

# Gye Sun Jeon

## List of Publications by Year in descending order

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Version: 2024-02-01

62  
papers

981  
citations

430874

18  
h-index

501196

28  
g-index

62  
all docs

62  
docs citations

62  
times ranked

1860  
citing authors

#	ARTICLE	IF	CITATIONS
1	Adipose-derived stem cell exosomes alleviate pathology of amyotrophic lateral sclerosis in vitro. <i>Biochemical and Biophysical Research Communications</i> , 2016, 479, 434-439.	2.1	105
2	Pathological Modification of TDP-43 in Amyotrophic Lateral Sclerosis with SOD1 Mutations. <i>Molecular Neurobiology</i> , 2019, 56, 2007-2021.	4.0	52
3	Potential Effect of S-Nitrosylated Protein Disulfide Isomerase on Mutant SOD1 Aggregation and Neuronal Cell Death in Amyotrophic Lateral Sclerosis. <i>Molecular Neurobiology</i> , 2014, 49, 796-807.	4.0	51
4	Intermittent Hypoxia Can Aggravate Motor Neuronal Loss and Cognitive Dysfunction in ALS Mice. <i>PLoS ONE</i> , 2013, 8, e81808.	2.5	47
5	Myelin Oligodendrocyte Glycoprotein-Immunoglobulin G in the CSF. <i>Neurology: Neuroimmunology and Neuroinflammation</i> , 2022, 9, .	6.0	43
6	Sensory Guillain-Barre syndrome following the ChAdOx1 nCov-19 vaccine: Report of two cases and review of literature. <i>Journal of Neuroimmunology</i> , 2021, 359, 577691.	2.3	42
7	The neuroprotective effect of the GSK-3 $\beta$ inhibitor and influence on the extrinsic apoptosis in the ALS transgenic mice. <i>Journal of the Neurological Sciences</i> , 2012, 320, 1-5.	0.6	35
8	High neutrophil-to-lymphocyte ratio predicts short survival duration in amyotrophic lateral sclerosis. <i>Scientific Reports</i> , 2020, 10, 428.	3.3	35
9	Capnography for Assessing Nocturnal Hypoventilation and Predicting Compliance with Subsequent Noninvasive Ventilation in Patients with ALS. <i>PLoS ONE</i> , 2011, 6, e17893.	2.5	32
10	Split-hand phenomenon in amyotrophic lateral sclerosis: A motor unit number index study. <i>Muscle and Nerve</i> , 2016, 53, 885-888.	2.2	31
11	Cerebrospinal fluid/serum gradient of IgG is associated with disability at acute attacks of neuromyelitis optica. <i>Journal of Neurology</i> , 2011, 258, 2176-2180.	3.6	25
12	Altered nucleocytoplasmic proteome and transcriptome distributions in an in vitro model of amyotrophic lateral sclerosis. <i>PLoS ONE</i> , 2017, 12, e0176462.	2.5	24
13	Myasthenia gravis seronegative for acetylcholine receptor antibodies in South Korea: Autoantibody profiles and clinical features. <i>PLoS ONE</i> , 2018, 13, e0193723.	2.5	23
14	Comparison between Flail Arm Syndrome and Upper Limb Onset Amyotrophic Lateral Sclerosis: Clinical Features and Electromyographic Findings. <i>Experimental Neurobiology</i> , 2014, 23, 253-257.	1.6	22
15	The Anti-Inflammatory Effect of Sulforaphane in Mice with Experimental Autoimmune Encephalomyelitis. <i>Journal of Korean Medical Science</i> , 2019, 34, e197.	2.5	21
16	Amyotrophic Lateral Sclerosis - Cell Based Therapy and Novel Therapeutic Development. <i>Experimental Neurobiology</i> , 2014, 23, 207-214.	1.6	20
17	A phosphomimetic mutant TDP-43 (S409/410E) induces Drosha instability and cytotoxicity in Neuro 2A cells. <i>Biochemical and Biophysical Research Communications</i> , 2015, 464, 236-243.	2.1	20
18	Takotsubo cardiomyopathy in amyotrophic lateral sclerosis. <i>Journal of the Neurological Sciences</i> , 2017, 375, 289-293.	0.6	20

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19	Neuroprotective effects of JGK-263 in transgenic SOD1-G93A mice of amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 2014, 340, 112-116.	0.6	17
20	Mutational spectrum of the SPAST and ATL1 genes in Korean patients with hereditary spastic paraplegia. Journal of the Neurological Sciences, 2015, 357, 167-172.	0.6	17
21	Repeated low-dose rituximab treatment based on the assessment of circulating B cells in patients with refractory myasthenia gravis. Therapeutic Advances in Neurological Disorders, 2019, 12, 175628641987118.	3.5	17
22	Dissociated leg muscle atrophy in amyotrophic lateral sclerosis/motor neuron disease: the "split-leg" sign. Scientific Reports, 2020, 10, 15661.	3.3	17
23	Diffusion Tensor Tractography Analysis of the Corpus Callosum Fibers in Amyotrophic Lateral		

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37	Electrodiagnostic data-driven clustering identifies a prognostically different subgroup of patients with chronic inflammatory demyelinating polyneuropathy. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 674-680.	1.9	7
38	A Novel <i>TFG</i> Mutation in a Korean Family with <i>α-Synucleinopathy</i> and Amyotrophic Lateral Sclerosis. <i>Movement Disorders</i> , 2022, 37, 384-391.	3.9	7
39	Adult onset Leigh syndrome with mitochondrial DNA 8344 A>G mutation. <i>Journal of Clinical Neuroscience</i> , 2014, 21, 2009-2011.	1.5	6
40	Pattern of Respiratory Deterioration in Sporadic Amyotrophic Lateral Sclerosis According to Onset Lesion by Using Respiratory Function Tests. <i>Experimental Neurobiology</i> , 2015, 24, 351-357.	1.6	6
41	Peripherally derived macrophages as major phagocytes in MOG encephalomyelitis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2019, 6, .	6.0	6
42	Distinctive patterns of peripheral neuropathy across the spectrum of plasma cell disorders. <i>Scientific Reports</i> , 2019, 9, 16769.	3.3	6
43	A case of MOG encephalomyelitis with T- cell lymphoma. <i>Multiple Sclerosis and Related Disorders</i> , 2020, 41, 102038.	2.0	6
44	Risk factors for developing post-thymectomy myasthenia gravis in patients with thymoma. <i>Muscle and Nerve</i> , 2021, 63, 531-537.	2.2	6
45	B Cell Immunophenotyping and Transcriptional Profiles of Memory B Cells in Patients with Myasthenia Gravis. <i>Experimental Neurobiology</i> , 2019, 28, 720-726.	1.6	6
46	HyperCKemia in Guillain-Barré Syndrome. <i>European Neurology</i> , 2020, 83, 415-420.	1.4	5
47	Progressive brain atrophy and white matter changes in MOG encephalomyelitis. <i>Neurology</i> , 2020, 95, 402-403.	1.1	4
48	A confusing case of multiple sclerosis and central nervous system graft versus host disease. <i>Korean Journal of Internal Medicine</i> , 2016, 31, 995-998.	1.7	4
49	Rapid Progression of Sporadic ALS in a Patient Carrying SOD1 p.Gly13Arg Mutation. <i>Experimental Neurobiology</i> , 2016, 25, 347-350.	1.6	3
50	Current perception threshold in diabetic sensory polyneuropathy with normal routine nerve conduction study. <i>Annals of Clinical Neurophysiology</i> , 2017, 19, 125.	0.2	3
51	FHL1 -mutated reducing body myopathy. <i>Neuropathology</i> , 2020, 40, 185-190.	1.2	3
52	Bone health in neuromyelitis optica: Bone mineral density and fractures. <i>Multiple Sclerosis and Related Disorders</i> , 2020, 42, 102080.	2.0	3
53	Multiple Symmetric Lipomatosis Presenting with Bilateral Brachial Plexopathy. <i>Journal of Clinical</i>		

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55	The Anti-Inflammatory Effects of Oral-Formulated Tacrolimus in Mice with Experimental Autoimmune Encephalomyelitis. <i>Journal of Korean Medical Science</i> , 2017, 32, 1502.	2.5	2
56	Rituximab-induced interstitial lung disease in a patient with aquaporin-4 immunoglobulin G-positive neuromyelitis optica spectrum disorder. <i>Multiple Sclerosis and Related Disorders</i> , 2018, 20, 192-193.	2.0	2
57	Spinobulbar muscular atrophy combined with atypical hereditary neuropathy with liability to pressure palsy. <i>Journal of Clinical Neuroscience</i> , 2018, 48, 90-92.	1.5	2
58	Previous psychiatric disorders in the multistep hypothesis of amyotrophic lateral sclerosis: a South Korean population study. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2022, 23, 601-607.	1.7	2
59	Reversible reddish skin color change in a patient with compressive radial neuropathy. <i>BMC Neurology</i> , 2018, 18, 120.	1.8	1
60	EBI2-expressing B cells in neuromyelitis optica spectrum disorder with AQP4-IgG: Association with acute attacks and serum cytokines. <i>Journal of Neuroimmunology</i> , 2021, 358, 577637.	2.3	1
61	Atypical initial manifestation of facioscapulohumeral muscular dystrophy mimicking neuralgic amyotrophy. <i>Neurology India</i> , 2016, 64, 173.	0.4	1
62	Response to the letter to the editor regarding an article, "Takotsubo cardiomyopathy in amyotrophic lateral sclerosis". <i>Journal of the Neurological Sciences</i> , 2017, 379, 341-342.	0.6	0