Han-Chih Hencher Lee

List of Publications by Citations

14 papers	205 citations	6 h-index	14 g-index
16 ext. papers	238 ext. citations	3.5 avg, IF	2.55 L-index

#	Paper	IF	Citations
14	Inborn errors of metabolism and expanded newborn screening: review and update. *Critical Reviews in Clinical Laboratory Sciences*, **2013**, 50, 142-62	9.4	99
13	Fatal viral infection-associated encephalopathy in two Chinese boys: a genetically determined risk factor of thermolabile carnitine palmitoyltransferase II variants. *Journal of Human Genetics*, **2011**, 56, 617-21	4.3	31
12	Non-invasive urinary screening for aromatic L-amino acid decarboxylase deficiency in high-prevalence areas: a pilot study. *Clinica Chimica Acta*, **2012**, 413, 126-30	6.2	16
11	The co-occurrence of serologically proven myasthenia gravis and Miller Fisher/Guillain Barr overlap syndrome: a case report. *Journal of the Neurological Sciences*, **2009**, 276, 187-8	3.2	13
10	2,4-Dinitrophenol: a threat to Chinese body-conscious groups. *Journal of the Chinese Medical Association*, **2014**, 77, 443-5	2.8	12
9	Role of postmortem genetic testing demonstrated in a case of glutaric aciduria type II. *Diagnostic Molecular Pathology*, **2010**, 19, 184-6		10
8	Gelsemium poisoning mediated by the non-toxic plant Cassytha filiformis parasitizing Gelsemium elegans. *Toxicon*, **2018**, 154, 42-49	2.8	5
7	Founder Mutation c.1516A>C in KLHL40 Is a Frequent Cause of Nemaline Myopathy With Hyponatremia in Ethnic Chinese. *Journal of Neuropathology and Experimental Neurology*, **2019**, 78, 854-864	3.1	4
6	Practical Aspects in Genetic Testing for Cardiomyopathies and Channelopathies. *Clinical Biochemist Reviews*, **2019**, 40, 187-200	7.3	4
5	Clinical and pathological characterization of FLNC-related myofibrillar myopathy caused by founder variant c.8129G>A in Hong Kong Chinese. *Clinical Genetics*, **2020**, 97, 747-757	4	3
4	Correlation study between spot urine protein-to-creatinine ratio and 24-hour urine protein measurement in 174 patients for proteinuria assessment. *Hong Kong Journal of Nephrology*, **2011**, 13, 51-54		3
3	Performance evaluation of five commercial assays for detection of acetaminophen. *Journal of Clinical Laboratory Analysis*, **2019**, 33, e22683	3	3
2	Flexi-Myo Panel Strategy: Genomic Diagnoses of Myopathies and Muscular Dystrophies by Next-Generation Sequencing. *Genetic Testing and Molecular Biomarkers*, **2020**, 24, 99-104	1.6	2
1	Kearns-Sayre Syndrome Minus: Two Cases of Identical Large-Scale Mitochondrial DNA Deletions with Presentations outside the Classical Triad.. *Case Reports in Genetics*, **2022**, 2022, 4153357	0.7	