Identification of a novel AIFM1 variant from a Chinese family with auditory neuropathy

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Background: Auditory neuropathy (AN) is a specific type of hearing loss characterized by impaired language comprehension. Apoptosis inducing factor mitochondrion associated 1 (AIFM1) is the most common gene associated with late-onset AN. In this study, we aimed to screen the pathogenic variant of AIFM1 in a Chinese family with AN and to explore the molecular mechanism underlying the function of such variant in the development of AN.

Methods: One patient with AN and eight unaffected individuals from a Chinese family were enrolled in this study. A comprehensive clinical evaluation was performed on all participants. A targeted next-generation sequencing (NGS) analysis of a total of 406 known deafness genes was performed to screen the potential pathogenic variants in the proband. Sanger sequencing was used to confirm the variants identified in all participants. The pathogenicity of variant was predicted by bioinformatics analysis. Immunofluorescence and Western blot analyses were performed to evaluate the subcellular distribution and expression of the wild type (WT) and mutant AIFM1 proteins. Cell apoptosis was evaluated based on the TUNEL analyses.

Results: Based on the clinical evaluations, the proband in this family was diagnosed with AN. The results of NGS and Sanger sequencing showed that a novel missense mutation of AIFM1, i.e., c.1367A > G (p. D456G), was identified in this family. Bioinformatics analysis indicated that this variant was pathogenic. Functional analysis showed that in comparison with the WT, the mutation c.1367A > G of AIFM1 showed no effect on its subcellular localization and the ability to induce apoptosis, but changed its protein expression level.

Conclusion: A novel variant of AIFM1 was identified for the first time, which was probably the genetic cause of AN in a Chinese family with AN.

KEYWORDS auditory neuropathy, hearing loss, mutation, apoptosis inducing factor mitochondrion associated 1 (AIFM1), next-generation sequencing (NGS)
Introduction

Auditory neuropathy (AN) is a type of auditory disorder mainly characterized by impaired speech comprehension ability, which manifests as a group of specific syndromes in audiology, also known as the auditory neuropathy spectrum disorder (ANSKD) (Starr et al., 1996; El-Badry and McFadden, 2009; Talaat et al., 2009).

The pathophysiology of AN indicates that the outer hair cells show normal function, whereas the inner hair cells and auditory nerves as well as their synapses are dysfunctional. The clinical manifestations include: (1) the degree of impaired speech recognition rate is inconsistent with the degree of hearing impairment, and the word comprehension ability and speech discrimination ability are poor, especially in noisy environment (Elrharchi et al., 2020); (2) the audiological examination is characterized by normal cochlear microphonic (CM) and meaningful otoacoustic emission (OAEs), but the auditory brainstem response (ABR) waveform is either missing or illegible in the patient (Starr et al., 1996). Studies showed that the prevalence rate of AN in neonates is 0.009%, and about 6.5% of the congenital sensorineural hearing loss is caused by AN (Boudewyns et al., 2016). AN is also one of the refractory diseases leading to hearing and speech communication disorders in both infants and adolescents, accounting for 10% of permanent hearing loss in children (Chinese Multi-center Research Collaborative Group on Clinical Diagnosis and Intervention of Auditory Neuropathy, 2022).

According to the age of onset, cases of AN are divided into two groups, i.e., congenital AN and late-onset AN. Studies have shown that the etiology of AN is complex, with about 48% as idiopathic, 10% related to various factors, such as ototoxic drugs and metabolic, immune, and infectious factors, and about 42% as genetic (Moser and Starr, 2016). There are three types of inheritance pattern for AN: autosomal recessive, autosomal dominant, and X-linked recessive (Wang et al., 2020). To date, among the over 20 genes associated with AN, apoptosis inducing factor mitochondrion associated 1 (AIFM1) is considered as the most prevalent gene associated with the late-onset AN and X-linked recessive inheritance mode (Zong et al., 2015; Wang et al., 2020).

First cloned and named in 1999, AIFM1 is located in the Xq25-q26 region of human chromosomes (Susin et al., 1999). The AIFM1 gene is 36.471 kb in length, containing a total of 16 exons and encoding a 613-amino acid apoptosis-inducing factor (AIF). The AIF is a flavin protein with oxidoreductase activity, composed of two flavin adenine dinucleotide (FAD) domains, a nicotinamide adenine dinucleotide (NADH) binding domain, a C-terminal domain with oxidoreductase activity, and a mitochondrial localization signal (MLS) at the N-terminal (Yuste et al., 2011). The AIFM1 has two main functions. First, it is involved in the caspase-independent death effect when mitochondria are transferred to nuclei under apoptosis stimulation, resulting in caspase-independent programmed cell death (Joza et al., 2001). Second, as a NADH oxidoreductase that is dependent on FAD, AIFM1 is crucial for oxidative phosphorylation, redox control, and respiratory chain activity in the organism (Ghezzi et al., 2010; Itai et al., 2011).

To date, a total of 19 AIFM1 variants have been identified as pathogenic factors for AN (https://www.hgmd.cf.ac.uk/; accessed on 03 October 2022). It is conceivable that there are other AIFM1 variants still remaining uncovered responsible for pathogenesis of AN. Therefore, in this study, we attempted to detect the pathogenic variants of AIFM1 in a Chinese family with AN enrolled in our hospital and to further investigate the effects of this variant on AN development.

Materials and methods

Subjects

One patient and eight unaffected individuals from a Chinese family as well as a total of 200 Chinese individuals with no genealogical relationships (i.e., controls) were included in this work. A comprehensive clinical evaluation, including the disease history and the audiological tests, was performed on all participants. The audiological tests included the pure tone threshold (PTA), distortion product otoacoustic emission (DPOAE), auditory brainstem response (ABR), speech discrimination score (SDS), and auditory steady-state response (ASSR). The informed consent was signed by each participant prior to the clinical evaluation. The study was conducted in accordance with the Declaration of Helsinki with all procedures performed in accordance with the ethical standards of Shandong Provincial Hospital (Approval # SWYX: NO. 2021-511).

Targeted next-generation sequencing

The genomic DNA was extracted from 1 ml whole blood using the Genomic Blood DNA Extraction Kit (Axygen, San Francisco, CA, United States) according to the manufacturer’s instruction. The proband’s DNA was diluted and then fragmented, ligated, amplified, and purified. GenCap® deafness gene capture probe V4.0 and GenCap® mitochondrial ring gene capture probe V1.0 (MyGenostics, Beijing, China) were used to capture the exon regions and the adjacent 20-bp intron regions of 406 genes of the proband. After elution, amplification, and purification, the captured regions were double-ended sequenced using Illumina HiSeq X TEN high-throughput sequencing platform (Illumina, United States). The detailed
procedures of library preparation, sequencing, and bioinformatics analysis were described in our previous study (Xiao et al., 2016).

### Polymerase chain reaction (PCR) amplification and Sanger sequencing

PCR was performed in a 50-μl reaction mixture with the specific primers (forward primer 5′-TCCCTTTGTATGAAAGCTAACTGG-3′ and reverse primer 5′-CCATTACAAGTGTTTCTTGAGCC-3′), as previously described (Bai et al., 2014). After the purification, the PCR products were sequenced by the ABI 3500 Genetic Analyzer (Thermo Scientific, Applied Biosystem, CA, United States). Nucleotide variations were identified by aligning the sequences with the AIFM1 available (GenBank accession NM_004208.4) using the Chrome software.

### Pathogenicity assessment of AIFM1 variant

The harmfulness of variant was evaluated with pathogenicity prediction tools and databases, including REVEL (Ioannidis et al., 2016), SIFT (http://sift.jcvi.org/), PolyPhen-2 (http://genetics.bwh.harvard.edu/pph2/), and Mutation Taster (http://www.mutationtaster.org/). The novelty of variant was determined by screening based on the 1000 Genomes (1000G) and Exome Aggregation Consortium (ExAC) databases as well as the literature and publicly available databases, i.e., the Deafness Variation Database (DVD) (http://deafnessvariationdatabase.org/; accessed on 03 October 2022) and the ClinVar database (http://www.ncbi.nlm.nih.gov/clinvar/; accessed on 03 October 2022).

A total of eight AIFM1 protein sequences of eight animal species, including *Homo sapiens* (Q95831), *Pan troglodytes* (K7BY76), *Bos taurus* (A0A4W217F3), *Macaca mulatta* (F7C728), *Canis lupus familiaris* (A0A8COJLH6), *Rattus norvegicus* (Q9JM53), *Mus musculus* (Q9Z0X1), and *Danio rerio* (Q5XFY2), were downloaded from the UniProt database to perform the multiple sequence alignment by the Clustal Omega online tools (https://www.ebi.ac.uk/Tools/msa/clustalo/; accessed on 03 October 2022) to assess the conservation of the AIFM1 proteins.

### Modeling analysis of AIFM1 protein structure

To determine the effects of AIFM1 variant on the protein structure, the 3-dimensional (3D) molecular structures of human wild type (WT) and mutant AIFM1 proteins were simulated using an automated homology modeling program I-TASSER (http://zhanglab.ccmb.med.umich.edu/; accessed on 03 September 2022) with the protein structure visualized by Swiss-PdbViewer 4.1.

### Cell culture

The Shandong Provincial Key Laboratory of Otology provided the HEK293 cells, which were cultured in the Dulbecco’s modified Eagle’s medium (DMEM) (Gibco, Grand Island, NY, United States) containing 10% fetal bovine serum (FBS) (Gibco, Grand Island, NY, United States) and 1% penicillin/streptomycin (Macgene, Beijing, China) with 5% CO₂ at 37°C.

### Transient transfection and immunofluorescence analysis

The WT and mutant AIFM1 plasmids expressing GFP-tagged (pCMV3-C-GFPSpark) were synthesized by BioSune Biotech (Jinan, China). Sequences of WT and mutant AIFM1 plasmids were verified by Sanger sequencing. The HEK293 cells were planted onto the 24-well plates and grown to 60% confluence. The expression plasmids of WT and mutant AIFM1 were transfected into cells by using Lipofectamine 3000 transfection reagent (Invitrogen, Waltham, MA, United States). In 48 h, the cells were fixed in 4% paraformaldehyde for 15 min, washed thrice with PBS, and then incubated with DAPI (D9542, Sigma-Aldrich, St. Louis, MO, United States) for 10 min in dark. Finally, the samples were washed three times with PBS. After sealing, the samples were observed and imaged through a Leica TCS SP8 confocal fluorescence microscope (Leica Microsystems, Biberach, Germany). The relative fluorescence intensity was quantified with ImageJ software.

### Western blot

Total protein was extracted from the HEK293 cells transfected with the WT and mutant AIFM1 plasmids. The proteins were denatured and separated by 10% SDS-PAGE electrophoresis and then transferred to polyvinylidene fluoride (PVDF) membranes (ISEQ00010, Merck Millipore, China). After blocking, the membranes were incubated with mouse monoclonal anti-GFP antibody (Proteintech, Wuhan, China) and mouse monoclonal anti-β-actin antibody (ZSGB-Bio, Beijing, China), and followed by anti-mouse IgG conjugated horseradish peroxidase (ZSGB-Bio, Beijing, China). Finally, the immunoblots were detected with an Immobilon Western HRP Substrate kit (WBKLS0100, Millipore, Schaffhausen, Switzerland) using the enhanced chemiluminescence system.
(Tanon5200, Shanghai, China). The gray scale of each band was quantified with ImageJ software and normalized by β-actin.

**Tunel assay**

The human HEK293 cells seeded in 24-well culture plates were transfected with plasmids of WT and mutant AIFM1 by using Lipofectamine 3000 transfection reagent (Invitrogen, Waltham, MA, United States). In 48 h, the cells were fixed in 4% paraformaldehyde for 30 min and permeabilized with proteinase K (20 μg/ml) at room temperature for 5 min. Each sample was stained with 50 μl TUNEL TdT Enzyme working solution (Meilunbio, Dalian, China) for 60 min at 37°C. The fluorescence intensity of the sample was detected with EX at 546 nm and EM at 570 nm.

**Statistical analysis**

Data presented as mean ± standard deviation (SD) of three biological replicates for each experiment were analyzed using one-way ANOVA (nonparametric or mixed) with Dunnett’s multiple comparison test and two-tailed unpaired t-test by Graphpad Prism (V8.3.0). The significant difference was set at the two-tailed p value less than 0.05.

**Results**

**Participants and clinical evaluations**

A total of nine members of a Chinese family were included in this study, including one affected (III-2) and eight unaffected (Figure 1A). The proband III-2 was a 21-year-old male and complained of hearing loss in both ears for 6 years, capable of hearing sounds but not clearly, with occasional tinnitus, which was more pronounced in noisy environments. The results of ABR showed that there was no significant waveform change observed in both ears of the proband with the stimulated of 96 db (Figure 1B). The PTA test results showed that the proband had low-frequency mild hearing loss in both ears (Figure 1C) with the SDS in the left and right ears of 45% and 43%, respectively, showing a disproportionate decline in pure tone hearing. The ASSR test showed that its threshold was significantly higher than...
that of PTA (Figure 1D). Notably, the DPOAE response was completely preserved (Figure 1E). These results of audiological tests indicated that the proband was presented with typical symptoms of AN.

Identification of a novel AIFM1 variant

The NGS analysis was performed based on the proband. The result showed that there was a base substitution (A to G) at position 1367 in the coding region of AIFM1 gene. Then, the candidate variant was confirmed by Sanger sequencing (Figure 2), showing that the proband carried the c.1367A > G hemizygous variant, the proband’s sister and mother had the c.1367A > G heterozygous variant, and the proband’s father did not have the variant, indicating that the variant of c.1367A > G came from the mother, and the WT allele came from the father. We then performed Sanger sequencing on other members of the family and found that the variant was co-segregated with the phenotypes of AN in this family (Figure 1A) with the X-linked recessive inheritance pattern in this family. The results of Sanger sequencing also showed that this variant was absent in the 200 controls. Furthermore, this variant was not detected in neither ExAC nor 1000G databases and was not reported in the literature and publicly available databases, i.e., the DVD and the ClinVar databases (Table 1). These results indicated that the novel variant of AIFM1, i.e., the missense mutation c.1367A > G, was identified in the proband of this Chinese family in this study.

Bioinformatics analysis of the pathogenicity of the AIFM1 variant

The results of bioinformatics analysis showed that the variant c.1367A > G was predicted to be deleterious by REVEL, SIFT, Polyphen-2, and Mutation Taster (Table 1). As showed in Figure 3A, the variant c.1367A > G was located in exon 13 of the FAD region of AIFM1, altering the triplet codon to substitute an aspartic (Asp, D, GAT) with the glycine (Gly, G, GGT) at position 456 (p. D456G). The results of the multiple sequence alignment showed that the AIFM1 p. D456G occurred at the evolutionarily highly conserved amino acids among eight species of vertebrates (Figure 3B). The 3D structures of WT and p. D456G AIFM1 were modeled based on the crystal structure of AIFM1 to reveal that compared with the WT, the structure of p. D456G AIFM1 protein was evidently changed (Figure 3C) with the polar, acidic amino acid Asp replaced by the non-polar amino acid Gly at position 456 and the secondary structure of the protein changed significantly posterior to the amino acid at position 456.

Functional analysis of the AIFM1 variant

After the transfection of WT and mutant AIFM1 plasmids into HEK293 cells, the subcellular localization of AIFM1 was detected by GFP fluorescence based on the fusion between the GFPSpark-tag and the C-terminal of AIFM1 protein. The results showed that green fluorescence was present in the entire cell transfected with vector plasmids and in the cytoplasm of the cells transfected with mutant and WT plasmids, indicating that both the WT and mutant AIFM1 proteins were localized in the cytoplasm (Figure 4A). Furthermore, immunofluorescence and quantitative results showed that the fluorescence intensity of the p. D456G AIFM1 was significantly lower than that of the WT

| Gene | Variant | REVEL | SIFT | Polyphen-2 | Mutation | ExAC | 1000 G | DVD | ClinVar |
|------|---------|-------|------|------------|-----------|------|--------|-----|---------|
| AIFM1 | c.1367A > G (p. D456G) | Probably damaging | Damaging | Damaging | Disease causing | Novel | Novel | Novel | Novel |

TABLE1 Characteristics of AIFM1 variant, analysis of predicted protein structure and disease-causing effects based on various databases.

Abbreviations: AIFM1, Apoptosis inducing factor mitochondrion associated 1; c, variation at cDNA level; p, variation at protein level; ExAC, Exome Aggregation Consortium; 1000 G, 1000 Genomes; DVD, Deafness Variation Database.
AIFM1 (Figures 4A,B). Western Blot analysis was conducted to further investigate the effects of AIFM1 variant on the protein expression. The results showed that the AIFM1 protein expression was not detected in the vector group transfected with vector plasmids, while the WT AIFM1 protein was detected with the expected molecular weight, suggesting that the staining reaction observed in the immunocytochemical analysis was specifically derived from the GFPSpark-tagged AIFM1 proteins. The molecular weight of p. D456G AIFM1 protein was the same as that of the WT AIFM1 protein but with weaker band than that of the WT AIFM1 protein (Figure 4C). The quantitative results showed that the expression level of p. D456G AIFM1 protein was significantly decreased compared with the WT AIFM1 protein (Figure 4D). In order to investigate the effect of p. D456G on cell survival, we performed the TUNEL staining assay (Figure 5). The results showed that compared with the vector group, HEK293 cells transfected with WT AIFM1 plasmid and c.1367A > G AIFM1 plasmid showed enhanced apoptosis, showing no significant difference between these two treatments.

Discussion

As a specific type of hearing loss, AN is characterized by impaired language comprehension. In most cases, AIFM1 gene is responsible for the cases of late-onset AN (Li et al., 2021), which contributes to the maintenance of normal mitochondrial morphology and physiology as well as caspase-independent apoptosis (Rance and Starr, 2015). In the present study, the possible genetic cause and molecular mechanisms underlying the development of AN were explored in a Chinese family with AN. The onset ages of patients with late-onset AN are mostly in the range of 5–20 years old with diverse clinical manifestations (Wang et al., 2020). Studies have shown that about 92.5% (360/389) of AN patients are presented with bilateral hearing loss.
Bing et al., 2019) with audiogram mainly of low-frequency ascending type and diverse pure tone hearing manifestations which are identified as normal or mild to severe hearing loss (Wang et al., 2003; Wang et al., 2006; Zong et al., 2015; Wang and Starr, 2018), accompanied by significant decline in speech recognition ability (Starr et al., 1996). The results of our study were consistent with the above characteristics reported in the literature, showing that the proband of the family in this study presented with typical symptoms of late-onset AN, i.e., the onset age of the proband was 15 years old, the audiogram showed low-frequency ascending type (Figure 1C), the PTA showed low-frequency mild hearing loss in both left and right ears with a disproportionate decrease in SDS (Figure 1C), the ABR waveform was illegible (Figure 1B), and the DPOAE was normal (Figure 1E). It was worth noting that although low-frequency ascending type was the most common type of hearing pattern of AN patients, including the patients involved in this study, there are still great differences in hearing patterns among AN patients. For example, a follow-up study found that some AN patients initially showed a low-frequency ascending hearing pattern, but with the progress of the disease, their high-frequency hearing was decreased rapidly to show a descending hearing pattern (Wang et al., 2020). Furthermore, some AN patients present with a descending hearing pattern involving the full frequency were accompanied by severe motor development impairment and mental retardation (Wang et al., 2019).

**FIGURE 4**
Subcellular localization and protein expression of wild type (WT) and p. D456G AIFM1 proteins in HEK293 cells. (A) Subcellular localization of WT and p. D456G AIFM1 proteins in HEK293 cells shown in DAPI, GFP-tagged, and merged images. GFP fluorescence is detected after the transient transfection in HEK293 cells. Scale bar = 40 μm. (B) Quantitative analysis of fluorescence intensity based on images shown in (A). The data are presented as mean ± standard deviation (SD) of three biological replicates. The significant difference is set at p < 0.01 (**) based on the two-tailed unpaired t-test. (C) Expressions of WT and p. D456G AIFM1 in transfected HEK293 cells based on Western blot using anti-GFP and anti-β-actin (i.e., internal control) antibodies. (D) Quantitative analysis of proteins shown in (C). The data are presented as mean ± standard deviation (SD) of three biological replicates. The significant difference is set at p < 0.0001 (****) based on the one-way ANOVA with Dunnett’s multiple comparison tests.
Furthermore, consistent with previous studies (Wang and Starr, 2018), the threshold of ASSR results in our study was also significantly higher than that of PTA (Figure 1D). Moreover, another important clinical manifestation revealed in our study was that the proband complained that his speech recognition ability was worsened in the noisy environment, which was in accordance with the results of a controlled study of AN patients (Lan et al., 2019), suggesting that the speech recognition ability of AN patients with sound speech recognition rate in quiet environment was significantly decreased in the noisy environment. Meanwhile, the proband also showed tinnitus and aggravation under noise environment, which was consistent with the previous study (Xie et al., 2019).

Since the genetic factors account for 42% of the etiology of AN (Moser and Starr, 2016), the genomic DNA of the proband was extracted for NGS analysis. Combined with Sanger sequencing technology, the results showed that the c.1367A>G variant of AIFM1 carried by the proband came from the mother, and this variant was co-segregated with the phenotype of the family and was not identified in a total of 200 healthy controls. Base on the literature and publicly available databases, the variant c. 1367A > G has not been reported, indicating that this variant of AIFM1 gene identified in this study was a novel one, expanding the pool of the mutations of AIFM1 gene. Meanwhile, this variant was predicted as deleterious by REVEL, SIFT, Polyphen-2, and Mutation Taster (Table 1), suggesting the potentially pathogenic property of this variant. To date, a total of 19 pathogenic variants of AIFM1 related to AN have been reported in literature (Elrharchi et al., 2020; Wang et al., 2020) and databases (Figure 6). The missense mutation (c.1367A > G) revealed in our study provided additional powerful support for molecular diagnosis of AN in clinical settings. These results evidently showed that a novel pathogenic AIFM1 variant, i.e., a missense mutation (c.1367A > G) in our study was first identified.

The variant of c.1367A > G was located in exon 13 of AIFM1 gene, causing the amino acid change of p. D456G in the second FAD domain of AIFM1 protein. Asp is an acidic polar aliphatic amino acid (Agius et al., 2018), which could be used as one of excitation-related neurotransmitter receptors in mammals (Kondoh et al., 2010), while Gly is a kind of non-polar aliphatic amino acid and an inhibitory neurotransmitter in the
as an apoptosis-inducing factor, the AIFM1 is a flavin protein localized in the inner membrane space of mitochondria and is transported from mitochondria to the nucleus to induce cell apoptosis upon the occurrence of the apoptotic injury (Joza et al., 2009). The immunofluorescence analysis showed that the subcellular localization of p.D456G AIFM1 was consistent with that of the wild-type AIFM1, probably due to the MLS (Yuste et al., 2011) and nuclear localization signal (NLS) (Wang et al., 2016) of AIFM1.

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Furthermore, the immunofluorescence analysis also showed that the fluorescence intensity of p. D456G AIFM1 was weaker than that of WT AIFM1, indicating that the variant of p. D456G changed the expression level of AIFM1. To further investigate the effect of p. D456G mutation on AIFM1 protein expression, western blot analysis was performed. The results were consistent with those of the immunofluorescence analysis, showing that the protein expression of p. D456G AIFM1 was significantly lower than that of wild-type AIFM1. As predicted by our 3D protein model, the p. D456G mutation affected the secondary/tertiary structure of the protein, resulting in degradation or low expression of the protein, which was further supported by the results of western blot. Previously, many scholars have studied the effect of AIFM1 mutation on protein expression with varied results. For example, patients with p. G338E (DiDato et al., 2016) and p. F210S (Sancho et al., 2017) variants all showed motor nerve lesions, and the expression level of AIFM1 was decreased. These results were consistent with the findings revealed in our study. Interestingly, in a study on axonal polyneuropathy caused by AIFM1 p. F210L variant (Hu et al., 2017), no significant change was detected in the AIFM1 protein expression, independent of the changes of different amino acids at the same site. Furthermore, in a study of mitochondrial encephalomyopathy caused by R201del variant in AIFM1, the protein expression was not significantly altered (Ghezzi et al., 2010).

Under normal circumstances, AIFM1 is first transcribed and translated into a 67-kD precursor molecule (Liu et al., 2018), which is then transported to mitochondria through the N-terminal domain MLS sequence. The mitochondrial processing peptidase (MPP) cleaves AIFM1 in mitochondria into 62-kD mature AIFM1, which acts as the FAD-dependent NADH oxidoreductase to contribute to the stabilization and maturation of mitochondrial oxidation of respiratory chain complex I as well as the removal of peroxide from the mitochondria (Susin et al., 1999). When cells are damaged by apoptosis, mature AIFM1 is cleaved into soluble apoptotic precursor protein, i.e., the truncated AIF (tAIF) of about 57 kD (Susin et al., 1999; Joza et al., 2001). The tAIF is released from mitochondria into cytosol and nucleus to induce two typical caspase-independent apoptosis phenomena: chromatin condensation and fragmentation of large DNA fragments of approximately 50 kD (Yuste et al., 2011).

A study of AIFM1 p. P488L variants in auditory neuropathy and peripheral neuropathy showed that AIFM1 p. P488L variants caused a mild increase in the rate of caspase-independent apoptosis in cells (Wang et al., 2019). Because the immunofluorescence and WB assays showed that the p. D456G variant reduced AIFM1 protein expression, the Tunel assay was performed to evaluate its adverse effect on cell survival. The results showed that the ability of AIFM1 to induce apoptosis was not significantly affected by the p. D456G mutation, showing the same effect on cell survival as that of WT AIFM, indicating that the p. D456G variant did not affect cell survival. This may also be the reason for the generation of mild symptoms of AN patients. Previous studies showed that the AIFM1 R201del variant caused severe mitochondrial encephalomyopathy and significantly increased parthanatos-linked cell death (Ghezzi et al., 2010). In the study of Cowchock Syndrome caused by AIFM1 p. E493V variant, Tunel assay of muscle biopsy showed a large number of apoptotic cells in the samples of the mutated individuals, while little or no staining was detected in the muscles of healthy controls (Rinaldi et al., 2012). However, in patients with distal motor neuropathy carrying the p. F210S mutation, the mutation did not enhance the apoptosis (Sancho et al., 2017), which was consistent with our study. Similarly, as mentioned above, the p. F210S mutation also reduced the protein expression level with mitochondrial fragmentation also observed in fibroblasts of this patient, suggesting that the development of distal motor neuropathy in this patient may be related to the defective mitochondrial respiration (Sancho et al., 2017). Therefore, we hypothesized that this mutation might affect the role of AIFM1 in oxidative phosphorylation, redox control, and respiratory chain activity. Study showed that knockdown of AIFM1 could attenuate mitochondrial respiration and ATP production, ultimately affecting cell functions (Zong et al., 2020). Furthermore, some studies showed that in the auditory pathway, including inner hair cells, glial cells in neural pathways, and spiral ganglion cell (SGC), the normal energy metabolism was the key to maintain its physiological activity (Yang et al., 2015). Therefore, as suggested in the previous studies (Zong et al., 2015), the variant revealed in our study may cause AN by affecting the function of AIFM1 protein in mitochondria, ultimately affecting the mitochondrial respiration and ATP synthesis in inner ear. Moreover, as mentioned above, the 3D prediction results showed that the p. D456G mutation affected both the secondary and tertiary structures of the C-terminal, while the binding of the N-terminal of CHCHD4 and the C-terminal of AIFM1 played an important role in the biogenesis of mitochondrial respiratory chain complex I. The decreased expression level of AIFM1 could affect this interaction, and then affect the function of the mitochondrial respiratory chain. These functional experiments further demonstrated that the novel mutation identified in this study could cause damage to AIFM1 protein, which could be the underlying genetic etiology of this family with AN, and further experiments of mitochondrial respiratory chain activity and oxidative phosphorylation are necessary to explore the pathological mechanism of this mutation.

In summary, we identified a novel c.1367A > G AIFM1 variant in a Chinese family with AN by targeted capture sequencing, expanding the AN-related mutation spectrum of AIFM1. Bioinformatics prediction and functional analysis showed that AIFM1 c.1367A > G was a pathogenic mutation and may be the genetic cause of AN in this Chinese family, providing additional molecular and clinical evidence to support the establishment of a strong genotype-phenotype correlation for AN.
Data availability statement

The data presented in the study are deposited in the the NCRI Sequence Read Archive (SRA) repository, accession number PRJNA891130.

Ethics statement

The studies involving human participants were reviewed and approved by the ethics committee of Shandong Provincial Hospital. Written informed consent to participate in this study was provided by the participants’ legal guardian/next of kin.

Author contributions

ZL and SY designed the research; RW analyzed the data and wrote the paper; XB, HY, and JM performed the research; RW, XB, and HY performed the clinical work.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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