Wolfgang P Ruf

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6274229/publications.pdf

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13	1,246	933447 10	1125743 13
papers	citations	h-index	g-index

13	13	13	2972
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Haploinsufficiency of TBK1 causes familial ALS and fronto-temporal dementia. Nature Neuroscience, 2015, 18, 631-636.	14.8	652
2	Hot-spot KIF5A mutations cause familial ALS. Brain, 2018, 141, 688-697.	7.6	167
3	Age-dependent defects of alpha-synuclein oligomer uptake in microglia and monocytes. Acta Neuropathologica, 2016, 131, 379-391.	7.7	140
4	Peripheral monocytes are functionally altered and invade the CNS in ALS patients. Acta Neuropathologica, 2016, 132, 391-411.	7.7	116
5	Impaired activation of ALS monocytes by exosomes. Immunology and Cell Biology, 2017, 95, 207-214.	2.3	39
6	Î±-synuclein interacts with SOD1 and promotes its oligomerization. Molecular Neurodegeneration, 2015, 10, 66.	10.8	29
7	LRRK2 contributes to monocyte dysregulation in Parkinsonâ€™s disease. Acta Neuropathologica Communications, 2016, 4, 123.	5.2	29
8	Screening forCHCHD10mutations in a large cohort of sporadic ALS patients: no evidence for pathogenicity of the p.P34S variant: Table 1. Brain, 2016, 139, e8-e8.	7.6	20
9	A biallelic mutation linksMYORGto autosomal-recessive primary familial brain calcification. Brain, 2019, 142, e4-e4.	7.6	17
10	Exacerbation of chronic inflammatory demyelinating polyneuropathy in concomitance with COVID-19. Journal of the Neurological Sciences, 2020, 418, 117106.	0.6	17
11	Protein Binding Partners of Dysregulated miRNAs in Parkinsonâ€™s Disease Serum. Cells, 2021, 10, 791.	4.1	11
12	Methylome analysis of ALS patients and presymptomatic mutation carriers in blood cells. Neurobiology of Aging, 2022, 116, 16-24.	3.1	8
13	Reply: Adult-onset distal spinal muscular atrophy: a new phenotype associated with KIF5A mutations. Brain, 2019, 142, e67-e67.	7.6	1