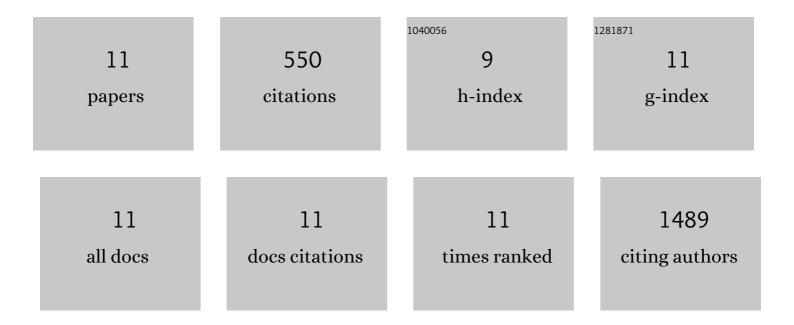
Silvia Testi

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7308333/publications.pdf Version: 2024-02-01



SILVIA TESTI

#	Article	IF	CITATIONS
1	RFC1 AAGGG repeat expansion masquerading as Chronic Idiopathic Axonal Polyneuropathy. Journal of Neurology, 2021, 268, 4280-4290.	3.6	22
2	Charcot-Marie-Tooth disease: experience from a large Italian tertiary neuromuscular center. Neurological Sciences, 2020, 41, 1239-1243.	1.9	16
3	<i>TBK1</i> Mutation Spectrum in an Extended European Patient Cohort with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. Human Mutation, 2017, 38, 297-309.	2.5	87
4	Co-Occurrence of the C9ORF72 Expansion and a Novel GRN Mutation in a Family with Alternative Expression of Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. Journal of Alzheimer's Disease, 2015, 44, 49-56.	2.6	11
5	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. Acta Neuropathologica, 2014, 128, 397-410.	7.7	93
6	A Novel PSEN1 Mutation in a Patient with Sporadic Early-Onset Alzheimer's Disease and Prominent Cerebellar Ataxia. Journal of Alzheimer's Disease, 2014, 41, 709-714.	2.6	10
7	A Panâ€ <scp>E</scp> uropean Study of the <i>C9orf72</i> Repeat Associated with <scp>FTLD</scp> : Geographic Prevalence, Genomic Instability, and Intermediate Repeats. Human Mutation, 2013, 34, 363-373.	2.5	247
8	Unusual features of central nervous system involvement in <scp>CMTX</scp> associated with a novel mutation of <scp><i>GJB1</i></scp> gene. Journal of the Peripheral Nervous System, 2012, 17, 407-411.	3.1	13
9	Autosomal Dominant Alzheimer's Disease with Early Frontal Lobe Involvement Associated with the Met239lle Mutation of Presenilin 2 Gene. Journal of Alzheimer's Disease, 2012, 31, 7-11.	2.6	8
10	Déjerineâ€Sottas syndrome with a silent nucleotide change of myelin protein zero gene. Journal of the Peripheral Nervous System, 2011, 16, 59-64.	3.1	8
11	Pitfalls in diagnosing mitochondrial neurogastrointestinal encephalomyopathy. Journal of Inherited Metabolic Disease, 2011, 34, 1199-1203.	3.6	35