

Hirofumi Maruyama

List of Publications by Year in descending order

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151
papers

5,139
citations

94269

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102304

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all docs

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docs citations

151
times ranked

7669
citing authors

#	ARTICLE	IF	CITATIONS
1	Autopsy Validation of the Diagnostic Accuracy of ¹²³ I-Metaiodobenzylguanidine Myocardial Scintigraphy for Lewy Body Disease. <i>Neurology</i> , 2022, 98, .	1.5	10
2	Clinical Factors Predicting Voluntary Driving Cessation among Patients with Parkinson's Disease. <i>Behavioural Neurology</i> , 2022, 2022, 1-6.	1.1	0
3	Impact of Previous Stroke on Clinical Outcome in Elderly Patients With Nonvalvular Atrial Fibrillation: ANAFIE Registry. <i>Stroke</i> , 2022, 53, 2549-2558.	1.0	4
4	An autopsy report of a familial amyotrophic lateral sclerosis case carrying VCP Arg487His mutation with a unique TDP ⁴³ proteinopathy. <i>Neuropathology</i> , 2021, 41, 118-126.	0.7	11
5	Analysis of genetic risk factors in Japanese patients with Parkinson's disease. <i>Journal of Human Genetics</i> , 2021, 66, 957-964.	1.1	4
6	Socio-economic impact on epilepsy outside of the nation-wide COVID-19 pandemic area. <i>Epilepsy and Behavior</i> , 2021, 117, 107886.	0.9	7
7	Impact of anatomical variations on ultrasonographic reference values of lower extremity peripheral nerves. <i>Muscle and Nerve</i> , 2021, 63, 890-896.	1.0	3
8	Study Protocol for a Randomized, Double-Blind, Placebo-Controlled, Phase-II Trial: AdrenoMedullin for Ischemic Stroke Study. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2021, 30, 105761.	0.7	4
9	Utility of Magnetic Resonance Spectroscopy for the Progression of Neurological Symptoms in Lenticulostriate Artery Territory Infarction. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2021, 30, 105747.	0.7	0
10	Nationwide online EEG education during coronavirus disease 2019 pandemic. <i>Clinical Neurophysiology</i> , 2021, 132, 2763-2765.	0.7	3
11	Histone deacetylase 10 knockout activates chaperone-mediated autophagy and accelerates the decomposition of its substrate. <i>Biochemical and Biophysical Research Communications</i> , 2020, 523, 246-252.	1.0	18
12	Effect of tooth loss and nutritional status on outcomes after ischemic stroke. <i>Nutrition</i> , 2020, 71, 110606.	1.1	9
13	Retinitis pigmentosa prior to familial ALS caused by a homozygous cilia and flagella-associated protein 410 mutation. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020, 91, 220-222.	0.9	6
14	Focal hyperperfusion and elevated lactate in the cerebral lesions with anti-GABA _A R encephalitis: A serial MRI study. <i>Journal of Neuroradiology</i> , 2020, 47, 243-246.	0.6	5
15	Conus Medullaris Infarction Involving the Paraspinal Muscles and Nerve Roots. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2020, 29, 104983.	0.7	3
16	Delayed Swallowing Reflex is Overlooked in Swallowing Screening Among Acute Stroke Patients. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2020, 29, 105303.	0.7	3
17	Nicotine-induced upregulation of miR-132-5p enhances cell survival in PC12 cells by targeting the anti-apoptotic protein Bcl-2. <i>Neurological Research</i> , 2020, 42, 405-414.	0.6	8
18	Safety and efficacy of rituximab in neuromyelitis optica spectrum disorders (RIN-1 study): a multicentre, randomised, double-blind, placebo-controlled trial. <i>Lancet Neurology</i> , The, 2020, 19, 298-306.	4.9	218

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19	Serum immunoglobulin G antibody titer to <i>Fusobacterium nucleatum</i> is associated with unfavorable outcome after stroke. <i>Clinical and Experimental Immunology</i> , 2020, 200, 302-309.	1.1	8
20	Anti-HMGCR Antibody-Positive Myopathy Shows Bcl-2-Positive Inflammation and Lymphocytic Accumulations. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 448-457.	0.9	13
21	Increased blood pressure variability during the subacute phase in patients with ischemic stroke presenting with a low ankle-brachial index. <i>Geriatrics and Gerontology International</i> , 2020, 20, 448-454.	0.7	1
22	A case of seronegative longitudinally extensive transverse myelitis with possible neuro sweet disease. <i>ENeurologicalSci</i> , 2020, 18, 100227.	0.5	0
23	Biallelic mutation of <i>HSD17B4</i> induces middle age-onset spinocerebellar ataxia. <i>Neurology: Genetics</i> , 2020, 6, e396.	0.9	6
24	Increased blood pressure variability during the subacute phase of ischemic stroke is associated with poor functional outcomes at 3 months. <i>Scientific Reports</i> , 2020, 10, 811.	1.6	12
25	Measurement of the length of vertebrobasilar arteries: A three-dimensional approach. <i>Journal of the Neurological Sciences</i> , 2020, 414, 116818.	0.3	1
26	Usefulness of Histogram-Profile Analysis in Ring-Enhancing Intracranial Lesions. <i>World Neurosurgery</i> , 2019, 131, e226-e236.	0.7	2
27	Steroid-responsive Nivolumab-induced Involuntary Movement with Anti-thyroid Antibodies. <i>Internal Medicine</i> , 2019, 58, 3577-3581.	0.3	6
28	A score to map the lateral nonprimary motor area: Multispectrum intrinsic brain activity versus cortical stimulation. <i>Epilepsia</i> , 2019, 60, 2294-2305.	2.6	1
29	Low-dose perampanel improves refractory cortical myoclonus by the dispersed and suppressed paroxysmal depolarization shifts in the sensorimotor cortex. <i>Clinical Neurophysiology</i> , 2019, 130, 1804-1812.	0.7	38
30	Prevalence of <i>RNF213</i> p.R4810K Variant in Early-Onset Stroke With Intracranial Arterial Stenosis. <i>Stroke</i> , 2019, 50, 1561-1563.	1.0	32
31	Warm Front Passage on the Previous Day Increased Ischemic Stroke Events. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2019, 28, 1873-1878.	0.7	3
32	A mutant <i>MATR3</i> mouse model to explain multisystem proteinopathy. <i>Journal of Pathology</i> , 2019, 249, 182-192.	2.1	16
33	Baseline Carotid Intima-Media Thickness and Stroke Recurrence During Secondary Prevention With Pravastatin. <i>Stroke</i> , 2019, 50, 1586-1589.	1.0	13
34	Mitochondrial localization of PABPN1 in oculopharyngeal muscular dystrophy. <i>Laboratory Investigation</i> , 2019, 99, 1728-1740.	1.7	6
35	Utility of Minimum Apparent Diffusion Coefficient Ratios in Alberta Stroke Program Early CT Score Regions for Deciding on Stroke Therapy. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2019, 28, 1371-1380.	0.7	1
36	A rational, multispectral mapping algorithm for primary motor cortex: A primary step before cortical stimulation. <i>Epilepsia</i> , 2019, 60, 547-559.	2.6	3

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37	The effect of a portable electrical muscle stimulation device at home on muscle strength and activation patterns in locomotive syndrome patients: A randomized control trial. <i>Journal of Electromyography and Kinesiology</i> , 2019, 45, 46-52.	0.7	22
38	Metastatic Malignant Lymphoma Mimicking Cerebral Toxoplasmosis with the "Target Sign". <i>Internal Medicine</i> , 2019, 58, 1157-1162.	0.3	3
39	Treatment of intractable resting tremor of spinocerebellar ataxia 42 with zonisamide. <i>Journal of the Neurological Sciences</i> , 2019, 396, 119-120.	0.3	3
40	Absence of an Autonomic Sign Assists in the Diagnosis of Extratemporal Lobe Epilepsy Manifesting Generalized Convulsion with Retained Awareness. <i>Internal Medicine</i> , 2019, 58, 1151-1155.	0.3	2
41	Synphilin-1 has neuroprotective effects on MPP ⁺ -induced Parkinson's disease model cells by inhibiting ROS production and apoptosis. <i>Neuroscience Letters</i> , 2019, 690, 145-150.	1.0	31
42	Cumulative Effects of LDL Cholesterol and CRP Levels on Recurrent Stroke and TIA. <i>Journal of Atherosclerosis and Thrombosis</i> , 2019, 26, 432-441.	0.9	26
43	Amyotrophic lateral sclerosis of long clinical course clinically presenting with progressive muscular atrophy. <i>Neuropathology</i> , 2019, 39, 47-53.	0.7	3
44	Desirable Low-Density Lipoprotein Cholesterol Levels for Preventing Stroke Recurrence. <i>Stroke</i> , 2018, 49, 865-871.	1.0	18
45	Impact of D-dimer levels for short-term or long-term outcomes in cryptogenic stroke patients. <i>Journal of Neurology</i> , 2018, 265, 628-636.	1.8	27
46	Blood Pressure Variability in Acute Ischemic Stroke: Influence of Infarct Location in the Insular Cortex. <i>European Neurology</i> , 2018, 79, 90-99.	0.6	15
47	Controlling nutritional status score for predicting 3-mo functional outcome in acute ischemic stroke. <i>Nutrition</i> , 2018, 55-56, 1-6.	1.1	54
48	Long-Term Effect of Pravastatin on Carotid Intima-Media Complex Thickness. <i>Stroke</i> , 2018, 49, 107-113.	1.0	16
49	Screening for Fabry Disease in Japanese Patients with Young-Onset Stroke by Measuring α -Galactosidase A and Globotriaosylsphingosine. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2018, 27, 3563-3569.	0.7	9
50	Factors Associated with Intima-Media Complex Thickness of the Common Carotid Artery in Japanese Noncardioembolic Stroke Patients with Hyperlipidemia: The J-STARS Echo Study. <i>Journal of Atherosclerosis and Thrombosis</i> , 2018, 25, 359-373.	0.9	20
51	Antithrombotic Therapy Strategy for Cancer-Associated Ischemic Stroke: A Case Series of 26 Patients. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2018, 27, e206-e211.	0.7	9
52	The effect of medication on vastus lateralis muscle activation patterns in Parkinson's disease patients. <i>Journal of Electromyography and Kinesiology</i> , 2018, 42, 66-73.	0.7	6
53	Alpha-2-macroglobulin as a Promising Biological Marker of Endothelial Function. <i>Journal of Atherosclerosis and Thrombosis</i> , 2018, 25, 350-358.	0.9	17
54	Effects of controlled abnormal joint movement on the molecular biological response in intra-articular tissues during the acute phase of anterior cruciate ligament injury in a rat model. <i>BMC Musculoskeletal Disorders</i> , 2018, 19, 175.	0.8	8

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55	4- ² ,6-Diamidino-2-Phenylindole Distinctly Labels Tau Deposits. <i>Journal of Histochemistry and Cytochemistry</i> , 2018, 66, 737-751.	1.3	2
56	Muscle-dominant wild-type TDP-43 expression induces myopathological changes featuring tubular aggregates and TDP-43-positive inclusions. <i>Experimental Neurology</i> , 2018, 309, 169-180.	2.0	11
57	Multi-component intrinsic brain activities as a safe alternative to cortical stimulation for sensori-motor mapping in neurosurgery. <i>Clinical Neurophysiology</i> , 2018, 129, 2038-2048.	0.7	6
58	Effects of Cilnidipine, an L/N-Type Calcium Channel Blocker, on Carotid Atherosclerosis in Japanese Post-Stroke Hypertensive Patients: Results from the CA-ATTEND Study. <i>Journal of Atherosclerosis and Thrombosis</i> , 2018, 25, 490-504.	0.9	9
59	Flip-Flop Phenomenon: Swallowing-Induced Arterial Displacement as an Indicator of Carotid Artery Disease. <i>Cerebrovascular Diseases</i> , 2018, 45, 258-262.	0.8	8
60	Pravastatin Reduces the Risk of Atherothrombotic Stroke when Administered within Six Months of an Initial Stroke Event. <i>Journal of Atherosclerosis and Thrombosis</i> , 2018, 25, 262-268.	0.9	7
61	Sex differences in variances of multi-channel surface electromyography distribution of the vastus lateralis muscle during isometric knee extension in young adults. <i>European Journal of Applied Physiology</i> , 2017, 117, 583-589.	1.2	19
62	Ischemic Stroke Mortality Is More Strongly Associated with Anemia on Admission Than with Underweight Status. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2017, 26, 1369-1374.	0.7	14
63	Maximum Tongue Pressure is Associated with Swallowing Dysfunction in ALS Patients. <i>Dysphagia</i> , 2017, 32, 542-547.	1.0	61
64	Blood pressure control with cilnidipine treatment in Japanese post-stroke hypertensive patients: The CA-ATTEND study. <i>Clinical and Experimental Hypertension</i> , 2017, 39, 225-234.	0.5	3
65	The origins of rimmed vacuoles and granulovacuolar degeneration bodies are associated with the Wnt signaling pathway. <i>Neuroscience Letters</i> , 2017, 638, 55-59.	1.0	7
66	Spatial electromyography distribution pattern of the vastus lateralis muscle during ramp up contractions in Parkinson's disease patients. <i>Journal of Electromyography and Kinesiology</i> , 2017, 37, 125-131.	0.7	20
67	Pyrimethamine-induced transient leukoencephalopathy: Similarity to methotrexate encephalopathy. <i>Neurology and Clinical Neuroscience</i> , 2017, 5, 75-75.	0.2	1
68	Modifications of tau protein after cerebral ischemia and reperfusion in rats are similar to those occurring in Alzheimer's disease - Hyperphosphorylation and cleavage of 4- and 3-repeat tau. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2017, 37, 2441-2457.	2.4	32
69	Ability of the Ankle Brachial Index and Brachial-Ankle Pulse Wave Velocity to Predict the 3-Month Outcome in Patients with Non-Cardioembolic Stroke. <i>Journal of Atherosclerosis and Thrombosis</i> , 2017, 24, 1167-1173.	0.9	14
70	First report of a Japanese family with spinocerebellar ataxia type 10: The second report from Asia after a report from China. <i>PLoS ONE</i> , 2017, 12, e0177955.	1.1	17
71	Various meteorological conditions exhibit both immediate and delayed influences on the risk of stroke events: The HEWS's stroke study. <i>PLoS ONE</i> , 2017, 12, e0178223.	1.1	13
72	The Multidisciplinary Swallowing Team Approach Decreases Pneumonia Onset in Acute Stroke Patients. <i>PLoS ONE</i> , 2016, 11, e0154608.	1.1	47

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73	Translocated in liposarcoma regulates the distribution and function of mammalian enabled, a modulator of actin dynamics. <i>FEBS Journal</i> , 2016, 283, 1475-1487.	2.2	1
74	Warfarin-Resistant Deep Vein Thrombosis during the Treatment of Acute Ischemic Stroke in Lung Adenocarcinoma. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2016, 25, e141-e145.	0.7	5
75	The identification of raft-derived tau-associated vesicles that are incorporated into immature tangles and paired helical filaments. <i>Neuropathology and Applied Neurobiology</i> , 2016, 42, 639-653.	1.8	18
76	Deviation in the recovery of the lower limb and respiratory muscles of patients with polymyositis: a preliminary clinical study. <i>Journal of Physical Therapy Science</i> , 2016, 28, 2652-2655.	0.2	2
77	Temporal Trends in Stroke Severity and Prior Antithrombotic Use Among Acute Ischemic Stroke Patients in Japan. <i>Circulation Journal</i> , 2016, 80, 2033-2036.	0.7	6
78	Tongue thickness evaluation using ultrasonography can predict swallowing function in amyotrophic lateral sclerosis patients. <i>Clinical Neurophysiology</i> , 2016, 127, 1669-1674.	0.7	45
79	Prediction of Pneumonia in Acute Stroke Patients Using Tongue Pressure Measurements. <i>PLoS ONE</i> , 2016, 11, e0165837.	1.1	57
80	Immediate effect of passive and active stretching on hamstrings flexibility: a single-blinded randomized control trial. <i>Journal of Physical Therapy Science</i> , 2015, 27, 3167-3170.	0.2	30
81	A mutation in the low voltage-gated calcium channel CACNA1G alters the physiological properties of the channel, causing spinocerebellar ataxia. <i>Molecular Brain</i> , 2015, 8, 89.	1.3	80
82	Endothelial dysfunction is associated with the severity of cerebral small vessel disease. <i>Hypertension Research</i> , 2015, 38, 291-297.	1.5	57
83	Telomere G-tail Length is a Promising Biomarker Related to White Matter Lesions and Endothelial Dysfunction in Patients With Cardiovascular Risk: A Cross-sectional Study. <i>EBioMedicine</i> , 2015, 2, 960-967.	2.7	15
84	A Case of Recurrent Ischemic Stroke Involving Subacute, Progressive Intracranial Cerebral Arterial Sclerosis Prior to Diagnosis with JAK2-mutated Polycythemia Vera. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2015, 24, e4-e6.	0.7	11
85	The Japan Statin Treatment Against Recurrent Stroke (J-STARS): A Multicenter, Randomized, Open-label, Parallel-group Study. <i>EBioMedicine</i> , 2015, 2, 1071-1078.	2.7	100
86	Clinicopathologic features of autosomal recessive amyotrophic lateral sclerosis associated with optineurin mutation. <i>Neuropathology</i> , 2014, 34, 64-70.	0.7	37
87	Exome sequencing reveals a novel <i>ANO10</i> mutation in a Japanese patient with autosomal recessive spinocerebellar ataxia. <i>Clinical Genetics</i> , 2014, 85, 296-297.	1.0	22
88	Neuropathological features of Japanese familial amyotrophic lateral sclerosis with p.N352S mutation in <i>TARDBP</i> . <i>Neuropathology and Applied Neurobiology</i> , 2014, 40, 231-236.	1.8	9
89	DYT6 in Japan—genetic screening and clinical characteristics of the patients. <i>Movement Disorders</i> , 2014, 29, 278-280.	2.2	2
90	Authors' reply to Drs M van Blijlitterswijk, R Rademakers and LH van den Berg. <i>Neuropathology and Applied Neurobiology</i> , 2014, 40, 359-360.	1.8	0

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91	Fiber Type-Specific Expression of Low-Density Lipoprotein Receptor-Related Protein 6 in Human Skeletal Muscles. <i>Pathobiology</i> , 2014, 81, 94-99.	1.9	5
92	Exome sequencing reveals a novel TTC19 mutation in an autosomal recessive spinocerebellar ataxia patient. <i>BMC Neurology</i> , 2014, 14, 5.	0.8	30
93	Exome sequencing reveals a novel MRE11 mutation in a patient with progressive myoclonic ataxia. <i>Journal of the Neurological Sciences</i> , 2014, 337, 219-223.	0.3	16
94	Mutations in Twinkle primase-helicase cause Perrault syndrome with neurologic features. <i>Neurology</i> , 2014, 83, 2054-2061.	1.5	86
95	An autopsy case of sporadic amyotrophic lateral sclerosis associated with the <sc>I113T</sc> mutation. <i>Neuropathology</i> , 2014, 34, 58-63.	0.7	19
96	A Japanese patient with familial <sc>ALS</sc> and a p.<sc>K510M</sc> mutation in the gene for <sc>FUS</sc> (<i><sc>FUS</sc></i>) resulting in the totally locked state. <i>Neuropathology</i> , 2014, 34, 504-509.	0.7	7
97	Detecting gene mutations in Japanese Alzheimer's patients by semiconductor sequencing. <i>Neurobiology of Aging</i> , 2014, 35, 1780.e1-1780.e5.	1.5	19
98	Ultrasonographic nerve enlargement of the median and ulnar nerves and the cervical nerve roots in patients with demyelinating Charcot-Marie-Tooth disease: distinction from patients with chronic inflammatory demyelinating polyneuropathy. <i>Journal of Neurology</i> , 2013, 260, 2580-2587.	1.8	77
99	Cerebellar ataxia with <i>SYNE1</i> mutation accompanying motor neuron disease. <i>Neurology</i> , 2013, 80, 600-601.	1.5	55
100	Optineurin and amyotrophic lateral sclerosis. <i>Geriatrics and Gerontology International</i> , 2013, 13, 528-532.	0.7	43
101	Optineurin suppression causes neuronal cell death via <sc>NF- κ B</sc> pathway. <i>Journal of Neurochemistry</i> , 2013, 126, 699-704.	2.1	80
102	Oromandibular dystonia associated with SCA36. <i>Movement Disorders</i> , 2013, 28, 558-559.	2.2	15
103	Molecular Markers for Granulovacuolar Degeneration Are Present in Rimmed Vacuoles. <i>PLoS ONE</i> , 2013, 8, e80995.	1.1	7
104	The clinical characteristics of spinocerebellar ataxia 36: A study of 2121 Japanese ataxia patients. <i>Movement Disorders</i> , 2012, 27, 1158-1163.	2.2	22
105	Investigation on circular asymmetry of geographical distribution in cancer mortality of Hiroshima atomic bomb survivors based on risk maps: analysis of spatial survival data. <i>Radiation and Environmental Biophysics</i> , 2012, 51, 133-141.	0.6	12
106	Optineurin with amyotrophic lateral sclerosis-related mutations abrogates inhibition of interferon regulatory factor-3 activation. <i>Neuroscience Letters</i> , 2011, 505, 279-281.	1.0	29
107	The UCHL1 S18Y polymorphism and Parkinson's disease in a Japanese population. <i>Parkinsonism and Related Disorders</i> , 2011, 17, 473-475.	1.1	5
108	Screening for OPTN mutations in amyotrophic lateral sclerosis in a mainly Caucasian population. <i>Neurobiology of Aging</i> , 2011, 32, 1923.e9-1923.e10.	1.5	20

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109	Optineurin is co-localized with FUS in basophilic inclusions of ALS with FUS mutation and in basophilic inclusion body disease. <i>Acta Neuropathologica</i> , 2011, 121, 555-557.	3.9	53
110	Clinicopathologic study on an ALS family with a heterozygous E478G optineurin mutation. <i>Acta Neuropathologica</i> , 2011, 122, 223-229.	3.9	60
111	Severe brain atrophy after long-term survival seen in siblings with familial amyotrophic lateral sclerosis and a mutation in the optineurin gene: a case series. <i>Journal of Medical Case Reports</i> , 2011, 5, 573.	0.4	6
112	Mutations of optineurin in amyotrophic lateral sclerosis. <i>Nature</i> , 2010, 465, 223-226.	13.7	1,097
113	Analysis on the Susceptibility Genes in Two Chinese Pedigrees with Familial Parkinson's Disease. <i>Neurology Research International</i> , 2010, 2010, 1-4.	0.5	2
114	<i>LRRK2</i> mutations and risk variants in Japanese patients with Parkinson's disease. <i>Movement Disorders</i> , 2009, 24, 1034-1041.	2.2	60
115	Screening for TARDBP mutations in Japanese familial amyotrophic lateral sclerosis. <i>Journal of the Neurological Sciences</i> , 2009, 284, 69-71.	0.3	45
116	Pathogenic expansions of the SCA6 locus are associated with a common CACNA1A haplotype across the globe: founder effect or predisposing chromosome?. <i>European Journal of Human Genetics</i> , 2008, 16, 841-847.	1.4	8
117	The CNTN4 c.4256C>T mutation is rare in Japanese with inherited spinocerebellar ataxia. <i>Journal of the Neurological Sciences</i> , 2008, 266, 180-181.	0.3	8
118	A polymorphism of LOC387715 gene is associated with age-related macular degeneration in the Japanese population. <i>Neuroscience Letters</i> , 2007, 414, 71-74.	1.0	43
119	Voltage-gated potassium channel antibodies associated limbic encephalitis in a patient with invasive thymoma. <i>Journal of the Neurological Sciences</i> , 2006, 250, 167-169.	0.3	31
120	Identification of a new family of spinocerebellar ataxia type 14 in the Japanese spinocerebellar ataxia population by the screening of PRKCG exon 4. <i>Movement Disorders</i> , 2006, 21, 1355-1360.	2.2	29
121	Identification and haplotype analysis of LRRK2 G2019S in Japanese patients with Parkinson disease. <i>Neurology</i> , 2006, 67, 697-699.	1.5	54
122	Possible Reduced Penetrance of Expansion of 44 to 47 CAG/CAA Repeats in the TATA-Binding Protein Gene in Spinocerebellar Ataxia Type 17. <i>Archives of Neurology</i> , 2004, 61, 209.	4.9	81
123	Dinucleotide repeat polymorphism in interferon- γ gene is not associated with sporadic Alzheimer's disease. <i>American Journal of Medical Genetics Part A</i> , 2004, 124B, 48-49.	2.4	3
124	A novel haplotype of spinocerebellar ataxia type 6 contributes to the highest prevalence in Western Japan. <i>Neuroscience Letters</i> , 2004, 358, 107-110.	1.0	4
125	Quantitative Assessment of Cerebral Blood Flow in Genetically Confirmed Spinocerebellar Ataxia Type 6. <i>Archives of Neurology</i> , 2004, 61, 933.	4.9	29
126	Influence of monocyte chemoattractant protein 1 gene polymorphism on age at onset of sporadic Parkinson's disease. <i>Movement Disorders</i> , 2003, 18, 953-955.	2.2	37

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127	SCA8 Repeat Expansion: Large CTA/CTG Repeat Alleles Are More Common in Ataxic Patients, Including Those with SCA6. <i>American Journal of Human Genetics</i> , 2003, 72, 704-709.	2.6	65
128	Dinucleotide repeat polymorphisms in the Neprilysin gene are not associated with sporadic Alzheimer's disease. <i>Neuroscience Letters</i> , 2002, 320, 105-107.	1.0	32
129	Difference in disease-free survival curve and regional distribution according to subtype of spinocerebellar ataxia: A study of 1,286 Japanese patients. <i>American Journal of Medical Genetics Part A</i> , 2002, 114, 578-583.	2.4	97
130	Contribution of the interleukin-1 β gene polymorphism in multiple system atrophy. <i>Movement Disorders</i> , 2002, 17, 808-811.	2.2	68
131	Citrullinemia type II in a 64-year-old man with fluctuating serum citrulline levels. <i>Journal of the Neurological Sciences</i> , 2001, 182, 167-170.	0.3	17
132	Diagnostic significance of tau protein in cerebrospinal fluid from patients with corticobasal degeneration or progressive supranuclear palsy. <i>Journal of the Neurological Sciences</i> , 2001, 183, 95-98.	0.3	99
133	Citrullinemia type II in a 64-year-old man with fluctuating serum citrulline levels: mutations in the SLC25A13 gene. <i>Journal of the Neurological Sciences</i> , 2001, 193, 63.	0.3	4
134	Genetic studies in Parkinson's disease with an α -synuclein/NACP gene polymorphism in Japan. <i>Neuroscience Letters</i> , 2001, 300, 125-127.	1.0	65
135	Influence of interleukin-1 β gene polymorphism on age-at-onset of spinocerebellar ataxia 6 (SCA6) in Japanese patients. <i>Neuroscience Letters</i> , 2001, 307, 128-130.	1.0	8
136	Lack of an association between cystatin C gene polymorphisms in Japanese patients with Alzheimer's disease. <i>Neurology</i> , 2001, 57, 337-339.	1.5	45
137	Lack of an Association of Estrogen Receptor α Gene Polymorphisms and Transcriptional Activity With Alzheimer Disease. <i>Archives of Neurology</i> , 2000, 57, 236.	4.9	144
138	Apolipoprotein E promoter polymorphism and sporadic Alzheimer's disease in a Japanese population. <i>Neuroscience Letters</i> , 1999, 259, 56-58.	1.0	44
139	Structure and regulation of the human NeuroD (BETA2/BHF1) gene. <i>Molecular Brain Research</i> , 1999, 69, 223-231.	2.5	21
140	Variation in the number of CAG repeats in the Machado-Joseph disease gene (MJD1) in the Japanese population. <i>Journal of the Neurological Sciences</i> , 1999, 166, 71-73.	0.3	2
141	Characteristic Magnetic Resonance Imaging Findings in Machado-Joseph Disease. <i>Archives of Neurology</i> , 1998, 55, 33.	4.9	142
142	Molecular features of the CAG repeats of spinocerebellar ataxia 6 (SCA6). <i>Human Molecular Genetics</i> , 1997, 6, 1283-1287.	1.4	129
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