

Emanuela Rotondo

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/543127/publications.pdf>

Version: 2024-02-01

10
papers

175
citations

1937685

4
h-index

1474206

9
g-index

10
all docs

10
docs citations

10
times ranked

359
citing authors

#	ARTICLE	IF	CITATIONS
1	Phenotypic Heterogeneity of the GRN Asp22fs Mutation in a Large Italian Kindred. <i>Journal of Alzheimer's Disease</i> , 2011, 24, 253-259.	2.6	62
2	Inflammatory molecules in Frontotemporal Dementia: Cerebrospinal fluid signature of progranulin mutation carriers. <i>Brain, Behavior, and Immunity</i> , 2015, 49, 182-187.	4.1	51
3	Social cognition impairment in genetic frontotemporal dementia within the GENFI cohort. <i>Cortex</i> , 2020, 133, 384-398.	2.4	26
4	The Gut Microbiome—Brain Crosstalk in Neurodegenerative Diseases. <i>Biomedicines</i> , 2022, 10, 1486.	3.2	20
5	Niemann-Pick Type C 1 (NPC1) and NPC2 Gene Variability in Demented Patients with Evidence of Brain Amyloid Deposition. <i>Journal of Alzheimer's Disease</i> , 2021, 83, 1313-1323.	2.6	5
6	Spontaneous confabulations in amnesic-mild cognitive impairment due to Alzheimer's disease: a new (yet old) atypical variant?. <i>Neurocase</i> , 2016, 22, 451-460.	0.6	4
7	Late-onset presentation and phenotypic heterogeneity of the rare R377W PSEN1 mutation. <i>European Journal of Neurology</i> , 2020, 27, 2630-2634.	3.3	3
8	Transmembrane Protein 106B Gene (TMEM106B) Variability and Influence on Progranulin Plasma Levels in Patients with Alzheimer's Disease. <i>Journal of Alzheimer's Disease</i> , 2014, 43, 757-761.	2.6	2
9	Detection of the SQSTM1 Mutation in a Patient with Early-Onset Hippocampal Amnesic Syndrome. <i>Journal of Alzheimer's Disease</i> , 2021, 79, 477-481.	2.6	2
10	Diogenes syndrome in dementia: a case report. <i>BJPsych Open</i> , 2021, 7, e43.	0.7	0