Hideshi Kawakami

List of Publications by Year
in descending order

162	10,164	61984 43	36028 97
papers	citations	h-index	g-index

166	166	166	11606
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	CAG expansions in a novel gene for Machado-Joseph disease at chromosome 14q32.1. Nature Genetics, 1994, 8, 221-228.	21.4	1,717
2	Genome-wide association study identifies common variants at four loci as genetic risk factors for Parkinson's disease. Nature Genetics, 2009, 41, 1303-1307.	21.4	1,217
3	Mutations of optineurin in amyotrophic lateral sclerosis. Nature, 2010, 465, 223-226.	27.8	1,097
4	Self-Organization of Polarized Cerebellar Tissue in 3D Culture of Human Pluripotent Stem Cells. Cell Reports, 2015, 10, 537-550.	6.4	531
5	Collaborative Analysis of Î±-Synuclein Gene Promoter Variability and Parkinson Disease. JAMA - Journal of the American Medical Association, 2006, 296, 661.	7.4	467
6	High frequency of open-angle glaucoma in Japanese patients with Alzheimer's disease. Journal of the Neurological Sciences, 2006, 246, 79-83.	0.6	216
7	Molecular features of the CAG repeats and clinical manifestation of Machado–Joseph disease. Human Molecular Genetics, 1995, 4, 807-812.	2.9	191
8	Lack of an Association of Estrogen Receptor Î± Gene Polymorphisms and Transcriptional Activity With Alzheimer Disease. Archives of Neurology, 2000, 57, 236.	4.5	144
9	Generation of transgenic mice with elevated blood pressure by introduction of the rat renin and angiotensinogen genes.. Proceedings of the National Academy of Sciences of the United States of America, 1990, 87, 5153-5157.	7.1	142
10	Characteristic Magnetic Resonance Imaging Findings in Machado-Joseph Disease. Archives of Neurology, 1998, 55, 33.	4.5	142
11	Sporadic ataxias in Japan â€“ a population-based epidemiological study. Cerebellum, 2008, 7, 189-197.	2.5	131
12	Molecular features of the CAG repeats of spinocerebellar ataxia 6 (SCA6). Human Molecular Genetics, 1997, 6, 1283-1287.	2.9	129
13	Linear ubiquitination is involved in the pathogenesis of optineurin-associated amyotrophic lateral sclerosis. Nature Communications, 2016, 7, 12547.	12.8	109
14	Direct Alteration of the P/Q-Type Ca²⁺Channel Property by Polyglutamine Expansion in Spinocerebellar Ataxia 6. Journal of Neuroscience, 1999, 19, RC14-RC14.	3.6	107
15	Structure and organization of the gene encoding human dopamine transporter. Gene, 1997, 195, 11-18.	2.2	98
16	Difference in disease-free survival curve and regional distribution according to subtype of spinocerebellar ataxia: A study of 1,286 Japanese patients. American Journal of Medical Genetics Part A, 2002, 114, 578-583.	2.4	97
17	Vulnerability of Purkinje Cells Generated from Spinocerebellar Ataxia Type 6 Patient-Derived iPSCs. Cell Reports, 2016, 17, 1482-1490.	6.4	91
18	Mutations in Twinkle primase-helicase cause Perrault syndrome with neurologic features. Neurology, 2014, 83, 2054-2061.	1.1	86

#	Article	IF	Citations
19	Lack of replication of thirteen single-nucleotide polymorphisms implicated in Parkinson's disease: a large-scale international study. Lancet Neurology, The, 2006, 5, 917-923.	10.2	83
20	Possible Reduced Penetrance of Expansion of 44 to 47 CAG/CAA Repeats in the TATA-Binding Protein Gene in Spinocerebellar Ataxia Type 17. Archives of Neurology, 2004, 61, 209.	4.5	81
21	Optineurin suppression causes neuronal cell death via <scp>NF</scp>â€ÎºB pathway. Journal of Neurochemistry, 2013, 126, 699-704.	3.9	80
22	A mutation in the low voltage-gated calcium channel CACNA1G alters the physiological properties of the channel, causing spinocerebellar ataxia. Molecular Brain, 2015, 8, 89.	2.6	80
23	Aortic Pulse Wave Velocity Predicts Cardiovascular Mortality in Middle-Aged and Elderly Japanese Men. Circulation Journal, 2009, 73, 549-553.	1.6	78
24	Characteristic Magnetic Resonance Imaging Findings in Spinocerebellar Ataxia 6. Archives of Neurology, 1998, 55, 1348.	4.5	77
25	Contribution of the interleukinâ€1Î² gene polymorphism in multiple system atrophy. Movement Disorders, 2002, 17, 808-811.	3.9	68
26	Genetic studies in Parkinson's disease with an Î±-synuclein/NACP gene polymorphism in Japan. Neuroscience Letters, 2001, 300, 125-127.	2.1	65
27	SCA8 Repeat Expansion: Large CTA/CTG Repeat Alleles Are More Common in Ataxic Patients, Including Those with SCA6. American Journal of Human Genetics, 2003, 72, 704-709.	6.2	65
28	Mutant Protein Kinase CÎ³ Found in Spinocerebellar Ataxia Type 14 Is Susceptible to Aggregation and Causes Cell Death. Journal of Biological Chemistry, 2005, 280, 29096-29106.	3.4	64
29	NO ASSOCIATION OF COMPLEMENT FACTOR H GENE POLYMORPHISM AND AGE-RELATED MACULAR DEGENERATION IN THE JAPANESE POPULATION. Retina, 2006, 26, 985-987.	1.7	62
30	LRRK2 mutations and risk variants in Japanese patients with Parkinson's disease. Movement Disorders, 2009, 24, 1034-1041.	3.9	60
31	Clinicopathologic study on an ALS family with a heterozygous E478G optineurin mutation. Acta Neuropathologica, 2011, 122, 223-229.	7.7	60
32	Cloning and Expression of a Human Glutamate Transporter. Biochemical and Biophysical Research Communications, 1994, 199, 171-176.	2.1	56
33	Homozygosity for Machado-Joseph disease gene enhances phenotypic severity.. Journal of Neurology, Neurosurgery and Psychiatry, 1996, 60, 354-356.	1.9	55
34	Cerebellar ataxia with SYNE1 mutation accompanying motor neuron disease. Neurology, 2013, 80, 600-601.	1.1	55
35	Identification and haplotype analysis of LRRK2 G2019S in Japanese patients with Parkinson disease. Neurology, 2006, 67, 697-699.	1.1	54
36	Optineurin is co-localized with FUS in basophilic inclusions of ALS with FUS mutation and in basophilic inclusion body disease. Acta Neuropathologica, 2011, 121, 555-557.	7.7	53

#	Article	IF	Citations
37	Genomeâ€wide association study in musician's dystonia: A risk variant at the arylsulfatase G locus?. Movement Disorders, 2014, 29, 921-927.	3.9	53
38	HLA and T-cell receptor gene polymorphisms in Guillain-Barrè•syndrome. Neurology, 1998, 51, 379-384.	1.1	52
39	Cerebral white matter lesions are not associated with apoE genotype but with age and female sex in Alzheimer's disease. Journal of Neurology, Neurosurgery and Psychiatry, 2000, 68, 653-656.	1.9	51
40	Influence of polymorphisms in the genes for cytokines and glutathione S-transferase omega on sporadic Alzheimerâ€™s disease. Neuroscience Letters, 2004, 368, 140-143.	2.1	51
41	Coagulation and vascular abnormalities in Crow-Fukase syndrome. , 1997, 20, 486-492.		50
42	Lack of an association between cystatin C gene polymorphisms in Japanese patients with Alzheimerâ€™s disease. Neurology, 2001, 57, 337-339.	1.1	45
43	Screening for TARDBP mutations in Japanese familial amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 2009, 284, 69-71.	0.6	45
44	Apolipoprotein E promoter polymorphism and sporadic Alzheimer's disease in a Japanese population. Neuroscience Letters, 1999, 259, 56-58.	2.1	44
45	Molecular Cloning and Characterization of a cDNA Encoding a Novel Basic Helix-Loop-Helix Protein Structurally Related to NeuroD/BHF1. Biochemical and Biophysical Research Communications, 1996, 220, 754-758.	2.1	43
46	A polymorphism of LOC387715 gene is associated with age-related macular degeneration in the Japanese population. Neuroscience Letters, 2007, 414, 71-74.	2.1	43
47	Optineurin and amyotrophic lateral sclerosis. Geriatrics and Gerontology International, 2013, 13, 528-532.	1.5	43
48	Expression of three glutamate transporter subtype mRNAs in human brain regions and peripheral tissues. Molecular Brain Research, 1996, 36, 189-192.	2.3	42
49	Unique features of the CAG repeats in Machadoâ€“Joseph disease. Nature Genetics, 1995, 9, 344-345.	21.4	41
50	HLA-DRB1 and tumor necrosis factor gene polymorphisms in Japanese patients with multiple sclerosis. Journal of Neuroimmunology, 1998, 92, 109-112.	2.3	40
51	Organization of the human orphan nuclear receptor Nurr1 gene. Gene, 1999, 230, 225-232.	2.2	40
52	Influence of a tumor necrosis factor gene polymorphism in Japanese patients with multiple system atrophy. Neuroscience Letters, 2005, 374, 218-221.	2.1	40
53	A single nucleotide polymorphism of dopamine transporter gene is associated with Parkinson's disease. Annals of Neurology, 2000, 47, 528-531.	5.3	39
54	Disinhibition of the somatosensory cortex in cervical dystoniaâ€”decreased amplitudes of high-frequency oscillations. Clinical Neurophysiology, 2004, 115, 1624-1630.	1.5	39

#	Article	IF	Citations
55	Glutathioneâ€Sâ€transferaseâ€1 and interleukinâ€1Î² gene polymorphisms in Japanese patients with Parkinson's disease. Movement Disorders, 2005, 20, 901-902.	3.9	39
56	Brainâ€derived neurotrophic factor gene polymorphisms in Japanese patients with sporadic Alzheimer's disease, Parkinson's disease, and multiple system atrophy. Movement Disorders, 2005, 20, 1031-1033.	3.9	39
57	Immunoreactivity of valosin-containing protein in sporadic amyotrophic lateral sclerosis and in a case of its novel mutant. Acta Neuropathologica Communications, 2014, 2, 172.	5.2	39
58	A new mitochondrial DNA mutation associated with mitochondrial myopathy: tRNA ^{Leu(UUR)} 3254C-to-G. Neurology, 1997, 49, 598-600.	1.1	37
59	Influence of monocyte chemoattractant protein 1 gene polymorphism on age at onset of sporadic Parkinson's disease. Movement Disorders, 2003, 18, 953-955.	3.9	37
60	Clinicopathologic features of autosomal recessive amyotrophic lateral sclerosis associated with optineurin mutation. Neuropathology, 2014, 34, 64-70.	1.2	37
61	Novel mutation in X-linked Charcot-Marie-Tooth disease associated with CNS impairment. Neurology, 2002, 59, 923-926.	1.1	34
62	Clinicopathologic investigation of a family with expanded SCA8 CTA/CTG repeats. Neurology, 2006, 67, 1479-1481.	1.1	34
63	Genetic contribution of the tumor necrosis factor region in guillainâ€barrÃ© syndrome. Annals of Neurology, 1998, 44, 815-818.	5.3	32
64	Dinucleotide repeat polymorphisms in the Neprilysin gene are not associated with sporadic Alzheimer's disease. Neuroscience Letters, 2002, 320, 105-107.	2.1	32
65	PLK1-mediated phosphorylation of WDR62/MCPH2 ensures proper mitotic spindle orientation. Human Molecular Genetics, 2017, 26, 4429-4440.	2.9	32
66	Voltage-gated potassium channel antibodies associated limbic encephalitis in a patient with invasive thymoma. Journal of the Neurological Sciences, 2006, 250, 167-169.	0.6	31
67	Exome sequencing reveals a novel TTC19 mutation in an autosomal recessive spinocerebellar ataxia patient. BMC Neurology, 2014, 14, 5.	1.8	30
68	Quantitative Assessment of Cerebral Blood Flow in Genetically Confirmed Spinocerebellar Ataxia Type 6. Archives of Neurology, 2004, 61, 933.	4.5	29
69	Identification of a new family of spinocerebellar ataxia type 14 in the japanese spinocerebellar ataxia population by the screening of PRKCG exon 4. Movement Disorders, 2006, 21, 1355-1360.	3.9	29
70	Optineurin with amyotrophic lateral sclerosis-related mutations abrogates inhibition of interferon regulatory factor-3 activation. Neuroscience Letters, 2011, 505, 279-281.	2.1	29
71	HLAs and genes in japanese patients with multiple sclerosis: evidence for increased frequencies of HLA-Cw3, HLA-DR2, and HLA-DQB1â^—0602. Human Immunology, 1992, 35, 116-124.	2.4	27
72	CAG repeat length and disease duration in Machado-Joseph disease: a new clinical classification. Journal of the Neurological Sciences, 1997, 152, 166-171.	0.6	27

#	Article	IF	Citations
73	Spinocerebellar ataxia type 6: MRI of three Japanese patients. Neuroradiology, 1998, 40, 222-227.	2.2	27
74	Effects of nerve growth factor and nicotine on the expression of nicotinic acetylcholine receptor subunits in PC12 cells. Neuroscience Research, 1999, 35, 175-181.	1.9	27
75	Tumor necrosis factor, tumor necrosis factor receptors type 1 and 2, lymphotoxin-Î±, and HLA-DRB1 gene polymorphisms in human T-Cell lymphotropic virus type I associated myelopathy. Human Immunology, 2000, 61, 1262-1269.	2.4	26
76	Localization of the gene (SLC1A3) encoding human glutamate transporter (GluT-1) to 5p13 by fluorescence in situ hybridization. Cytogenetic and Genome Research, 1995, 69, 209-210.	1.1	25
77	Association between interleukin-6 gene polymorphism and human T-Cell leukemia virus type I associated myelopathy. Human Immunology, 2002, 63, 696-700.	2.4	25
78	Multiple Proteinopathies in Familial ALS Cases With Optineurin Mutations. Journal of Neuropathology and Experimental Neurology, 2018, 77, 128-138.	1.7	25
79	The clinical characteristics of spinocerebellar ataxia 36: A study of 2121 Japanese ataxia patients. Movement Disorders, 2012, 27, 1158-1163.	3.9	22
80	Exome sequencing reveals a novel <scp>ANO10</scp> mutation in a Japanese patient with autosomal recessive spinocerebellar ataxia. Clinical Genetics, 2014, 85, 296-297.	2.0	22
81	Second derivative of the finger photoplethysmogram and cardiovascular mortality in middle-aged and elderly Japanese women. Hypertension Research, 2017, 40, 207-211.	2.7	22
82	Prediction Model of Amyotrophic Lateral Sclerosis by Deep Learning with Patient Induced Pluripotent Stem Cells. Annals of Neurology, 2021, 89, 1226-1233.	5.3	22
83	Reevaluation of the exact CAG repeat length in hereditary cerebellar ataxias using highly denaturing conditions and long PCR. Human Genetics, 1996, 97, 591-595.	3.8	21
84	Structure and regulation of the human NeuroD (BETA2/BHF1) gene. Molecular Brain Research, 1999, 69, 223-231.	2.3	21
85	FXTAS is difficult to differentiate from neuronal intranuclear inclusion disease through skin biopsy: a case report. BMC Neurology, 2021, 21, 396.	1.8	21
86	Screening for OPTN mutations in amyotrophic lateral sclerosis in a mainly Caucasian population. Neurobiology of Aging, 2011, 32, 1923.e9-1923.e10.	3.1	20
87	An autopsy case of sporadic amyotrophic lateral sclerosis associated with the <scp>I113T</scp>â€¦<scp>SOD1</scp> mutation. Neuropathology, 2014, 34, 58-63.	1.2	19
88	Detecting gene mutations in Japanese Alzheimer's patients by semiconductor sequencing. Neurobiology of Aging, 2014, 35, 1780.e1-1780.e5.	3.1	19
89	Influence of cytokine and mannose binding protein gene polymorphisms on human t-cell leukemia virus type i (hTLV-i) provirus load in HTLV-I asymptomatic carriers. Human Immunology, 2003, 64, 453-457.	2.4	17
90	First report of a Japanese family with spinocerebellar ataxia type 10: The second report from Asia after a report from China. PLoS ONE, 2017, 12, e0177955.	2.5	17

#	Article	IF	Citations
91	Inhibition by folded isomers of L-2-(carboxycyclopropyl) glycine of glutamate uptake via the human glutamate transporter hGluT-1. European Journal of Pharmacology, 1995, 289, 387-390.	2.6	16
92	Exome sequencing reveals a novel MRE11 mutation in a patient with progressive myoclonic ataxia. Journal of the Neurological Sciences, 2014, 337, 219-223.	0.6	16
93	Electrophysiological studies in spinocerebellar ataxia type 6. NeuroReport, 2000, 11, 969-972.	1.2	15
94	Oromandibular dystonia associated with SCA36. Movement Disorders, 2013, 28, 558-559.	3.9	15
95	Optineurin defects cause TDP43-pathology with autophagic vacuolar formation. Neurobiology of Disease, 2021, 148, 105215.	4.4	15
96	Neuroprotective mechanism of bromocriptine. Lancet, The, 1995, 346, 1305.	13.7	14
97	Effect of amino acid ergot alkaloids on glutamate transport via human glutamate transporter hGluT-1. Journal of the Neurological Sciences, 1998, 155, 31-36.	0.6	14
98	Compound heterozygote mutations in the SIGMAR1 gene in an oldestâ€old patient with amyotrophic lateral sclerosis. Geriatrics and Gerontology International, 2018, 18, 1519-1520.	1.5	14
99	Cloning and Expression of a Rat Brain Basic Helixâ€“Loopâ€“Helix Factor. Biochemical and Biophysical Research Communications, 1996, 221, 199-204.	2.1	13
100	A family with Machado-Joseph disease, previously diagnosed as dentatorubral-pallidoluysian atrophy. Neurology, 1996, 46, 1154-1156.	1.1	13
101	Nicotinic acetylcholine receptors and neurodegenerative disease. Alcohol, 2001, 24, 79-81.	1.7	13
102	Investigation on circular asymmetry of geographical distribution in cancer mortality of Hiroshima atomic bomb survivors based on risk maps: analysis of spatial survival data. Radiation and Environmental Biophysics, 2012, 51, 133-141.	1.4	12
103	An autopsy report of a familial amyotrophic lateral sclerosis case carrying VCP Arg487His mutation with a unique TDPâ€43 proteinopathy. Neuropathology, 2021, 41, 118-126.	1.2	11
104	Spinocerebellar ataxia type 6 in relation to CAG repeat length. Acta Neurologica Scandinavica, 1999, 99, 209-212.	2.1	10
105	Co-morbidity of progressive supranuclear palsy and amyotrophic lateral sclerosis: a clinical-pathological case report. BMC Neurology, 2019, 19, 168.	1.8	10
106	Middle-age-onset cerebellar ataxia caused by a homozygous TWNK variant: a case report. BMC Medical Genetics, 2020, 21, 68.	2.1	10
107	A necropsied case of Machado-Joseph disease with a hyperintense signal of transverse pontine fibres on long TR sequences of magnetic resonance images. Journal of Neurology, Neurosurgery and Psychiatry, 1998, 64, 140-141.	1.9	9
108	A quantitatively-modeled homozygosity mapping algorithm, qHomozygosityMapping, utilizing whole genome single nucleotide polymorphism genotyping data. BMC Bioinformatics, 2010, 11, S5.	2.6	9

#	Article	IF	Citations
109	Neuropathological features of <scp>J</scp>apanese familial amyotrophic lateral sclerosis with p.<scp>N</scp>352<scp>S</scp> mutation in <scp>TARDBP</scp>. Neuropathology and Applied Neurobiology, 2014, 40, 231-236.	3.2	9
110	Homozygosity Mapping on Homozygosity Haplotype Analysis to Detect Recessive Disease-Causing Genes from a Small Number of Unrelated, Outbred Patients. PLoS ONE, 2011, 6, e25059.	2.5	9
111	Structural Organization and Expression of the Gene for Bovine Myosin I Heavy Chain1. Journal of Biochemistry, 1992, 111, 302-309.	1.7	8
112	Influence of interleukin-1Î² gene polymorphism on age-at-onset of spinocerebellar ataxia 6 (SCA6) in Japanese patients. Neuroscience Letters, 2001, 307, 128-130.	2.1	8
113	Pathogenic expansions of the SCA6 locus are associated with a common CACNA1A haplotype across the globe: founder effect or predisposing chromosome?. European Journal of Human Genetics, 2008, 16, 841-847.	2.8	8
114	The CNTN4 c.4256C>T mutation is rare in Japanese with inherited spinocerebellar ataxia. Journal of the Neurological Sciences, 2008, 266, 180-181.	0.6	8
115	Autosomal recessive Andersen-Tawil syndrome with a novel mutation L94P in Kir2.1. Neurology and Clinical Neuroscience, 2013, 1, 131-137.	0.4	8
116	RELATIVE BIOLOGICAL EFFECTIVENESS OF NEUTRONS DERIVED FROM THE EXCESS RELATIVE RISK MODEL WITH THE ATOMIC BOMB SURVIVORS DATA MANAGED BY HIROSHIMA UNIVERSITY. Radiation Protection Dosimetry, 2018, 180, 346-350.	0.8	8
117	Genetic screening for potassium channel mutations in Japanese autosomal dominant spinocerebellar ataxia. Journal of Human Genetics, 2020, 65, 363-369.	2.3	8
118	Optineurin regulates osteoblastogenesis through STAT1. Biochemical and Biophysical Research Communications, 2020, 525, 889-894.	2.1	8
119	Cytokines and myelin antibodies in Crow-Fukase syndrome. , 1996, 19, 1620-1622.		7
120	A <scp>J</scp>apanese patient with familial <scp>ALS</scp> and a p.<scp>K510M</scp> mutation in the gene for <scp>FUS</scp> (<scp>FUS</scp>) resulting in the totally lockedâ€in state. Neuropathology, 2014, 34, 504-509.	1.2	7
121	Novel compound heterozygous mutations in the <scp>PARK</scp>2 gene identified in a Chinese pedigree with earlyâ€onset Parkinson's disease. Brain and Behavior, 2018, 8, e00901.	2.2	7
122	Aggressive periodontitis and NOD2 variants. Journal of Human Genetics, 2020, 65, 841-846.	2.3	7
123	Kv11 (etherâ€Âà€goâ€goâ€related gene) voltageâ€dependent K⁺ channels promote resonance and oscillation of subthreshold membrane potentials. Journal of Physiology, 2021, 599, 547-569.	2.9	7
124	No association between apolipoprotein E alleles and olivopontocerebellar atrophy. Journal of the Neurological Sciences, 1998, 158, 110-112.	0.6	6
125	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. Journal of Medical Case Reports, 2011, 5, 573.	0.8	6
126	Retinitis pigmentosa prior to familial ALS caused by a homozygous cilia and flagella-associated protein 410 mutation. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 220-222.	1.9	6

#	Article	IF	Citations
127	Biallelic mutation of HSD17B4 induces middle ageâ€“onset spinocerebellar ataxia. Neurology: Genetics, 2020, 6, e396.	1.9	6
128	Identification and analysis of the promoter region of the human NeuroD-related factor (NDRF). Biochimica Et Biophysica Acta Gene Regulatory Mechanisms, 1999, 1445, 142-147.	2.4	5
129	The UCHL1 S18Y polymorphism and Parkinsonâ€™s disease in a Japanese population. Parkinsonism and Related Disorders, 2011, 17, 473-475.	2.2	5
130	Clinical Manifestations of Autosomal Recessive Early-Onset Parkinsonism with Diurnal Fluctuation. Advances in Behavioral Biology, 1996, , 485-489.	0.2	5
131	The first Japanese case of primary familial brain calcification caused by an MYORG variant. Journal of Human Genetics, 2020, 65, 917-920.	2.3	5
132	Ciliary Neurotrophic Factor Induced Increase in Î²-Amyloid Precursor Protein mRNA in Rat C6 Glioma Cells. Biochemical and Biophysical Research Communications, 1994, 204, 391-398.	2.1	4
133	A novel haplotype of spinocerebellar ataxia type 6 contributes to the highest prevalence in Western Japan. Neuroscience Letters, 2004, 358, 107-110.	2.1	4
134	Activation of human SII cortex during exploratory finger movement and hand clenching tasks. NeuroReport, 2005, 16, 145-148.	1.2	4
135	C-terminal mutations in SYNE1 are associated with motor neuron disease in patients with SCAR8. Journal of the Neurological Sciences, 2019, 402, 118-120.	0.6	4
136	Long-term MRI findings of adult-onset neuronal intranuclear inclusion disease. Clinical Neurology and Neurosurgery, 2021, 201, 106456.	1.4	4
137	Analysis of genetic risk factors in Japanese patients with Parkinsonâ€™s disease. Journal of Human Genetics, 2021, 66, 957-964.	2.3	4
138	Serum deprivation alters the expression and the splicing at exons 7, 8 and 15 of the Î²-amyloid precursor protein in the C6 glioma cell line. Molecular Brain Research, 1996, 39, 12-22.	2.3	3
139	Dinucleotide repeat polymorphism in interferon-? gene is not associated with sporadic Alzheimer's disease. American Journal of Medical Genetics Part A, 2004, 124B, 48-49.	2.4	3
140	Autopsy Case of Later-Onset Pontocerebellar Hypoplasia Type 1: Pontine Atrophy and Pyramidal Tract Involvement. Journal of Child Neurology, 2010, 25, 1429-1434.	1.4	3
141	An autopsy case of familial amyotrophic lateral sclerosis withFUSR521G mutation. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2014, 15, 305-308.	1.7	3
142	Treatment of intractable resting tremor of spinocerebellar ataxia 42 with zonisamide. Journal of the Neurological Sciences, 2019, 396, 119-120.	0.6	3
143	Amyotrophic lateral sclerosis of long clinical course clinically presenting with progressive muscular atrophy. Neuropathology, 2019, 39, 47-53.	1.2	3
144	Zonisamide can ameliorate the voltage-dependence alteration of the T-type calcium channel CaV3.1 caused by a mutation responsible for spinocerebellar ataxia. Molecular Brain, 2020, 13, 163.	2.6	3

#	Article	IF	Citations
145	A single nucleotide polymorphism of dopamine transporter gene is associated with Parkinson's disease. Annals of Neurology, 2000, 47, 528-531.	5.3	3
146	Variation in the number of CAG repeats in the Machadoâ€"Joseph disease gene (MJD1) in the Japanese population. Journal of the Neurological Sciences, 1999, 166, 71-73.	0.6	2
147	Analysis on the Susceptibility Genes in Two Chinese Pedigrees with Familial Parkinson's Disease. Neurology Research International, 2010, 2010, 1-4.	1.3	2
148	DYT6 in Japanâ€"genetic screening and clinical characteristics of the patients. Movement Disorders, 2014, 29, 278-280.	3.9	2
149	Prospects and status of the dosimetry system for atomic bomb survivor cohort study conducted at Research Institute for Radiation Biology and Medicine of Hiroshima University. Journal of Radiation Research, 2021, 62, i107-i113.	1.6	2
150	Novel monoallelic variant in ERLIN2 causes spastic paraplegia converted to amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 2021, 430, 119984.	0.6	2
151	Knockdown of optineurin controls C2C12 myoblast differentiation via regulating myogenin and MyoD expressions. Differentiation, 2022, 123, 1-8.	1.9	2
152	Neurogenetics. Neurology, 2015, 84, 1070-1071.	1.1	1
153	Cytokines and myelin antibodies in Crowâ€Fukase syndrome. Muscle and Nerve, 1996, 19, 1620-1622.	2.2	1
154	Reevaluation of the exact CAG repeat length in hereditary cerebellar ataxias using highly denaturing conditions and long PCR. Human Genetics, 1996, 97, 591-595.	3.8	1
155	Localization of the gene encoding the human L-glutamate transporter (GLT-1) to 11p11.2-p13 by fluorescence in situ hybridization. Human Genetics, 1996, 97, 387-389.	3.8	1
156	Comparison of two families with and without ataxia harboring novel variants in PRKCG. Journal of Human Genetics, 0, , .	2.3	1
157	Authorsâ€™ reply to <scp>D</scp>rs <scp>M</scp> van <scp>B</scp>litterswijk, <scp>R R</scp>ademakers and <scp>LH</scp> van den <scp>B</scp>erg. Neuropathology and Applied Neurobiology, 2014, 40, 359-360.	3.2	0
158	ã,¢ãƒ«ãƒ„ãƒã,ãƒžãƒ¼ç—…ã«é–¢ã™ã,‹ã,°ãƒ«ã,¿ãƒŸãƒ³é…¸è¼¸é€è›‹ç™½ã®å^†å¢"Ÿç‰©å¦çš„ç"ç©¶. Japanese Journal of Geriatrics, 20	0.1	0
159	Risk factors for dementia. , 2001, , 279-286.		0
160	Dopamine Transporter and Parkinsonâ€™s Disease. Advances in Behavioral Biology, 2002, , 467-470.	0.2	0
161	Convenient diagnosis of spinal and bulbar muscular atrophy using a microchip electrophoresis system. American Journal of Neurodegenerative Disease, 2013, 2, 35-9.	0.1	0
162	Channelopathies and Cerebellar Disease. , 2022, , 1399-1413.		0