

Xianting Li

List of Publications by Year in descending order

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13
papers

1,753
citations

687363

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h-index

1125743

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14
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docs citations

14
times ranked

2847
citing authors

#	ARTICLE	IF	CITATIONS
1	Ser ¹²⁹² Autophosphorylation Is an Indicator of LRRK2 Kinase Activity and Contributes to the Cellular Effects of PD Mutations. <i>Science Translational Medicine</i> , 2012, 4, 164ra161.	12.4	324
2	Enhanced Striatal Dopamine Transmission and Motor Performance with LRRK2 Overexpression in Mice Is Eliminated by Familial Parkinson's Disease Mutation G2019S. <i>Journal of Neuroscience</i> , 2010, 30, 1788-1797.	3.6	309
3	Functional variants in the <i>LRRK2</i> gene confer shared effects on risk for Crohn's disease and Parkinson's disease. <i>Science Translational Medicine</i> , 2018, 10, .	12.4	273
4	Leucine-rich repeat kinase 2 (LRRK2)/PARK8 possesses GTPase activity that is altered in familial Parkinson's disease R1441C/G mutants. <i>Journal of Neurochemistry</i> , 2007, 103, 238-247.	3.9	200
5	Phosphorylation-Dependent 14-3-3 Binding to LRRK2 Is Impaired by Common Mutations of Familial Parkinson's Disease. <i>PLoS ONE</i> , 2011, 6, e17153.	2.5	141
6	Structural model of the dimeric Parkinson's protein LRRK2 reveals a compact architecture involving distant interdomain contacts. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E4357-E4366.	7.1	130
7	Autophagy is induced upon platelet activation and is essential for hemostasis and thrombosis. <i>Blood</i> , 2015, 126, 1224-1233.	1.4	106
8	Parkinson's Disease-Associated LRRK2 Hyperactive Kinase Mutant Disrupts Synaptic Vesicle Trafficking in Ventral Midbrain Neurons. <i>Journal of Neuroscience</i> , 2017, 37, 11366-11376.	3.6	103
9	A Founder Mutation in VPS11 Causes an Autosomal Recessive Leukoencephalopathy Linked to Autophagic Defects. <i>PLoS Genetics</i> , 2016, 12, e1005848.	3.5	50
10	Vitamin B12 modulates Parkinson's disease LRRK2 kinase activity through allosteric regulation and confers neuroprotection. <i>Cell Research</i> , 2019, 29, 313-329.	12.0	42
11	Synj1 haploinsufficiency causes dopamine neuron vulnerability and alpha-synuclein accumulation in mice. <i>Human Molecular Genetics</i> , 2020, 29, 2300-2312.	2.9	29
12	Selective autophagy of AKAP11 activates cAMP/PKA to fuel mitochondrial metabolism and tumor cell growth. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	7.1	27
13	Low-variance RNAs identify Parkinson's disease molecular signature in blood. <i>Movement Disorders</i> , 2015, 30, 813-821.	3.9	18