



Original Article

Prevalence of Trinucleotide Expansions in SCA17/TBP and JPH3 Genes and Octapeptide Insertion in PRNP Gene in Iranian Patients with Huntington-Disease like Syndrome

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ABSTRACT

Background and objective: Huntington's disease (HD) is an autosomal dominant disorder that mainly affects adults. Although mutations in the *IT15* gene have been known as the main cause of the disease, patients with HD like (HDL) syndrome have mutations in genes other than the *IT15* gene. In this study, we investigate the frequency of mutations in *SCA17/TBP*, *JPH3* and *PRNP* genes in patients with HDL syndrome.

Methods: The frequency of mutations in *SCA17/TBP*, *JPH3* and *PRNP* genes was studied in 56 patients with HDL phenotype but without trinucleotide expansion in the *IT15* gene. DNA was extracted from peripheral whole blood by the salting out method. PCR was performed using specific primers for each gene. PCR products were separated on polyacrylamide gel. Sequencing was performed on some samples to confirm the PCR results.

Results: We found neither trinucleotide expansion in the *JPH3* and *SCA17*, nor octapeptide insertion in the *PRNP* gene.

Conclusion: Based on the results, Iranian patients with HDL syndrome do not have mutation in the *TBP*, *JPH3* and *PRNP* genes. However, this result may be due to population differences, rarity of the mutations in the studied genes and the small number of study subjects. Therefore, studies with a larger study population that investigate other mutations, such as point mutations in the mentioned genes may help clarify the exact cause of HDL phenotype in Iranian patients.

Keywords: Huntington's Disease, HDL, JPH3, PRNP, SCA17

INTRODUCTION

Huntington's disease (HD) is a progressive neurodegenerative disease which mainly occurs in adults (1). The most important clinical symptoms of the disease include movement disorders, chorea, psychological disorders, epilepsy, mood changes and depression. The average age of onset is between 30 to 50 years; however, 5 to 10 percent of the patients develop the disease before the age of 20, a condition known as the juvenile HD (2).

Huntington's disease is caused by the CAG repeat expansion (poly-glutamine) in the coding region of *IT15* (HTT) gene located at 4p16.3. Patients with HD-like (HDL) syndromes have symptoms similar to HD but do not have the trinucleotide expansion in the *IT15* gene. There are three HDL syndromes (HDL1, HDL2 and HDL4) with autosomal dominant inheritance and one (HDL3) with recessive inheritance (3).

HDL1 disease or transmissible spongiform encephalopathy (prion disease) is a rare, progressive neurodegenerative disorder that affects one in every 1,000,000 people annually. The age of onset of HDL1 is 25-45 years. The symptoms include cognitive difficulties, ataxia and myoclonic seizures, personality changes, psychiatric symptoms, fatigue, emotional numbness, dysarthria, dementia and motor disturbance with chorea (4, 5). The disease is caused by the insertion of octapeptide repeats in the *PRNP* gene, which is located on the short arm of chromosome 20. The octapeptide repeat region lies *between codon 51 and 91*, which includes a nonapeptide (Pro-Gln- (Glu) 4-Trp-Gly-Gln) followed by an octapeptide (Pro-His- (Gly) 3-Trp -Gly-Gln). With each repeat, 24 nucleotides (8 amino acids) will be added to the gene. Normally, there are four octapeptide repeats in the *PRNP* gene. In HDL1 patients, the repeat numbers is increased to 11-13, which produces an abnormally long cellular prion protein. It is not clear that how abnormal proteins destroy neurons and cause characteristic features of HDL1 (6).

Huntington's disease-like 2 (HDL2) is caused by CAG/CTG expansion in the *JPH3* gene located at 16q23. Normal and pathogenic alleles have 6-28 and 44-57 repeats, respectively. Alleles with 29-40 repeats are unstable (7).

The disease usually appears at the age of 29 to 41 and has a 10 to 20 years survival time. Clinical symptoms of the disease include hypokinesia (rigidity, bradykinesia), tremor, dysarthria, psychological symptoms, emotional and cognitive abnormalities, convulsion and chorea. HDL2 is a rare and most commonly observed in Africa (8).

Huntington's disease-like 4 (HDL4), also known as spinal cerebellar ataxia type 17 (SCA17), is caused by a mutation in the TATA-binding protein (*TBP*) gene, which is located at 6q27 and contains eight exons. The CAG/CAA repeats are located in exon 3 of this gene. Normal alleles have 25-42 CAG/CAA repeats. The pathogenicity of alleles with 43 and 44 repeats is uncertain. Furthermore, alleles with up to 49 repeats have incomplete penetrance (9,10). Main symptoms of HDL4 include uncoordinated movement, dementia, involuntary movement, psychosis, seizure and rigidity. The age of onset (3–75 years) and clinical presentation are extremely variable (11).

HDL3 is mapped to 4p15.3 and has recessive inheritance. The responsible gene for this disease is not known (12).

In HDL syndromes, the repeat number is significantly correlated with early onset and disease severity (13). The aim of this study was to evaluate frequency of the mentioned mutations in the *TBP*, *JPH3* and *PRNP* genes of patients with HDL syndromes.

MATERIALS AND METHODS

In this study, we investigated the expansion of trinucleotide repeats in the *JPH3* and *SCA17/TBP* genes and the insertion of octapeptide repeats in the *PRNP* gene. We recruited 56 unrelated HDL patients (34 men and 22 women, mean age: 52 ± 5 years) with normal trinucleotide repeat numbers in the *IT15* gene, who were referred to the Tehran Medical Genetics Laboratory between 2006 and 2010. Written consent was obtained from patients or their guardians.

DNA was extracted from peripheral whole blood using the salting-out method. The quantity and quality of the extracted DNA was assessed by a spectrophotometer (Nanodrop 2000c, ThermoScientific) (14). Three primer pairs were designed to amplify the region of interest of each gene (Table 1).

Table 1. Sequence of the primers used for amplification of the *JPH3*, *PRNP* and *TBP* genes

Primer	Sequences (5'→3')	Product length
JPH3-F	AATCGATCTGTGCCTTCATTC	139 bp
JPH3-R	GTTCCCTGCACAGAAACCATC	
PRNP-F	AAGCCTGGAGGATGGAACAC	233 bp
PRNP-R	TTACTCGGCTTGTTCCACTGAC	
SCA17-F	CCTTATGGCACTGGACTGAC	245 bp
SCA17-R	GTTCCCTGTGTTGCCTGCTG	

Amplification was done using the Veriti 96 well Thermal Cycler (Applied Biosystems) with the following conditions: initial denaturation at 95 °C for 5 min, followed by 32 cycles of 94 °C for 40 sec, 65 °C for 40 sec, 72 °C for 40 sec and final extension at 72 °C for 10 min. PCR products were separated on 12% polyacrylamide gel stained with silver nitrate. Also, for determining the exact number of repeats, several samples from the studied genes were sequenced by MacroGen Co. (South Korea).

RESULTS

In this study, all patients had normal repeat numbers (Figures 1-3). The average repeat numbers for the *JPH3* and *TBP* genes was 13-15 and 33-35, respectively. In addition, the average number of octapeptide repeats in the *PRNP* gene was 3-4.

Figure 4 shows an example of the *PRNP* gene sequencing result in one of the patients using the forward primer.

Figure 1. PCR products of the *PRNP* gene amplification on 12% polyacrylamide gel. Columns 1 to 6: HDL patients, column 7: normal control sample, M: 50 bp size marker

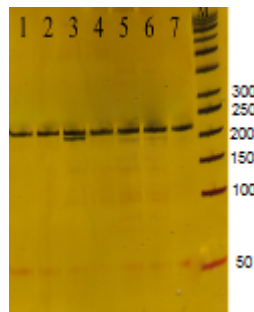


Figure 2. PCR products of the *JPH3* gene amplification on 12% polyacrylamide gel. Columns 1 to 6: HDL patients, column 7: normal control sample, M: 50 bp size marker.

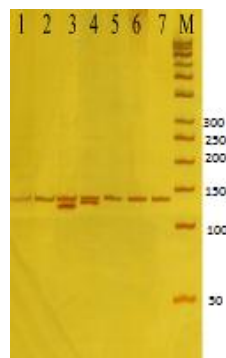


Figure 3. PCR products of the *TBP* gene amplification on 12% polyacrylamide gel. Columns 1 to 6: HDL patients, column 7: normal control sample, M: 50 bp size marker.

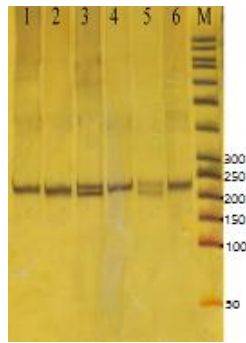
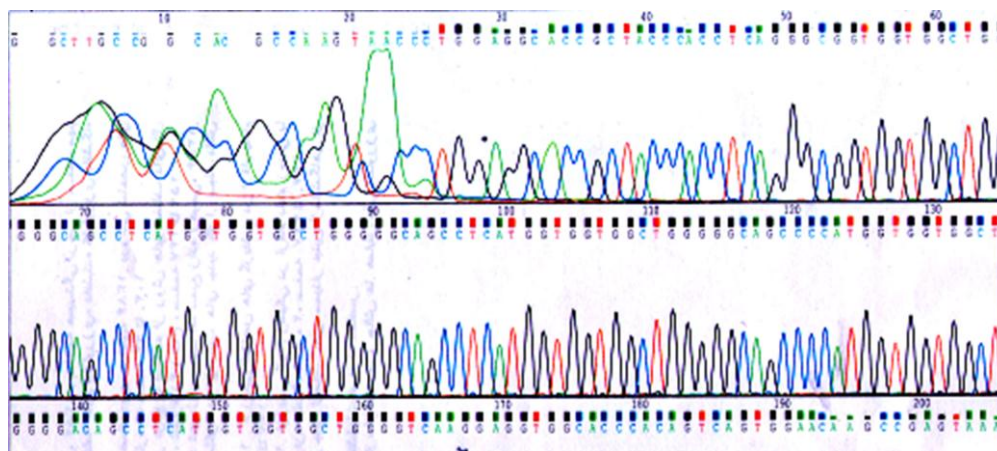


Figure 4. Sequencing result of octapeptide region in the *PRNP* gene using the forward primer.



DISCUSSION

Huntington's disease and HDL syndromes are autosomal dominant disorders (except for HDL3) characterized by progressive neurodegeneration, cognitive impairment and behavioral abnormalities that occur in adulthood. In 1993, abnormality in the *IT15* (HTT) gene was identified as the causative agent for HD (15, 16). However, it was found that a number of patients with clinical symptoms of HD do not have a three nucleotide expansion in their *IT15* gene. Later, other genes such as *PRNP*, *JPH3*, *SCA17* and *DRPLA* were found to be responsible for HDL symptoms.

In this study, 56 HDL patients with no mutation in the *IT15* gene were studied for *JPH3*, *PRNP* and *TBP* genes abnormality. No trinucleotide expansion or octapeptide insertion was found in the subjects. In a study on 106 Brazilian patients, 37 and 29 patients were affected by HD and HDL syndromes,

respectively. Moreover, 10.3% of the HDL patients had mutations in the *JPH3* gene and were therefore diagnosed with HDL2 (17). In a study in America on 538 patients with HDL phenotypes, six patients had mutations in the *JPH3* gene. In Japan, no patient had *JPH3* mutation (18). In Serbia and Montenegro, 198 HDL patients were studied for CTG repetition expansion in the *JPH3* gene using PCR and sequencing. It was revealed that alleles had 11-18 CTG repeats, so HDL2 is very rare in these populations (19). Some studies have proposed other genes as candidate for HDL diagnosis. For instance, *CREBBP* and *POU3F2* genes were studied in Portuguese patients and there was no mutation in the genes. In Japan, mutation in the *DRPLA* gene was common among HDL patients (20). Table 2 summarizes a number of studies that have been performed on these genes in different populations.

Table 2. Frequency of HDL syndromes in different populations

Year of study	Study population	Number of subjects	Studied genes	Frequency of mutations
1999	America and Britain (13)	15	<i>SCAI</i> , 2 and 3 and <i>DRPLA</i>	0
2001	African Americans (13)	330	<i>HDL2</i>	1.2%
2002	Germany and Austria (14)	1600	<i>SAC17</i> <i>HDL2</i>	9(0.005%) 0
2003	France (8)	252	<i>HDL1</i> , <i>HDL2</i> , <i>SCAI7</i> and <i>DRPLA</i>	<i>HDL2</i> =2(33%) <i>SCAI7</i> =2(33%)
2004	North America (13)	538	<i>HDL2</i>	6
2005	Yugoslavia (16)	48	<i>DRPLA</i> , 23, 17, <i>SCAI</i> , <i>HDLA1</i> , <i>HDL2</i> and <i>NFP</i>	0
2007	South Africa (17)	50	<i>HDL2</i>	15(30%)
2006	Portugal (18)	107	<i>SCAI7</i> , <i>DRPLA</i> , <i>NFP</i> , <i>HDL1</i> and <i>HDL2</i>	0
2008	England (5)	258	<i>DRPLA</i> , 17, 23, <i>SCAI</i> , <i>HDL1</i> , <i>HDL2</i> , <i>FRDA</i> and <i>NFP</i>	<i>SCAI7</i> =5(1.8%) <i>HDL1</i> =1(0.4%) <i>FRDA</i> =1(0.4%) <i>HDL2</i> =1(0.4%)
2008	Poland (19)	224	<i>DRPLA</i> , <i>SCAI7</i> and <i>HDL2</i>	<i>SCAI7</i> =1(0.44%)
2011	Brazil (12)	29	<i>ChAC</i> , <i>DRPLA</i> , 17, 23, <i>SCAI</i> and <i>HDL2</i>	<i>HDL2</i> =3(10.3%) <i>ChAC</i> =2(6.8%)
2014	Iran (the present study)	56	<i>HDL1</i> , <i>HDL2</i> and <i>HDL4</i>	<i>HDL1</i> =0 <i>HDL2</i> =0 <i>HDL4</i> =0

CONCLUSION

.....We found no mutation in the *TBP*, *JPH3* and *PRNP* genes of patients with HDL syndromes. However, this result may be due to population differences, rarity of the mutations in the studied genes and the small number of study subjects. Therefore, studies with a larger study population that investigate other mutations, such as point mutations in the mentioned genes may help clarify the exact cause of HDL phenotype in Iranian patients.

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CONFLICT OF INTEREST

.....There is no conflict of interest regarding publication of this article.

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