A Novel Splice Site Variant in the LDLRAP1 Gene Causes Familial Hypercholesterolemia

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ABSTRACT

Background: FH, a hereditary disorder, is caused by pathogenic variants in the LDLR, APOB, and PCSK9 genes. This study has assessed genetic variants in a family, clinically diagnosed with FH. Methods: A family was recruited from MASHAD study in Iran with possible FH based on the Simon Broom criteria. The DNA sample of an affected individual (proband) was analyzed using WES, followed by bioinformatics and segregation analyses. Results: A novel splice site variant (c.345-2A>G) was detected in the LDLRAP1 gene, which was segregated in all affected family members. Moreover, HMGCR rs3846662 g.23092A>G was found to be homozygous (G/G) in the proband, probably leading to reduced response to simvastatin and pravastatin. Conclusion: LDLRAP1 c.345-2A>G could alter the PTB, which acts as an important part of biological pathways related to lipid metabolism.

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List of Abbreviations:

FH, familial hypercholesterolemia; ES, exome sequencing; HSF, human splice finder; LDL-C, Low-density lipoprotein cholesterol; LDLRAP1, LDL receptor adaptor protein 1; MASHAD, Mashhad Stroke and Heart Atherosclerotic Disorders; PTB, phosphotyrosine-binding domain; WES, whole exome sequencing; WT, wild type
INTRODUCTION

Familial hypercholesterolemia is an inherited condition due to the pathogenic variants of LDLR, APOB, and PCSK9 genes\(^1\). The most significant complication for FH patient is the high risk of premature coronary heart disease\(^2\). Based on estimates, the prevalence rate of FH is 1:200 in Western populations; however, over 80% of FH patients remain undiagnosed\(^3\).

The World Health Organization has recommended large-scale screening for the identification of FH patients, who will benefit mostly from early treatment with lipid-lowering drugs. Early diagnosis and treatment of these patients can reduce the risk of cardiovascular disease, aiming at lowering the LDL-C concentration, and contribute to the proper management of other risk factors during the life\(^4\). Therefore, it is crucial to understand the molecular basis of FH to diagnose the disease and manage therapeutic approaches. About 5% of the cases before the age of 60, who have experienced myocardial infarction, is estimated to be heterozygous FH\(^5\).

Recent molecular techniques such as WES aim at finding novel genetic variants, which is mainly important in multiethnic populations\(^9\). The \textit{LDLRAP1} gene or ARH is an adapter protein that facilitates the endocytosis of LDLR into hepatocytes. Mutations in this gene have been demonstrated to induce a recessive type of FH\(^6\). It has also been reported that some heterozygous carriers of pathogenic variants in the \textit{LDLRAP1} gene show high LDL-C levels\(^8\). We describe a novel homozygous splice site variant c.345-2A>G in the \textit{LDLRAP1} gene, which was identified in an FH family.

MATERIALS AND METHODS

Subjects and clinical presentation

This survey is a family-based pedigree study, as a part of a larger research on the genetic assessment of dyslipidemia, MASHAD cohort study, using WES \(^9\). Peripheral blood samples were collected, and the sera were separated. Total serum levels of cholesterol, HDL-C, LDL-C, and triglyceride were measured according to a method described previously\(^9\).

DNA extraction and WES

DNA samples were extracted using the standard salting-out method. The quality and quantity of DNA sample were assessed by a Nanodrop (Thermo Scientific, USA) and the genomic DNA extracts were analyzed on the 0.7% agarose gel. WES was performed for the proband’s sample (II.1, Fig. 1) at the Persian Bayan Gene Research and Training Center (Shiraz, Iran). The ES condition was performed as follows: bidirectional sequencing of the complete coding region plus 2-kb upstream and 1-kb downstream, with 150x reads, on an Illumina HiSeq 2500 (Illumina, San Diego, CA, USA). The reads were aligned with the reference genome (hg38) sequences using WinterVar (http://wintervar.wglab.org)\(^10\). This process was followed by the detection of single nucleotide variants and small insertion and deletions, as well as by the identification of all other variants in the exons of the target genes. Moreover, the effect of nonsynonymous missense variants was predicted using VarSome\(^11\) and HSF \(^12\). Assessment of variants was carried out by computational prediction tools and genetic databases. For further analysis, variants with a minor allele frequency lower than 1% were selected. After the confirmation of the candidate variant, parents and siblings were screened to assess the co-segregation of the suggested variant using PCR and Sanger sequencing.

Ethical statement

The above-mentioned sampling protocol was approved by the Research Ethics Committee of Mashhad University of Medical Sciences, Mashhad, Iran (ethical code: IR.MUMS.MEDICAL.REC. 1386.250)\(^9\). All participants have signed the informed consent.

RESULTS

Clinical findings

The proband (II.1, Fig. 1) was a 25-year-old male (with complaints of obesity and skin lesions on his hands and elbows), who was referred to a geneticist. After a detailed physical examination, the proband was found to have a history of two episodes of myocardial infarction at ages 23 and 25 years. Moreover, he had bilateral xanthelasma (Fig. 1). According to Simon Broom Criteria \(^13\), he was diagnosed with FH, due to high total cholesterol and LDL-C serum levels before treatment. Other family members consisted of one brother and two sisters who had also high total cholesterol levels and clinical manifestations shown in Table 1. The proband was on treatment with rosuvastatin (40 mg) and ezetimibe (10 mg). Moreover, he took aspirin (80 mg) and bisoprolol (5 mg) for coronary artery disease and hypertension treatment, respectively. The proband responded to the treatment, though he did not reach the lipid target level based on the European Society of Cardiology recommendations.
Molecular findings

Preliminary analysis of the data for the F-10 family revealed that there was a potentially pathogenic variant within the \textit{LDLRAP1} gene: NG_008932.1 (NM_015627.2); c.345-2A>G (GRCh38). This variant caused a nucleotide change at c.345-2A>G, an acceptor splice site in IVS-3 in the \textit{LDLRAP1} gene. The c.345-2A>G variant was novel as no report was found for this variant in the genomic databases at the time of this study. Moreover, searching for the frequency of this variant in databases such as ExAC (http://exac.broadinstitute.org/), dbSNP (https://www.ncbi.nlm.nih.gov/snp/), and gnomAD (https://gnomad.broadinstitute.org/) yielded no results. To identify the clinical classification of the variant, mutation-related databases, such as ClinVar (https://www.ncbi.nlm.nih.gov/clinvar/), OMIM (https://www.omim.org/), and HGMD (http://www.hgmd.cf.ac.uk/ac/index.php), were also searched, which no report was found for the c.345-2A>G. The pathogenicity of the c.345-2A>G variant was assessed through HSF, which aims to help the assessment of the pre-mRNA splicing through 12 different algorithms to recognize and predict the effect of nucleotide changes on splicing sites consisting of the acceptor/donor splice sites and the branch point and auxiliary sequences\cite{11}. In the case of a genetic variant, if the WT score is more than the threshold and the score variation (between WT and Mutant) is less than -10% for HSF, it is considered that the breaking of the splice site has occurred. Since c.345-2A>G variation causes -30.94% difference, it is predicted as a breaking site variant. Moreover, this variant was assessed through the VarSome online tool, which allows users to search variants of interest in their genomic context and collects data from multiple databases in a central location, providing free and easy sharing knowledge on human genomic variations\cite{13}. According to all other available tools such as MutationTaster (http://www.mutationtaster.org/), FATHMM-MKL (http://fathmm.biocompute.org.uk/fathmmMKL.htm), and CADD phred score (https://cadd.gs.washington.edu/), c.345-2A>G was predicted as disease-causing variant. Pathogenicity was also described according to the American College of Medical Genetics and Genomics guidelines\cite{14}. Therefore, c.345-2A>G variant was classified as a ‘pathogenic’ variant.

Co-segregation analysis

The available family members of the proband were segregated for the candidate variant in the \textit{LDLRAP1} gene using Sanger sequencing. The proband (II.1), two sisters (II.2 & II.3) of the proband and one brother (II.4) were homozygous for the \textit{LDLRAP1} gene c.345-
A Novel Variant in the LDLRAP1 Gene

Ahangari et al.

Table 1. Characteristics of the investigated family carrying novel variant of LDLRAP1

| Variants & Parameters | Father | Mother | Proband | II.2 | II.3 | II.3 |
|-----------------------|--------|--------|---------|------|------|------|
| LDLRAP1 variant zygosity | Htz | Htz | Hmz | Hmz | Hmz | Hmz |
| Age (y) | 54 | 47 | 25 | 23 | 16 | 16 |
| Gender | M | F | M | F | F | M |
| Total cholesterol (mg/dL) | 177 | 195 | 670 | 610 | 588 | 567 |
| Triglyceride (mg/dL) | 200 | 210 | 126 | 150 | 132 | 161 |
| LDL-C (mg/dL) | 135 | 120 | 435 | 389 | 410 | 378 |
| HDL-C (mg/dL) | 45 | 51 | 86 | 56 | 59 | 62 |
| Symptoms | None | None | Xanthelasma, Xanthema, CAD | Xanthelasma, Xanthoma | Xanthelasma, Xanthoma | Xanthelasma, Xanthoma |

Hmz, homozygote; Htz, heterozygote; M, male; F, female

2A>G variants (Fig. 1D). The parents (I.1 & I.2) were heterozygous. The variant segregation results as well as the lipid profile and clinical symptoms are summarized in Table 1.

Pharmacogenetics study

It is well known that statins are the first-line therapy for hypercholesterolemia, though responses to these drugs have shown significant differences among patients. These differences in drug response are partly attributed to the variations in genes involved in pharmacokinetics, pharmacodynamics, and lipid metabolisms such as ABCG2, SLC01B1, CYP3A4, and HMGCR. In this regard, we analyzed WES data for the best known reported genetic variants associated with lipid-lowering therapeutic response. We found rs3846662 g.23092A>G, an intronic variant, as homozygous (G/G) in affected individuals.

DISCUSSION

In this clinically ascertained patient with FH who had severe hypercholesterolemia, we found an acceptor splice-site mutation (c.345-2A>G) in intron 3 of the LDLRAP1 gene. This variant was neither reported in any other hypercholesterolemic patients in MASHAD cohort study nor in the Iranome database. This variant was also absent in ExAC and 1000G databases. The rs781769339 represented as an A>T substitution has previously been reported as a splice acceptor variant. According to five applied algorithms, this variant was interpreted to alter splicing. The LDLRAP1 gene encodes a protein consisting of 308 amino acids that involve a PTB (170 amino acids). There are significantly similar sequences to the PTB domains in several adapter proteins. PTB domains, situated in the cytoplasmic domains of several cell surface receptors such as LDLR, bind to the NPXY consensus sequence. Exon 4 of LDLRAP1 is located in the PTB/PID interaction domain. Northern blot analysis has revealed that LDLRAP1 expression is typically occurred at high levels in the kidney, liver, and placenta, while it is expressed at lower levels in the brain, heart, muscle, colon, spleen, intestine, lung, and leukocytes.

Hypercholesterolemia may be occurred due to failure in the hepatic uptake of LDL in the patient. In a report, the genetic analysis of a Mexican FH family with two affected siblings indicated a new mutation (IVS4 + 2T > G) that affects the donor splice site in LDLRAP1 IVS-4, while parents and other siblings were heterozygous. Substitution of IVS4 + 2T>G caused another alternative transcript with 78 deleted nucleotides in mature mRNA in the template. Translation of this mRNA led to the production of ARH-26-a mutated protein without 26 amino acids and also the lack of the b6 and b7 strands of the PTB domain. This was the first report of a mutation leading to an altered PTB domain. Furthermore, increasing LDL uptake by lymphocytes has been reported in individuals carrying LDLRAP1 mutation. As described above, we found g.23092A>G (rs3846662, an intronic variant) within the HMGCR gene in the affected individuals, as well. This variant led to a probably reduced response to simvastatin and pravastatin. Exon 13 of pre-mRNA alternative splicing of HMGCR results in two transcripts, known as rs3846662 with full-length HMGCR and Δ13 HMGCR. HMGCR exon 13 encodes a part of the catalytic/statin-binding domain. The rs3846662 modifies the binding motif of heterozygous nuclear ribonucleoprotein A1, which regulates the alternative splicing of HMGCR. It has been suggested that the high amount of Δ13 HMGCR mRNA in carriers of the rs3846662 allele leads to probable lower activity in HMGCR, as well as lower levels of baseline LDL-C and reduced sensitivity and response to the statin.
A Novel Variant in the LDLRAP1 Gene

Ahangari et al.

inhibition[25].
Overall, we describe, herein, an Iranian FH pedigree with a novel splice site acceptor variant in the LDLRAP1 gene. This variant results in a breaking site in IVS-3 within the PTB domain, which may affect the LDLR and also other related receptors. Functional studies and validation of this variant may lead to a more comprehensive FH screening in the future.

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CONFLICT OF INTEREST. None declared.

REFERENCES
1. Humphries SE, Cranston T, Allen M, Middleton-Price H, Fernandez MC, Senior V, Hawe E, Iversen A, Wray R, Crook MA, Wierzbicki AS. Mutational analysis in UK patients with a clinical diagnosis of familial hypercholesterolaemia: relationship with plasma lipid traits, heart disease risk and utility in relative tracing. Journal of molecular medicine 2006; 84(3): 203-214.
2. Hopkins PN. Encouraging appropriate treatment for familial hypercholesterolemia. Clinical lipidology 2010; 5(3): 339-354.
3. Hopkins PN, Toth PP, Ballantyne CM, Rader DJ. Familial hypercholesterolemias: prevalence, genetics, diagnosis and screening recommendations from the National Lipid Association Expert Panel on Familial Hypercholesterolemia. Journal of clinical lipidology 2011; 5(3): S9-S17.
4. DeMott K, Nherera L, Shaw E, Minhas R, Humphries S, Kathoria M, Ritchie G, Nunes V, Davies D, Lee P, McDowell I, Neil A, Qureshi N, Rowlands P, Seed M, Stracey H, Thorogood M, Watson M. Clinical Guidelines and Evidence Review for Familial Hypercholesterolaemia: The identification and management of adults and children with familial hypercholesterolaemia. 2008. London: National Collaborating Centre for Primary Care and Royal College of General Practitioners. Retrieved from: http://www.safap.it/servizi_lineeguida_200809/CG071F ullGuideline.pdf
5. Goldstein JL, Brown MS. The LDL receptor. Arteriosclerosis, thrombosis, and vascular biology 2009; 29(4): 431-438.
6. Ahmad Z, Adams-Huet B, Chen C, Garg A. Low prevalence of mutations in known loci for autosomal dominant hypercholesterolemia in a multi-ethnic patient cohort. Circulation: cardiovascular genetics 2012; 5(6): 666-675.
7. K, Wilund K, Arca M, Zuliani G, Fellin R, Maioli M, Calandra S, Bertolini S, Cossu F, Grishin N, Barnes R, Cohen JC, Hobbs HH. Autosomal recessive hypercholesterolemia caused by mutations in a putative LDL receptor adaptor protein. Science 2001; 292(5520): 1394-1398.
8. Harada K, Miyamoto Y, Morisaki H, Ohta N, Yamanaka I, Kokubo Y, Makino H, Harada-Shiba M, Okayama Y, Tomoike H. A novel Thr56Met mutation of the autosomal recessive hypercholesterolemia gene associated with hypercholesterolemia. Journal of atherosclerosis and thrombosis 2010 ; 17(2): 131-140.
9. Ghayour-Mobarhan M, Mooshebati M, Esmaily H, Ebrahimi M, Parizadeh SMR, Heidari-Bakavoli AR, Safarian M, Mokhter N, Nematy M, Saber H, Mohammadi M, Andalibi MS, Ferns GA, Azarpazhooh MR, Mashhad stroke and heart atherosclerotic disorder (MASHAD) study: design, baseline characteristics and 10-year cardiovascular risk estimation. International journal of public health 2015; 60(5): 561-572.
10. Li Q, Wang K. InterVar: clinical interpretation of genetic variants by the 2015 ACMG-AMP guidelines. The American Journal of Human Genetics 2017; 100(2): 267-280.
11. Kopanos C, Tsiofakas V, Kouris A, Chapple CE, Aguiller MA, Meyer R, Massouras A. VarSome: The human genomic variant search engine. Bioinformatics 2019; 35(11): 1978-1980.
12. Desmet F-O, Hamrson D, Lalande M, Collod-Bèroud G, Claustres M, Bèroud C. Human splicing finder: an online bioinformatics tool to predict splicing signals. Nucleic acids research 2009; 37(9): e67.
13. Scientific Steering Committee on behalf of the Simon Broome Register Group. Mortality in treated heterozygous familial hypercholesterolaemia: implications for clinical management. Atherosclerosis 1999; 142(1): 105-112.
14. Richards S, Aziz N, Bale S, Bick D, Das S, Gastier-Foster J, Grody WW, Hegde M, Lyon E, Spector E, Voelkerding K, Rehm HL, ACMG Laboratory Quality Assurance Committee. Standards and guidelines for the interpretation of sequence variants: a joint consensus recommendation of the American College of Medical Genetics and Genomics and the Association for Molecular Pathology. Genetics in medicine 2015; 17(5): 405-423.
15. Nieminen T, Kähönen M, Viiri LE, Grönroos P, Lehtimäki T. Pharmacogenetics of apolipoprotein E gene during lipid-lowering therapy: lipid levels and prevention of coronary heart disease. Pharmacogenomis 2008; 9(10):1475-86.
16. Taghizadeh E, Mirzaei F, Jalilian N, Ghayour Mobarhan M, Ferns GA, Pasdar A. A novel mutation in USP1 gene is associated with familial combined hyperlipidemia. IUBMB life 2019; 72(4): 616-623.
17. Taghizadeh E, Ghayour-Mobarhan M, Ferns GA, Pasdar A. A novel variant in LPL gene is associated with familial combined hyperlipidemia. Biofactors 2020; 46(1): 94-99.
18. Dvir H, Shah M, Girardi E, Guo L, Farquhar MR,
Zajonc DM. Atomic structure of the autosomal recessive hypercholesterolemia phosphotyrosine-binding domain in complex with the LDL-receptor tail. *Proceedings of the National Academy of Sciences of the United States of America* 2012; 109(18): 6916-6921.

19. Canizales-Quinteros S, Aguilar-Salinas CA, Huertas-Vázquez A, Ordóñez-Sánchez ML, Rodríguez-Torres M, Venturas-Gallegos JL, Riba L, Ramírez-Jimenez S, Salas-Montiel R, Medina-Palacios G, Robles-Osorio L, Miliar-García A, Miliar-García A, Rosales-León L, Ruiz-Ordaz BH, Zentella-Dehesa A, Ferré-D’Amare A, Gómez-Pérez FJ, Tusié-Luna MT. A novel ARH splice site mutation in a Mexican kindred with autosomal recessive hypercholesterolemia. *Human genetics* 2005; 116(1-2):114-120.

20. Thedrez A, Sjouke B, Passard M, Prampart-Fauvet S, Guédon A, Croyal M, Dallinga-Thie G, Peter J, Blom D, Ciccarese M, Cefalu AB, Pisciotta L, Santos RD, Averna M, Riba L, Lo K, Mangravite LM, Naidoo D, Kutilova M, Medina MW, HNRNPA1 regulates HMGCR alternative splicing and modulates cellular cholesterol metabolism. *Human molecular genetics* 2014; 23(2):319-322.

21. Burkhardt R, Kenny EE, Lowe JK, Birkeland A, Josowitz R, Noel M, Salit J, Maller JB, Pe’er I, Daly MJ, Altshuler D, Stoffel M, Friedman JM, Breslow JL. Common SNPs in HMGCR in micronesians and whites associated with LDL-cholesterol levels affect alternative splicing of exon13. *Arteriosclerosis, thrombosis, and vascular biology* 2008; 28(11): 2078-2084.

22. Johnson JM, Castle J, Garrett-Engele P, Kan Z, Loerch PM, Armour CD, Santos R, Schadt EE, Stoughton R, Shoemaker DD. Genome-wide survey of human alternative pre-mRNA splicing with exon junction microarrays. *Science* 2003; 302(5653): 2141-2144.

23. Istvan ES, Deisenhofer J. Structural mechanism for statin inhibition of HMG-CoA reductase. *Science* 2001; 292(5519): 1160-1164.

24. Yu C-Y, Theusch E, Lo K, Mangravite LM, Naidoo D, Kutilova M, Medina MW, HNRNPA1 regulates HMGCR alternative splicing and modulates cellular cholesterol metabolism. *Human molecular genetics* 2014; 23(2):319-322.

25. Chung JY, Cho SK, Oh ES, Lee DH, Lim LA, Jang SB, Lee YJ, Park K, Park MS. Effect of HMGCR variant alleles on low-density lipoprotein cholesterol—lowering response to atorvastatin in healthy korean subjects. *Journal of clinical pharmacology* 2012; 52(3): 339-346.

Iran. Biomed. J. 25 (5): 374-379 379