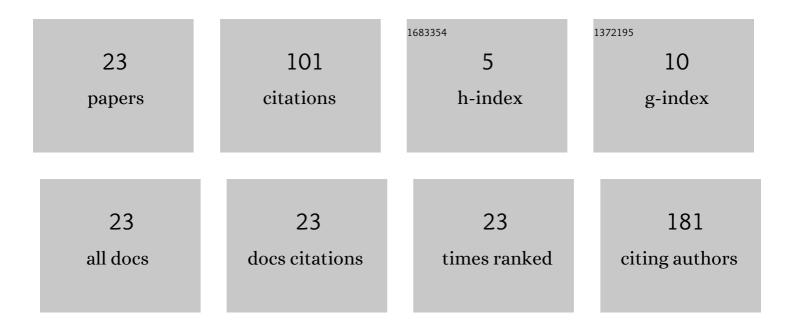
Yuko Kawahara

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Simultaneous assessment of cognitive and affective functions in multiple system atrophy and cortical cerebellar atrophy in relation to computerized touch-panel screening tests. Journal of the Neurological Sciences, 2015, 351, 24-30.	0.3	20
2	Early detection of cognitive decline in mild cognitive impairment and Alzheimer's disease with a novel eye tracking test. Journal of the Neurological Sciences, 2021, 427, 117529.	0.3	20
3	Cognitive and Affective Assessments of Multiple Sclerosis (MS) and Neuromyelitis Optica (NMO) Patients Utilizing Computerized Touch Panel-type Screening Tests. Internal Medicine, 2014, 53, 2281-2290.	0.3	16
4	Improvement of a decreased anti-oxidative activity by edaravone in amyotrophic lateral sclerosis patients. Journal of the Neurological Sciences, 2020, 415, 116906.	0.3	13
5	A patient develops transient unique cerebral and cerebellar lesions after unruptured aneurysm coiling. BMC Neurology, 2015, 15, 49.	0.8	10
6	4-Hydroxyl-2-Nonenal Localized Expression Pattern in Retrieved Clots is Associated with Large Artery Atherosclerosis in Stroke Patients. Journal of Stroke and Cerebrovascular Diseases, 2021, 30, 105583.	0.7	5
7	Marked hypertriglyceridemia induced by interferon- \hat{I}^2 1a therapy in a clinically isolated syndrome patient. Journal of the Neurological Sciences, 2017, 373, 144-146.	0.3	4
8	Aggressive Garcin's syndrome by acquired cystic disease of kidney-related renal cell carcinoma in a long-term hemodialytic patient. Journal of the Neurological Sciences, 2015, 355, 216-218.	0.3	3
9	Cognitive and affective functions of aged subacute myeloâ€optico neuropathy patients in Japan. Neurology and Clinical Neuroscience, 2015, 3, 173-178.	0.2	2
10	Asymmetrical and Isolated Hypoglossal Nerve Palsy Accompanied by a New Subset of Anti-ganglioside Antibodies in a Patient with Diffuse Large B Cell Lymphoma. Internal Medicine, 2019, 58, 283-286.	0.3	2
11	Type 2 Alexander disease with a novel glial fibrillary acidic protein gene mutation and its unique clinical features. Neurology and Clinical Neuroscience, 2017, 5, 183-185.	0.2	1
12	A Japanese patient with a VCP mutation c.290GÂ>ÂA (p.G97E) presenting a rapid progressive respiratory failure. Neurology and Clinical Neuroscience, 2019, 7, 361-364.	0.2	1
13	A unique case with positive antiâ€myelin oligodendrocyte glycoprotein antibody presenting multiple brain lesions. Neurology and Clinical Neuroscience, 2020, 8, 92-95.	0.2	1
14	The first case of chronic inflammatory demyelinating polyneuropathy after transsexualism and continuous testosterone administration. Neurology and Clinical Neuroscience, 2021, 9, 346-348.	0.2	1
15	A case of rheumatoid meningoencephalitis induced by pembrolizumab. Neurology and Clinical Neuroscience, 2021, 9, 470.	0.2	1
16	Three cases of GFAP astrocytopathy, one with bilateral ovarian teratoma. Neurology and Clinical Neuroscience, 0, , .	0.2	1
17	Multiâ€nodal combination therapy rescued a frequent ischemic stroke patient due to giant cell arteritis. Neurology and Clinical Neuroscience, 2019, 7, 132-135.	0.2	0
18	Very rare solitary primary peripheral nerve onset cytotoxic moleculeâ€positive peripheral Tâ€cell lymphoma (<scp>PTCL</scp>). Neurology and Clinical Neuroscience, 2019, 7, 146-149.	0.2	0

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#	Article	IF	CITATIONS
19	A unique case of myasthenia gravis mimicking Garcin's syndrome. Neurology and Clinical Neuroscience, 2020, 8, 399-402.	0.2	0
20	Successful treatment of antiâ€GAD antibodyâ€associated autoimmune cerebellar ataxia with combined immunotherapies. Neurology and Clinical Neuroscience, 0, , .	0.2	0
21	The Oldest Japanese Case of Combined Central and Peripheral Demyelination, which Developed Nine Years After the First Instance of Optic Neuritis. Internal Medicine, 2021, 60, 305-308.	0.3	0
22	A case of a heterozygous ABCC6 mutation showing recurrent ischemic strokes and intracranial hemorrhages. Neurology and Clinical Neuroscience, 0, , .	0.2	0
23	A case of successful renal transplantation of <scp>Charcotâ€Marieâ€Tooth</scp> disease associated with <scp>FSCS</scp> due to mutation of the <scp> <i>INF2</i> </scp> gene. Neurology and Clinical Neuroscience, 0, , .	0.2	0