Chih-Chao Yang

List of Publications by Year
in descending order

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49	3,694	394421 19	233421 45
papers	citations	h-index	g-index

51	51	51	5786
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Real-world evidence on the safety and effectiveness of fingolimod in patients with multiple sclerosis from Taiwan. Journal of the Formosan Medical Association, 2021, 120, 542-550.	1.7	7
2	Thymidine Kinase 2 Deficiencyâ€“Induced Adult-Onset Ptosis and Proximal Weakness. Neurology: Clinical Practice, 2021, 11, e379-e382.	1.6	0
3	Novel c.435delC mutation in XK gene found in a Taiwanese patient with McLeod syndrome. Transfusion, 2021, 61, E28-E30.	1.6	1
4	A systematic review of late-onset and very-late-onset multiple acyl-coenzyme A dehydrogenase deficiency: Cohort analysis and patient report from Taiwan. Neuromuscular Disorders, 2021, 31, 218-225.	0.6	3
5	The Frequency and Perceived Effectiveness of Pain Self-Management Strategies Used by Individuals With Migraine. The Journal of Nursing Research: JNR, 2021, 29, e154.	1.7	5
6	Analysis of site-specific glycan profiles of serum proteins in patients with multiple sclerosis or neuromyelitis optica spectrum disorderâ€”a pilot study. Glycobiology, 2021, 31, 1230-1238.	2.5	2
7	Lactate peak in muscle disclosed by magnetic resonance spectroscopy in a patient with CPEO-plus syndrome. ENeurologicalSci, 2021, 24, 100360.	1.3	1
8	Clinical and genetic characterization of adultâ€onset leukoencephalopathy caused by CSF1R mutations. Annals of Clinical and Translational Neurology, 2021, 8, 2121-2131.	3.7	9
9	Cardiac manifestations and prognostic implications of hereditary transthyretin amyloidosis associated with transthyretin Ala97Ser. Journal of the Formosan Medical Association, 2020, 119, 693-700.	1.7	13
10	Autosomal dominant cerebellar ataxia, deafness, and narcolepsy with amenorrhea, subclinical optic atrophy, and electroencephalographic abnormality: A case report. ENeurologicalSci, 2020, 21, 100271.	1.3	0
11	Reply to â€œUse DN4-T to rule out non-neuropathic painâ€. Journal of the Chinese Medical Association, 2020, 83, 511-511.	1.4	0
12	Multiple sclerosis and neuromyelitis optica after optic neuritis: A nationwide cohort study in Taiwan. Multiple Sclerosis and Related Disorders, 2020, 44, 102379.	2.0	3
13	Modeling spinocerebellar ataxias 2 and 3 with iPSCs reveals a role for glutamate in disease pathology. Scientific Reports, 2019, 9, 1166.	3.3	29
14	Migraine-Specific Quality of Life QuestionnaireÂChinese version 2.1 (MSQv2.1-C): psychometric evaluation in patients with migraine. Health and Quality of Life Outcomes, 2019, 17, 108.	2.4	12
15	Development and validation of a Taiwan version of the DN4-T questionnaire. Journal of the Chinese Medical Association, 2019, 82, 623-627.	1.4	11
16	Differentiation of remitting neuromyelitis optica spectrum disorders from multiple sclerosis by integrating parameters from serum proteins and lymphocyte subsets. Journal of Neuroimmunology, 2018, 318, 45-52.	2.3	7
17	Development and validation of a Taiwan version of the ID Pain questionnaire (ID Pain-T). Journal of the Chinese Medical Association, 2018, 81, 12-17.	1.4	12
18	Patisiran, an RNAi Therapeutic, for Hereditary Transthyretin Amyloidosis. New England Journal of Medicine, 2018, 379, 11-21.	27.0	1,944

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19	Sleep Disordered Breathing Mimicking Myasthenia Crisis in a Patient with Myasthenia Gravis. Journal of Clinical Sleep Medicine, 2016, 12, 767-769.	2.6	3
20	History of neurology in Taiwan. Neurology, 2015, 84, 1803-1804.	1.1	0
21	The Temporal Profiles of Changes in Nerve Excitability Indices in Familial Amyloid Polyneuropathy. PLoS ONE, 2015, 10, e0141935.	2.5	13
22	A novel XK gene mutation in a Taiwanese family with McLeod syndrome. Journal of the Neurological Sciences, 2014, 340, 221-224.	0.6	7
23	Cardioembolic stroke related to limb-girdle muscular dystrophy 1B. BMC Research Notes, 2013, 6, 32.	1.4	13
24	Fabry Disease: A Rare Cause of Fever of Unknown Origin. American Journal of Kidney Diseases, 2012, 59, 161-162.	1.9	6
25	Carbamazepine-Induced Toxic Effects and HLA-B*1502 Screening in Taiwan. New England Journal of Medicine, 2011, 364, 1126-1133.	27.0	631
26	Neuroimaging Findings in a Brain With Niemannâ€“Pick Type C Disease. Journal of the Formosan Medical Association, 2011, 110, 537-542.	1.7	36
27	Carrier Screening for Spinal Muscular Atrophy (SMA) in 107,611 Pregnant Women during the Period 2005â€“2009: A Prospective Population-Based Cohort Study. PLoS ONE, 2011, 6, e17067.	2.5	72
28	The Mutational Spectrum in a Cohort of Charcot-Marie-Tooth Disease Type 2 among the Han Chinese in Taiwan. PLoS ONE, 2011, 6, e29393.	2.5	84
29	The effect of interferon Î²-1a on optic neuritis relapse in patients with multiple sclerosis. Graefe's Archive for Clinical and Experimental Ophthalmology, 2010, 248, 231-235.	1.9	2
30	Mutation analysis and characterization of alternative splice variants of the Wilson disease gene ATP7B. Hepatology, 2010, 52, 1662-1670.	7.3	27
31	Probable variant Creutzfeldtâ€“Jakob disease in Asia: A case report from Taiwan and review of two prior cases. Psychiatry and Clinical Neurosciences, 2010, 64, 652-658.	1.8	4
32	Neurosarcoidosis Affecting the Spinal Cord. Journal of the Formosan Medical Association, 2010, 109, 676-679.	1.7	2
33	Correlation of Survival Motor Neuron Expression in Leukocytes and Spinal Cord in Spinal Muscular Atrophy. Journal of Pediatrics, 2009, 154, 303-305.	1.8	9
34	Predictors for Outcome and Treatment Delay in Patients With Tuberculous Meningitis. American Journal of the Medical Sciences, 2009, 338, 134-139.	1.1	53
35	Identification of deletion and duplication genotypes of thePMP22gene using PCRâ€RFLP, competitive multiplex PCR, and multiplex ligationâ€dependent probe amplification: A comparison. Electrophoresis, 2008, 29, 618-625.	2.4	13
36	Transthyretin Ala97Ser in Chineseâ€“Taiwanese patients with familial amyloid polyneuropathy: Genetic studies and phenotype expression. Journal of the Neurological Sciences, 2008, 267, 91-99.	0.6	26

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37	Comparison of two PCR-based molecular methods in the diagnosis of CMT 1A and HNPP diseases in Chinese. Clinical Neurology and Neurosurgery, 2008, 110, 466-471.	1.4	5
38	Nephrogenic Systemic Fibrosis Associated with Gadolinium Use. Journal of the Formosan Medical Association, 2008, 107, 270-274.	1.7	21
39	Cutaneous Polyarteritis Nodosa in a Patient With Fabry Disease. Archives of Dermatology, 2008, 144, 122-3.	1.4	3
40	Myopathy in Gaucher disease. Journal of Inherited Metabolic Disease, 2008, 31, 489-491.	3.6	4
41	Nerve function and dysfunction in acute intermittent porphyria. Brain, 2008, 131, 2510-2519.	7.6	75
42	Characterization of a familial case with primary erythromelalgia from Taiwan. Journal of Neurology, 2007, 254, 210-214.	3.6	51
43	Multiple Sclerosis With Childhood Onset: Report of 21 Cases in Taiwan. Pediatric Neurology, 2006, 35, 327-334.	2.1	22
44	Levamisole-Induced Multifocal Inflammatory Leukoencephalopathy. Medicine (United States), 2006, 85, 203-213.	1.0	47
45	The spectrum ofWRNmutations in Werner syndrome patients. Human Mutation, 2006, 27, 558-567.	2.5	198
46	Identification of forty-five novel and twenty-three knownNF1 mutations in Chinese patients with neurofibromatosis type 1. Human Mutation, 2006, 27, 832-832.	2.5	30
47	Efficacy, safety, and tolerability of pramipexole in untreated and levodopa-treated patients with Parkinson's disease. Journal of the Neurological Sciences, 2003, 216, 81-87.	0.6	46
48	Quantitative pathology of cutaneous nerve terminal degeneration in the human skin. Acta Neuropathologica, 2001, 102, 455-461.	7.7	87
49	Immunolocalization of transcription factor NF-Î°B in inclusion-body myositis muscle and at normal human neuromuscular junctions. Neuroscience Letters, 1998, 254, 77-80.	2.1	44