Yuichi Hayashi

List of Publications by Year in descending order

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84 papers	1,354 citations	430754 18 h-index	395590 33 g-index
92 all docs	92 docs citations	92 times ranked	2168 citing authors

#	Article	IF	Citations
1	Progressive supranuclear palsy with predominant frontal presentation exhibiting progressive nonfluent aphasia due to crossed aphasia. Neuropathology, 2022, , .	0.7	1
2	Specific electroencephalogram features in the very early phases of sporadic Creutzfeldtâ€“Jakob disease. Journal of the Neurological Sciences, 2022, 437, 120265.	0.3	1
3	Clinicopathological findings of a mitochondrial encephalopathy, lactic acidosis, and strokeâ€like episodes/Leigh syndrome overlap patient with a novel m. 3482A>G mutation in MTâ€ND1. Neuropathology, 2021, 41, 84-90.	0.7	4
4	Long-term preservation of pharyngeal swallowing function in MM2-cortical-type sporadic Creutzfeldt-Jakob disease. Prion, 2021, 15, 82-86.	0.9	1
5	Clinical characteristics of intractable or persistent hiccups and nausea associated with herpes zoster. Clinical Neurology and Neurosurgery, 2021, 207, 106751.	0.6	2
6	Albuminâ€bilirubin score for predicting neuropsychiatric symptoms in patients receiving ifosfamideâ€based chemotherapy. Journal of Clinical Pharmacy and Therapeutics, 2021, 46, 794-799.	0.7	1
7	Brain Abscess Presenting as Prolonged Headache in a Patient with Amyotrophic Lateral Sclerosis under Mechanical Ventilation. Internal Medicine, 2020, 59, 581-583.	0.3	0
8	Focal sharp waves are a specific early-stage marker of the MM2-cortical form of sporadic Creutzfeldt-Jakob disease. Prion, 2020, 14, 207-213.	0.9	3
9	Clinicopathological findings of a long-term survivor of V180I genetic Creutzfeldt-Jakob disease. Prion, 2020, 14, 109-117.	0.9	11
10	Serial evaluation of swallowing function in a long-term survivor of V180I genetic Creutzfeldt-Jakob disease. Prion, 2020, 14, 180-184.	0.9	6
11	Autoimmune Glial Fibrillary Acidic Protein Astrocytopathy Presenting with Slowly Progressive Myelitis and Longitudinally Extensive Spinal Cord Lesions. Internal Medicine, 2020, 59, 2777-2781.	0.3	7
12	Appropriate Medication in Elderly Patients. The Journal of the Japanese Society of Internal Medicine, 2020, 109, 2206-2214.	0.0	0
13	Cytokines and biological markers in autoimmune GFAP astrocytopathy: The potential role for pathogenesis and therapeutic implications. Journal of Neuroimmunology, 2019, 334, 576999.	1.1	18
14	Adverse event profiles of ifosfamide-induced encephalopathy analyzed using the Food and Drug Administration Adverse Event Reporting System and the Japanese Adverse Drug Event Report databases. Cancer Chemotherapy and Pharmacology, 2019, 84, 1097-1105.	1.1	10
15	Clinicopathological findings of an MM2-cortical-type sporadic Creutzfeldt-Jakob disease patient with cortical blindness during a course of glaucoma and age-related macular degeneration. Prion, 2019, 13, 124-131.	0.9	5
16	Functional evaluation of PDGFB-variants in idiopathic basal ganglia calcification, using patient-derived iPS cells. Scientific Reports, 2019, 9, 5698.	1.6	8
17	Authors' Reply: Mixed Alzheimer's and Lewy-related Pathology Can Cause Corticobasal Syndrome with Visual Hallucinations. Internal Medicine, 2019, 58, 1815-1815.	0.3	1
18	Clinical characteristics of autoimmune GFAP astrocytopathy. Journal of Neuroimmunology, 2019, 332, 91-98.	1.1	94

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19	Pathological and/or clinical workâ€up are required in atypical Creutzfeldtâ€“Jakob disease cases with periodic lateralised epileptiform discharge. Psychogeriatrics, 2019, 19, 519-520.	0.6	1
20	Relationship between number of drugs and duration of hospital stay in older patients with neuromuscular diseases. Geriatrics and Gerontology International, 2018, 18, 1018-1024.	0.7	3
21	Long-term clinical follow-up of a patient with non-paraneoplastic cerebellar ataxia associated with anti-mGluR1 autoantibodies. Journal of Neuroimmunology, 2018, 319, 63-67.	1.1	13
22	Higher levels of progranulin in cerebrospinal fluid of patients with lymphoma and carcinoma with CNS metastasis. Journal of Neuro-Oncology, 2018, 137, 455-462.	1.4	9
23	Inorganic phosphorus (Pi) in CSF is a biomarker for SLC20A2-associated idiopathic basal ganglia calcification (IBGC1). Journal of the Neurological Sciences, 2018, 388, 150-154.	0.3	32
24	Time-Dependent Decline in Serum Phenytoin Concentration With Heightened Convulsive Seizure Risk by Prolonged Administration of Fosphenytoin in Japanese: A Retrospective Study. Therapeutic Drug Monitoring, 2018, 40, 507-511.	1.0	2
25	Drug-induced Pressure Ulcers in a Middle-aged Patient with Early-stage Parkinson's Disease. Internal Medicine, 2018, 57, 1483-1486.	0.3	4
26	Diagnosing Corticobasal Syndrome Based on the Presence of Visual Hallucinations and Imaging with Amyloid Positron Emission Tomography. Internal Medicine, 2018, 57, 605-611.	0.3	2
27	Reduction in the numbers of drugs administered to elderly inâ€patients with polypharmacy by a multidisciplinary review of medication using electronic medical records. Geriatrics and Gerontology International, 2017, 17, 653-658.	0.7	12
28	Reversible splenial lesion of the corpus callosum associated with meningococcal meningitis. Journal of the Neurological Sciences, 2017, 373, 81-82.	0.3	7
29	Comparison of cerebrospinal fluid profiles in Alzheimerâ€™s disease with multiple cerebral microbleeds and cerebral amyloid angiopathy-related inflammation. Journal of Neurology, 2017, 264, 373-381.	1.8	11
30	Neuropathological findings from an autopsied case showing posterior reversible encephalopathy syndrome-like neuroradiological findings associated with premedication including tacrolimus for autologous peripheral blood stem cell transplantation. Journal of the Neurological Sciences, 2017, 375, 382-387.	0.3	4
31	Reply to: Amyotrophic lateral sclerosis with frontotemporal dementia (ALS-FTD) syndrome as a phenotype of Creutzfeldt-Jakob disease (CJD)? A case report. Journal of the Neurological Sciences, 2017, 375, 489.	0.3	3
32	Association between naturally occurring anti-amyloid Î² autoantibodies and medial temporal lobe atrophy in Alzheimer's disease. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 126-131.	0.9	12
33	Case of adult-onset neuronal intranuclear hyaline inclusion disease with negative electroretinogram. Documenta Ophthalmologica, 2017, 134, 221-226.	1.0	17
34	IVIG treatment for repeated hypothermic attacks associated with LGI1 antibody encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2017, 4, e348.	3.1	1
35	Central hypothermia associated with Alexander disease. A case report. Clinical Neurology and Neurosurgery, 2017, 157, 31-33.	0.6	4
36	Clinical findings of a probable case of MM2-cortical-type sporadic Creutzfeldt-Jakob disease with antibodies to anti-N-terminus of Î±-enolase. Prion, 2017, 11, 454-464.	0.9	9

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37	Anti-C1q autoantibodies in patients with neuromyelitis optica spectrum disorders. Journal of Neuroimmunology, 2017, 310, 150-157.	1.1	6
38	Acuteâ€Onset Severe Occipital Neuralgia Associated With High Cervical Lesion in Patients With Neuromyelitis Optica Spectrum Disorder. Headache, 2017, 57, 1145-1151.	1.8	9
39	A multidisciplinary medical network approach is crucial for increasing the number of autopsies for prion disease [Reply to: How can we increase the numbers of autopsies for prion disease? A model system in Japan]. Journal of the Neurological Sciences, 2017, 377, 95-96.	0.3	4
40	An autopsy-verified case of steroid-responsive encephalopathy with convulsion and a false-positive result from the real-time quaking-induced conversion assay. Prion, 2017, 11, 284-292.	0.9	17
41	Pathological examination is required for the case of rapidly progressive dementia with only positive result of RT-QUIC assay. Prion, 2017, 11, 469-470.	0.9	2
42	Cerebrospinal Fluid C-C Motif Chemokine Ligand 2 Correlates with Brain Atrophy and Cognitive Impairment in Alzheimerâ€™s Disease. Journal of Alzheimer's Disease, 2017, 61, 581-588.	1.2	18
43	Sarcopenia. Journal of the Japanese Society for Food Science and Technology, 2017, 64, 515-516.	0.1	0
44	Increased cerebrospinal fluid progranulin correlates with interleukin-6 in the acute phase of neuromyelitis optica spectrum disorder. Journal of Neuroimmunology, 2017, 305, 175-181.	1.1	21
45	Frailty. Journal of the Japanese Society for Food Science and Technology, 2017, 64, 446-446.	0.1	0
46	Polypharmacy. Journal of the Japanese Society for Food Science and Technology, 2017, 64, 517-518.	0.1	0
47	Improvement of memory recall by quercetin in rodent contextual fear conditioning and human early-stage Alzheimerâ€™s disease patients. NeuroReport, 2016, 27, 671-676.	0.6	36
48	An autopsy-verified case of FTLD-TDP type A with upper motor neuron-predominant motor neuron disease mimicking MM2-thalamic-type sporadic Creutzfeldt-Jakob disease. Prion, 2016, 10, 492-501.	0.9	13
49	Preserved regional cerebral blood flow in the occipital cortices, brainstem, and cerebellum of patients with V180I-129M genetic Creutzfeldt-Jakob disease in serial SPECT studies. Journal of the Neurological Sciences, 2016, 370, 145-151.	0.3	12
50	Identification of target antigens of naturally occurring autoantibodies in cerebrospinal fluid. Journal of Proteomics, 2015, 128, 450-457.	1.2	12
51	B-cell-activating factor belonging to the tumor necrosis factor family (BAFF) and a proliferation-inducing ligand (APRIL) levels in cerebrospinal fluid of patients with meningoencephalitis. Journal of the Neurological Sciences, 2015, 352, 79-83.	0.3	14
52	Decreased regional cerebral blood flow in the bilateral thalami and medulla oblongata determined by an easy Z-score (eZIS) analysis of 99mTc-ECD-SPECT images in a case of MM2-thalamic-type sporadic Creutzfeldtâ€“Jakob disease. Journal of the Neurological Sciences, 2015, 358, 447-452.	0.3	12
53	Evaluation of SLC20A2 mutations that cause idiopathic basal ganglia calcification in Japan. Neurology, 2014, 82, 705-712.	1.5	71
54	Corticosteroid therapy in a patient with cerebral amyloid angiopathy-related inflammation. Journal of Neuroinflammation, 2013, 10, 39.	3.1	27

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55	Effects of a polymorphism in the GFAP promoter on the age of onset and ambulatory disability in late-onset Alexander disease. Journal of Human Genetics, 2013, 58, 635-638.	1.1	11
56	Cardiac sympathetic function in the patients with amyotrophic lateral sclerosis: analysis using cardiac [123I] MIBG scintigraphy. Journal of Neurology, 2013, 260, 2380-2386.	1.8	35
57	Is there a difference in gastric emptying between myotonic dystrophy type 1 patients with and without gastrointestinal symptoms?. Journal of Neurology, 2013, 260, 1611-1616.	1.8	6
58	High frequency of calcification in basal ganglia on brain computed tomography images in <scp>J</scp>apanese older adults. Geriatrics and Gerontology International, 2013, 13, 706-710.	0.7	67
59	Decreased Bioelements Content in the Hair of Patients with Fahrâ€™s Disease (Idiopathic Bilateral) Tj ETQq1 1 0.784314 rgBT /Overlock	1.9	5
60	Intracranial Calcification in a Patient with HDR Syndrome and a <i>GATA3</i> Mutation. Internal Medicine, 2013, 52, 161-162.	0.3	3
61	Peduncular hallucinations in brainstem encephalitis drawn by a patient. Neurology, 2012, 79, 1625-1625.	1.5	3
62	Anti-Endothelial Cell Antibodies in Patients with Cerebral Small Vessel Disease. Current Neurovascular Research, 2012, 9, 296-301.	0.4	20
63	Antibodies Against the Tom40 Subunit of the Translocase of the Outer Mitochondrial Membrane Complex and Cognitive Impairment in Alzheimer's Disease. Journal of Alzheimer's Disease, 2012, 29, 373-377.	1.2	6
64	Late-onset Patients with Sporadic Amyotrophic Lateral Sclerosis in Japan have a Higher Progression Rate of ALSFRS-R at the Time of Diagnosis. Internal Medicine, 2012, 51, 579-584.	0.3	16
65	Elevated Antiâ€“Heat Shock Protein 60 Antibody Titer is Related to White Matter Hyperintensities. Journal of Stroke and Cerebrovascular Diseases, 2012, 21, 305-309.	0.7	7
66	Is there delayed gastric emptying in patients with multiple system atrophy? An analysis using the 13C-acetate breath test. Journal of Neurology, 2012, 259, 1448-1452.	1.8	21
67	Patterns of levels of biological metals in CSF differ among neurodegenerative diseases. Journal of the Neurological Sciences, 2011, 303, 95-99.	0.3	213
68	Diffuse Skeletal Muscles Uptake of [18F] Fluorodeoxyglucose on Positron Emission Tomography in Primary Muscle Peripheral T-cell Lymphoma. Internal Medicine, 2011, 50, 2021-2024.	0.3	5
69	Neuromyelitis Optica in Japanese Sisters. Internal Medicine, 2011, 50, 2829-2832.	0.3	8
70	Identification of antibodies as biological markers in serum from multiple sclerosis patients by immunoproteomic approach. Journal of Neuroimmunology, 2011, 233, 175-180.	1.1	14
71	Is there a delayed gastric emptying of patients with early-stage, untreated Parkinsonâ€™s disease? An analysis using the 13C-acetate breath test. Journal of Neurology, 2011, 258, 421-426.	1.8	69
72	Markedly Ring-enhanced Optic Nerves Due to Metastasis of Signet-ring Cell Gastric Carcinoma. Internal Medicine, 2010, 49, 517-517.	0.3	4

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73	Serial Monitoring of Basal Metabolic Rate for Therapeutic Evaluation in an Isaacs' Syndrome Patient with Chronic Fluctuating Symptoms. Internal Medicine, 2010, 49, 475-477.	0.3	3
74	Motor-dominant chronic inflammatory demyelinating polyneuropathy. Journal of Neurology, 2010, 257, 621-629.	1.8	34
75	High Levels of Copper, Zinc, Iron and Magnesium, but not Calcium, in the Cerebrospinal Fluid of Patients with Fahrâ€™s Disease. Case Reports in Neurology, 2010, 2, 46-51.	0.3	20
76	Nuclear changes in skeletal muscle extend to satellite cells in autosomal dominant Emery-Dreifuss muscular dystrophy/limb-girdle muscular dystrophy 1B. Neuromuscular Disorders, 2009, 19, 29-36.	0.3	57
77	DNA microarray analysis of transcriptional responses of mouse spinal cords to physical exercise. Journal of Toxicological Sciences, 2009, 34, 445-448.	0.7	7
78	Relationship between ribosomal RNA gene transcription activity and motoneuron death: Observations of avulsion and axotomy of the facial nerve in rats. Journal of Neuroscience Research, 2008, 86, 435-442.	1.3	3
79	Selective cauda equina hypertrophy with idiopathic inflammation. Muscle and Nerve, 2008, 38, 1065-1069.	1.0	1
80	Progressive multifocal leukoencephalopathy and CD4+ T-lymphocytopenia in a patient with SjÃ¶gren syndrome. Journal of the Neurological Sciences, 2008, 268, 195-198.	0.3	23
81	Adenoviral gene transfer of hepatocyte growth factor prevents death of injured adult motoneurons after peripheral nerve avulsion. Brain Research, 2006, 1111, 187-195.	1.1	18
82	Association between metallothionein genes polymorphisms and sporadic amyotrophic lateral sclerosis in a Japanese population. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2006, 7, 22-26.	2.3	24
83	Peripheral nerve avulsion injuries as experimental models for adult motoneuron degeneration. Neuropathology, 2005, 25, 371-380.	0.7	20
84	Motoneuron degeneration after facial nerve avulsion is exacerbated in presymptomatic transgenic rats expressing human mutant Cu/Zn superoxide dismutase. Journal of Neuroscience Research, 2005, 82, 63-70.	1.3	12