Rosanna Tortelli

List of Publications by Year
in descending order

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Version: 2024-02-01

54	2,395	304602	223716
papers	citations	22 h-index	46 g-index

59	59	59	5072
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The diagnostic accuracy of late-life depression is influenced by subjective memory complaints and educational level in an older population in Southern Italy. Psychiatry Research, 2022, 308, 114346.	1.7	3
2	A Remote Digital Monitoring Platform to Assess Cognitive and Motor Symptoms in Huntington Disease: Cross-sectional Validation Study. Journal of Medical Internet Research, 2022, 24, e32997.	2.1	15
3	The Modified Five-Point Test (MFPT): normative data for a sample of Italian elderly. Neurological Sciences, 2021, 42, 2431-2440.	0.9	2
4	The use of wearable/portable digital sensors in Huntington's disease: A systematic review. Parkinsonism and Related Disorders, 2021, 83, 93-104.	1.1	28
5	Brain-derived neurotrophic factor in cerebrospinal fluid and plasma is not a biomarker for Huntingtonâ€™s disease. Scientific Reports, 2021, 11, 3481.	1.6	12
6	Comparison of the ability of the Kingâ€™s and MiToS staging systems to predict disease progression and survival in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, 22, 1-9.	1.1	6
7	Kynurenine pathway metabolites in cerebrospinal fluid and blood as potential biomarkers in Huntington's disease. Journal of Neurochemistry, 2021, 158, 539-553.	2.1	18
8	Reduction of Sniff Nasal Inspiratory Pressure (SNIP) as an Early Indicator of the Need of Enteral Nutrition in Patients with Amyotrophic Lateral Sclerosis. Brain Sciences, 2021, 11, 1091.	1.1	1
9	Clinical features and outcomes of the flail arm and flail leg and pure lower motor neuron MND variants: a multicentre Italian study. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1001-1003.	0.9	14
10	Cerebrospinal fluid endo-lysosomal proteins as potential biomarkers for Huntingtonâ€™s disease. PLoS ONE, 2020, 15, e0233820.	1.1	8
11	Mutant huntingtin and neurofilament light have distinct longitudinal dynamics in Huntingtonâ€™s disease. Science Translational Medicine, 2020, 12, .	5.8	64
12	Plasma Inflammatory Cytokines Are Elevated in ALS. Frontiers in Neurology, 2020, 11, 552295.	1.1	48
13	Ageâ€Related Central Auditory Processing Disorder, MCI, and Dementia in an Older Population of Southern Italy. Otolaryngology - Head and Neck Surgery, 2020, 163, 348-355.	1.1	39
14	The Italian Version of the Test Your Memory (TYM-I): A Tool to Detect Mild Cognitive Impairment in the Clinical Setting. Frontiers in Psychology, 2020, 11, 614920.	1.1	4
15	Cerebrospinal fluid endo-lysosomal proteins as potential biomarkers for Huntingtonâ€™s disease. , 2020, 15, e0233820.		0
16	Cerebrospinal fluid endo-lysosomal proteins as potential biomarkers for Huntingtonâ€™s disease. , 2020, 15, e0233820.		0
17	Cerebrospinal fluid endo-lysosomal proteins as potential biomarkers for Huntingtonâ€™s disease. , 2020, 15, e0233820.		0
18	Cerebrospinal fluid endo-lysosomal proteins as potential biomarkers for Huntingtonâ€™s disease. , 2020, 15, e0233820.		0

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19	Diagnostic Value of Cerebrospinal Fluid Neurofilament Light Protein in Neurology. JAMA Neurology, 2019, 76, 1035.	4.5	455
20	Associations of Cognitive Function and Education Level With All-Cause Mortality in Adults on Hemodialysis: Findings From the COGNITIVE-HD Study. American Journal of Kidney Diseases, 2019, 74, 452-462.	2.1	24
21	Comparative Analysis of C9orf72 and Sporadic Disease in a Large Multicenter ALS Population: The Effect of Male Sex on Survival of C9orf72 Positive Patients. Frontiers in Neuroscience, 2019, 13, 485.	1.4	35
22	Incidence of frontotemporal lobar degeneration in Italy. Neurology, 2019, 92, e2355-e2363.	1.5	35
23	Episodic memory and learning rates in amyotrophic lateral sclerosis without dementia. Cortex, 2019, 117, 257-265.	1.1	9
24	Loss of Swallow Tail Sign on Susceptibility-Weighted Imaging in Dementia with Lewy Bodies. Journal of Alzheimer's Disease, 2019, 67, 61-65.	1.2	15
25	Cardiovascular diseases may play a negative role in the prognosis of amyotrophic lateral sclerosis. European Journal of Neurology, 2018, 25, 861-868.	1.7	29
26	Prevalence and patterns of cognitive impairment in adult hemodialysis patients: the COGNITIVE-HD study. Nephrology Dialysis Transplantation, 2018, 33, 1197-1206.	0.4	52
27	Pseudobulbar affect as a negative prognostic indicator in amyotrophic lateral sclerosis. Acta Neurologica Scandinavica, 2018, 138, 55-61.	1.0	8
28	The potential of solanezumab and gantenerumab to prevent Alzheimerâ€™s disease in people with inherited mutations that cause its early onset. Expert Opinion on Biological Therapy, 2018, 18, 25-35.	1.4	34
29	Social Dysfunction in Older Age and Relationships with Cognition, Depression, and Apathy: The GreatAGE Study. Journal of Alzheimer's Disease, 2018, 65, 989-1000.	1.2	42
30	Plasma Î²-amyloid 1â€“42 reference values in cognitively normal subjects. Journal of the Neurological Sciences, 2018, 391, 120-126.	0.3	19
31	Clinical features and genetic characterization of two dizygotic twins with C9orf72 expansion. Neurobiology of Aging, 2018, 69, 293.e1-293.e8.	1.5	1
32	Clinical and genetic analyses of familial and sporadic frontotemporal dementia patients in Southern Italy. Alzheimer's and Dementia, 2017, 13, 858-869.	0.4	24
33	Midlife Metabolic Profile and the Risk of Late-Life Cognitive Decline. Journal of Alzheimer's Disease, 2017, 59, 121-130.	1.2	41
34	Reversible Cognitive Frailty, Dementia, and All-Cause Mortality. The Italian Longitudinal Study on Aging. Journal of the American Medical Directors Association, 2017, 18, 89.e1-89.e8.	1.2	126
35	Comorbidity of dementia with amyotrophic lateral sclerosis (ALS): insights from a large multicenter Italian cohort. Journal of Neurology, 2017, 264, 2224-2231.	1.8	19
36	Primary progressive aphasia: a review of neuropsychological tests for the assessment of speech and language disorders. Aphasiology, 2017, 31, 1359-1378.	1.4	16

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37	Factors predicting survival in ALS: a multicenter Italian study. Journal of Neurology, 2017, 264, 54-63.	1.8	96
38	Classification of Single Normal and Alzheimer's Disease Individuals from Cortical Sources of Resting State EEG Rhythms. Frontiers in Neuroscience, 2016, 10, 47.	1.4	73
39	Time to generalization and prediction of survival in patients with amyotrophic lateral sclerosis: a retrospective observational study. European Journal of Neurology, 2016, 23, 1117-1125.	1.7	19
40	Validity of the Geriatric Depression Scale-30 against the gold standard diagnosis of depression in older age: The GreatAGE Study. European Psychiatry, 2016, 33, S85-S85.	0.1	0
41	Pseudobulbar affect (PBA) in an incident ALS cohort: results from the Apulia registry (SLAP). Journal of Neurology, 2016, 263, 316-321.	1.8	18
42	Tau-based therapeutics for Alzheimerâ€™s disease: active and passive immunotherapy. Immunotherapy, 2016, 8, 1119-1134.	1.0	61
43	Genome-wide association analyses identify new risk variants and the genetic architecture of amyotrophic lateral sclerosis. Nature Genetics, 2016, 48, 1043-1048.	9.4	494
44	Time to generalisation as a predictor of prognosis in amyotrophic lateral sclerosis: TableÂ1. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 678-679.	0.9	15
45	Classification of Healthy Subjects and Alzheimer's Disease Patients with Dementia from Cortical Sources of Resting State EEG Rhythms: A Study Using Artificial Neural Networks. Frontiers in Neuroscience, 2016, 10, 604.	1.4	51
46	Targeting Cognitive Frailty: Clinical and Neurobiological Roadmap for a Single Complex Phenotype. Journal of Alzheimer's Disease, 2015, 47, 793-813.	1.2	108
47	Frontal assessment battery for detecting executive dysfunction in amyotrophic lateral sclerosis without dementia: a retrospective observational study. BMJ Open, 2015, 5, e007069.	0.8	25
48	COGNITIVE-HD study: protocol of an observational study of neurocognitive functioning and association with clinical outcomes in adults with end-stage kidney disease treated with haemodialysis. BMJ Open, 2015, 5, e009328.	0.8	10
49	Epidemiology of age related hearing loss: A review. Hearing, Balance and Communication, 2015, 13, 77-81.	0.1	20
50	Cerebrospinal fluid neurofilament light chain levels: marker of progression to generalized amyotrophic lateral sclerosis. European Journal of Neurology, 2015, 22, 215-218.	1.7	60
51	Sniff nasal inspiratory pressure as a prognostic factor of tracheostomy or death in amyotrophic lateral sclerosis. Journal of Neurology, 2015, 262, 593-603.	1.8	56
52	Amyotrophic Lateral Sclerosis: An Aging-Related Disease. Current Geriatrics Reports, 2015, 4, 142-153.	1.1	17
53	Prevention of Late-life Cognitive Disorders: Diet-Related Factors, Dietary Patterns, and Frailty Models. Current Nutrition Reports, 2014, 3, 110-129.	2.1	9
54	A novel KIF5A mutation in an Italian family marked by spastic paraparesis and congenital deafness. Journal of the Neurological Sciences, 2014, 343, 218-220.	0.3	9