

Dae-Seong Kim

List of Publications by Year in descending order

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Version: 2024-02-01

16
papers

206
citations

1163117

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1372567

10
g-index

16
all docs

16
docs citations

16
times ranked

406
citing authors

#	ARTICLE	IF	CITATIONS
1	Muscle and Nerve Biopsy in Various Neuromuscular Disorders. Daehan Sin-gyeong Geunyuk Jilhwon Hakoeji, 2022, 14, 6-15.	0.0	0
2	Nomenclature of emerging therapeutics in neurology. Annals of Clinical Neurophysiology, 2021, 23, 29-34.	0.2	0
3	Nogo-A regulates myogenesis via interacting with Filamin-C. Cell Death Discovery, 2021, 7, 1.	4.7	62
4	A case of X-linked Charcot-Marie-tooth disease type 1 manifesting as recurrent alternating hemiplegia with transient cerebral white matter lesions. Annals of Clinical Neurophysiology, 2021, 23, 130-133.	0.2	0
5	Diagnostic Approaches to Various Muscle Diseases Based on Muscle Pathology. Journal of the Korean Neurological Association, 2021, 39, 274-286.	0.1	3
6	Focal eosinophilic myositis presenting with leg pain and tenderness. Annals of Clinical Neurophysiology, 2020, 22, 125-128.	0.2	1
7	Muscle pathology in neuromuscular disorders. Annals of Clinical Neurophysiology, 2020, 22, 51-60.	0.2	2
8	Characterization of congenital myopathies at a Korean neuromuscular center. Muscle and Nerve, 2018, 58, 235-244.	2.2	8
9	Myasthenia gravis seronegative for acetylcholine receptor antibodies in South Korea: Autoantibody profiles and clinical features. PLoS ONE, 2018, 13, e0193723.	2.5	23
10	NUDT15 p.R139C variant is common and strongly associated with azathioprine-induced early leukopenia and severe alopecia in Korean patients with various neurological diseases. Journal of the Neurological Sciences, 2017, 378, 64-68.	0.6	29
11	Clinical and genetic diversity of nemaline myopathy from a single neuromuscular center in Korea. Journal of the Neurological Sciences, 2017, 383, 61-68.	0.6	13
12	A family with dynamin 2-related centronuclear myopathy without ocular involvement. Journal of Genetic Medicine, 2016, 13, 51-54.	0.2	0
13	Mutation analysis of <i>SPAST</i> , <i>ATL1</i> , and <i>REEP1</i> in Korean Patients with Hereditary		