Qiuju Wang

List of Publications by Year
in descending order

11 papers	256 citations	1163117 8 h-index	1281871 11 g-index
11 all docs	11 docs citations	11 times ranked	499 citing authors

#	Article	IF	Citations
1	Generation of a gene corrected human isogenic iPSC line (CPGHi001-A-1) from a hearing loss patient with the TMC1 p.M418K mutation using CRISPR/Cas9. Stem Cell Research, 2022, 60, 102736.	0.7	4
2	Clinical characteristics of patients with unilateral auditory neuropathy. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2021, 42, 103143.	1.3	4
3	High Frequency of AIFM1 Variants and Phenotype Progression of Auditory Neuropathy in a Chinese Population. Neural Plasticity, 2020, 2020, 1-12.	2.2	10
4	Generation of a human induced pluripotent stem cell line (CPGHi001-A) from a hearing loss patient with the TMC1 p.M418K mutation. Stem Cell Research, 2020, 49, 101982.	0.7	7
5	Recurrent de novo WFS1 pathogenic variants in Chinese sporadic patients with nonsyndromic sensorineural hearing loss. Molecular Genetics & Genomic Medicine, 2020, 8, e1367.	1.2	9
6	Phenotype prediction of Mohr-Tranebjaerg syndrome (MTS) by genetic analysis and initial auditory neuropathy. BMC Medical Genetics, 2019, 20, 11.	2.1	12
7	Identification of four <scp>TMC</scp>1 variations in different Chinese families with hereditary hearing loss. Molecular Genetics & Genomic Medicine, 2018, 6, 504-513.	1.2	17
8	Novel recessive PDZD7 biallelic mutations in two Chinese families with nonâ€syndromic hearing loss. American Journal of Medical Genetics, Part A, 2018, 176, 99-106.	1.2	15
9	Mutations in apoptosis-inducing factor cause X-linked recessive auditory neuropathy spectrum disorder. Journal of Medical Genetics, 2015, 52, 523-531.	3.2	92
10	A Novel DFNA36 Mutation in TMC1 Orthologous to the Beethoven (Bth) Mouse Associated with Autosomal Dominant Hearing Loss in a Chinese Family. PLoS ONE, 2014, 9, e97064.	2.5	61
11	Targeted High-Throughput Sequencing Identifies Pathogenic Mutations in KCNQ4 in Two Large Chinese Families with Autosomal Dominant Hearing Loss. PLoS ONE, 2014, 9, e103133.	2.5	25