

# Manting Xu

## List of Publications by Citations

**Source:** <https://exaly.com/author-pdf/3906479/manting-xu-publications-by-citations.pdf>

**Version:** 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

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. <i>American Journal of Human Genetics</i> , <b>2018</b> , 103, 817-825	11	24
8	Clinical Assessments and EEG Analyses of Encephalopathies Associated With Dynamin-1 Mutation. <i>Frontiers in Pharmacology</i> , <b>2019</b> , 10, 1454	5.6	4
7	Identification of a novel m.3955G>A variant in MT-ND1 associated with Leigh syndrome. <i>Mitochondrion</i> , <b>2021</b> , 62, 13-23	4.9	2
6	Whole genome and exome sequencing identify mutations as a new cause of progressive cavitating leukoencephalopathy. <i>Journal of Medical Genetics</i> , <b>2021</b> ,	5.8	1
5	Biallelic -Variants Leading to Developmental Regression With Progressive Spasticity and Brain Atrophy in a Chinese Patient. <i>Frontiers in Genetics</i> , <b>2021</b> , 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. <i>Frontiers in Pharmacology</i> , <b>2021</b> , 12, 605803	5.6	1
3	Clinical Attributes and Electroencephalogram Analysis of Patients With Varying AlpersaSyndrome Genotypes. <i>Frontiers in Pharmacology</i> , <b>2021</b> , 12, 669516	5.6	0
2	Identification of a Novel Variant in Causing MELAS. <i>Frontiers in Genetics</i> , <b>2021</b> , 12, 638749	4.5	0
1	Generation of an iPSC line from a patient with early-onset epileptic encephalopathy carrying CARS2 (p.G476R) mutation.. <i>Stem Cell Research</i> , <b>2021</b> , 59, 102633	1.6	