A likely pathogenic POLD1 variant associated with mandibular hypoplasia, deafness, progeroid features, and lipodystrophy syndrome in a Chinese patient

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Abstract

Background: Mandibular hypoplasia, deafness, progeroid features, and lipodystrophy syndrome (MDPL; OMIM# 615381) is a rare autosomal dominant disorder, with only a few reported cases worldwide. Herein, we describe the clinical features and underlying molecular etiology of MDPL syndrome in an 8-year-old Chinese patient.

Methods: We performed otological, endocrine, ultrasound, and radiological examinations, as well as genetic testing. Additionally, the literature concerning MDPL was reviewed to do a retrospective analysis of the pathogenesis, genotype–phenotype correlation, and clinical management.

Results: The proband was diagnosed with MDPL, presenting with mandibular hypoplasia, a characteristic facial appearance, lipodystrophy, and sensorineural hearing loss (SNHL). Whole-exome sequencing and bioinformatics analysis revealed a de novo missense variant in the POLD1 gene, NM_002691.4:c.3185A>G (NP_002682.2:p.(Gln1062Arg)). The retrospective analysis showed wide variation in the MDPL phenotype, but the most frequent features included mandibular hypoplasia, characteristic facial appearance, lipodystrophy, and SNHL.

Conclusions: This study supplements the mutational spectrum of POLD1. The genetic analysis contributes to the diagnosis of syndromic deafness, and it has a vital role in clinical management and future genetic consultation.

Keywords: Mandibular hypoplasia, progeroid features, and lipodystrophy syndrome, MDPL, Sensorineural hearing loss, POLD1

Introduction

Mandibular hypoplasia, deafness, progeroid features, and lipodystrophy syndrome (MDPL; OMIM# 615381) is a rare disease characterized by mandibular hypoplasia, a characteristic facial appearance, lipodystrophy and sensorineural hearing loss (SNHL). Shastry et al. [1] first described MDPL in seven unrelated patients from mandibulofacial dysplasia (MAD) patients without pathogenic variants in known causal genes. In four unrelated patients with MDPL, Weedon et al. [2] first reported a recurrent heterozygous in-frame deletion of the POLD1 gene, c.1812_1814del (p.(Ser605del)) through exome sequencing. The clinical manifestations of MDPL also include prominent eyes, crowded teeth, poor breast development in females, hepatomegaly, a small mouth, hepatic steatosis, cryptorchidism and hypogonadism in

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males, osteoporosis, joint contractures, kyphosis, scoliosis, a beaked nose, tight skin, scleroderma-like changes, telangiectases, skin atrophy, a high-pitched voice, insulin resistance, diabetes mellitus, abnormal liver function tests, and hypertriglyceridemia. MDPL should be differentiated from MAD (OMIM# 248370, 608612) and Werner syndrome (WS; OMIM# 277700). MAD was transmitted in an autosomal recessive pattern, and the etiology was attributed to LMNA or ZMPSTE24 alterations. MAD patients presented mainly with growth retardation, craniofacial anomalies, skeletal anomalies, and skin changes. Compared to MAD patients, MDPL patients have a better overall prognosis, normal hair, no alopecia, clavicle dysplasia, or acroosteolysis [1]. WS, which results from pathogenic variants in the WRN gene, is an autosomal recessive disorder characterized by premature aging and an increased risk of various forms of cancer. Compared to WS patients, MDPL patients develop a facial appearance characteristic of premature aging at a younger age, but do not exhibit premature loss or graying of hair [3].

The present study reports an MDPL case in a Chinese family. Whole-exome sequencing identified a likely pathogenic variant in the POLD1 gene. A literature review was performed to examine the phenotypic variability of MDPL.

Materials and methods

Patients and clinical investigations

An 8-year-old girl who suffered from hearing loss presented to the Department of Otorhinolaryngology, Affiliated Hospital of Zhengzhou University, China. Audiological evaluation revealed severe bilateral SNHL, with no indication of a conductive hearing loss. Data related to the pedigree of the patient were collected from her parents. The clinical assessment was performed, including audiological, endocrine, ultrasound, and radiological examinations. The project was approved by the Ethics Committee of the Affiliated Hospital of Zhengzhou University (No.: 2018008) and all described procedures were complied with the Helsinki Declaration. A signed informed consent form was obtained from the proband’s parents.

Genetic testing and bioinformatics analysis

Whole blood (3 ml) was collected from the affected proband, her parents, and her younger sister. Genomic DNA was isolated from the blood following the manufacturer’s protocol of the Genomic DNA Purification kit (GenMagBio, Changzhou, China). The whole-exome sequencing procedures, including fragmentation, end-repair of genomic DNA, targeted enrichment, and sequencing, were identical to those described in previous studies [4]. After trimming the adapters and low-quality reads, cleaned reads were mapped to the human reference genome (version GRCh37). Bioinformatics analyses, including variant calling, annotation, and filtering, were performed as those described previously [4]. The ACMG (American College of Medical Genetics) standards and guidelines were used to interpret sequence variants [5]. Variant nomenclature was based on POLD1 canonical transcript NM_002691.4. To verify the candidate variants and test for co-segregation in the kindred, Primer-BLAST was used to design primers for c.3185A>G (forward primer 5'-CAGGAGCCGTGTGTAGTTG-3' and reverse primer 5'-TCAACAGCTGGAAGGGATG-3'). Then PCR and Sanger sequencing were carried out.

Literature review

Using MDPL and POLD1 as keywords, PubMed was searched for articles published between 2010 and 2021 on the genotype and phenotype of MDPL.

Results

Case presentation

The proband (II-1, Fig. 1a), aged 8-year-old, from a non-consanguineous Chinese parent. She was born at 32 weeks of gestation with a birth weight of 2.1 kg. She exhibited poor growth and thin limbs since the age of 4 years. She is allergic to fish, eggs, milk, and wheat. She had prominent eyes, a beaked nose, a small mouth, kyphosis, tight skin, and a high-pitched voice. She was diagnosed with severe bilateral SNHL on the basis of pure-tone audiometry tested at six years of age. Her bone age was delayed, and was estimated at three years (Fig. 2a, b). Distortion product otoacoustic emissions were absent at all frequencies on both sides. The auditory brainstem response was evaluated to determine the average (right, 100 dB nHL normal hearing level; left, 90 dB nHL) and bone conduction (right, 50 dB nHL; left, 50 dB nHL) thresholds. The auditory steady-state response thresholds at 500,1,000, 2,000, and 4000 Hz were 70, 90, 90, and 90 dB nHL in the left ear, and were 70, 90, 90, and 80 dB nHL in the right ear, respectively. X-rays showed crowded teeth and mandibular hypoplasia (Fig. 2b). Temporal bone CT was normal. Osteoporosis, joint contractures, and insulin resistance were not observed. The levels of glucose, triglycerides, insulin, aspartate transaminase, alanine transaminase, and total bilirubin were within the normal limits. No internal organ abnormalities were observed by ultrasound. At the age of 8 years, the patient’s height and weight were 1.26 m and 16 kg, respectively. The follicle-stimulating hormone, luteinizing hormone, estradiol, and prolactin levels were 25.7 mIU/mL, 0.92 mIU/mL, 8.44 pg/mL, and
Fig. 1 Pedigree and Sanger sequencing of the MDPL family. a Pedigree of the MDPL family b Sanger sequencing

Fig. 2 Facial phenotype, audiogram, and X-ray of the proband. a Bilateral severe sensorineural hearing loss revealed by pure-tone audiometry showed. The use of hearing aids led to a successful outcome. b When she was 6 years old, the bone age was only 3 years. X-ray showing crowded teeth and mandibular hypoplasia
41.2 ng/mL, respectively. She had a normal intelligence level. The use of hearing aids in both ears provided good results. The proband’s family members were unaffected and had normal hearing (Fig. 1a).

**Molecular diagnosis**

The proband carried a missense variant of the *POLD1* gene, identified by whole-exome sequencing: c.3185A>G (Fig. 1b). The variant was validated in the family members using Sanger sequencing; the results showed that the proband's parents and younger sister did not carry this variant, suggesting a de novo status in the proband (both maternity and paternity confirmed, PS2). The variant was not found previously in public databases (PM2_Supporting). The proband's phenotype is highly specific for MDPL with a single genetic etiology (PP4). Multiple lines of computational evidence, including SIFT, Polyphen2, and MutationTaster, support its deleterious effect on the *POLD1* protein (PP3). This variant has been submitted to the Clinvar database (SCV001477293) and classified as variant of uncertain significance (VUS). In summary, the variant c.3185A>G was reclassified as likely pathogenic.

**Discussions**

**Mechanism**

MDPL is caused by pathogenic variants in the *POLD1* gene, which encodes for the catalytic subunit (p125) of DNA polymerase delta (Polδ). The catalytic subunit is responsible for synthesizing the lagging strand DNA during DNA replication with both 5′- to 3′-polymerase activity and 3′- to 5′-exonuclease activity [6, 7]. Polδ is involved in DNA replication and maintains genomic stability. Downregulation of the p125 subunit causes genomic instability and ultimately DNA replication errors. The proofreading ability of Polδ results from its exonuclease activity, which is essential to ensure replication fidelity [6]. The molecular mechanism is still unclear. The p125 mutation leads to reduced genomic stability, cellular senescence, and apoptosis, which may cause deafness and progeroid features in MDPL patients.

**Molecular mapping**

Encoded by the *POLD1* gene with a size of 34 kb and 29 exons, POLD1 has polymerase and exonuclease activities on its p125 catalytic subunit. POLD1 consists of putative nuclear localization signal (amino acids 4–19), exonuclease domain (amino acids 306–519), polymerase active site (amino acids 581–910), and ZnF domain (amino acids 1012–1083), which contains two conserved cysteine-rich metal-binding motifs: CysA and CysB [7]. The six previously reported *POLD1* pathogenic variants in MDPL patients are summarized in Table 1. Most variants were de novo. The hotspot of *POLD1* variants associated with MDPL is the c.1812_1814del. Weedon et al. [2] showed using in vitro functional studies that the c.1812_1814del resulted in inefficient interaction of DNA polymerase with dNTPs and incorporation into the extended DNA strand. However, the nucleic acid exonuclease activity was retained, implying that the proofreading function was normal and did not increase the rate of

| Table 1: *POLD1* pathogenic variants in MDPL |
|---------------------------------------------|
| Nucleotide changes | Amino acid changes | Type of variants | Publications |
| c.[589_589+1del; 3298G>A] | p.(Gly1100Arg) | Missense | Oh et al. [8] |
| c.1519G>T | p.(Arg507Cys) | Missense | Pelosini et al. [9] |
| c.1812_1814del | p.(Ser605del) | In-Frame | Lessel et al. [3] |
| c.3185A>G | p.(Gln1062Arg) | Missense | Weedon et al. [2] |
| c.3199G>A | p.(Glu1067Lys) | Missense | Lessel et al. [3] |
| c.3209T>A | p.(Ile1070Asn) | Missense | Reinier et al. [10] |
| | | | Elouej et al. [7] |
| | | | Okada et al. [11] |
| | | | Wang et al. [12] |
| | | | Fiorillo et al. [13] |
| | | | Sasaki et al. [14] |
| | | | Wang et al. [15] |
| | | | Murdocca et al. [6] |
| | | | Yu et al. [16] |
| | | | Zhou et al. [17] |
| | | | This study |
| | | | Ajluni et al. [18] |
| | | | Elouej et al. [7] |

Nucleotide numbering is based on GenBank reference sequence NM_002691.4

MDPL, mandibular hypoplasia, deafness, progeroid features, and lipoatrophy syndrome

*The first variant was originally described as "NM_001256849.1:c.584_585del" in the study
| Site                      | Features                                | Frequency | Shastry et al. [1], Spo N | Weedon et al. [2], Spo | Pelosi et al. [9], Spo | Reinier et al. [10], Spo | Lessel et al. [3] | Ajluni et al. [18] | Okada et al. [11], Spo | Wang et al. [12], Spo | Fiorillo et al. [13], Spo | Sasaki et al. [14], Spo | Wang et al. [15], Spo | Oh et al. [8] | Yu et al. [16], Spo | Zhou et al. [17], Spo | Murdocca et al. [6] | This study Proband |
|--------------------------|-----------------------------------------|-----------|---------------------------|------------------------|------------------------|------------------------|-------------------|---------------------|-----------------------|---------------------|------------------------|------------------------|---------------------|----------------|-------------------|------------------|-------------------|------------------|
| Weight                   | Normal birth weight                     | 23/23     | 7/7                       | 2/2                    | +                      | +                      | 4/4               | NA                  | 2/2                   | +                  | +                      | NA                     | NA                  | +               | +                 | +                | NA                | NA               |
| Face                     | Mandibular hypoplasia                   | 32/33     | 7/7                       | 2/2                    | +                      | +                      | 7/8               | 2/2                 | 2/2                   | +                  | +                      | +                      | +                  | –                | +                 | +                | +                 | +                |
|                          | Progeroid appearance                   | 8/11      | NA                        | NA                     | +                      | +                      | NA                | 0/2                 | NA                   | NA                  | +                      | NA                     | +                  | +                | NA               | NA                | +                |
| Ears                     | Sensorineural hearing loss              | 26/32     | 7/7                       | 2/2                    | +                      | +                      | 5/8               | 0/2                 | 1/2                   | +                  | +                      | +                      | +                  | +                | +                 | +                | +                 | +                |
| Eyes                     | Prominent eyes                          | 14/16     | 7/7                       | NA                     | –                      | +                      | NA                | NA                  | NA                   | –                  | +                      | NA                     | –                  | +                | +                 | +                | +                 | +                |
| Nose                     | Beaked nose                             | 29/30     | 7/7                       | 2/2                    | +                      | +                      | 8/8               | NA                  | 2/2                   | +                  | +                      | +                      | +                  | –                | +                 | +                | +                 | +                |
| Mouth                    | Small mouth                             | 18/19     | 7/7                       | NA                     | +                      | +                      | NA                | NA                  | NA                   | +                  | +                      | +                      | –                  | +                | +                 | +                | +                 | +                |
| Teeth                    | Crowded teeth                           | 19/28     | 2/7                       | 2/2                    | +                      | +                      | 5/8               | NA                  | 2/2                   | +                  | +                      | NA                     | NA                 | +                | +                 | +                | +                 | +                |
| Breasts                  | Poor breast development (female)        | 8/8       | 2/2                       | M                      | +                      | +                      | NA                | NA                  | 1/1                   | M                  | M                      | NA                     | NA                 | NA               | M                 | +                | +                 | +                |
| Liver                    | Hepatomegaly                            | 6/14      | 3/5                       | 0/2                    | +                      | +                      | NA                | NA                  | 0/2                   | NA                  | NA                     | NA                     | NA                 | –                | NA               | –                | NA                | –                |
|                          | Hepatic steatosis                       | 12/20     | 2/5                       | 1/1                    | +                      | +                      | 1/2               | NA                  | 0/2                   | +                  | +                      | NA                     | +                  | –                | +                 | NA               | +                |
| Internal genitalia       | Cryptorchidism (male)                   | 7/10      | 3/5                       | 2/2                    | F                      | F                      | NA                | NA                  | 1/1                   | +                  | NA                     | F                      | F                  | –                | –                 | –                | –                |
| Skeleton                 | Osteoporosis                            | 9/24      | 1/7                       | 2/2                    | NA                     | NA                      | 2/7               | NA                  | 1/2                   | NA                  | +                      | –                      | NA                 | –                | NA               | NA              |
|                          | Joint contractures                     | 17/30     | 5/7                       | 2/2                    | –                      | +                      | 1/8               | 2/2                 | 2/2                   | +                  | +                      | NA                     | NA                 | –                | –                 | +                | –                |
| Spine                    | Kyphosis                                | 6/10      | NA                        | 2/2                    | +                      | –                      | NA                | NA                  | 1/2                   | –                  | NA                     | +                      | NA                 | –                | NA               | NA              |
|                          | Scoliosis                               | 4/9       | NA                        | 2/2                    | +                      | –                      | NA                | 1/2                 | –                     | NA                  | NA                     | NA                     | NA                 | –                | NA               | NA              |
| Skin                     | Tight skin                              | 18/26     | 3/7                       | 2/2                    | –                      | –                      | 8/8               | NA                  | NA                   | +                  | +                      | NA                     | –                  | +                | +                 | NA               |
|                          | Scleroderma-like changes                | 5/15      | 1/7                       | NA                     | +                      | +                      | NA                | NA                  | NA                   | –                  | NA                     | –                      | NA                 | –                | +                 | +                |
|                          | Telangiectases                          | 17/26     | 4/6                       | 2/2                    | –                      | +                      | 5/8               | NA                  | 1/2                   | +                  | NA                     | NA                     | +                  | –                | –                 | +                |

**Table 2** A review of physical features of MDPL
Table 2 (continued)

| Site Features | Frequency | Shastry et al. [1], Spo | Weedon et al. [2], Spo | Peleśni et al. [9], Spo | Reinier et al. [10], Spo | Ajiuni et al. [18], Spo | Okada et al. [11], Spo | Wang et al. [12], Spo | Fiorillo et al. [13], Spo | Sasaki et al. [14], Spo | Wang et al. [15], Spo | Oh et al. [8], Spo | Yu et al. [16], Spo | Zhou et al. [17], Spo | Murdocca et al. [6] | This study Proband |
|---------------|-----------|-------------------------|------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|
| Skin atrophy  | 9/15      | 7/7                     | NA                     | –          | +                      | NA                      | NA                     | –                      | –                      | NA                     | +                      | NA                     | –                      | –                      | NA                     | NA                     |
| Soft tissues  | 32/33     | 7/7                     | 2/2                    | +          | +                      | 8/8                     | 2/2                    | 2/2                    | +                      | +                      | +                      | +                      | –                      | –                      | +                      | +                      |
| Lipodystrophy | 23/31     | 3/7                     | NA                     | –          | +                      | 8/8                     | 2/2                    | 1/2                    | +                      | +                      | +                      | +                      | –                      | –                      | +                      | +                      |
| Loss of subcutaneous fat, particularly affecting the limbs | 7/31 | 4/7 | NA | – | 0/8 | 0/2 | 1/2 | – | – | – | – | – | – | – | – | – |
| Increased visceral fat | 14/22 | 2/5 | 2/2 | + | + | 1/2 | NA | 0/2 | + | + | + | + | – | + | NA | + | – |
| Voice High-pitched voice | 13/25 | 1/7 | 2/2 | NA | + | 5/8 | NA | 0/2 | + | + | NA | + | NA | – | NA | NA | NA | + |
| Endocrine Insulin resistance | 11/22 | 1/7 | 2/2 | + | + | NA | 2/2 | NA | – | – | + | + | + | – | – | NA | + | – |
| Diabetes mellitus | 13/33 | 2/7 | 2/2 | – | + | 2/8 | 2/2 | 0/2 | + | – | – | + | – | – | – | + | + | – |
| Hypogonadism (male) | 5/8 | NA | 2/2 | F | F | 1/2 | NA | 0/1 | + | + | F | F | F | F | F | F | F | F |
| Others Abnormal liver function tests | 7/15 | NA | 2/2 | NA | + | 0/3 | 2/2 | 0/1 | + | – | NA | + | NA | – | – | NA | NA | – |
| Hypertriglyceridemia | 20/29 | 6/7 | 2/2 | + | + | 5/6 | 2/2 | 0/2 | – | + | NA | + | – | – | – | – | + | NA | – |
| Onset of lipodystrophy in early childhood | 24/25 | 5/5 | 2/2 | + | + | 4/4 | NA | 2/2 | + | + | + | + | – | + | + | + | + | + |
| Onset of hearing loss in first or second decade | 23/28 | 7/7 | 1/2 | + | + | 5/8 | NA | 1/1 | + | + | + | + | NA | – | + | NA | + | + |

MDPL, mandibular hypoplasia, deafness, progeroid features, and lipodystrophy syndrome; N, none of molecular genetic diagnosis; Spo, Sporadic; F, female; M, male; NA, not available
base substitution errors, resulting only in abnormal DNA synthesis. In our study, the missense variant c.3185A>G was found on the CysB motif (amino acids 1058–1076).

Phenotypic variability
Lessel et al. [3] described eight POLD1 pathogenic variants carriers who did not have hearing loss or mandibular hypoplasia; these patients had a lower incidence of metabolic abnormalities and joint contractures, indicating that POLD1 pathogenic variants could result in a variable expression. The phenotypic variability of MDPL syndrome remains unclear. There are 33 reported cases of MDPL worldwide, with ~70% from Europe and the Americas and ~30% from Asia. Almost all reported cases of MDPL have mandibular hypoplasia, a characteristic facial appearance, lipodystrophy, and SNHL (Table 2). Additionally, all female patients have poor breast development. More than half of all patients have hepatic steatosis, joint contractures, kyphosis, skin changes, a high-pitched voice, and hypertriglyceridemia, with cryptorchidism and hypogonadism also seen in males. Lipodystrophy almost always occurs in early childhood, while hearing loss occurs in the first or second decade.

Treatment and prevention
Audiological, ophthalmological, endocrine, ultrasound, and radiological examinations should be regularly performed for MDPL patients. Additionally, levels of glucose, triglycerides, insulin, and liver enzymes should be measured. SNHL is usually bilateral and sometimes progressive, with onset in the first or second decade of life. The most common intervention for SNHL is a hearing aid (Table 3). The proband in our study presented with severe SNHL and had a successful outcome with the use of a hearing aid. Cochlear implants can be used for patients with profound deafness who have poor outcomes with hearing aids. However, the effectiveness of cochlear implants in MDPL patients remains to be studied.

MDPL is an autosomal dominant trait that has a 50% probability of being inherited by the offspring. Genetic counseling should be recommended for the proband and her future spouse. A preimplantation genetic diagnosis may help prevent MDPL in the offspring. Whole-exome sequencing contributes to MDPL diagnosis and is beneficial for genetic counseling of MDPL patients.

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Author contributions
WL and WT designed the study and performed genetic counseling. BZ and HX collected clinical material, drafted and revised the manuscript. LM and BZ carried out whole-exome sequencing, and data analysis. ZP and HF performed phenotypic and audiometric analysis, and helped to write the manuscript. All authors have read and approved of the manuscript.

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Availability of data and materials
All data supporting the findings of this study are available on request from the corresponding author. The likely pathogenic variant have been submitted to

### Table 3 Interventions of SNHL in reported MDPL

| Previous publications | Age at SNHL onset (years) | Interventions |
|-----------------------|---------------------------|---------------|
| Shastry et al. [1], patient 100.3 | 18 | NA |
| Shastry et al. [1], patient 200.5 | 15 | NA |
| Shastry et al. [1], patient 300.4 | 7 | NA |
| Shastry et al. [1], patient 400.3 | 8 | NA |
| Shastry et al. [1], patient 500.4 | 16 | NA |
| Shastry et al. [1], patient 600.4 | 14 | NA |
| Shastry et al. [1], patient 700.3 | 6 | NA |
| Weedon et al. [2], patient 1 | 12 | Bilateral hearing aids |
| Weedon et al. [17], patient 2 | 33 | NA |
| Pelosini et al. [9] | 25 | Hearing aid |
| Reinier et al. [10] | 10 | NA |
| Lessel et al. [3], patient 1 | N | NA |
| Lessel et al. [3], patient 2 | 12 | NA |
| Lessel et al. [3], patient 3 | 11 | NA |
| Lessel et al. [3], patient 4 | N | NA |
| Lessel et al. [3], patient 5 | 14 | NA |
| Lessel et al. [3], patient 6 | 14 | NA |
| Lessel et al. [3], patient 7 | N | NA |
| Lessel et al. [3], patient 8 | NA | NA |
| Ajluni et al. [18], patient 6 | N | N |
| Ajluni et al. [18], patient 7 | N | N |
| Eloeue et al. [7], patient 1 | N | N |
| Eloeue et al. [7], patient 2 | 4 | Bilateral hearing aids |
| Okada et al. [11] | 7 | Hearing aids |
| Wang et al. [12] | 10 | Hearing aids |
| Fiorillo et al. [13] | 2 | NA |
| Sasaki et al. [14] | Puberty | NA |
| Wang et al. [15] | 15 | NA |
| Oh et al. [8], SB127-219 | NA | NA |
| Oh et al. [8], SB127-277 | NA | NA |
| Yu et al. [16] | 25 | NA |
| Zhou et al. [17] | 20 | NA |
| Murdocca et al. [6] | NA | NA |
| This study proband | 6 | Bilateral hearing aids |

SNHL, sensorineural hearing loss; MDPL, mandibular hypoplasia, deafness, progeroid features, and lipodystrophy syndrome; N, no; NA, not available.
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