

# Cinzia Tiloca

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

10  
papers

1,436  
citations

10  
h-index

10  
g-index

10  
ext. papers

1,786  
ext. citations

18.7  
avg, IF

2.29  
L-index

#	Paper	IF	Citations
10	The LRRK2 Variant E193K Prevents Mitochondrial Fission Upon MPP+ Treatment by Altering LRRK2 Binding to DRP1. <i>Frontiers in Molecular Neuroscience</i> , <b>2018</b> , 11, 64	6.1	25
9	ALS-associated missense and nonsense TBK1 mutations can both cause loss of kinase function. <i>Neurobiology of Aging</i> , <b>2018</b> , 71, 266.e1-266.e10	5.6	44
8	Mutations in the vesicular trafficking protein annexin A11 are associated with amyotrophic lateral sclerosis. <i>Science Translational Medicine</i> , <b>2017</b> , 9,	17.5	74
7	Poly(GP) proteins are a useful pharmacodynamic marker for -associated amyotrophic lateral sclerosis. <i>Science Translational Medicine</i> , <b>2017</b> , 9,	17.5	128
6	Genome-wide association analyses identify new risk variants and the genetic architecture of amyotrophic lateral sclerosis. <i>Nature Genetics</i> , <b>2016</b> , 48, 1043-8	36.3	328
5	NEK1 variants confer susceptibility to amyotrophic lateral sclerosis. <i>Nature Genetics</i> , <b>2016</b> , 48, 1037-42	36.3	149
4	Exome-wide rare variant analysis identifies TUBA4A mutations associated with familial ALS. <i>Neuron</i> , <b>2014</b> , 84, 324-31	13.9	229
3	C9orf72 repeat expansions are restricted to the ALS-FTD spectrum. <i>Neurobiology of Aging</i> , <b>2014</b> , 35, 936.e13-7	5.6	24
2	Oligoclonal bands in the cerebrospinal fluid of amyotrophic lateral sclerosis patients with disease-associated mutations. <i>Journal of Neurology</i> , <b>2013</b> , 260, 85-92	5.5	19
1	Mutations in the profilin 1 gene cause familial amyotrophic lateral sclerosis. <i>Nature</i> , <b>2012</b> , 488, 499-503	50.4	416