# Manting Xu

## List of Publications by Citations



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|---|---|---|---|
| 9 papers | 34 citations | 2 h-index | 5 g-index |
| 11 ext. papers | 55 ext. citations | 5.2 avg, IF | 0.39 L-index |

| # | Paper | IF | Citations |
|---|---|---|---|
| 9 | Bi-allelic ADPRHL2 Mutations Cause Neurodegeneration with Developmental Delay, Ataxia, and Axonal Neuropathy. *American Journal of Human Genetics*, **2018**, 103, 817-825 | 11 | 24 |
| 8 | Clinical Assessments and EEG Analyses of Encephalopathies Associated With Dynamin-1 Mutation. *Frontiers in Pharmacology*, **2019**, 10, 1454 | 5.6 | 4 |
| 7 | Identification of a novel m.3955G▯▯A variant in MT-ND1 associated with Leigh syndrome. *Mitochondrion*, **2021**, 62, 13-23 | 4.9 | 2 |
| 6 | Whole genome and exome sequencing identify mutations as a new cause of progressive cavitating leukoencephalopathy. *Journal of Medical Genetics*, **2021**, | 5.8 | 1 |
| 5 | Biallelic -Variants Leading to Developmental Regression With Progressive Spasticity and Brain Atrophy in a Chinese Patient. *Frontiers in Genetics*, **2021**, 12, 685035 | 4.5 | 1 |
| 4 | Cinical, Metabolic, and Genetic Analysis and Follow-Up of Eight Patients With Mutations Presenting With Leigh/Leigh-Like Syndrome. *Frontiers in Pharmacology*, **2021**, 12, 605803 | 5.6 | 1 |
| 3 | Clinical Attributes and Electroencephalogram Analysis of Patients With Varying AlpersaSyndrome Genotypes. *Frontiers in Pharmacology*, **2021**, 12, 669516 | 5.6 | 0 |
| 2 | Identification of a Novel Variant in Causing MELAS. *Frontiers in Genetics*, **2021**, 12, 638749 | 4.5 | 0 |
| 1 | Generation of an iPSC line from a patient with early-onset epileptic encephalopathy carrying CARS2 (p.G476R) mutation.. *Stem Cell Research*, **2021**, 59, 102633 | 1.6 | |