

# Chuan Zhang

## List of Publications by Year in descending order

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60  
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#	ARTICLE	IF	CITATIONS
1	Variant analysis of 92 Chinese Han families with hearing loss. BMC Medical Genomics, 2022, 15, 12.	1.5	6
2	Mutation analysis of <i>TCOF1</i> gene in Chinese Treacher Collins syndrome patients. Journal of Clinical Laboratory Analysis, 2021, 35, e23567.	2.1	10
3	Detailed pedigree analyses and prenatal diagnosis for a family with mucopolysaccharidosis type II. BMC Medical Genomics, 2021, 14, 175.	1.5	2
4	Maternal UPD of chromosome 7 in a patient with Silver-Russell syndrome and Pendred syndrome. Journal of Clinical Laboratory Analysis, 2020, 34, e23407.	2.1	5
5	Mutation analysis, treatment and prenatal diagnosis of Chinese cases of methylmalonic acidemia. Scientific Reports, 2020, 10, 12509.	3.3	10
6	Compound heterozygous variants of the <i>FBXO7</i> gene resulting in infantile-onset Parkinsonian-pyramidal syndrome in siblings of a Chinese family. Journal of Clinical Laboratory Analysis, 2020, 34, e23324.	2.1	8
7	A novel <i>LOXHD1</i> variant in a Chinese couple with hearing loss. Journal of International Medical Research, 2019, 47, 6082-6090.	1.0	6