Dario Saracino

List of Publications by Year
in descending order

30 papers	365 citations	840585 11 h-index	839398 18 g-index
30 all docs	30 docs citations	30 times ranked	676 citing authors

#	Article	IF	Citations
1	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. Alzheimer's and Dementia, 2022, 18, 1408-1423.	0.4	24
2	Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. Neurobiology of Aging, 2022, , .	1.5	1
3	Systematic Review on the Role of Lobar Cerebral Microbleeds in Cognition. Journal of Alzheimer's Disease, 2022, 86, 1025-1035.	1.2	2
4	How can we define the presymptomatic C9orf72 disease in 2022? An overview on the current definitions of preclinical and prodromal phases. Revue Neurologique, 2022, 178, 426-436.	0.6	2
5	Hyperkinetic manifestations in superficial siderosis: evidence for pathogenic network disruption. Neurological Sciences, 2021, 42, 719-722.	0.9	0
6	Plasma microRNA signature in presymptomatic and symptomatic subjects with C9orf72-associated frontotemporal dementia and amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 485-493.	0.9	25
7	Primary Progressive Aphasia Associated With GRN Mutations. Neurology, 2021, 97, e88-e102.	1.5	23
8	Plasma NfL levels and longitudinal change rates in C9orf72 and GRN-associated diseases: from tailored references to clinical applications. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 1278-1288.	0.9	25
9	Implication of folate deficiency in CYP2U1 loss of function. Journal of Experimental Medicine, 2021, 218, .	4.2	13
10	Primary progressive aphasias associated with C9orf72 expansions: Another side of the story. Cortex, 2021, 145, 145-159.	1.1	9
11	The missense p.Trp7Arg mutation in GRN gene leads to progranulin haploinsufficiency. Neurobiology of Aging, 2020, 85, 154.e9-154.e11.	1.5	3
12	Delayed postâ€hypoxic leukoencephalopathy with a peculiar autoantibody association. Neurology and Clinical Neuroscience, 2020, 8, 86-88.	0.2	1
13	Homozygous GRN mutations: new phenotypes and new insights into pathological and molecular mechanisms. Brain, 2020, 143, 303-319.	3.7	54
14	One novel GRN null mutation, two different aphasia phenotypes. Neurobiology of Aging, 2020, 87, 141.e9-141.e14.	1.5	6
15	The Rise of the GRN C157KfsX97 Mutation in Southern Italy: Going Back to the Fall of the Western Roman Empire. Journal of Alzheimer's Disease, 2020, 78, 387-394.	1.2	1
16	Plasma progranulin levels for frontotemporal dementia in clinical practice: a 10-year French experience. Neurobiology of Aging, 2020, 91, 167.e1-167.e9.	1.5	24
17	Case Report: Histopathology and Prion Protein Molecular Properties in Inherited Prion Disease With a De Novo Seven-Octapeptide Repeat Insertion. Frontiers in Cellular Neuroscience, 2020, 14, 150.	1.8	4
18	Isolated parkinsonism is an atypical presentation of GRN and C9orf72 gene mutations. Parkinsonism and Related Disorders, 2020, 80, 73-81.	1.1	13

#	Article	IF	Citations
19	Dramatic neurological debut in a case of KÃ¶hlmeier-Degos disease. Neurological Sciences, 2019, 40, 2201-2203.	0.9	1
20	Presymptomatic spinal cord pathology in c9orf72 mutation carriers: A longitudinal neuroimaging study. Annals of Neurology, 2019, 86, 158-167.	2.8	71
21	Neuro-BehÃ§etâ€™s disease presenting as an isolated progressive cognitive and behavioral syndrome. Neurocase, 2018, 24, 238-241.	0.2	2
22	Novel VCP mutations expand the mutational spectrum of frontotemporal dementia. Neurobiology of Aging, 2018, 72, 187.e11-187.e14.	1.5	19
23	A cluster of progranulin C157KfsX97 mutations in Southern Italy: clinical characterization and genetic correlations. Neurobiology of Aging, 2017, 49, 219.e5-219.e13.	1.5	4
24	Diagnostic contribution of magnetic resonance imaging in an atypical presentation of motor neuron disease. Quantitative Imaging in Medicine and Surgery, 2017, 7, 727-731.	1.1	1
25	Sphenoidal pneumosinus dilatans due to anterior skull base meningiomas â€“ CT and MRI aspects: Report of two new cases and literature review. Neuroradiology Journal, 2016, 29, 295-297.	0.6	2
26	A novel diagnostic method to detect truncated neurofibromin in neurofibromatosis 1. Journal of Neurochemistry, 2015, 135, 1123-1128.	2.1	13
27	A case of progressive frontal lobe syndrome in a sporadic form of Cerebral Amyloid Angiopathy: A singular overlap with fronto-temporal dementia?. Journal of the Neurological Sciences, 2015, 359, 247-249.	0.3	2
28	Foix-Chavany-Marie syndrome in a 17-year-old female with congenital cytomegalovirus infection. Neuropsychiatric Disease and Treatment, 2014, 10, 2249.	1.0	3
29	New-Onset Refractory Status Epilepticus Mimicking Herpes Virus Encephalitis. Case Reports in Neurology, 2013, 5, 162-167.	0.3	6
30	A progranulin mutation associated with cortico-basal syndrome in an Italian family expressing different phenotypes of fronto-temporal lobar degeneration. Neurological Sciences, 2012, 33, 93-97.	0.9	11