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	Patisiran, an RNAi Therapeutic, for Hereditary Transthyretin Amyloidosis. New England Journal of Medicine, 2018, 379, 11-21.	27.0	1,944
2	Carbamazepine-Induced Toxic Effects and HLA-B*1502 Screening in Taiwan. New England Journal of Medicine, 2011, 364, 1126-1133.	27.0	631
3	The spectrum ofWRNmutations in Werner syndrome patients. Human Mutation, 2006, 27, 558-567.	2.5	198
4	Quantitative pathology of cutaneous nerve terminal degeneration in the human skin. Acta Neuropathologica, 2001, 102, 455-461.	7.7	87
5	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
6	Nerve function and dysfunction in acute intermittent porphyria. Brain, 2008, 131, 2510-2519.	7.6	75
7	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
8	Predictors for Outcome and Treatment Delay in Patients With Tuberculous Meningitis. American Journal of the Medical Sciences, 2009, 338, 134-139.	1.1	53
9	Characterization of a familial case with primary erythromelalgia from Taiwan. Journal of Neurology, 2007, 254, 210-214.	3.6	51
10	Levamisole-Induced Multifocal Inflammatory Leukoencephalopathy. Medicine (United States), 2006, 85, 203-213.	1.0	47
11	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
12	Immunolocalization of transcription factor NF-ÎºB in inclusion-body myositis muscle and at normal human neuromuscular junctions. Neuroscience Letters, 1998, 254, 77-80.	2.1	44
13	Neuroimaging Findings in a Brain With Niemannâ€“Pick Type C Disease. Journal of the Formosan Medical Association, 2011, 110, 537-542.	1.7	36
14	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
15	Modeling spinocerebellar ataxias 2 and 3 with iPSCs reveals a role for glutamate in disease pathology. Scientific Reports, 2019, 9, 1166.	3.3	29
16	Mutation analysis and characterization of alternative splice variants of the Wilson disease gene ATP7B. Hepatology, 2010, 52, 1662-1670.	7.3	27
17	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
18	Multiple Sclerosis With Childhood Onset: Report of 21 Cases in Taiwan. Pediatric Neurology, 2006, 35, 327-334.	2.1	22

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19	Nephrogenic Systemic Fibrosis Associated with Gadolinium Use. Journal of the Formosan Medical Association, 2008, 107, 270-274.	1.7	21
20	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
21	Cardioembolic stroke related to limb-girdle muscular dystrophy 1B. BMC Research Notes, 2013, 6, 32.	1.4	13
22	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
23	The Temporal Profiles of Changes in Nerve Excitability Indices in Familial Amyloid Polyneuropathy. PLoS ONE, 2015, 10, e0141935.	2.5	13
24	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
25	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
26	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
27	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
28	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
29	A novel XK gene mutation in a Taiwanese family with McLeod syndrome. Journal of the Neurological Sciences, 2014, 340, 221-224.	0.6	7
30	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
31	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
32	Fabry Disease: A Rare Cause of Fever of Unknown Origin. American Journal of Kidney Diseases, 2012, 59, 161-162.	1.9	6
33	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
34	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
35	Myopathy in Gaucher disease. Journal of Inherited Metabolic Disease, 2008, 31, 489-491.	3.6	4
36	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

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37	Cutaneous Polyarteritis Nodosa in a Patient With Fabry Disease. Archives of Dermatology, 2008, 144, 122-3.	1.4	3
38	Sleep Disordered Breathing Mimicking Myasthenia Crisis in a Patient with Myasthenia Gravis. Journal of Clinical Sleep Medicine, 2016, 12, 767-769.	2.6	3
39	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
40	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
41	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
42	Neurosarcoidosis Affecting the Spinal Cord. Journal of the Formosan Medical Association, 2010, 109, 676-679.	1.7	2
43	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
44	Novel c.435delC mutation in XK gene found in a Taiwanese patient with McLeod syndrome. Transfusion, 2021, 61, E28-E30.	1.6	1
45	Lactate peak in muscle disclosed by magnetic resonance spectroscopy in a patient with CPEO-plus syndrome. ENeurologicalSci, 2021, 24, 100360.	1.3	1
46	History of neurology in Taiwan. Neurology, 2015, 84, 1803-1804.	1.1	0
47	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
48	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
49	Thymidine Kinase 2 Deficiencyâ€“Induced Adult-Onset Ptosis and Proximal Weakness. Neurology: Clinical Practice, 2021, 11, e379-e382.	1.6	0