Yuta Ichinose

List of Publications by Year
in descending order

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30 papers	318 citations	1040056 9 h-index	940533 16 g-index
31 all docs	31 docs citations	31 times ranked	629 citing authors

#	Article	IF	Citations
1	The identified clinical features of Parkinson's disease in homo-, heterozygous and digenic variants of PINK1. Neurobiology of Aging, 2021, 97, 146.e1-146.e13.	3.1	14
2	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
3	Palpebral ptosis as the initial symptom of amyotrophic lateral sclerosis. Neurological Sciences, 2020, 41, 211-212.	1.9	2
4	VPS13Dâ€related disorders presenting as a pure and complicated form of hereditary spastic paraplegia. Molecular Genetics & Genomic Medicine, 2020, 8, e1108.	1.2	29
5	RFC1 repeat expansion in Japanese patients with late-onset cerebellar ataxia. Journal of Human Genetics, 2020, 65, 1143-1147.	2.3	32
6	A Japanese SPG4 Patient with a Confirmed De Novo Mutation of the SPAST Gene. Internal Medicine, 2020, 59, 2311-2315.	0.7	1
7	Variants in saposin D domain of prosaposin gene linked to Parkinsonâ€™s disease. Brain, 2020, 143, 1190-1205.	7.6	72
8	Sympathetic outflow to skin predicts central autonomic dysfunction in multiple system atrophy. Neurological Sciences, 2020, 41, 2241-2248.	1.9	5
9	Conjugal multiple system atrophy: Computing chance or investigating real patients?. Parkinsonism and Related Disorders, 2020, 75, 122-123.	2.2	1
10	Conjugal multiple system atrophy: Be cautious when calculating numbers of probability. Parkinsonism and Related Disorders, 2020, 77, 178-179.	2.2	0
11	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
12	Clinical and Genetic Study of the First Japanese FTDP-17 Patient with a Mutation of +3 in Intron 10 in the <i>MAPT</i> Gene. Internal Medicine, 2019, 58, 2397-2400.	0.7	2
13	Conjugal cerebellar type of multiple system atrophy: Person-to-person transmission?. Parkinsonism and Related Disorders, 2019, 69, 68-70.	2.2	5
14	UBAP1 mutations cause juvenile-onset hereditary spastic paraplegias (SPG80) and impair UBAP1 targeting to endosomes. Journal of Human Genetics, 2019, 64, 1055-1065.	2.3	15
15	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
16	Spinocerebellar ataxia type 31 associated with REM sleep behavior disorder: a case report. BMC Neurology, 2019, 19, 9.	1.8	7
17	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
18	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

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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	Novel GARS mutation presenting as autosomal dominant intermediate Charcotâ€Marieâ€Tooth disease. Journal of the Peripheral Nervous System, 2019, 24, 156-160.	3.1	8
21	Exome Sequencing Reveals a Novel Homozygous Frameshift Mutation in the <i>CYP7B1</i> Gene in a Japanese Patient with SPG5. Internal Medicine, 2019, 58, 719-722.	0.7	2
22	PLA2G6-associated neurodegeneration presenting as a complicated form of hereditary spastic paraplegia. Journal of Human Genetics, 2019, 64, 55-59.	2.3	17
23	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
24	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
25	Novel mutations in the ALDH18A1 gene in complicated hereditary spastic paraplegia with cerebellar ataxia and cognitive impairment. Journal of Human Genetics, 2018, 63, 1009-1013.	2.3	18
26	Vasomotor regulation in patients with multiple system atrophy. Journal of Neural Transmission, 2017, 124, 477-481.	2.8	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	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
29	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
30	Characteristic MRI findings in beta-propeller protein-associated neurodegeneration (BPAN). Neurology: Clinical Practice, 2014, 4, 175-177.	1.6	27