A Novel Homozygous Frameshift Mutation in CCN6 Causing Progressive Pseudorheumatoid Dysplasia (PPRD) in a Consanguineous Yemeni Family

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Background: Progressive pseudorheumatoid dysplasia (PPRD) inherited in an autosomal recessive fashion, is a disabling disease, characterized by platyspondyly, irregularities of the vertebral bodies, narrowing of the intervertebral discs and intraarticular spaces, widening of the epiphysis-metaphysis, multiple joint contractures, and disproportionate short stature. A number of studies have been performed on this deformity in various populations around the globe, including the Arab population. Mutations in CCN6, located on 6q22, are reported to cause this anomaly.

Case Presentation: The present study describes the investigation of a consanguineous family of Yemeni origin. Clinical examination of the patient revealed short stature with progressive skeletal abnormalities, stiffness and enlargement of small joints of the hands along with restriction of movements of proximal interphalangeal (PIP) and distal interphalangeal (DIP) joints with weakness and gait disturbance. Sanger sequencing revealed a novel homozygous frameshift deletion mutation (c.746delT; p.Val249Glyfs*10) in CCN6 which may lead to NMD (Nonsense mediated decay). This mutation expands the spectrum of pathogenic variants in CCN6 causing PPRD.

Keywords: progressive pseudorheumatoid dysplasia (PPRD), consanguinity, novel frameshift mutation, CCN6, Nonsense Mediated Decay
INTRODUCTION

Spondylo-epi-metaphyseal dysplasias (SEMDs) are a heterogeneous group of disorders carrying different inheritance patterns (1). SEMDs are described as a set of epiphyseal, metaphyseal, and vertebral abnormalities. SEMDs are diagnosed based on typical skeletal presentations and/or associated extra clinical features of skeletal defects (2, 3). PPRD (OMIM 208230); Spondyloepiphysial dysplasia tarda with progressive arthropathy (SEDT-PA), is described as a progressive stiffness of joints, remarkably reduced mobility of the cervical spine, and swollen fingers (4). Early development among patients with PPRD is usually normal but during the initial years of childhood the formation of the joint contractures occurs in the hands which spreads to the hips, knees, and the spine (5).

As PPRD is a progressive disorder, all the joints (small and large, including spine) will have progressively limited movement. Stature is usually normal at the onset of the disease but have been reported below percentile 3 at the time of diagnosis in most of the cases (6). Garcia Segarra et al. (6) have also reported that there are variations in early symptoms of the patients but about 50% of the patients had gait anomalies and fatigability. In 30% of the cases the first sign was IP (interphalangeal joints) swelling and some degree of knee deformity in about 20% of the cases while pain was reported as the initial symptom in 15% of the patients (6). Radiological surveys indicate compressed vertebral bodies, defects in ossifications, and malformed acetabular portion of pelvis (7). PPRD has an autosomal recessive mode of inheritance which seems to be more frequent in the Middle East and Gulf states because of the high consanguinity rate and large family size, with an incidence of ~1:1,000,000 in Arab countries, (7, 8), however a few cases from non-Mediterranean origin (Germany, United Kingdom, Belgium, Italy, Poland, France UK, USA, Morocco, Ecuador, Japan, India, Pakistan, and China) have been reported as well (5, 6, 9).

PPRD phenotype is often confused with skeletal deformities of ankylosing spondylitis, rheumatoid arthritis (RA), and mucopolysaccharidosis (1, 6) but the characteristic features of RA, such as the destruction of bones, is missing in PPRD (7). Though PPRD is clinically indistinguishable from juvenile idiopathic arthritis (JIA) (7), the results regarding RA factor and inflammation during diagnostic testing are always negative and PPRD patients also show a reduced response to antirheumatic drugs (5, 7).

PPRD is caused by mutations in cellular communication network factor 6 (CCN6), previously known as Wnt1 inducible signaling pathway protein 3 (WISP3), which maps to chromosome 6q22 (10), and encodes a connective tissue growth factor involved in cell growth and differentiation (11). The human cartilage homeostasis and bone growth are highly dependent on the expression of CCN6 (12, 13).

Two homologs CCN4 and CCN5 (also known as WISP1, WISP2) of CCN6 have been described in the literature, which are expressed in WNT1 transformed cell (14, 15). The role of WNT-signaling proteins in cell fate determination and the regulation of cell morphology and proliferation is extraordinary (15–18). CCN6, along with its orthologs (CCN4, CCN5), shows a differential expression in primary human colon cancer (14, 15). Hurvitz et al. pointed out that CNN6 might have a consistent role in human skeletal homeostasis and the regulation of skeletal development in adults (15).

In the present study, we clinically diagnosed PPRD in a Yemeni patient, born from a consanguineous marriage. Sanger sequencing of coding regions of CCN6 has revealed a novel deletion mutation (c.746delT; p.Val249Glyfs*10) in this patient.

CASE PRESENTATION

Methods

The study was conceived according to the principals of the Declaration of Helsinki. Both adult and minor participants were completely informed about the clinical procedures and molecular diagnosis. Informed written consents of parents were obtained on the institutional patient recruitment and consent proformas and research protocols were approved under the reference number 24/14 by the medical ethics and research committee of King Abdulaziz University, Jeddah, Saudi Arabia. Affected member (IV-4) underwent thorough clinical, radiological and other relevant laboratory investigations. Venous blood samples were collected from affected (IV-4) and available unaffected individuals (III-1, III-2, IV-3) of the family (Figure 1). Genomic DNA was extracted following a standard protocol.

Sanger Sequencing

Chain termination DNA sequencing technique was used for the identification of pathogenic variant in CCN6 and co-segregation

![Image](https://via.placeholder.com/150)

FIGURE 1 | Pedigree of a Yemeni family segregating an autosomal recessive PPRD. Arrow indicates the index patient (IV-4). The samples which were available for the genetic analysis are marked with asterisks (*). wt/delT represents the heterozygous genotype of the carriers, wt/wt represents the wild type genotype of an unaffected member while delT/delT represents the homozygous deletion in the affected member of the family.
The Family was subject to DNA sequencing of CCN6 to know the genetic cause of this anomaly and for appropriate genetic counseling.

**Clinical Findings**

The 13-year-old female (IV-4) is the 5th offspring of a consanguineous Yemeni couple (III-1, III-2) (Figure 1). She was born to a healthy 33-year-old mother (III-2) and a 37-year-old father (III-1) as a result of an uncomplicated pregnancy with a limited range of active and passive movements mainly at intraphalangeal, elbow, knee, hip, ankle, shoulder, and wrist, associated with progressive difficulty in walking and abnormal gait was observed. She exhibited no inflammatory signs in her joints.

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**RESULTS**

A 13-year-old patient (IV-4) was referred to the genetic clinic, King Abdulaziz University Hospital, Jeddah, for the genetic consultation. After review of the family history, clinical symptoms, and laboratory investigations of the patient, there were several clues which guided us to the diagnosis of PPRD (Figure 2a). The progressive restriction of movements of the elbow, knee, hip, ankle, shoulder, and wrist, associated with progressive difficulty in walking and abnormal gait was observed. She exhibited no inflammatory signs in her joints.

**Investigations**

Cytogenetic analysis of the patient (IV-4) revealed a 46, XX karyotype. Rheumatoid factor, antinuclear antibody, and anti-dsDNA tests were negative. Erythrocyte sedimentation rate, C-reactive protein, complete blood count, coagulation profile, urea and electrolytes, thyroid function tests, parathyroid hormone, and liver enzymes were all within normal range. Bone Marrow Density was −2SD for her age. N-acetylgalactosamine-6-sulfate and arylsulfatase-B were within normal range. Bone Marrow Density was −2SD for her age. N-acetylgalactosamine-6-sulfate and arylsulfatase-B were within normal range. Bone Marrow Density was −2SD for her age. N-acetylgalactosamine-6-sulfate and arylsulfatase-B were within normal range. Bone Marrow Density was −2SD for her age. N-acetylgalactosamine-6-sulfate and arylsulfatase-B were within normal range. Bone Marrow Density was −2SD for her age. N-acetylgalactosamine-6-sulfate and arylsulfatase-B were within normal range. Bone Marrow Density was −2SD for her age. N-acetylgalactosamine-6-sulfate and arylsulfatase-B were within normal range. Bone Marrow Density was −2SD for her age. N-acetylgalactosamine-6-sulfate and arylsulfatase-B were within normal range. Bone Marrow Density was −2SD for her age. N-acetylgalactosamine-6-sulfate and arylsulfatase-B were within normal range.

**Radiological Examination**

Radiological examination revealed a diffuse decreased density of the visualized bones, irregular articular surface of the PIP joints of hands and feet with relative decrease in joint space and enlargement of the epi-metaphysis of the interphalangeal joints (Figure 2e). This was associated with prominent distal epiphysis of the radius and ulna (Figures 2c,f). Regarding elbow joints, there were prominent capitulum and trochlea of the distal humerus and irregularity of the articular surface (Figure 2f). Hips showed bilateral fine irregularities of articular surfaces (Figure 2g). An X-ray of the spine showed anterior beaking of the lumbar vertebral bodies, mild irregularities of the anterior aspect of the superior end plates along with a mild form of platyspondyly involving most of the examined vertebrae (Figure 2b).

**Mutation Identification**

All five exons of CCN6 were amplified in the affected individual (IV-4) DNA. Sanger sequencing analysis revealed a novel homozygous deletion mutation (c.746delT;
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CCN6 Deletion in a Yemeni Family

FIGURE 2 | Clinical pictures of the patient IV-4 (a) showing camptodactyly of all fingers, with prominent PIP and DIP joints (b) represents the inability of the patient to make a fist (c) shows the prominent knee joints (d) shows short trunk. Radiological findings of the patient IV-4. (e) an X-rays of the right hand shows the irregular articular surface of the PIP joints with relative decrease in joint space and enlargement of the epi-metaphysis of the interphalangeal joints (f) shows elbow joints with prominent capitulum and trochlea of the distal humerus and irregularity of the articular surface (g) shows hips with bilateral fine irregularities of articular surfaces (h) shows the anterior beaking of the vertebral bodies with mild form of platyspondyly.

p.Val249Glyfs*10) in exon-4 of CCN6 (Figure 3). Considering that the identified variant is pathogenic, Sanger sequencing of both parents (III-1, III-2) and an unaffected sibling (IV-3) was carried out using the same set of primers for exon-4 (Table 1), which confirmed the co-segregation of this variant within the family. The allele frequency of this variant was not found in the Genome Aggregation Database (gnomAD) (http://gnomad.broadinstitute.org/gene/ENSG00000112761). An online prediction algorithm, MutationTaster (http://www.mutationtaster.org) has predicted this variant as a frameshift mutation likely to suffer Nonsense Mediated Decay (NMD). This variant has been submitted to ClinVar database (https://submit.ncbi.nlm.nih.gov/clinvar/) under an accession ID SCV000902272.

DISCUSSION

Here, we report a consanguineous Yemeni family with one affected member (IV-4) showing typical features of PPRD. The clinical laboratory findings coincided with the apparent clinical phenotypes. The thorough examination of parents (III-1, III-2) and an unaffected brother (IV-3) revealed the absence of PPRD symptoms. Sanger sequencing revealed a homozygous deletion mutation (c.746delT; p.Val249Glyfs*10) in CCN6 which co-segregates within the family. Previously, CCN6 gene mutations have been reported in PPRD patients from various geographical regions of the world.

CCN6 encodes a protein of 372 amino acids which is composed of a signal peptide domain encoded by exon-1 and four conserved cysteine-rich domains; IGFBP N-terminal (Insulin-like growth factor binding protein N-terminal) domain, vWFC (von Willebrand factor type-C module) domain, TSP type-1 (thrombospondin) domain, CTCK (C-terminal cystine knot) domain, encoded by exon-2, 3, 4, and 5 of CCN6, respectively.
well-elaborated (5). Missense, nonsense, deletions, insertions and splice site mutations have been reported in this gene causing PPRD phenotype (HGMD, Professional, 2018.3; http://www.hgmd.cf.ac.uk/ac/gene.php?gene=CCN6).

Connective tissue growth factor/cysteine-rich protein 61/nephroblastoma overexpressed (CNN) family member CCN6 is a WNT-signaling pathway protein which plays a vital role in the expression regulation of a transcriptional factor SOX9, collagen II, and aggrecan in chondrocyte cell lines (13, 19). CCN6 plays another role in controlling the activity of superoxide dismutase. The number of reactive oxygen species is increased in the absence of CCN6 expression (6, 20). A study on a single PPRD affected individual conducted by Zhou et al. (21) has provided information about the increased cell proliferation and abnormal matrix metalloproteinase processing in human chondrocytes (21). Certain in vivo experiments in zebrafish have established the role of CCN6 in the development of cartilages. It has been proposed that the malfunction of signaling pathways governed by bone morphogenetic protein (BMP) and/or Wnt-signaling protein, in the case of a nonfunctional mutant CCN6, may lead to human cartilage development failure in PPRD patients (22). However, the causative pathomechanism of malformation of cartilage in human PPRD patients is still not well-elaborated (6).

In the present study, we have identified a novel homozygous frameshift variant (c.746delT; p.Val249Glyfs*10) in CCN6 causing PPRD in a Yemeni patient. In a review article, Hentze and Kulozik, have explained that the imperfect messages which are the products of frameshift mutations are eliminated by conserved mammalian surveillance mechanisms like NMD (23). MutationTaster (http://www.mutationtaster.org/) has also predicted that this novel variant may lead to frameshift and eventually to NMD, which is anticipated to cause PPRD in our patient.

CONCLUSION

The present work aims to report a clinical study of a Yemeni patient who is a product of a consanguineous union and the identification of pathogenic mutation causing PPRD phenotype. Our data is novel and increases the mutational spectrum of CCN6 gene.

DATA AVAILABILITY

No datasets were generated or analyzed for this study.

ETHICS STATEMENT

Informed written consents were obtained from parents about the clinical and molecular diagnosis and for publishing the clinical photographs and mutation data. The study was approved under the reference number 24/14 by the medical ethics and research committee of King Abdulaziz University, Jeddah, Saudi Arabia.

AUTHOR CONTRIBUTIONS

NG, MA, and JA-A enrolled the patients, made the clinical diagnosis, and wrote the clinical synopsis. AE did radiological analysis and wrote a report about that. NW and AP designed the study and wrote the initial draft of this manuscript. NW contributed to Sanger sequencing. NW, AP, and NG contributed to critical revisions of the manuscript. All authors have thoroughly reviewed and approved the final draft.

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**Conflict of Interest Statement:** 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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