Short Communication

Novel mutation of \textit{IFT140} in an infant with Mainzer-Saldino syndrome presenting with retinal dystrophy

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ABSTRACT

A seven-month-old girl presented with bilateral roving nystagmus, hyperopia, and retinal dystrophy, and was brought to our ophthalmology clinic. Visual-evoked potentials (VEPs) were non-recordable in both the eyes. No other systemic symptoms or abnormalities were observed. Whole exome sequencing (WES) identified a compound heterozygous mutation in the \textit{IFT140} gene: c.1990G \textgreater{} A (p. Glu664Lys) and c.2214_2217del (p. Asp738GlufsTer47). The genetic results support a diagnosis of Mainzer-Saldino syndrome (MSS). Importantly, c.2214_2217del is a novel mutation in the \textit{IFT140} gene. Although the patient presents with isolated retinal dystrophy, it is crucial to monitor renal function over time. Taken together, our results reinforce the role of \textit{IFT140} in syndromic ciliopathies. This report also highlights the role of combined WES approaches in identifying underlying mutations in infants presenting with isolated retinal dystrophy, considering MSS may present differently over time.

1. Introduction

Ciliopathies are a heterogeneous group of diseases that affect intraflagellar transport (IFT) and alter cilial function [1]. Given the widespread presence of IFT in the cellular cilia and flagella of eukaryotic organisms, its mutations can give rise to a wide range of symptoms in multiple systems. IFT particles are multiprotein complexes that can be biochemically grouped into two subcomplexes, IFTA and IFTB [2], having distinct functions. IFTA is crucial in the retrograde transport whereas IFTB is crucial in the anterograde trafficking, along ciliary axonomes [3]. Mutations in IFTA form ciliary bulges due to protein accumulation at the distal tip, whereas mutations in IFTB often lead to failure in cilia formation due to defects in importing cilia-building components [4,5].

Mainzer-Saldino syndrome (MSS) is a rare autosomal recessive ciliopathy, clinically characterized by cone-shaped phalangeal epiphyses, chronic renal disease, and retinal dystrophy. Less common characteristics, such as short stature, cerebellar ataxia, and hepatic fibrosis [6]. MSS is most commonly attributed to the dysfunction of the \textit{IFT140} protein, which encodes a subunit of IFTA [7]. Pathogenic variation in \textit{IFT140} has not only been reported in patients with MSS, but isolated retinitis pigmentosa (RP), cranioectodermal dysplasia (CED), Jeune syndrome and optiz trigonocephaly syndrome (OTCS) [8,9].

Here, we report a pediatric patient carrying a novel pathogenic mutation in the \textit{IFT140} gene, presenting with isolated retinal dystrophy. Although there was no evident systemic manifestation on initial presentation, the genetic results supported a diagnosis of MSS.

2. Materials and methods

We collected the patient’s medical records, family history, and clinical presentation, along with blood samples for hemogram and
biochemistry profiles. Bone age study, renal ultrasound, liver ultrasound, and detailed ophthalmic examinations were performed. Whole exome sequencing (WES) was performed to detect possible DNA alterations. First, 100 ng genomic DNA from the sample to be tested was fragmented to an average size of 180–280 bp. Library preparation and enrichment capture were performed using Illumina platform-compatible Roche KAPA HyperExome kit. Enriched captured DNA fragments were amplified and sequenced using 2 × 150 bp Paired-End formats on a NovaSeq sequencer. Low-quality bases (Q < 30) were trimmed. Sequence data were aligned based on GRCh38 using DRAGEN (SW:05.021.595.3.7.5) and variant site searching was performed using DRAGEN (SW:05.021.595.3.7.5). Confirmed variant sites were annotated using the Variant Effect Predictor (v101) and Jannovar (0.35). Human phenotype ontology and Online Mendelian Inheritance in Man (OMIM) databases were used to identify candidate genes based on patient phenotypes. Variants previously reported as benign or likely to be benign in the ClinVar database and variants with a prevalence of >5% in the Common Variation Databases (dbSNP version 150 and Taiwan Biobank) were excluded. Genetic variants in the CDS region were analyzed for changes in protein structure using prediction softwares, including SIFT, PolyPhen-2, and MutationTaster. The variants were then clinically interpreted according to the ACMG/AMP 2015 guidelines. We also performed Sanger sequencing with further trio analysis, for the patient and her parents. All collected data were de-identified. The corresponding author has full access to all the data and bears the final responsibility for the decision to submit the data for publication.

3. Case presentation

A seven-month-old female infant was born at full term as the first child of a non-consanguineous couple. Her parents reported that she was unable to follow objects and she was referred to our ophthalmology department. On initial examination, she was blinking with light but not following objects. Roving nystagmus presented in all positions of gaze was evident, and Hirschberg test showed orthotropia. Hyperopic refractive errors of approximately +6.0 Diopter were noted in both the eyes. Anterior segment evaluation and intraocular pressure measurements were insignificant. Fundus examination revealed attenuated vessels with vascular sheathing and marbleized diffuse retinal pigment dispersion (Fig. 1C). Visual evoked potentials (VEPs) were non-recordable bilaterally (Fig. 2). Brain magnetic resonance imaging (MRI) showed no structural abnormalities. Ultrasonography at the initial presentation revealed no significant abnormalities in the kidneys, liver, or pancreas. No clinical or radiographic evidence of skeletal dysplasia was observed. We noted that this patient had some facial features, like frontal bossing, high forehead and broad nasal bridge (Fig. 1A). However, she did not have narrow thorax, shorter ribs or any other skeletal abnormalities. No hearing impairment was observed.

Notably, WES illustrated two important compound heterozygous
mutations affecting the IFT140 (intraflagellar transport 140) gene. A paternal missense mutation NM_014714.4:c.1990G>A, NP_055529.2: p. Glu664Lys (located at chr16:1564074) and a maternal frameshift mutation NM_014714.4:c.2214_2217del, NP_055529.2: p.Asp738GlufsTer47 (located at chr16:1558117–1558120) are shown in Fig. 1B and Fig. 3. The former mutation has been reported to be pathogenic in the ClinVar database (variant ID:31679) with a global allele frequency of 0.004% (gnomAD–Exomes) and an East Asian frequency of 0.004% (gnomAD–Genomes). This mutation has been correlated with Mainzer-Saldino syndrome (MSS) in previous studies (PMID: 26359340, PMID: 26216056, PMID:22503633). The c.1990G > A (p.Glu664Lys) changes showed approximately 80% disorganization of IFT140 and loss of basal body localization associated with an increase in the cytoplasmic portion (PMID:22503633). The second mutation observed, c. 2214_2217del (Asp738GlufsTer47), was a novel mutation. Currently, there is no incidence record in commonly used databases, including ExAC, 1000 genomes, gnomAD, and the Taiwan Biobank. This variant has neither been documented in ClinVar nor been reported in PubMed. Since this mutation causes premature termination of protein translation, the prediction software MutationTaster predicts that the effect of this mutation on the protein results is “Disease causing,” and the point mutation is classified as “Pathogenic” according to the ACMG guidelines. Based on these findings, the patient was diagnosed with MSS.

4. Results and discussion

Ciliopathies are genetic disorders associated with the dysfunction of primary cilia which affect multiple organ systems [10]. Primary cilia
operate as cellular signaling centers that modulate various extracellular stimuli, and they are essential in regulating cellular responses during early vertebrate and specific tissue development [11]. Ciliopathies can be organ-specific, as in the case of polycystic kidney disease, nephronophthisis, or LCA [12,13], or can be syndromic, as in the case of Jeune syndrome, Bardet-Biedl syndrome, and MSS. As research indicates that the cumulative prevalence of ciliopathies is approximately 1 in every 2000 individuals, accurate and timely diagnosis is essential to avoid significant morbidity and mortality [14].

Currently, IFT140 mutations have been identified in several syndromic ciliopathies, including MSS. Khan et al. studied 11 subjects with confirmed homozygous IFT140-related retinopathy and found that although some patients had frank skeletal or renal disease, not all patients showed obvious extraocular findings [15]. This is consistent with the presentation in our patient, where ophthalmic phenotypes including infantile nystagmus, hyperopia, and retinal dystrophy were apparent with no systemic manifestations observed to date. LCA and retinitis pigmentosa are commonly described in a series of MSS cases. In these reports, early onset severe visual impairment may appear earlier than systemic symptoms [7,16,17]. This finding can partly be explained by the distinct role of IFT140 in maintaining cilia function in photoreceptors [18]. In light of these findings, it is important to note that variable expressivity for ciliopathies is common, and that careful systemic work-up and monitor need to be emphasized.

In this study, the frameshift mutation c.2214_2217del (p. Asp738GlufsTer47) detected in our patient is a novel variant located in the IFT140 gene cause Opitz trigonocephaly C syndrome in a patient with typical clinical picture, and prenatal diagnosis, Hum. Genet. 68 (1984) 104–115.

In summary, we identified compound heterozygous IFT140 variants in two polish families with Sensenbrenner syndrome and early onset end-stage renal disease, Orphanet. J. Rare Dis. 15 (2020) 36.

In this study, we reported a novel frameshift mutation in the IFT140 gene in a patient with MSS. Given the pathogenic variation in IFT140, MSS must be considered as a differential diagnosis in patients with isolated retinal dystrophy. The present study strengthens the rationale that early monitoring of systemic diseases including renal function is crucial in patients with MSS.

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Declaration of Competing Interest

There are no conflicts of interest to disclose.

Data availability

The data that has been used is confidential.

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