Ling M Wong

List of Publications by Year
in descending order

15 papers	289 citations	1040056 9 h-index	1058476 14 g-index
16 all docs	16 docs citations	16 times ranked	322 citing authors

#	Article	IF	Citations
1	Young adult female fragile X premutation carriers show age- and genetically-modulated cognitive impairments. Brain and Cognition, 2011, 75, 255-260.	1.8	72
2	Adult Female Fragile X Premutation Carriers Exhibit Age- and CGG Repeat Length-Related Impairments on an Attentionally Based Enumeration Task. Frontiers in Human Neuroscience, 2011, 5, 63.	2.0	59
3	Enhanced Manual and Oral Motor Reaction Time in Young Adult Female Fragile X Premutation Carriers. Journal of the International Neuropsychological Society, 2011, 17, 746-750.	1.8	28
4	Abnormal N400 word repetition effects in fragile X-associated tremor/ataxia syndrome. Brain, 2010, 133, 1438-1450.	7.6	24
5	A Cross-Sectional Analysis of the Development of Response Inhibition in Children with Chromosome 22q11.2 Deletion Syndrome. Frontiers in Psychiatry, 2013, 4, 81.	2.6	22
6	Children With Chromosome 22q11.2 Deletion Syndrome Exhibit Impaired Spatial Working Memory. American Journal on Intellectual and Developmental Disabilities, 2014, 119, 115-132.	1.6	22
7	Young adult male carriers of the fragile X premutation exhibit genetically modulated impairments in visuospatial tasks controlled for psychomotor speed. Journal of Neurodevelopmental Disorders, 2012, 4, 26.	3.1	14
8	Eye movements reveal impaired inhibitory control in adult male fragile X premutation carriers asymptomatic for FXTAS.. Neuropsychology, 2014, 28, 571-584.	1.3	14
9	Next-generation strategies for gene-targeted therapies of central nervous system disorders: A workshop summary. Molecular Therapy, 2021, 29, 3332-3344.	8.2	12
10	Neuropsychological measurement of inhibitory control in posttraumatic stress disorder: An exploratory antisaccade paradigm. Journal of Clinical and Experimental Neuropsychology, 2017, 39, 1002-1012.	1.3	8
11	Temporal dynamics of attentional selection in adult male carriers of the fragile X premutation allele and adult controls. Frontiers in Human Neuroscience, 2015, 9, 37.	2.0	4
12	Funding community collaboration to develop effective therapies for neurofibromatosis type 1 tumors. EMBO Molecular Medicine, 2020, 12, e11656.	6.9	3
13	NINDS launches network to develop treatments for ultra-rare neurological diseases. Nature Biotechnology, 2021, 39, 1497-1499.	17.5	3
14	Quantifying the resolution of spatial and temporal representation in children with 22q11.2 deletion syndrome. Journal of Neurodevelopmental Disorders, 2019, 11, 40.	3.1	2
15	A cross-sectional analysis of orienting of visuospatial attention in child and adult carriers of the fragile X premutation. Journal of Neurodevelopmental Disorders, 2014, 6, 45.	3.1	0