Christina Liang

List of Publications by Year
in descending order

25 papers	813 citations	759233 12 h-index	713466 21 g-index
25 all docs	25 docs citations	25 times ranked	1643 citing authors

#	Article	IF	Citations
1	Patient care standards for primary mitochondrial disease in Australia: an Australian adaptation of the Mitochondrial Medicine Society recommendations. Internal Medicine Journal, 2022, 52, 110-120.	0.8	3
2	Outcome of patient with myasthenia gravis with the use of immunotherapy in metastatic Merkel cell carcinoma. Oxford Medical Case Reports, 2022, 2022, omac012.	0.4	1
3	Late adult-onset spinal muscular atrophy with lower extremity predominance (SMALED). BMJ Case Reports, 2022, 15, e248297.	0.5	1
4	Use of Whole-Genome Sequencing for Mitochondrial Disease Diagnosis. Neurology, 2022, 99, .	1.1	33
5	080â€…The diagnostic journey of mitochondrial disease patients. , 2021, , .		2
6	Neuropathy in sporadic inclusion body myositis: A multi-modality neurophysiological study. Clinical Neurophysiology, 2020, 131, 2766-2776.	1.5	8
7	Rehabilitation for ataxia study: protocol for a randomised controlled trial of an outpatient and supported home-based physiotherapy programme for people with hereditary cerebellar ataxia. BMJ Open, 2020, 10, e040230.	1.9	14
8	Safety and efficacy of intravenous bimagrumab in inclusion body myositis (RESILIENT): a randomised, double-blind, placebo-controlled phase 2b trial. Lancet Neurology, The, 2019, 18, 834-844.	10.2	91
9	Sarcolemmal depolarization in sporadic inclusion body myositis assessed with muscle velocity recovery cycles. Clinical Neurophysiology, 2019, 130, 2272-2281.	1.5	9
10	024â€…Resistance exercises with blood flow restriction in patients with sporadic inclusion body myositis. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, A9.1-A9.	1.9	1
11	Sarcolemmal excitability in the myotonic dystrophies. Muscle and Nerve, 2018, 57, 595-602.	2.2	12
12	Mitochondrial diseases. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 147, 125-141.	1.8	30
13	N-of-1 trial of thymoquinone and vorinostat in a patient with sialidosis type 1. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, e1.98-e1.	1.9	1
14	Serum metabolomic profiling for diagnosis of mitochondrial diseases. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, e1.7-e1.	1.9	0
15	Muscle membrane dysfunction in inclusion body myopathy studied by muscle velocity recovery cycles. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, e1.38-e1.	1.9	0
16	Sarcolemmal excitability attributes of the myotonic dystrophies as assessed by muscle velocity recovery cycles (mvrcs). Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, e1.52-e1.	1.9	0
17	A comparison of current serum biomarkers as diagnostic indicators of mitochondrial diseases. Neurology, 2016, 86, 2010-2015.	1.1	89
18	Rare variants in SQSTM1 and VCP genes and risk of sporadic inclusion body myositis. Neurobiology of Aging, 2016, 47, 218.e1-218.e9.	3.1	40

#	Article	IF	Citations
19	Systematic review of cardiac electrical disease in Kearnsâ€“Sayre syndrome and mitochondrial cytopathy. International Journal of Cardiology, 2015, 181, 303-310.	1.7	81
20	Expanding the phenotype of GMPPB mutations. Brain, 2015, 138, 836-844.	7.6	54
21	Use of Whole-Exome Sequencing for Diagnosis of Limb-Girdle Muscular Dystrophy. JAMA Neurology, 2015, 72, 1424.	9.0	164
22	The broadening spectrum of mitochondrial disease: Shifts in the diagnostic paradigm. Biochimica Et Biophysica Acta - General Subjects, 2014, 1840, 1360-1367.	2.4	48
23	Comparing axonal excitability in past polio to amyotrophic lateral sclerosis. Muscle and Nerve, 2014, 50, 602-604.	2.2	1
24	Axonal excitability in X-linked dominant Charcot Marie Tooth disease. Clinical Neurophysiology, 2014, 125, 1261-1269.	1.5	12
25	Necrotizing autoimmune myopathy. Current Opinion in Rheumatology, 2011, 23, 612-619.	4.3	118