

Dario Saracino

List of Publications by Year in descending order

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Version: 2024-02-01

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	Presymptomatic spinal cord pathology in <i>C9orf72</i> mutation carriers: A longitudinal neuroimaging study. <i>Annals of Neurology</i> , 2019, 86, 158-167.	2.8	71
2	Homozygous GRN mutations: new phenotypes and new insights into pathological and molecular mechanisms. <i>Brain</i> , 2020, 143, 303-319.	3.7	54
3	Plasma microRNA signature in presymptomatic and symptomatic subjects with C9orf72-associated frontotemporal dementia and amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2021, 92, 485-493.	0.9	25
4	Plasma NFL levels and longitudinal change rates in <i>C9orf72</i> and <i>GRN</i> -associated diseases: from tailored references to clinical applications. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2021, 92, 1278-1288.	0.9	25
5	Plasma progranulin levels for frontotemporal dementia in clinical practice: a 10-year French experience. <i>Neurobiology of Aging</i> , 2020, 91, 167.e1-167.e9.	1.5	24
6	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. <i>Alzheimer's and Dementia</i> , 2022, 18, 1408-1423.	0.4	24
7	Primary Progressive Aphasia Associated With <i>GRN</i> Mutations. <i>Neurology</i> , 2021, 97, e88-e102.	1.5	23
8	Novel VCP mutations expand the mutational spectrum of frontotemporal dementia. <i>Neurobiology of Aging</i> , 2018, 72, 187.e11-187.e14.	1.5	19
9	A novel diagnostic method to detect truncated neurofibromin in neurofibromatosis 1. <i>Journal of Neurochemistry</i> , 2015, 135, 1123-1128.	2.1	13
10	Implication of folate deficiency in CYP2U1 loss of function. <i>Journal of Experimental Medicine</i> , 2021, 218, .	4.2	13
11	Isolated parkinsonism is an atypical presentation of GRN and C9orf72 gene mutations. <i>Parkinsonism and Related Disorders</i> , 2020, 80, 73-81.	1.1	13
12	A progranulin mutation associated with cortico-basal syndrome in an Italian family expressing different phenotypes of fronto-temporal lobar degeneration. <i>Neurological Sciences</i> , 2012, 33, 93-97.	0.9	11
13	Primary progressive aphasia associated with C9orf72 expansions: Another side of the story. <i>Cortex</i> , 2021, 145, 145-159.	1.1	9
14	New-Onset Refractory Status Epilepticus Mimicking Herpes Virus Encephalitis. <i>Case Reports in Neurology</i> , 2013, 5, 162-167.	0.3	6
15	One novel GRN null mutation, two different aphasia phenotypes. <i>Neurobiology of Aging</i> , 2020, 87, 141.e9-141.e14.	1.5	6
16	A cluster of progranulin C157KfsX97 mutations in Southern Italy: clinical characterization and genetic correlations. <i>Neurobiology of Aging</i> , 2017, 49, 219.e5-219.e13.	1.5	4
17	Case Report: Histopathology and Prion Protein Molecular Properties in Inherited Prion Disease With a De Novo Seven-Octapeptide Repeat Insertion. <i>Frontiers in Cellular Neuroscience</i> , 2020, 14, 150.	1.8	4
18	Foix-Chavany-Marie syndrome in a 17-year-old female with congenital cytomegalovirus infection. <i>Neuropsychiatric Disease and Treatment</i> , 2014, 10, 2249.	1.0	3

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19	The missense p.Trp7Arg mutation in GRN gene leads to progranulin haploinsufficiency. <i>Neurobiology of Aging</i> , 2020, 85, 154.e9-154.e11.	1.5	3
20	A case of progressive frontal lobe syndrome in a sporadic form of Cerebral Amyloid Angiopathy: A singular overlap with fronto-temporal dementia?. <i>Journal of the Neurological Sciences</i> , 2015, 359, 247-249.	0.3	2
21	Sphenoidal pneumosinus dilatans due to anterior skull base meningiomas – CT and MRI aspects: Report of two new cases and literature review. <i>Neuroradiology Journal</i> , 2016, 29, 295-297.	0.6	2
22	Neuro-Behçet's disease presenting as an isolated progressive cognitive and behavioral syndrome. <i>Neurocase</i> , 2018, 24, 238-241.	0.2	2
23	Systematic Review on the Role of Lobar Cerebral Microbleeds in Cognition. <i>Journal of Alzheimer's Disease</i> , 2022, 86, 1025-1035.	1.2	2
24	How can we define the presymptomatic C9orf72 disease in 2022? An overview on the current definitions of preclinical and prodromal phases. <i>Revue Neurologique</i> , 2022, 178, 426-436.	0.6	2
25	Diagnostic contribution of magnetic resonance imaging in an atypical presentation of motor neuron disease. <i>Quantitative Imaging in Medicine and Surgery</i> , 2017, 7, 727-731.	1.1	1
26	Dramatic neurological debut in a case of Klüver-Huemer-Degos disease. <i>Neurological Sciences</i> , 2019, 40, 2201-2203.	0.9	1
27	Delayed post-hypoxic leukoencephalopathy with a peculiar autoantibody association. <i>Neurology and Clinical Neuroscience</i> , 2020, 8, 86-88.	0.2	1
28	The Rise of the GRN C157KfsX97 Mutation in Southern Italy: Going Back to the Fall of the Western Roman Empire. <i>Journal of Alzheimer's Disease</i> , 2020, 78, 387-394.	1.2	1
29	Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. <i>Neurobiology of Aging</i> , 2022, , .	1.5	1
30	Hyperkinetic manifestations in superficial siderosis: evidence for pathogenic network disruption. <i>Neurological Sciences</i> , 2021, 42, 719-722.	0.9	0