

# Silvia Testi

## List of Publications by Year in descending order

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11  
papers

550  
citations

1040056

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1281871

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g-index

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11  
docs citations

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times ranked

1489  
citing authors

#	ARTICLE	IF	CITATIONS
1	A Pan-European Study of the <i>C9orf72</i> Repeat Associated with FTLD: Geographic Prevalence, Genomic Instability, and Intermediate Repeats. <i>Human Mutation</i> , 2013, 34, 363-373.	2.5	247
2	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. <i>Acta Neuropathologica</i> , 2014, 128, 397-410.	7.7	93
3	<i>TBK1</i> Mutation Spectrum in an Extended European Patient Cohort with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. <i>Human Mutation</i> , 2017, 38, 297-309.	2.5	87
4	Pitfalls in diagnosing mitochondrial neurogastrointestinal encephalomyopathy. <i>Journal of Inherited Metabolic Disease</i> , 2011, 34, 1199-1203.	3.6	35
5	RFC1 AAGGG repeat expansion masquerading as Chronic Idiopathic Axonal Polyneuropathy. <i>Journal of Neurology</i> , 2021, 268, 4280-4290.	3.6	22
6	Charcot-Marie-Tooth disease: experience from a large Italian tertiary neuromuscular center. <i>Neurological Sciences</i> , 2020, 41, 1239-1243.	1.9	16
7	Unusual features of central nervous system involvement in CMTX associated with a novel mutation of <i>GJB1</i> gene. <i>Journal of the Peripheral Nervous System</i> , 2012, 17, 407-411.	3.1	13
8	Co-Occurrence of the C9ORF72 Expansion and a Novel GRN Mutation in a Family with Alternative Expression of Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. <i>Journal of Alzheimer's Disease</i> , 2015, 44, 49-56.	2.6	11
9	A Novel PSEN1 Mutation in a Patient with Sporadic Early-Onset Alzheimer's Disease and Prominent Cerebellar Ataxia. <i>Journal of Alzheimer's Disease</i> , 2014, 41, 709-714.	2.6	10
10	Djerine-Sottas syndrome with a silent nucleotide change of myelin protein zero gene. <i>Journal of the Peripheral Nervous System</i> , 2011, 16, 59-64.	3.1	8
11	Autosomal Dominant Alzheimer's Disease with Early Frontal Lobe Involvement Associated with the Met239Ile Mutation of Presenilin 2 Gene. <i>Journal of Alzheimer's Disease</i> , 2012, 31, 7-11.	2.6	8