Chiara Cupidi

List of Publications by Year
in descending order

36 papers	3,939 citations	394286 19 h-index	330025 37 g-index
38 all docs	38 docs citations	38 times ranked	7654 citing authors

#	Article	IF	Citations
1	Gene Expression Imputation Across Multiple Tissue Types Provides Insight Into the Genetic Architecture of Frontotemporal Dementia and Its Clinical Subtypes. Biological Psychiatry, 2021, 89, 825-835.	0.7	10
2	Mendelian randomization implies no direct causal association between leukocyte telomere length and amyotrophic lateral sclerosis. Scientific Reports, 2020, 10, 12184.	1.6	4
3	C9orf72, age at onset, and ancestry help discriminate behavioral from language variants in FTLD cohorts. Neurology, 2020, 95, e3288-e3302.	1.5	7
4	Refining the Spectrum of Neuronal Intranuclear Inclusion Disease: A Case Report. Journal of Neuropathology and Experimental Neurology, 2019, 78, 665-670.	0.9	21
5	Genetic meta-analysis of diagnosed Alzheimerâ€™s disease identifies new risk loci and implicates AÎ², tau, immunity and lipid processing. Nature Genetics, 2019, 51, 414-430.	9.4	1,962
6	The largest caucasian kindred with dentatorubralâ€pallidoluysian atrophy: A founder mutation in italy. Movement Disorders, 2019, 34, 1919-1924.	2.2	5
7	Frequency of Cardiovascular Genetic Risk Factors in a Calabrian Population and Their Effects on Dementia. Journal of Alzheimer's Disease, 2018, 61, 1179-1187.	1.2	5
8	Brainâ€derived neurotrophic factor modulates cholesterol homeostasis and Apolipoprotein E synthesis in human cell models of astrocytes and neurons. Journal of Cellular Physiology, 2018, 233, 6925-6943.	2.0	33
9	A C6orf10/LOC101929163 locus is associated with age of onset in C9orf72 carriers. Brain, 2018, 141, 2895-2907.	3.7	39
10	Effects of Multiple Genetic Loci on Age atÂOnset in Frontotemporal Dementia. Journal of Alzheimer's Disease, 2017, 56, 1271-1278.	1.2	4
11	The novel PSEN1 M84V mutation associated to frontal dysexecutive syndrome, spastic paraparesis, and cerebellar atrophy in a dominant Alzheimer's disease family. Neurobiology of Aging, 2017, 56, 213.e7-213.e12.	1.5	19
12	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. Nature Genetics, 2017, 49, 1373-1384.	9.4	783
13	Prevalence of Delirium in a Population of Elderly Outpatients with Dementia: A Retrospective Study. Journal of Alzheimer's Disease, 2017, 61, 251-257.	1.2	12
14	Role of Niemann-Pick Type C Disease Mutations in Dementia. Journal of Alzheimer's Disease, 2016, 55, 1249-1259.	1.2	24
15	Angela R.: a familial Alzheimerâ€™s disease case in the days of Auguste D.. Journal of Neurology, 2016, 263, 2494-2498.	1.8	3
16	NGF controls APP cleavage by downregulating APP phosphorylation at Thr668: relevance for Alzheimer's disease. Aging Cell, 2016, 15, 661-672.	3.0	57
17	Italian Frontotemporal Dementia Network (FTD Group-SINDEM): sharing clinical and diagnostic procedures in Frontotemporal Dementia in Italy. Neurological Sciences, 2015, 36, 751-757.	0.9	9
18	Marinescoâ€“SjÃ¶gren syndrome caused by a new SIL1 frameshift mutation. Journal of the Neurological Sciences, 2015, 354, 112-113.	0.3	4

#	Article	IF	Citations
19	Homozygous carriers of APP A713T mutation in an autosomal dominant Alzheimer disease family. Neurology, 2015, 84, 2266-2273.	1.5	30
20	Haptoglobin Interacts with Apolipoprotein E and Beta-Amyloid and Influences Their Crosstalk. ACS Chemical Neuroscience, 2014, 5, 837-847.	1.7	39
21	Novel N-terminal domain mutation in prion protein detected in 2 patients diagnosed with frontotemporal lobar degeneration syndrome. Neurobiology of Aging, 2014, 35, 2657.e7-2657.e11.	1.5	15
22	Frontotemporal dementia and its subtypes: a genome-wide association study. Lancet Neurology, The, 2014, 13, 686-699.	4.9	302
23	Association of the Variant Cys139Arg at GRN Gene to the Clinical Spectrum of Frontotemporal Lobar Degeneration. Journal of Alzheimer's Disease, 2014, 40, 679-685.	1.2	11
24	Frontal dementia related to thalamic stroke: a case report. Neurological Sciences, 2013, 34, 999-1001.	0.9	1
25	Autosomal Dominant Frontotemporal Lobar Degeneration Due to the C9ORF72 Hexanucleotide Repeat Expansion: Late-Onset Psychotic Clinical Presentation. Biological Psychiatry, 2013, 74, 384-391.	0.7	105
26	Sleep quality in caregivers of patients with Alzheimer's disease and Parkinson's disease and its relationship to quality of life â€“ CORRIGENDUM. International Psychogeriatrics, 2013, 25, 1211-1211.	0.6	1
27	Contribution of Cerebrospinal Fluid Thymosin Î²4 Levels to the Clinical Differentiation of Creutzfeldt-Jakob Disease. Archives of Neurology, 2012, 69, 868-72.	4.9	11
28	Tumor Diagnosis Preceding Alzheimer's Disease Onset: Is There a Link Between Cancer and Alzheimer's Disease?. Journal of Alzheimer's Disease, 2012, 31, 177-182.	1.2	45
29	Sleep quality in caregivers of patients with Alzheimer's disease and Parkinson's disease and its relationship to quality of life. International Psychogeriatrics, 2012, 24, 1827-1835.	0.6	64
30	Epidemiology and genetics of frontotemporal dementia: a door-to-door survey in Southern Italy. Neurobiology of Aging, 2012, 33, 2948.e1-2948.e10.	1.5	40
31	REM sleep behavior disorder in a patient with frontotemporal dementia. Neurological Sciences, 2012, 33, 371-373.	0.9	21
32	Lipofuscin Hypothesis of Alzheimerâ€™s Disease. Dementia and Geriatric Cognitive Disorders Extra, 2011, 1, 292-296.	0.6	30
33	Neocortical Variation of AÎ² Load in Fully Expressed, Pure Alzheimer's Disease. Journal of Alzheimer's Disease, 2010, 19, 57-68.	1.2	19
34	Hereditary Cerebral Hemorrhage With Amyloidosis Associated With the E693K Mutation of APP. Archives of Neurology, 2010, 67, 987-95.	4.9	87
35	Acute reversible parkinsonism in a diabetic-uremic patient. Clinical Neurology and Neurosurgery, 2006, 108, 601-603.	0.6	29
36	Neuropathological and Clinical Phenotype of an Italian Alzheimer Family with M239V Mutation of Presenilin 2 Gene. Journal of Neuropathology and Experimental Neurology, 2004, 63, 199-209.	0.9	39