Raffaele Dubbioso

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8372001/publications.pdf

Version: 2024-02-01

75 papers 1,515 citations

331670 21 h-index 377865 34 g-index

80 all docs 80 docs citations

80 times ranked 2246 citing authors

#	Article	IF	CITATIONS
1	Effect of SARS-CoV-2 mRNA vaccination in MS patients treated with disease modifying therapies. EBioMedicine, 2021, 72, 103581.	6.1	184
2	Functional involvement of central cholinergic circuits and visual hallucinations in Parkinson's disease. Brain, 2009, 132, 2350-2355.	7.6	115
3	Diagnostic contribution and therapeutic perspectives of transcranial magnetic stimulation in dementia. Clinical Neurophysiology, 2021, 132, 2568-2607.	1.5	85
4	The effects of prolonged cathodal direct current stimulation on the excitatory and inhibitory circuits of the ipsilateral and contralateral motor cortex. Journal of Neural Transmission, 2012, 119, 1499-1506.	2.8	71
5	A cross-sectional study investigating frequency and features of definitely diagnosed diabetic painful polyneuropathy. Pain, 2018, 159, 2658-2666.	4.2	49
6	Centre-surround organization of fast sensorimotor integration in human motor hand area. Neurolmage, 2017, 158, 37-47.	4.2	47
7	The therapeutic use of non-invasive brain stimulation in multiple sclerosis – a review. Restorative Neurology and Neuroscience, 2017, 35, 497-509.	0.7	46
8	Association of Variants in the <i>SPTLC1</i> Gene With Juvenile Amyotrophic Lateral Sclerosis. JAMA Neurology, 2021, 78, 1236.	9.0	46
9	Anodal transcranial direct current stimulation of motor cortex does not ameliorate spasticity in multiple sclerosis. Restorative Neurology and Neuroscience, 2015, 33, 487-492.	0.7	39
10	The Effect of Cerebellar Degeneration on Human Sensori-motor Plasticity. Brain Stimulation, 2015, 8, 1144-1150.	1.6	37
11	The Italian multicenter experience with edaravone in amyotrophic lateral sclerosis. Journal of Neurology, 2020, 267, 3258-3267.	3.6	37
12	Subclinical neurological involvement does not develop if Wilson's disease is treated early. Parkinsonism and Related Disorders, 2016, 24, 15-19.	2.2	34
13	Fast Intracortical Sensory-Motor Integration: A Window Into the Pathophysiology of Parkinson's Disease. Frontiers in Human Neuroscience, 2019, 13, 111.	2.0	34
14	Electrophysiological characterisation in hereditary spastic paraplegia type 5. Clinical Neurophysiology, 2011, 122, 819-822.	1.5	31
15	<scp>The neuropathy in hereditary transthyretin amyloidosis</scp> : A <scp>narrative review</scp> . Journal of the Peripheral Nervous System, 2021, 26, 155-159.	3.1	30
16	Executive functions are impaired in heterozygote patients with oculopharyngeal muscular dystrophy. Journal of Neurology, 2012, 259, 833-837.	3 . 6	27
17	The combined treatment with orbital and pretarsal botulinum toxin injections in the management of poorly responsive blepharospasm. Neurological Sciences, 2014, 35, 397-400.	1.9	26
18	Multimodal evoked potentials follow up in multiple sclerosis patients under fingolimod therapy. Journal of the Neurological Sciences, 2016, 365, 143-146.	0.6	26

#	Article	IF	Citations
19	Central cholinergic dysfunction in the adult form of Niemann Pick disease type C: a further link with Alzheimer's disease?. Journal of Neurology, 2014, 261, 804-808.	3.6	24
20	The emotional impact of COVID-19 outbreak in amyotrophic lateral sclerosis patients: evaluation of depression, anxiety and interoceptive awareness. Neurological Sciences, 2020, 41, 2339-2341.	1.9	24
21	The Myelin Content of the Human Precentral Hand Knob Reflects Interindividual Differences in Manual Motor Control at the Physiological and Behavioral Level. Journal of Neuroscience, 2021, 41, 3163-3179.	3.6	24
22	Anti-GAD antibody ocular flutter: expanding the spectrum of autoimmune ocular motor disorders. Journal of Neurology, 2013, 260, 2675-2677.	3.6	23
23	Short-latency afferent inhibition in patients with Parkinson's disease and freezing of gait. Journal of Neural Transmission, 2015, 122, 1533-1540.	2.8	22
24	Electrophysiological characterization of adult-onset Niemann–Pick type C disease. Journal of the Neurological Sciences, 2015, 348, 262-265.	0.6	22
25	Autoimmune Autonomic Ganglionopathy. Archives of Neurology, 2011, 68, 504.	4.5	19
26	Somatosensory Temporal Discrimination Threshold Is Increased in Patients with Cerebellar Atrophy. Cerebellum, 2013, 12, 456-459.	2.5	19
27	Atypical clinical and radiological presentation of cryptococcal choroid plexitis in an immunocompetent woman. Journal of the Neurological Sciences, 2013, 334, 180-182.	0.6	19
28	Cognitive correlates of prospective memory in dystonia. Parkinsonism and Related Disorders, 2019, 66, 51-55.	2.2	19
29	Spasmodic dysphonia follow-up with videolaryngoscopy and voice spectrography during treatment with botulinum toxin. Neurological Sciences, 2015, 36, 1679-1682.	1.9	18
30	Thalamic and cortical hyperexcitability in juvenile myoclonic epilepsy. Clinical Neurophysiology, 2020, 131, 2041-2046.	1.5	18
31	Dealing with immune-mediated neuropathies during COVID-19 outbreak: practical recommendations from the task force of the Italian Society of Neurology (SIN), the Italian Society of Clinical Neurophysiology (SINC) and the Italian Peripheral Nervous System Association (ASNP). Neurological Sciences. 2020. 41. 1345-1348.	1.9	17
32	Motor performance deterioration accelerates after 50 years of age in Charcotâ€Marieâ€Tooth type 1A patients. European Journal of Neurology, 2018, 25, 301-306.	3.3	16
33	Case of acute motor conduction block neuropathy (AMCBN). Muscle and Nerve, 2009, 39, 224-226.	2.2	15
34	Abnormal sensorimotor cortex and thalamo-cortical networks in familial adult myoclonic epilepsy type 2: pathophysiology and diagnostic implications. Brain Communications, 2022, 4, fcac037.	3.3	15
35	Neurophysiological Signatures of Motor Impairment in Patients with Rett Syndrome. Annals of Neurology, 2020, 87, 763-773.	5.3	14
36	Cervical dystonia patients display subclinical gait changes. Parkinsonism and Related Disorders, 2017, 43, 97-100.	2.2	13

#	Article	IF	CITATIONS
37	Early predictive factors of disability in CIDP. Journal of Neurology, 2017, 264, 1939-1944.	3.6	11
38	One-year follow up of three Italian patients with Duchenne muscular dystrophy treated with ataluren: is earlier better?. Therapeutic Advances in Neurological Disorders, 2018, 11, 175628641880958.	3.5	11
39	Six-minute walk test is reliable and sensitive in detecting response to therapy in CIDP. Journal of Neurology, 2019, 266, 860-865.	3.6	11
40	Electrophysiological comparison between males and females in HNPP. Neurological Sciences, 2013, 34, 1429-1432.	1.9	10
41	Long-term therapy with miglustat and cognitive decline in the adult form of Niemann-Pick disease type C: a case report. Neurological Sciences, 2018, 39, 1015-1019.	1.9	10
42	Brain Stimulation as a Therapeutic Tool in Amyotrophic Lateral Sclerosis: Current Status and Interaction With Mechanisms of Altered Cortical Excitability. Frontiers in Neurology, 2020, 11, 605335.	2.4	10
43	In vivo evidence of cortical amyloid deposition in the adult form of Niemann Pick type C. Heliyon, 2019, 5, e02776.	3.2	9
44	Teaching Video Neuro <i>Images</i> : Clonus of the lower jaw. Neurology, 2014, 82, e96.	1.1	8
45	Postganglionic Sudomotor Assessment in Early Stage of Multiple System Atrophy and Parkinson Disease. Neurology, 2022, 98, .	1.1	8
46	Muscle fiber type disproportion (FTD) in a family with mutations in the <i>LMNA</i> gene. Muscle and Nerve, 2015, 51, 604-608.	2.2	7
47	Does motor cortex plasticity depend on the type of mutation in the leucine-rich repeat kinase 2 gene?. Movement Disorders, 2017, 32, 947-948.	3.9	7
48	Disruption of GABA(A)-mediated intracortical inhibition in patients with chorea-acanthocytosis. Neuroscience Letters, 2017, 654, 107-110.	2.1	7
49	Cortical tremor: a tantalizing conundrum between cortex and cerebellum. Brain, 2020, 143, e87-e87.	7.6	7
50	Clinical and Molecular Spectrum of Myotonia and Periodic Paralyses Associated With Mutations in SCN4A in a Large Cohort of Italian Patients. Frontiers in Neurology, 2020, 11, 646.	2.4	7
51	Brain Plasticity in Charcot-Marie-Tooth Type 1A Patients? A Combined Structural and Diffusion MRI Study. Frontiers in Neurology, 2020, 11, 795.	2.4	7
52	Cutaneous sensory and autonomic denervation in progressive supranuclear palsy. Neuropathology and Applied Neurobiology, 2021, 47, 653-663.	3.2	7
53	Neurophysiological and behavioural correlates of ocrelizumab therapy on manual dexterity in patients with primary progressive multiple sclerosis. Journal of Neurology, 2022, 269, 4791-4801.	3.6	7
54	The Treatment of Hypersalivation in Rett Syndrome with Botulinum Toxin: Efficacy and Clinical Implications. Neurology and Therapy, 2019, 8, 155-160.	3.2	6

#	Article	lF	CITATIONS
55	Different cortical excitability profiles in hereditary brain iron and copper accumulation. Neurological Sciences, 2020, 41, 679-685.	1.9	6
56	<i>BDNF</i> polymorphism and interhemispheric balance of motor cortex excitability: a preliminary study. Journal of Neurophysiology, 2022, 127, 204-212.	1.8	6
57	Prognostic Markers of Ocrelizumab Effectiveness in Multiple Sclerosis: A Real World Observational Multicenter Study. Journal of Clinical Medicine, 2022, 11, 2081.	2.4	6
58	Relationship between high-frequency activity in the cortical sensory and the motor hand areas, and their myelin content. Brain Stimulation, 2022, 15, 717-726.	1.6	6
59	Isolated intracranial Mycobacterium avium complex granulomas in an immune-competent man. Journal of the Neurological Sciences, 2015, 349, 264-265.	0.6	4
60	Multimodal evaluation of an Italian family with a hereditary spastic paraplegia and <i>POLR3A</i> mutations. Annals of Clinical and Translational Neurology, 2020, 7, 2326-2331.	3.7	4
61	Probing Context-Dependent Modulations of Ipsilateral Premotor-Motor Connectivity in Relapsing-Remitting Multiple Sclerosis. Frontiers in Neurology, 2020, 11, 193.	2.4	4
62	Alteration of interoceptive sensitivity: expanding the spectrum of behavioural disorders in amyotrophic lateral sclerosis. Neurological Sciences, 2022, 43, 5403-5410.	1.9	4
63	Mutilating fingertip ulcers in uncontrolled type 1 diabetes mellitus. Neurological Sciences, 2014, 35, 123-124.	1.9	3
64	The occurrence of lateral shift in cervical dystonia. Neurological Sciences, 2017, 38, 683-686.	1.9	3
65	The impact of symptoms on daily life as perceived by patients with Charcot-Marie-Tooth type 1A disease. Neurological Sciences, 2022, 43, 559-563.	1.9	3
66	Validation of the DYALS (dysphagia in amyotrophic lateral sclerosis) questionnaire for the evaluation of dysphagia in ALS patients. Neurological Sciences, 2022, 43, 3195-3200.	1.9	3
67	Thermosensitive hereditary neuropathy with liability to pressure palsy. Muscle and Nerve, 2011, 43, 448-449.	2.2	2
68	A case of congenital cataracts, facial dysmorphisms, neuropathy, and hyperkinetic movement disorder. Movement Disorders, 2013, 28, 559-560.	3.9	1
69	Chronic inflammatory demyelinating polyneuropathy mimicking an acute painful diabetic neuropathy. Neurological Sciences, 2015, 36, 1509-1510.	1.9	1
70	Need to find a signature of abnormal brain oscillations in task-specific focal dystonia. Clinical Neurophysiology, 2019, 130, 1025-1026.	1.5	1
71	Primary Progressive Multiple Sclerosis Under Anti-TNFα Treatment: A Case Report. Journal of Central Nervous System Disease, 2020, 12, 117957352097382.	1.9	1
72	A Case of Late-Onset Pompe Disease Occurring with a Muscle Weakness Pattern Similar to that of Facioscapulohumeral Muscular Dystrophy. Journal of Neuromuscular Diseases, 2015, 2, S52-S52.	2.6	0

#	Article	IF	CITATIONS
73	Treatment of hypersalivation in rett syndrome with botulinum toxin: Efficacy and clinical implications. Toxicon, 2018, 156, S28-S29.	1.6	O
74	Transcranial brain stimulation and structural MRI., 0,,.		0
75	Ocrelizumab treatment in multiple sclerosis: Prospective real world observational multi-center study in Campania, Italy. Journal of the Neurological Sciences, 2021, 429, 118129.	0.6	O