

# Shi-Ge Wang

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5662934/publications.pdf>

Version: 2024-02-01

11  
papers

107  
citations

1478505

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1372567

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docs citations

11  
times ranked

172  
citing authors

#	ARTICLE	IF	CITATIONS
1	The Phenotypic and Genetic Spectrum of Paroxysmal Kinesigenic Dyskinesia in China. <i>Movement Disorders</i> , 2020, 35, 1428-1437.	3.9	28
2	Expanding the clinical spectrum of adult-onset neuronal intranuclear inclusion disease. <i>Acta Neurologica Belgica</i> , 2022, 122, 647-658.	1.1	15
3	Primary familial brain calcification presenting as paroxysmal kinesigenic dyskinesia: Genetic and functional analyses. <i>Neuroscience Letters</i> , 2020, 714, 134543.	2.1	13
4	Altered structural and functional connectivity in CSF1R-related leukoencephalopathy. <i>Brain Imaging and Behavior</i> , 2020, 15, 1655-1666.	2.1	11
5	Advances in hyperekplexia and other startle syndromes. <i>Neurological Sciences</i> , 2021, 42, 4095-4107.	1.9	9
6	Neurodevelopmental disorder caused by a truncating de novo variant of IRF2BPL. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2021, 84, 47-52.	2.0	7
7	Variants in LAMC3 Causes Occipital Cortical Malformation. <i>Frontiers in Genetics</i> , 2021, 12, 616761.	2.3	7
8	Teaching Video NeuroImages: Cautious walking gait in siblings with hereditary hyperekplexia. <i>Neurology</i> , 2019, 92, e2068-e2069.	1.1	5
9	Clinical and genetic analyses of 150 patients with paroxysmal kinesigenic dyskinesia. <i>Journal of Neurology</i> , 2022, 269, 4717-4728.	3.6	5
10	c.1263+1G>A Is a Latent Hotspot for CYP27A1 Mutations in Chinese Patients With Cerebrotendinous Xanthomatosis. <i>Frontiers in Genetics</i> , 2020, 11, 682.	2.3	4
11	Excessive Startle with Novel <i>GLRA1</i> Mutations in 4 Chinese Patients and a Literature Review of		