	2
50	Rare Variants in Autophagy and Non-Autophagy Genes in Late-Onset Pompe Disease: Suggestions of Their Disease-Modifying Role in Two Italian Families. International Journal of Molecular Sciences, 2021, 22, 3625.	1.8	2
51	Diagnostic Accuracy of Dysphagia Screening in Stroke Care: Answer to the Letter by Toscano etÂal.. Journal of Stroke and Cerebrovascular Diseases, 2021, 30, 105666.	0.7	0
52	Nodular morphea in a patient with Steinert disease. Giornale Italiano Di Dermatologia E Venereologia, 2019, 154, 209-210.	0.8	0