A genetic assay of three patients in the same family with Holt-Oram syndrome; a case report

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Abstract

Holt-Oram syndrome (HOS) is a developmental disorder inherited in an autosomal-dominant pattern. Affected organs are the heart and forelimbs with upper extremity skeletal defects and congenital heart malformation. In this study we present three cases of HOS in the same family. In one of these three individuals we detected a transition of C to T (CTG - GTT, V205V) in exon 7 of the TBX5 gene. This nucleotide change causes no amino acid change and potential pathologic effects remain unknown.

Keywords: Holt-Oram syndrome, Congenital heart malformation, TBX5 gene

Introduction

Holt-Oram syndrome (HOS) is a developmental disorder inherited in an autosomal-dominant pattern. Affected organs are the heart and forelimbs with upper extremity skeletal defects and congenital heart malformation. This syndrome was first described by Holt and Oram (1), who observed atrial septal defect in members of four generations of a family associated with a congenital anomaly of the thumbs.

Haploinsufficiency of TBX5 on the human chromosome 12 (12q24.1) is the cause of HOS (2, 3). Holt-Oram syndrome is the most common heart-hand syndrome with an estimated frequency of about one in 100,000 live births (4). This syndrome has been reported from various racial and ethnic groups worldwide (5).

The upper-limb malformations in HOS are fully penetrant, but congenital heart malformations occur in approximately 75% of affected individuals (6).

Materials and Methods

In this study we present three cases of HOS in the same family:

Patient 1 was 23-year-old female who presented with palpitations. The electrocardiogram revealed sinus pauses with junctional escape rhythm. The echocardiogram revealed a 45 mm atrial septal defect of the ostium secundum type with enlarged right chambers, moderate pulmonary arterial hypertension (systolic pressure: 53 mmHg), mild tricuspid regurgitation, and inferior vena cava (IVC) dilation of 24 mm. These findings were confirmed during surgery.

Patient 2 was a 25-year-old male who presented with palpitations and syncope with onset at age nine. The echocardiogram revealed a large atrial septal defect of the ostium secundum type and multiple ventricular septal defects. Surgery was first performed at age nine.

Patient 3 was a 27-year-old male who presented with palpitations, dyspnea, and syncope. The echocardiogram revealed two atrial septal defects of the ostium secundum type, enlarged right chambers, and mild tricuspid and mitral regurgitation.

Peripheral blood samples of these patients were collected in EDTA anticoagulant tubes and DNA
was extracted using the salting out method, with modifications described previously (Fig. 1) (7).

Genomic DNA of exons 2-9 of the TBX5 gene, with NC_000012.11 as the NCBI reference sequence, was sequenced using the primers listed in Table 1.

Each exon was amplified under the following conditions: one cycle at 94 °C for 5 min, followed by 30 cycles of denaturation at 94 °C for 30 sec, annealing at 60 °C for 30 sec, and extension at 72 °C for 30 sec, and one cycle of post-elongation at 72 °C for 7 min. Each 25 µL reaction contained 20 mmol/L Tris-HCl pH 8.4, 50 mmol/L KCl, 2 mmol/L MgCl2, 10 pmol of each primer, 0.2 mmol/L of each dNTP mix, 1.25 units of Smart Taq DNA polymerase, and 100-200 ng of genomic DNA.

Table 1. Primers used to amplify exons 2-9 of the TBX5 gene.

<table>
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<tr>
<th>Exon</th>
<th>Primer</th>
<th>Size (bp)</th>
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| 2    | Forward 5'-GCTTCTTTGCTCAGAGACAGAACCT-3’  
     | Reverse 5'-GGAAGAGAACGGGAGGAAAAGCCA-3’ | 277       |
| 3    | Forward 5'-AGTTTGGGGAAGGAATGCCCACTAC-3’  
     | Reverse 5'-TCTCTTGGTCCCCTCTACA-3’ | 200       |
| 4    | Forward 5'-AACGGGGCTAGTTTCCGCTTCCACG-3’  
     | Reverse 5'-CTTTTTGGGAGAAGGTTCCACTTTT-3’ | 307       |
| 5    | Forward 5'-CCCTAAAATGGATGGAGGC-3’  
     | Reverse 5'-CTTTTTGGGAGAAGGTTCCACTTTT-3’ | 261       |
| 6    | Forward 5'-CTCTGTCGCTGAACTGAAGCACGCT-3’  
     | Reverse 5'-CTGGAAGCTGCTGCTGCTGCT-3’ | 340       |
| 7    | Forward 5'-GAGGGAGACAAGGCGGGGAATCCAG-3’  
     | Reverse 5'-GAGGGAGACGGCGAGGCTCC-3’ | 250       |
| 8    | Forward 5'-CATTTTGCTGCGTTCCTCACACC-3’  
     | Reverse 5'-GGGATAGGCACTGCAAGAAAGGACT-3’ | 515       |
| 9    | Forward 5'-GAGGGAGACAAGGCGGGGAATCCAG-3’  
     | Reverse 5'-GAGGGAGACGGCGAGGCTCC-3’ | 802       |

Results
Sequence analysis of the TBX5 gene in two of our three patients identified no nucleotide changes, but in one patient we detected a C to T transition (CTG-GTT, V205V) in exon 7 of the TBX5 gene (Fig. 2). This nucleotide change is at codon 205 of TBX5. This transition causes no amino acid change and potential pathologic effects remain unknown.

Discussion
TBX5 is a member of the T-Box gene family required in vertebrate cardiogenesis and normal heart development (8, 9). Holt-Oram syndrome is a developmental disorder with full penetrance caused by haploinsufficiency in the TBX5 gene (3). This syndrome shows variable expression as a spectrum of mutations and thereby genotype-phenotype correlation. In this syndrome cardiac involvement may be absent in patients with upper limb defects. Immunohistochemical studies with anti-human TBX5 protein antibodies demonstrated cardiac expression during embryogenesis throughout the epicardium and in cardiomyocyte nuclei in the myocardia of all four cardiac chambers (10, 11).
Clinical manifestations generally present as malformations of the upper limbs, cardiac septation defects such as atrial and ventricular septal defects (ASD and VSD, respectively), and disruption of normal cardiac conduction system development (12). Wide varieties of skeletal defects and congenital heart disease were observed; however, the severity of skeletal involvement did not parallel that of cardiac disease. Smith et al. (1979) reported five of 39 affected patients had normal EKGs despite typical limb defects (13). Basson et al. (1997) reported a nonsense mutation (Glut69-stop codon) in affected members of one family that caused a truncated protein without a T-box domain (2), and in another study Yang et al. (2000) identified three novel mutations that included a frame shift in exons 3–9 with breakpoints within introns 1–2 and 9–10 (17).

Presently, modern high-throughput DNA sequencing technologies such as pyro sequencing and next generation sequencing (NGS) can be used to search for nucleotide changes in entire candidate genes in patients with genetic disorders. These methods are improvements over exon-by-exon PCR and subsequent DNA sequencing.

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References