Takamura Nagasaka

List of Publications by Year
in descending order

26	129	1478505 6	1372567 10
papers	citations	h-index	g-index

29	29	29	171
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical and histopathological features of progressive-type familial amyloidotic polyneuropathy with TTR Lys54. Journal of the Neurological Sciences, 2009, 276, 88-94.	0.6	25
2	Cerebral Hypermetabolism Demonstrated by FDG PET in Familial Creutzfeldt-Jakob Disease. Clinical Nuclear Medicine, 2011, 36, 725-727.	1.3	13
3	Exome sequencing reveals a novel missense mutation in the KIAA0196 gene in a Japanese patient with SPG8. Clinical Neurology and Neurosurgery, 2016, 144, 36-38.	1.4	11
4	Non-convulsive status epilepticus associated with neuronal intranuclear inclusion disease: A case report and literature review. Epilepsy & Behavior Case Reports, 2019, 11, 103-106.	1.5	10
5	Localization of lipocaline-type prostaglandinï¿½D synthase in rat brain: immunoelectron microscopic study. Histochemistry and Cell Biology, 2004, 121, 483-91.	1.7	8
6	Ageâ€related changes in blood pressure and heart rates of patients with Parkinson's disease. Journal of Clinical Hypertension, 2021, 23, 175-178.	2.0	7
7	Familial Amyloidotic Polyneuropathy and Transthyretin. Sub-Cellular Biochemistry, 2012, 65, 565-607.	2.4	6
8	Muscle sympathetic nerve activity in frontotemporal lobar degeneration is similar to amyotrophic lateral sclerosis. Clinical Autonomic Research, 2016, 26, 1-5.	2.5	5
9	Sympathetic outflow to skin predicts central autonomic dysfunction in multiple system atrophy. Neurological Sciences, 2020, 41, 2241-2248.	1.9	5
10	Vasomotor regulation in patients with multiple system atrophy. Journal of Neural Transmission, 2017, 124, 477-481.	2.8	4
11	Thymoma-associated anti-LGI1 encephalitis and myasthenia gravis: A unique combination with autoantibodies. ENeurologicalSci, 2022, 27, 100395.	1.3	4
12	Pre- and postganglionic vasomotor dysfunction causes distal limb coldness in multiple system atrophy. Journal of the Neurological Sciences, 2017, 380, 191-195.	0.6	3
13	Pathological findings in a patient with non-dystrophic myotonia with a mutation of the SCN4A gene; a case report. BMC Neurology, 2019, 19, 125.	1.8	3
14	Japanese amyotrophic lateral sclerosis patient with learning disabilities with a deletion mutation in the Câ€terminal of the <scp>FUS</scp>/<scp>TLS</scp> gene. Neurology and Clinical Neuroscience, 2015, 3, 192-193.	0.4	2
15	Decreasing 123I-ioflupane SPECT accumulation and 123I-MIBG myocardial scintigraphy uptake in a patient with a novel homozygous mutation in the ZFYVE26 gene. Neurological Sciences, 2019, 40, 429-431.	1.9	2
16	Paraneoplastic sensorimotor neuropathy associated with mediastinal germ cell tumor: favorable outcome after high-dose intravenous immunoglobulin therapy. Neurological Sciences, 2020, 41, 723-725.	1.9	2
17	Sympathetic nerve outflow to skin in a case with dentatorubral-pallidoluysian atrophy. Journal of Clinical Neuroscience, 2021, 87, 80-83.	1.5	2
18	Morphological Alterations of the Sarcotubular System in Permanent Myopathy of Hereditary Hypokalemic Periodic Paralysis with a Mutation in the CACNA1S Gene. Journal of Neuropathology and Experimental Neurology, 2020, 79, 1276-1292.	1.7	2

#	Article	IF	Citations
19	A concise overview of recent breakthroughs in imaging of ALS. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders: Official Publication of the World Federation of Neurology, Research Group on Motor Neuron Diseases, 2000, 1, 3-6.	1.2	1
20	Opsoclonus-myoclonus syndrome associated with multiple system atrophy. Cerebellum and Ataxias, 2014, 1, 15.	1.9	1
21	No relation between sympathetic outflow to muscles and respiratory function in amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 2015, 358, 66-71.	0.6	1
22	Changes in sympathetic thermoregulatory function with aging. Clinical Autonomic Research, 2016, 26, 461-464.	2.5	1
23	Thalamic and Thalamic Reticular Nucleus Abnormalities in a Patient With Chromosome 22q11.2 Deletion Syndrome. Journal of Neuropsychiatry and Clinical Neurosciences, 2015, 27, e81-e82.	1.8	0
24	Sympathetic neurograms showing characteristics of both muscle and skin sympathetic nerve activity in a case with pure autonomic failure. Clinical Autonomic Research, 2018, 28, 347-349.	2.5	0
25	Coexistence of Charcot-Marie-Tooth 1A and nondystrophic myotonia due to PMP22 duplication and SCN4A pathogenic variants: a case report. BMC Neurology, 2022, 22, 17.	1.8	0
26	A MOG antibodyâ€related disorder associated with peripheral facial nerve palsy. Clinical and Experimental Neuroimmunology, 0, , .	1.0	0