

# Young-Chul Choi

## List of Publications by Year in descending order

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Version: 2024-02-01

120  
papers

1,895  
citations

270111

25  
h-index

406436

35  
g-index

126  
all docs

126  
docs citations

126  
times ranked

3102  
citing authors

#	ARTICLE	IF	CITATIONS
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1	Incidence, Disability, and Mortality in Patients With Guillain-Barré Syndrome in Korea: A Nationwide		
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#	ARTICLE	IF	CITATIONS
19	Gender differences influence over insomnia in Korean population: A cross-sectional study. , 2020, 15, e0227190.		0
20	Gender differences influence over insomnia in Korean population: A cross-sectional study. , 2020, 15, e0227190.		0
21	Gender differences influence over insomnia in Korean population: A cross-sectional study. , 2020, 15, e0227190.		0
22	Prevalence and Socioeconomic Status of Patients with Genetic Myopathy in Korea: A Nationwide, Population-Based Study. Neuroepidemiology, 2019, 53, 115-120.	1.1	4
23	Risk of osteoporosis in patients with chronic inflammatory neuropathy- a population-based cohort study. Scientific Reports, 2019, 9, 9131.	1.6	2
24	Neurological Manifestations of Myeloneuropathy in Patients with Nitrous Oxide Intoxication. Journal		

#	ARTICLE	IF	CITATIONS
37	Lower-extremity magnetic resonance imaging in patients with hyperkalemic periodic paralysis carrying the SCN4A mutation T704M: 30-month follow-up of seven patients. <i>Neuromuscular Disorders</i> , 2018, 28, 837-845.	0.3	9
38	<i>FAT1</i> Gene Alteration in Facioscapulohumeral Muscular Dystrophy Type 1. <i>Yonsei Medical Journal</i> , 2018, 59, 337.	0.9	10
39	Pathogenic Variant of REEP1 in a Korean Family with Autosomal-Dominant Hereditary Spastic		

#	ARTICLE	IF	CITATIONS
55	Clinical and Pathological Heterogeneity of Korean Patients with <i>CAPN3</i> Mutations. Yonsei Medical Journal, 2016, 57, 173.	0.9	7

56 Recurrent Episodes of Rhabdomyolysis after Seizures in a Patient with Glycogen Storage Disease Type



#	ARTICLE	IF	CITATIONS
73	Clinical and Genetic Aspects in Twelve Korean Patients with Adrenomyeloneuropathy. Yonsei Medical Journal, 2014, 55, 676.	0.9	17
74	The Role of Insulin Resistance in Diabetic Neuropathy in Koreans with Type 2 Diabetes Mellitus: A 6-Year Follow-Up Study. Yonsei Medical Journal, 2014, 55, 700.	0.9	24
75	The Significance of Clinical and Laboratory Features in the Diagnosis of Glycogen Storage Disease Type V: A Case Report. Journal of Korean Medical Science, 2014, 29, 1021.	1.1	3
76	Congenital muscular dystrophy type 1A with residual merosin expression. Korean Journal of Pediatrics, 2014, 57, 149.	1.9	6
77	Subdural Hemorrhage Mimicking Peripheral Neuropathy. Journal of Korean Neurosurgical Society, 2014, 56, 166.	0.5	3
78	Clinical, immunohistochemical, Western blot, and genetic analysis in dystrophinopathy. Journal of Clinical Neuroscience, 2013, 20, 1099-1105.	0.8	15
79	Clinical and electromyographic features of radiation-induced lower cranial neuropathy. Clinical Neurophysiology, 2013, 124, 598-602.	0.7	27
80	Fluid-Attenuated Inversion Recovery Hyperintense Vessels in Posterior Cerebral Artery Infarction. Cerebrovascular Diseases Extra, 2013, 3, 46-54.	0.5	57
81	A Case of GNE Myopathy Presenting a Rapid Deterioration during Pregnancy. Journal of Clinical		

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91	The Assessment of Routine Electroencephalography in Patients with Altered Mental Status. Yonsei Medical Journal, 2011, 52, 933.	0.9	2
92	Treatment of Spontaneous Cervical Spinal Subdural Hematoma with Methylprednisolone Pulse Therapy. Yonsei Medical Journal, 2011, 52, 692.	0.9	17
93	Three Cases of Manifesting Female Carriers in Patients with Duchenne Muscular Dystrophy. Yonsei Medical Journal, 2011, 52, 192.	0.9	32
94	Elevated serum level of interleukin-32 $\pm$ in the patients with myasthenia gravis. Journal of Neurology, 2011, 258, 1865-1870.	1.8	29
95	Serum neuron-specific enolase level as a biomarker in differential diagnosis of seizure and syncope. Journal of Neurology, 2010, 257, 1708-1712.	1.8	34
96	Clinical Heterogeneity in Korean Patients with Nemaline Myopathy. Yonsei Medical Journal, 2010, 51, 225.	0.9	3

97

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109	A Case of Adult Polyglucosan Body Disease. <i>Yonsei Medical Journal</i> , 2007, 48, 701.	0.9	3
110	MuSK antibody-positive, seronegative myasthenia gravis in Korea. <i>Journal of Clinical Neuroscience</i> , 2006, 13, 353-355.	0.8	35
111	Comparison of Clinical Characteristics Between Congenital Fiber Type Disproportion Myopathy and Congenital Myopathy with Type 1 Fiber Predominance. <i>Yonsei Medical Journal</i> , 2006, 47, 513.	0.9	11
112	Mutation analysis of the GNE gene in Korean patients with distal myopathy with rimmed vacuoles. <i>Journal of Human Genetics</i> , 2006, 51, 137-140.	1.1	33
113	Phenotypic variability in Kennedy's disease: implication of the early diagnostic features. <i>Acta Neurologica Scandinavica</i> , 2005, 112, 57-63.	1.0	107
114	Pneumomediastinum Due to Intractable Hiccup as the Presenting Symptom of Multiple Sclerosis. <i>Yonsei Medical Journal</i> , 2005, 46, 292.	0.9	9
115	Magnetic Resonance Tractography in a Patient with Alexia without Agraphia. <i>European Neurology</i> , 2005, 54, 174-176.	0.6	5
116	Clinical and Pathological Characteristics of Four Korean Patients with Limb-Girdle Muscular Dystrophy type 2B. <i>Journal of Korean Medical Science</i> , 2004, 19, 447.	1.1	5
117	Increase in Transglutaminase 2 in Idiopathic Inflammatory Myopathies. <i>European Neurology</i> , 2004, 51, 10-14.	0.6	29
118	Prediction of Early Clinical Severity and Extent of Neuronal Damage in Anterior-Circulation Infarction Using the Initial Serum Neuron-Specific Enolase Level. <i>Archives of Neurology</i> , 2003, 60, 37.	4.9	61
119	Sporadic Inclusion Body Myositis Correlates with Increased Expression and Cross-linking by Transglutaminases 1 and 2. <i>Journal of Biological Chemistry</i> , 2000, 275, 8703-8710.	1.6	49
120	LGMD2E with a novel nonsense variant in <i>SGCB</i> gene: a case of LGMD2E with a novel variant. <i>Annals of Clinical Neurophysiology</i> , 0, 22, 29.	0.1	0