Serena Nannucci

List of Publications by Year
in descending order

17	349	933447 10	940533 16
papers	citations	h-index	g-index

17	17	17	827
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The Cerebral Autosomal-Dominant Arteriopathy With Subcortical Infarcts and Leukoencephalopathy (CADASIL) Scale. Stroke, 2012, 43, 2871-2876.	2.0	68
2	White Matter Microstructural Damage in Small Vessel Disease Is Associated With Montreal Cognitive Assessment But Not With Mini Mental State Examination Performances. Stroke, 2015, 46, 262-264.	2.0	47
3	Heterozygous mutations of <scp>HTRA</scp>1 gene in patients with familial cerebral small vessel disease. CNS Neuroscience and Therapeutics, 2017, 23, 759-765.	3.9	46
4	Operationalizing mild cognitive impairment criteria in small vessel disease: the VMCI-Tuscany Study. , 2016, 12, 407-418.		34
5	Cerebral microbleeds in patients with mild cognitive impairment and small vessel disease: The Vascular Mild Cognitive Impairment (VMCI)-Tuscany study. Journal of the Neurological Sciences, 2016, 368, 195-202.	0.6	27
6	Acetazolamide for the prophylaxis of migraine in CADASIL: a preliminary experience. Journal of Headache and Pain, 2012, 13, 299-302.	6.0	26
7	White matter microstructural damage and depressive symptoms in patients with mild cognitive impairment and cerebral small vessel disease: the VMCIâ€Tuscany Study. International Journal of Geriatric Psychiatry, 2016, 31, 611-618.	2.7	15
8	The VAS-COG clinic: an out-patient service for patients with cognitive and behavioral consequences of cerebrovascular diseases. Neurological Sciences, 2012, 33, 1277-1283.	1.9	13
9	Application of the DSM-5 Criteria for Major Neurocognitive Disorder to Vascular MCI Patients. Dementia and Geriatric Cognitive Disorders Extra, 2018, 8, 104-116.	1.3	13
10	Circulating Biomarkers in Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy Patients. Journal of Stroke and Cerebrovascular Diseases, 2017, 26, 823-833.	1.6	12
11	Familial cerebral cavernous malformation: report of a further Italian family. Neurological Sciences, 2009, 30, 143-147.	1.9	11
12	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.	2.6	11
13	Progranulin Genetic Screening in Frontotemporal Lobar Degeneration Patients From Central Italy. Cellular and Molecular Neurobiology, 2012, 32, 13-16.	3.3	10
14	The Florence VAS-COG Clinic: A Model for the Care of Patients with Cognitive and Behavioral Disturbances Consequent to Cerebrovascular Diseases. Journal of Alzheimer's Disease, 2014, 42, S453-S461.	2.6	10
15	Inherited leukoencephalopathies with clinical onset in middle and old age. Journal of the Neurological Sciences, 2014, 347, 1-13.	0.6	3
16	Response to Letter Regarding Article, â€œThe Cerebral Autosomal-Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy Scale. A Screening Tool to Select Patients for NOTCH3 Gene Analysisâ€• Stroke, 2013, 44, e19.	2.0	2
17	High lipoprotein(a) serum levels in three CADASIL families. Journal of Neurology, 2012, 259, 379-380.	3.6	1