Monica Bandettini di Poggio

List of Publications by Year
in descending order

32	1,166	361413 20	414414 32
papers	citations	h-index	g-index

32	32	32	2415
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Endovascular treatment of patients with acute ischemic stroke and tandem occlusion due to internal carotid artery dissection: A multicenter experience. Neuroradiology Journal, 2023, 36, 86-93.	1.2	3
2	Complications of mechanical thrombectomy for acute ischemic stroke: Incidence, risk factors, and clinical relevance in the Italian Registry of Endovascular Treatment in acute stroke. International Journal of Stroke, 2021, 16, 818-827.	5.9	32
3	Coverage of the requirements of first and second level stroke unit in Italy. Neurological Sciences, 2021, 42, 1073-1079.	1.9	1
4	Carotid Stenting and Mechanical Thrombectomy in Patients with Acute Ischemic Stroke and Tandem Occlusions: Antithrombotic Treatment and Functional Outcome. American Journal of Neuroradiology, 2020, 41, 2088-2093.	2.4	20
5	A longitudinal clinical and MRI evaluation of the treatment with erenumab. Neurological Sciences, 2020, 41, 463-464.	1.9	1
6	Management of acute ischemic stroke, thrombolysis rate, and predictors of clinical outcome. Neurological Sciences, 2019, 40, 319-326.	1.9	24
7	Trauma and amyotrophic lateral sclerosis: a european population-based case-control study from the EURALS consortium. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2018, 19, 118-125.	1.7	26
8	Clinical epidemiology of amyotrophic lateral sclerosis in Liguria, Italy: An update of LIGALS register. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2016, 17, 535-542.	1.7	29
9	Molecular Chaperones in the Pathogenesis of Amyotrophic Lateral Sclerosis: The Role of HSPB1. Human Mutation, 2016, 37, 1202-1208.	2.5	45
10	A novel Arg147Trp MATR3 missense mutation in a slowly progressive ALS Italian patient. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2015, 16, 530-531.	1.7	27
11	Whole-blood global DNA methylation is increased in amyotrophic lateral sclerosis independently of age of onset. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2014, 15, 98-105.	1.7	54
12	Brown-Vialetto-Van Laere syndrome: Clinical and neuroradiological findings of a genetically proven patient. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2014, 15, 141-144.	1.7	11
13	Plasma amino acids patterns and age of onset of amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2014, 15, 371-375.	1.7	8
14	Physical activity and amyotrophic lateral sclerosis: A European populationâ€based caseâ€“control study. Annals of Neurology, 2014, 75, 708-716.	5.3	79
15	Validation of the Italian version of the Movement Disorder Societyâ€”Unified Parkinsonâ€™s Disease Rating Scale. Neurological Sciences, 2013, 34, 683-687.	1.9	123
16	Myasthenia gravis developing in an HIV-negative patient with Kaposiâ€™s sarcoma. Neurological Sciences, 2013, 34, 1249-1250.	1.9	4
17	The FIG4 gene does not play a major role in causing ALS in Italian patients. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2013, 14, 228-229.	1.7	8
18	Dopamine-agonist responsive Parkinsonism in a patient with the SANDO syndrome caused by POLG mutation. BMC Medical Genetics, 2013, 14, 105.	2.1	25

#	Article	IF	Citations
19	Neuropathy and levodopa in Parkinson's disease: Evidence from a multicenter study. Movement Disorders, 2013, 28, 1391-1397.	3.9	114
20	Clinical epidemiology of ALS in Liguria, Italy. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2013, 14, 52-57.	1.7	26
21	A novel compound heterozygous mutation of <scp>C20orf54</scp> gene associated with <scp>B</scp>rownâ€“<scp>V</scp>ialettoâ€“<scp>V</scp>an <scp>L</scp>aere <scp>s</scp>yndrome in an <scp>I</scp>talian family. European Journal of Neurology, 2013, 20, e94-5.	3.3	9
22	Discrepancies in reporting the CAG repeat lengths for Huntington's disease. European Journal of Human Genetics, 2012, 20, 20-26.	2.8	20
23	Trauma and amyotrophic lateral sclerosis: a caseâ€“control study from a populationâ€based registry. European Journal of Neurology, 2012, 19, 1509-1517.	3.3	63
24	Whole body cholesterol metabolism is impaired in Huntington's disease. Neuroscience Letters, 2011, 494, 245-249.	2.1	75
25	Clarithromycin-induced neurotoxicity in adults. Journal of Clinical Neuroscience, 2011, 18, 313-318.	1.5	63
26	Whole Body and Cardiac Metaiodobenzylguanidine Kinetics in Parkinson Disease and Multiple System Atrophy. Clinical Nuclear Medicine, 2010, 35, 311-316.	1.3	7
27	A case of secondary syphilis presenting as optic neuritis. Neurological Sciences, 2010, 31, 365-367.	1.9	24
28	Antiphospholipid Syndrome and Reversible Posterior Leukoencephalophaty Syndrome. Seminars in Arthritis and Rheumatism, 2010, 40, e9-e10.	3.4	7
29	Enlarging clinical spectrum of FALS with TARDBP gene mutations: S393L variant in an Italian family showing phenotypic variability and relevance for genetic counselling. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2010, 11, 223-227.	2.1	29
30	Observing Huntingtonâ€™s Disease: the European Huntingtonâ€™s Disease Networkâ€™s REGISTRY. PLOS Currents, 2010, 2, RRN1184.	1.4	124
31	Normal and mutant HTT interact to affect clinical severity and progression in Huntington disease. Neurology, 2009, 73, 1280-1285.	1.1	84
32	Carcinoma of the tongue mimicking bulbar amyotrophic lateral sclerosis. Neurological Sciences, 2008, 29, 127-127.	1.9	1