Kazumasa Shindo

List of Publications by Year
in descending order

48	422	933447	839539
papers	citations	10 h-index	18 g-index

51	51	51	623
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Variants in saposin D domain of prosaposin gene linked to Parkinsonâ€™s disease. Brain, 2020, 143, 1190-1205.	7.6	72
2	Increased sympathetic outflow to muscles in patients with amyotrophic lateral sclerosis: a comparison with other neuromuscular patients. Journal of the Neurological Sciences, 1995, 134, 57-60.	0.6	43
3	Characteristic MRI findings in beta-propeller protein-associated neurodegeneration (BPAN). Neurology: Clinical Practice, 2014, 4, 175-177.	1.6	27
4	Sympathetic sudomotor and vasoconstrictive neural function in patients with Parkinson's disease. Parkinsonism and Related Disorders, 2008, 14, 548-552.	2.2	26
5	Chronological changes of sympathetic outflow to muscles in amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 2004, 227, 79-84.	0.6	22
6	A Japanese SCA5 family with a novel three-nucleotide in-frame deletion mutation in the SPTBN2 gene: a clinical and genetic study. Journal of Human Genetics, 2014, 59, 569-573.	2.3	20
7	Sympathetic sudomotor neural function in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2011, 12, 39-44.	2.1	18
8	Aging Effects of Sympathetic Reflex Activities on Skin Nerves. Gerontology, 2003, 49, 366-373.	2.8	17
9	Analysis of the relationship between muscle sympathetic nerve activity and cardiac123I-metaiodobenzylguanidine uptake in patients with Parkinson's disease. Movement Disorders, 2005, 20, 1419-1424.	3.9	15
10	Novel mutations in the PNPLA6 gene in Boucher-NeuhÃuser syndrome. Journal of Human Genetics, 2015, 60, 217-220.	2.3	15
11	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
12	Temporal prolongation of decreased skin blood flow causes cold limbs in Parkinsonâ€™s disease. Journal of Neural Transmission, 2013, 120, 445-451.	2.8	10
13	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
14	Prolonged Sympathetic Reflex Latency on Skin Nerves in Sporadic Cerebellar Degeneration. Archives of Neurology, 1999, 56, 462.	4.5	9
15	A comparison of sympathetic outflow to muscles between cervical spondylotic amyotrophy and ALS. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders: Official Publication of the World Federation of Neurology, Research Group on Motor Neuron Diseases, 2002, 3, 233-238.	1.2	7
16	Hot cross bun sign in a late-onset SCA1 patient. Neurological Sciences, 2016, 37, 1873-1874.	1.9	7
17	Spinocerebellar ataxia type 31 associated with REM sleep behavior disorder: a case report. BMC Neurology, 2019, 19, 9.	1.8	7
18	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

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19	Neuroimaging, genetic, and enzymatic study in a Japanese family with a GBA gross deletion. Parkinsonism and Related Disorders, 2019, 61, 57-63.	2.2	6
20	Muscle sympathetic nerve activity in frontotemporal lobar degeneration is similar to amyotrophic lateral sclerosis. Clinical Autonomic Research, 2016, 26, 1-5.	2.5	5
21	Conjugal cerebellar type of multiple system atrophy: Person-to-person transmission?. Parkinsonism and Related Disorders, 2019, 69, 68-70.	2.2	5
22	Sympathetic outflow to skin predicts central autonomic dysfunction in multiple system atrophy. Neurological Sciences, 2020, 41, 2241-2248.	1.9	5
23	A treatable case of autoimmune GFAP astrocytopathy presenting chronic progressive cognitive impairment. Neurological Sciences, 2020, 41, 2999-3002.	1.9	5
24	Vasomotor regulation in patients with multiple system atrophy. Journal of Neural Transmission, 2017, 124, 477-481.	2.8	4
25	Camptocormia as an onset symptom of myasthenia gravis. Neurological Sciences, 2017, 38, 515-516.	1.9	4
26	Thymoma-associated anti-LGI1 encephalitis and myasthenia gravis: A unique combination with autoantibodies. ENeurologicalSci, 2022, 27, 100395.	1.3	4
27	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
28	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
29	A patient with ossification of the yellow ligament and ventriculomegaly with 22q11.2 deletion syndrome undiagnosed until adulthood. Heliyon, 2020, 6, e05600.	3.2	3
30	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
31	Novel SLC20A2 mutation in primary familial brain calcification with disturbance of sustained phonation and orofacial apraxia. Journal of the Neurological Sciences, 2018, 390, 1-3.	0.6	2
32	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
33	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
34	Palpebral ptosis as the initial symptom of amyotrophic lateral sclerosis. Neurological Sciences, 2020, 41, 211-212.	1.9	2
35	Sympathetic nerve outflow to skin in a case with dentatorubral-pallidoluysian atrophy. Journal of Clinical Neuroscience, 2021, 87, 80-83.	1.5	2
36	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

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37	Repeated brain Magnetic Resonance Imaging Provides Clues for the Diagnosis of Autoimmune Glial Fibrillary Acid Protein Astrocytopathy. Internal Medicine, 2022, , .	0.7	2
38	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
39	Opsoclonus-myoclonus syndrome associated with multiple system atrophy. Cerebellum and Ataxias, 2014, 1, 15.	1.9	1
40	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
41	Changes in sympathetic thermoregulatory function with aging. Clinical Autonomic Research, 2016, 26, 461-464.	2.5	1
42	A Japanese family with a novel nonsense mutation in the spastin gene associated with both cerebellar ataxia and cognitive impairment. Journal of the Neurological Sciences, 2019, 397, 114-116.	0.6	1
43	Orthostatic hypotension as a core symptom in a Japanese family harboring SNCA duplication. Parkinsonism and Related Disorders, 2020, 81, 28-30.	2.2	1
44	A case of recurrent polymyalgia rheumatica-like complications with pregnancy. Rheumatology International, 2010, 30, 541-542.	3.0	0
45	Active vasodilation by sympathetic outflow to limb skin in a patient with progressive aphasia. NeuroReport, 2014, 25, 303-306.	1.2	0
46	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
47	Muscle Sympathetic Nerve Activity in Neurological Disorders. , 2017, , 13-30.		0
48	A MOG antibodyâ€related disorder associated with peripheral facial nerve palsy. Clinical and Experimental Neuroimmunology, 0, , .	1.0	0