

Delia Gagliardi

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1658293/publications.pdf>

Version: 2024-02-01

27
papers

420
citations

1039406

9
h-index

752256

20
g-index

28
all docs

28
docs citations

28
times ranked

778
citing authors

#	ARTICLE	IF	CITATIONS
1	Structural Brain Connectome and Cognitive Impairment in Parkinson Disease. <i>Radiology</i> , 2017, 283, 515-525.	3.6	77
2	Time Is Motor Neuron: Therapeutic Window and Its Correlation with Pathogenetic Mechanisms in Spinal Muscular Atrophy. <i>Molecular Neurobiology</i> , 2018, 55, 6307-6318.	1.9	53
3	Diagnostic and Prognostic Role of Blood and Cerebrospinal Fluid and Blood Neurofilaments in Amyotrophic Lateral Sclerosis: A Review of the Literature. <i>International Journal of Molecular Sciences</i> , 2019, 20, 4152.	1.8	47
4	Nusinersen treatment and cerebrospinal fluid neurofilaments: An explorative study on Spinal Muscular Atrophy type 3 patients. <i>Journal of Cellular and Molecular Medicine</i> , 2020, 24, 3034-3039.	1.6	47
5	Extracellular vesicles and amyotrophic lateral sclerosis: from misfolded protein vehicles to promising clinical biomarkers. <i>Cellular and Molecular Life Sciences</i> , 2021, 78, 561-572.	2.4	42
6	MicroRNA's as regulators of cell death mechanisms in amyotrophic lateral sclerosis. <i>Journal of Cellular and Molecular Medicine</i> , 2019, 23, 1647-1656.	1.6	24
7	Estimating the impact of COVID-19 pandemic on services provided by Italian Neuromuscular Centers: an Italian Association of Myology survey of the acute phase. <i>Acta Myologica</i> , 2020, 39, 57-66.	1.5	24
8	Can Intestinal Pseudo-Obstruction Drive Recurrent Stroke-Like Episodes in Late-Onset MELAS Syndrome? A Case Report and Review of the Literature. <i>Frontiers in Neurology</i> , 2019, 10, 38.	1.1	17
9	Insights into the identification of a molecular signature for amyotrophic lateral sclerosis exploiting integrated microRNA profiling of iPSC-derived motor neurons and exosomes. <i>Cellular and Molecular Life Sciences</i> , 2022, 79, 189.	2.4	12
10	Diagnostic and prognostic value of CSF neurofilaments in a cohort of patients with motor neuron disease: A cross-sectional study. <i>Journal of Cellular and Molecular Medicine</i> , 2021, 25, 3765-3771.	1.6	10
11	Stathmins and Motor Neuron Diseases: Pathophysiology and Therapeutic Targets. <i>Biomedicines</i> , 2022, 10, 711.	1.4	9
12	Impact of COVID-19 on the quality of life of patients with neuromuscular disorders in the Lombardy area, Italy. <i>Muscle and Nerve</i> , 2021, 64, 474-482.	1.0	7
13	Central Nervous System Involvement in Common Variable Immunodeficiency: A Case of Acute Unilateral Optic Neuritis in a 26-Year-Old Italian Patient. <i>Frontiers in Neurology</i> , 2018, 9, 1031.	1.1	6
14	Bilateral Cavernous Carotid Aneurysms: Atypical Presentation of a Rare Cause of Mass Effect. A Case Report and a Review of the Literature. <i>Frontiers in Neurology</i> , 2018, 9, 619.	1.1	6
15	Targeting PTB for Glia-to-Neuron Reprogramming In Vitro and In Vivo for Therapeutic Development in Neurological Diseases. <i>Biomedicines</i> , 2022, 10, 399.	1.4	6
16	Homozygous SOD1 Variation L144S Produces a Severe Form of Amyotrophic Lateral Sclerosis in an Iranian Family. <i>Neurology: Genetics</i> , 2022, 8, e645.	0.9	6
17	Insights into disease mechanisms and potential therapeutics for C9orf72-related amyotrophic lateral sclerosis/frontotemporal dementia. <i>Ageing Research Reviews</i> , 2020, 64, 101172.	5.0	5
18	Ophthalmoplegia Due to Miller Fisher Syndrome in a Patient With Myasthenia Gravis. <i>Frontiers in Neurology</i> , 2019, 10, 823.	1.1	4

#	ARTICLE	IF	CITATIONS
19	Subclinical Leber's hereditary optic neuropathy with pediatric acute spinal cord onset: more than meets the eye. <i>BMC Neurology</i> , 2018, 18, 220.	0.8	3
20	Secondary prevention of cryptogenic stroke in patients with patent foramen ovale: a systematic review and meta-analysis. <i>Internal and Emergency Medicine</i> , 2018, 13, 1287-1303.	1.0	3
21	Hyperacute extensive spinal cord infarction and negative spine magnetic resonance imaging: a case report and review of the literature. <i>Medicine (United States)</i> , 2020, 99, e22900.	0.4	3
22	Herpes Simplex virus type 2 myeloradiculitis with a pure motor presentation in a liver transplant recipient. <i>Transplant Infectious Disease</i> , 2020, 22, e13236.	0.7	2
23	Sodium Levels Predict Disability at Discharge in Guillain-Barré Syndrome: A Retrospective Cohort Study. <i>Frontiers in Neurology</i> , 2021, 12, 729252.	1.1	2
24	Expanding the clinical spectrum of the mitochondrial mutation A13084T in the <i>ND5</i> gene. <i>Neurology: Genetics</i> , 2020, 6, e511.	0.9	1
25	Frailties and critical issues in neuromuscular diseases highlighted by SARS-CoV-2 pandemic: how many patients are still "invisible"? <i>Acta Myologica</i> , 2022, 41, 24-29.	1.5	1
26	New Insights into Cerebral Vessel Disease Landscapes at Single-Cell Resolution: Pathogenetic and Therapeutic Perspectives. <i>Biomedicines</i> , 2022, 10, 1693.	1.4	1
27	How to define and enhance diagnostic and assistance pathways in neuromuscular diseases during the COVID-19 pandemic: the concept of network.. <i>Acta Myologica</i> , 2021, 40, 172-176.	1.5	0