Douglas Sato

List of Publications by Year in descending order

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Ποιιείλο δάτο

#	Article	IF	CITATIONS
1	Distinction between MOG antibody-positive and AQP4 antibody-positive NMO spectrum disorders. Neurology, 2014, 82, 474-481.	1.1	743
2	MRI characteristics of neuromyelitis optica spectrum disorder. Neurology, 2015, 84, 1165-1173.	1.1	523
3	MOG antibody–positive, benign, unilateral, cerebral cortical encephalitis with epilepsy. Neurology: Neuroimmunology and NeuroInflammation, 2017, 4, e322.	6.0	334
4	MOG cell-based assay detects non-MS patients with inflammatory neurologic disease. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e89.	6.0	322
5	Demographic and clinical features of neuromyelitis optica: A review. Multiple Sclerosis Journal, 2015, 21, 845-853.	3.0	278
6	Myelin-oligodendrocyte glycoprotein antibody-associated disease. Lancet Neurology, The, 2021, 20, 762-772.	10.2	261
7	Neuromyelitis optica and multiple sclerosis: Seeing differences through optical coherence tomography. Multiple Sclerosis Journal, 2015, 21, 678-688.	3.0	209
8	Myasthenia gravis and neuromyelitis optica spectrum disorder. Neurology, 2012, 78, 1601-1607.	1.1	177
9	MOG-IgG-Associated Optic Neuritis, Encephalitis, and Myelitis: Lessons Learned From Neuromyelitis Optica Spectrum Disorder. Frontiers in Neurology, 2018, 9, 217.	2.4	156
10	Current concept of neuromyelitis optica (NMO) and NMO spectrum disorders. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 922-930.	1.9	149
11	MRI and retinal abnormalities in isolated optic neuritis with myelin oligodendrocyte glycoprotein and aquaporin-4 antibodies: a comparative study: TableÂ1. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 446-448.	1.9	134
12	CSF cytokine profile in MOG-IgG+ neurological disease is similar to AQP4-IgG+ NMOSD but distinct from MS: a cross-sectional study and potential therapeutic implications. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 927-936.	1.9	116
13	Bilateral frontal cortex encephalitis and paraparesis in a patient with anti-MOG antibodies. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 534-536.	1.9	113
14	Treatment of MOG-IgG-associated disorder with rituximab: An international study of 121 patients. Multiple Sclerosis and Related Disorders, 2020, 44, 102251.	2.0	110
15	Update on biomarkers in neuromyelitis optica. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e134.	6.0	104
16	Aquaporin-4 antibody–positive cases beyond current diagnostic criteria for NMO spectrum disorders. Neurology, 2013, 80, 2210-2216.	1.1	98
17	Th17 Cells Pathways in Multiple Sclerosis and Neuromyelitis Optica Spectrum Disorders: Pathophysiological and Therapeutic Implications. Mediators of Inflammation, 2016, 2016, 1-11.	3.0	92
18	Myelin injury without astrocytopathy in neuroinflammatory disorders with MOG antibodies. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 1257-1259.	1.9	89

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19	Lesion length of optic neuritis impacts visual prognosis in neuromyelitis optica. Journal of Neuroimmunology, 2016, 293, 28-33.	2.3	81
20	Neuromyelitis optica should be classified as an astrocytopathic disease rather than a demyelinating disease. Clinical and Experimental Neuroimmunology, 2012, 3, 58-73.	1.0	79
21	Fulminant demyelinating encephalomyelitis. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e175.	6.0	75
22	Treatment of MOG antibody associated disorders: results of an international survey. Journal of Neurology, 2020, 267, 3565-3577.	3.6	64
23	Severe demyelination but no astrocytopathy in clinically definite neuromyelitis optica with anti-myelin-oligodendrocyte glycoprotein antibody. Multiple Sclerosis Journal, 2015, 21, 656-659.	3.0	63
24	Different etiologies and prognoses of optic neuritis in demyelinating diseases. Journal of Neuroimmunology, 2016, 299, 152-157.	2.3	63
25	Use of Advanced Magnetic Resonance Imaging Techniques in Neuromyelitis Optica Spectrum Disorder. JAMA Neurology, 2015, 72, 815.	9.0	59
26	Highly encephalitogenic aquaporin 4-specific T cells and NMO-IgG jointly orchestrate lesion location and tissue damage in the CNS. Acta Neuropathologica, 2015, 130, 783-798.	7.7	55
27	Severely exacerbated neuromyelitis optica rat model with extensive astrocytopathy by high affinity anti-aquaporin-4 monoclonal antibody. Acta Neuropathologica Communications, 2015, 3, 82.	5.2	54
28	Neuromyelitis Optica Spectrum Disorders. Neuroimaging Clinics of North America, 2017, 27, 251-265.	1.0	53
29	Changes in Th17 and regulatory T cells after fingolimod initiation to treat multiple sclerosis. Journal of Neuroimmunology, 2014, 268, 95-98.	2.3	50
30	Cerebrospinal fluid aquaporinâ€4 antibody levels in neuromyelitis optica attacks. Annals of Neurology, 2014, 76, 305-309.	5.3	49
31	Persistent MOG-IgG positivity is a predictor of recurrence in MOG-IgG-associated optic neuritis, encephalitis and myelitis. Multiple Sclerosis Journal, 2019, 25, 1907-1914.	3.0	45
32	Viral encephalitis: a practical review on diagnostic approach and treatment. Jornal De Pediatria, 2020, 96, 12-19.	2.0	44
33	Treatment of neuromyelitis optica: an evidence based review. Arquivos De Neuro-Psiquiatria, 2012, 70, 59-66.	0.8	43
34	Latin American consensus recommendations for management and treatment of neuromyelitis optica spectrum disorders in clinical practice. Multiple Sclerosis and Related Disorders, 2020, 45, 102428.	2.0	42
35	Influenza-associated MOG antibody-positive longitudinally extensive transverse myelitis: a case report. BMC Neurology, 2014, 14, 224.	1.8	40
36	Treatment and outcome of aquaporin-4 antibody–positive NMOSD. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	37

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37	Clinical Spectrum and Treatment of Neuromyelitis Optica Spectrum Disorders: Evolution and Current Status. Brain Pathology, 2013, 23, 647-660.	4.1	36
38	Clinical spectrum of inflammatory central nervous system demyelinating disorders associated with antibodies against myelin oligodendrocyte glycoprotein. Neurochemistry International, 2019, 130, 104319.	3.8	35
39	The clinical spectrum associated with myelin oligodendrocyte glycoprotein antibodies (anti-MOG-Ab) in Thai patients. Multiple Sclerosis Journal, 2016, 22, 964-968.	3.0	31
40	Clinical features and long-term outcome of a group of Japanese children with inflammatory central nervous system disorders and seropositivity to myelin-oligodendrocyte glycoprotein antibodies. Brain and Development, 2015, 37, 849-852.	1.1	30
41	Atypical presentations of neuromyelitis optica. Arquivos De Neuro-Psiquiatria, 2011, 69, 824-828.	0.8	29
42	Seronegative Neuromyelitis Optica Spectrum - The challenges on disease definition and pathogenesis. Arquivos De Neuro-Psiquiatria, 2014, 72, 445-450.	0.8	25
43	Idiopathic aquaporin-4 antibody negative longitudinally extensive transverse myelitis. Multiple Sclerosis Journal, 2015, 21, 710-717.	3.0	25
44	Serological markers associated with neuromyelitis optica spectrum disorders in South India. Annals of Indian Academy of Neurology, 2016, 19, 505.	0.5	24
45	Brazilian Consensus for the Treatment of Multiple Sclerosis: Brazilian Academy of Neurology and Brazilian Committee on Treatment and Research in Multiple Sclerosis. Arquivos De Neuro-Psiquiatria, 2018, 76, 539-554.	0.8	22
46	Correlation between the corpus callosum index and brain atrophy, lesion load, and cognitive dysfunction in multiple sclerosis. Multiple Sclerosis and Related Disorders, 2018, 20, 154-158.	2.0	20
47	Post-vaccination MDEM associated with MOG antibody in a subclinical Chlamydia infected boy. Brain and Development, 2016, 38, 690-693.	1.1	19
48	Role of Glutamatergic Excitotoxicity in Neuromyelitis Optica Spectrum Disorders. Frontiers in Cellular Neuroscience, 2019, 13, 142.	3.7	19
49	Anti– <scp>N</scp> â€methylâ€ <scp>D</scp> â€aspartate receptor encephalitis with multiphasic demyelination. Annals of Neurology, 2014, 76, 462-464.	5.3	18
50	MOG-antibody-associated disease is different from MS and NMOSD and should be considered as a distinct disease entity – Yes. Multiple Sclerosis Journal, 2020, 26, 272-274.	3.0	18
51	Status of the neuromyelitis optica spectrum disorder in Latin America. Multiple Sclerosis and Related Disorders, 2021, 53, 103083.	2.0	18
52	Distinction between MOG antibody-positive and AQP4 antibody-positive NMO spectrum disorders. Neurology, 2014, 83, 1122-1123.	1.1	17
53	CSF levels of glutamine synthetase and GFAP to explore astrocytic damage in seronegative NMOSD. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 605-611.	1.9	17
54	Distinction between MOG antibody–positive and AQP4 antibody–positive NMO spectrum disorders. Neurology, 2014, 83, 475-476.	1.1	16

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55	Characterization of pain syndromes in patients with neuromyelitis optica. European Journal of Pain, 2020, 24, 1548-1568.	2.8	16
56	Relapsing optic neuritis and isolated transverse myelitis are the predominant clinical phenotypes for patients with antibodies to myelin oligodendrocyte glycoprotein in India. Multiple Sclerosis Journal - Experimental, Translational and Clinical, 2016, 2, 205521731667563.	1.0	15
57	Neutralizing Antibodies Are Associated with a Reduction of Interferon-β Efficacy during the Treatment of Japanese Multiple Sclerosis Patients. Tohoku Journal of Experimental Medicine, 2012, 228, 85-92.	1.2	14
58	Detection of MOG-IgG by cell-based assay: moving from discovery to clinical practice. Neurological Sciences, 2021, 42, 73-80.	1.9	14
59	Myelin oligodendrocyte glycoprotein immunoglobulin Gâ€associated disease: An overview. Clinical and Experimental Neuroimmunology, 2018, 9, 48-55.	1.0	13
60	AQP4 antibody serostatus. Neurology, 2013, 81, 1186-1188.	1.1	12
61	Anti-MOG (Myelin Oligodendrocyte Glycoprotein)–Positive Severe Optic Neuritis with Optic Disc Ischaemia and Macular Star. Neuro-Ophthalmology, 2015, 39, 285-288.	1.0	12
62	Experimental Models of Neuroimmunological Disorders: A Review. Frontiers in Neurology, 2020, 11, 389.	2.4	11
63	Aquaporin-4 antibody-positive myelitis initially biopsied for suspected spinal cord tumors: Diagnostic considerations. Multiple Sclerosis Journal, 2014, 20, 621-626.	3.0	10
64	Unraveling B lymphocytes in CNS inflammatory diseases. Neurology, 2020, 95, 733-744.	1.1	10
65	Treatment of MOG-IgG associated disease in paediatric patients: A systematic review. Multiple Sclerosis and Related Disorders, 2021, 56, 103216.	2.0	10
66	Neuromyelitis optica without typical opticospinal phenotype. Multiple Sclerosis Journal, 2010, 16, 1154-1155.	3.0	9
67	MOG-IgG serological status matters in paediatric ADEM. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 242-242.	1.9	8
68	MOG-IgG associated optic neuritis is not multiple sclerosis. Arquivos De Neuro-Psiquiatria, 2017, 75, 687-691.	0.8	8
69	Detection of autoantibodies in central nervous system inflammatory disorders: Clinical application of cell-based assays. Multiple Sclerosis and Related Disorders, 2020, 38, 101858.	2.0	8
70	Consensus recommendations for the diagnosis and treatment of primary progressive multiple sclerosis in Latin America. Journal of the Neurological Sciences, 2018, 393, 4-13.	0.6	7
71	Viral encephalitis: a practical review on diagnostic approach and treatment. Jornal De Pediatria (Versão Em Português), 2020, 96, 12-19	0.2	7
72	Safety indicators of a novel multi supplement based on guarana, selenium, and L-carnitine: Evidence from human and red earthworm immune cells. Food and Chemical Toxicology, 2021, 150, 112066.	3.6	6

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73	Long-term safety of azathioprine for treatment of neuromyelitis optica spectrum disorders. Arquivos De Neuro-Psiquiatria, 2021, 79, 229-232.	0.8	5
74	Decreased convulsive threshold and memory loss after anti-NMDAR positive CSF injection in zebrafish. Journal of Neuroimmunology, 2021, 359, 577689.	2.3	5
75	Mechanisms of myelin repair, MRI techniques and therapeutic opportunities in multiple sclerosis. Multiple Sclerosis and Related Disorders, 2022, 58, 103407.	2.0	5
76	Detection of MOG-IgG in Clinical Samples by Live Cell-Based Assays: Performance of Immunofluorescence Microscopy and Flow Cytometry. Frontiers in Immunology, 2021, 12, 642272.	4.8	4
77	Case of autoantibodies against Nâ€methylâ€Dâ€aspartate receptor+/antibodies against myelinâ€oligodendrocyte glycoprotein+ multiphasic acute disseminated encephalomyelitis (<scp>ADEM</scp>). Clinical and Experimental Neuroimmunology, 2014, 5, 49-51.	1.0	3
78	Reduced quality of life in a pediatric-onset Neuromyelitis optica spectrum disorders cohort. Multiple Sclerosis and Related Disorders, 2021, 56, 103252.	2.0	3
79	Time to target brain atrophy and neurodegeneration in multiple sclerosis. Arquivos De Neuro-Psiquiatria, 2016, 74, 181-182.	0.8	2
80	Oral fingolimod to treat multiple sclerosis: see your cardiologist first. Arquivos De Neuro-Psiquiatria, 2014, 72, 651-652.	0.8	2
81	Aquaporin-4 antibody-positive cases beyond current diagnostic criteria for NMO spectrum disorders. Neurology, 2014, 82, 372-372.	1.1	1
82	Leukoencephalopathy resolution after atypical mycobacterial treatment: a case report. BMC Neurology, 2015, 15, 159.	1.8	1
83	Healthy aging during the COVID-19 pandemic. PAJAR - Pan-American Journal of Aging Research, 2021, 9, e41087.	0.1	1
84	Simultaneous bilateral optic neuritis and longitudinally extensive transverse myelitis following vaccination against COVID-19: A case report. Neuroimmunology Reports, 2021, 1, 100041.	0.4	1
85	2014 Joint Americas Committee for Treatment and Research in Multiple Sclerosis–European Committee for Treatment and Research in Multiple Sclerosis Meeting. Clinical and Experimental Neuroimmunology, 2015, 6, 100-102.	1.0	0
86	A Regional Analysis of Relapsing Remitting MS (RRMS) Patients Treated with Natalizumab Suggests a Need to Diagnose and Treat Early in Latin Americans. Value in Health, 2016, 19, A437.	0.3	0
87	Autoimmune encephalitis in patients with anti-myelin oligodendrocyte glycoprotein-antibody. Journal of the Neurological Sciences, 2017, 381, 792.	0.6	0
88	Discrimination of spinal cord sarcoidosis from neuromyelitis optica spectrum disorer or spondylotic myelopathy. Journal of the Neurological Sciences, 2017, 381, 449-450.	0.6	0
89	Sometimes less is more in multiple sclerosis drug switching. Arquivos De Neuro-Psiquiatria, 2016, 74, 605-606.	0.8	0
90	Quality of life goes beyond the medical priorities in multiple sclerosis: assessing the impact of social support network. Arquivos De Neuro-Psiquiatria, 2017, 75, 263-264.	0.8	0

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91	Isolated rhombencephalitis with good clinical recovery. Arquivos De Neuro-Psiquiatria, 2017, 75, 757-757.	0.8	0