

Familial LCAT Deficiency and Low HDL-C Levels: In silico Characterization of Two Rare LCAT Missense Mutations

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Abstract: Mutations in the lecithin-cholesterol acyltransferase (*LCAT*) gene, which catalyzes the esterification of cholesterol, result in two types of autosomal recessive disorders: Familial *LCAT* deficiency (FLD) and Fish Eye Disease (FED). While both phenotypes are characterized by corneal opacities and different forms of dyslipidemia, such as low levels of high-density lipoprotein-cholesterol (HDL-C), FLD exhibits more severe clinical manifestations like splenomegaly, anemia, and renal failure. We describe the first clinically and genetically confirmed case of FLD in Colombia which corresponds to a 46-year-old woman with corneal opacity, hypothyroidism, and dyslipidemia, who does not have any manifestations of renal failure, with two pathogenic heterozygous missense variants in the *LCAT* gene: *LCAT* (NM_000229.2):c.803G>A (p.Arg268His) and *LCAT* (NM_000229.2):c.368G>C (p.Arg123Pro). In silico analysis of the mutations predicted the physicochemical properties of the mutated protein, causing instability and potentially decreased *LCAT* function. These compound mutations highlight the clinical heterogeneity of the phenotypes associated with *LCAT* gene mutations.

Keywords: eye, *LCAT*, cholesterol/trafficking, genomics, VLDL, lecithin cholesterol acyltransferase deficiency, *LCAT* deficiency, alpha-*LCAT* deficiency, fish eye disease

Introduction

Familial *LCAT* deficiency (FLD; MIM 245900) and Fish Eye Disease (FED; MIM 136120) are two autosomal recessive disorders caused by mutations in the lecithin-cholesterol acyltransferase (*LCAT*) gene which is located in the q22.1 region of chromosome 16 and is made up of 6 exons that code for a 440 amino acid residue glycoprotein. *LCAT* is expressed mainly in the liver, although it is also found in smaller amounts in the brain, testicles and plasma.¹

LCAT catalyzes the esterification of unesterified cholesterol (UC) in plasma, the maturation of high-density lipoproteins (HDL) and is essential for the reverse cholesterol transport from peripheral tissues to the liver.² The enzyme reversibly binds to lipoproteins and is responsible for transferring the acyl chain from the second position of lecithin to the hydroxyl group of UC housed within plasma lipoproteins, thus generating esterified cholesterol (EC) and lysolecithin.³ Since EC is significantly more hydrophobic than UC in plasma, the molecule moves into the core of lipoproteins allowing their maturation.¹

The prevalence of mutations in the *LCAT* gene is estimated to be below 1/1,000,000 and of the population with extremely low HDL levels, approximately 2–9% is related to some level of *LCAT* deficiency.^{4,5} Decreased *LCAT* enzymatic activity is characterized clinically by bilateral corneal opacity due to the accumulation of cholesterol deposits

in the corneal stroma and decreased levels of cholesterol housed in HDL in an esterified form known as high-density lipoprotein-cholesterol (HDL-C), which is the result of the decrease in the maturation of HDL and the low capacity of *LCAT* to esterify UC inside lipoproteins.⁶

LCAT enzymatic activity is classified into two groups: alpha *LCAT* and beta *LCAT* enzymatic activity. Alpha *LCAT* activity esterifies the UC in HDL, whereas beta *LCAT* activity catalyzes the reaction in low-density lipoproteins (LDL) and very low-density lipoproteins (VLDL).^{2,4} Patients that retain only de beta *LCAT* enzymatic activity will present low levels of HDL-C and milder clinical manifestations because beta *LCAT* is still active on VLDL and LDL: the characteristic bilateral corneal opacity, low HDL-C levels, high LDL-C and TG levels, and normal to elevated plasma EC levels.⁶ However, if both groups of enzymatic activity are affected, cholesterol esterification will be almost null in lipoproteins and will lead to familial *LCAT* deficiency (FLD, MIM: 25900) which is a severe pathology with clinical manifestations such as low HDL-C levels, corneal opacities, elevated plasma TG levels, hemolytic anemia, splenomegaly, proteinuria and progressive renal failure lead by the accumulation of UC in tissues like the glomeruli and erythrocytes' membrane.³

Here we describe the first clinically and genetically confirmed case of FLD in Colombia, which corresponds to a 46-year-old woman with corneal opacity, hypothyroidism, dyslipidemia, and episodes of anemia, with no splenomegaly and no manifestations of renal failure, with two compound heterozygous variants in the *LCAT* gene: *LCAT* (NM_000229.2): c.803G>A (p.Arg268His) and *LCAT* (NM_000229.2):c.368G>C (p.Arg123Pro). This research contributes to enriching the spectrum of variants of this rare disease, as well as highlighting the clinical heterogeneity of this phenotype.

Experimental Procedures

Sample Processing

Blood samples were collected in 4mL EDTA tubes and genomic DNA extraction was performed using the QIAamp DNA Mini Kit (QIAGEN, Germany) following the manufacturer's protocol. The concentration and purity (260/280 and 260/230 ratios) of the nucleic acids were evaluated using a NanoDrop 2000 spectrophotometer (Thermo Fisher Scientific, USA). Long PCR was performed using the *Promega GoTaq® Long PCR Master Mix* with the primers 5'-GGTTGCCCCGTTGATTCTGTTG-3' and 3'-ACTGAACAACTCGGGTCCT-5', generating a 6336 bp amplicon covering all the exons of the *LCAT* gene. The PCR conditions were as follows: an initial denaturation at 94°C for 2 minutes, followed by 40 cycles of denaturation at 93°C for 30 seconds, annealing at 60.5°C for 45 seconds, and extension at 72°C for 6 minutes, with a final extension step at 72°C for 5 minutes. To ensure proper amplification, PCR products were separated by gel electrophoresis with 1% agarose at 100V for 40 minutes, stained with ethidium bromide, and visualized using UV light. Since non-specific amplifications were obtained, it was necessary to perform amplicon purification by band excision using the E.Z.N.A.® Cycle Pure Kit (Omega bio-tek, USA).

Purified amplicons were used for single-stranded Sanger sequencing of all the exons of the *LCAT* gene using the BigDye Terminator v.3.1 (Applied Biosystems, Foster City, USA) following the manufacturer's instructions. The primers used for each exon sequencing are described in Table 1. The Sanger sequencing products were purified with the BigDye®

Table 1 Sequencing Primers

Type	Primer	Exon
Forward	5'- CCCACTCCCACACCAGATAA-3'	1
Forward	5'-GTGTAAGCAGGGGAGGGTAA-3'	2-3
Forward	5'-CACCCCTAGCCCCAACACG-3'	3
Forward	5'-GAGTACCTGGACAGCAGCA-3'	4-5
Forward	5'-ACAGCTCCACCCAACAGA-3'	6
Reverse	5'-CCACGCCGTAAAGACAGT-3'	6

Xterminator™ Purification Kit (Applied Biosystems, Thermo Fisher Scientific, USA) and loaded in the 3500 Genetic Analyzer (Thermo Fisher Scientific, USA). Sequence data were analyzed using MEGA X software⁷ and the GenBank reference sequence of the *LCAT* gene (NG_009778.1).

Mutation Analysis

In order to predict the potential impact of a gene variant in protein structure or function, in silico bioinformatic tools that analyze the stability and functionality of the mutated protein were used: SIFT predictor (<https://sift.bii.a-star.edu.sg/>) which uses sequence homology to predict whether an amino acid substitution will affect protein function;⁸ PolyPhen-2 (<http://genetics.bwh.harvard.edu/pph2/>) which uses annotated UniProt entries to predict whether the protein variation occurs within an important structural or functional site of the protein based mainly on a well annotated crystal structure or modeled protein;⁹ I-mutant 2.0 (<https://folding.biofold.org/i-mutant/i-mutant2.0.html>) which from an experimental thermodynamic database predicts changes in protein stability based on changes in free energy¹⁰ and CADD (<https://cadd.gs.washington.edu/>) which analyzes multiple parameters built through other predictors and databases and integrates them into a single global score.¹¹ Finally, to review the impact that the change of amino acids has from the physical-chemical point of view on the interactions and functionality of the protein, the HOPE platform (www.cmbi.umcn.nl/hope) was used, which based on the properties of the wild-type amino acid contrasted with the mutant, predicts the changes and alterations that could occur.¹²

Case Presentation

A 46-year-old Colombian woman, daughter of non-consanguineous parents, was referred to the genetic area due to a differential diagnosis between *LCAT* deficiency phenotypes and Tangier disease because of corneal opacity without visual acuity alteration, but with progressive deterioration of night vision, since the age of 10 years. The patient appeared phenotypically healthy, except for the corneal opacity (Figure 1). At the time of consultation, the patient did not present hepatomegaly or splenomegaly, proteinuria, anemia or other clinical characteristics. However, the patient had presented anemia episodes in 2018 and 2019. Electrolytes information revealed hypercalciuria and a significant hypercalcemia, 31.86 mg/dL and 10.30 mg/dL, respectively. The patient had no symptoms of coronary atherosclerosis. The results of the tests of liver and renal function were normal. Hemogram without evidence of leukocytosis, leukopenia, or any white cellular abnormality. No evidence of relevant alteration in hemoglobin levels. Platelets in adequate ranges without evidence of thrombocytosis or thrombocytopenia. Vitamin D, serum and urine creatinine, ferritin and phosphorus levels in serum are within the standard parameters.

The patient was diagnosed with primary hypothyroidism at the age of 30 years (TSH 9.09mUI/mL). Throughout her life she presented with chronic dyslipidemia characterized by persistently low HDL-C levels stood out, even down to 2.8 mg/dL (Figure 2), high levels of triglycerides (216 mg/dL) and low levels of Apo-A1 (34.6 mg/dL). Total cholesterol



Figure 1 Patient's corneal opacity.

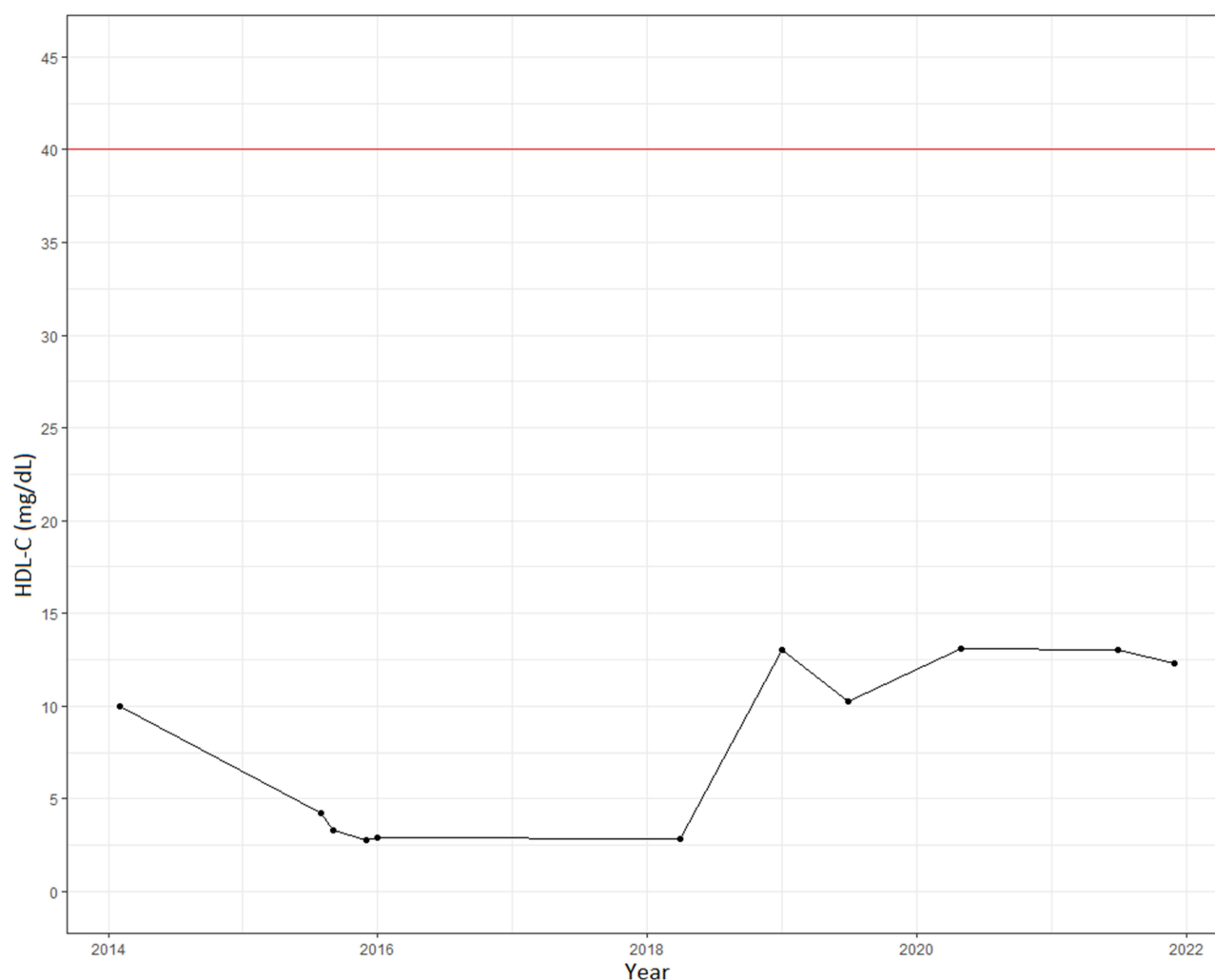


Figure 2 Temporal record of the patient's HDL-C levels. Red line corresponds to the minimum value of the clinical reference.

of 95.0 mg/dL and non-HDL cholesterol of 92.2 mg/dL are within the standard population. VLDL levels have been above the reference value (30 mg/dL) and LDL-C levels have been continuously low, 43 mg/dL and 38.80mg/dL, respectively.

At the age of 39 years, lipoprotein electrophoresis revealed hypoalphalipoproteinemia (14.4% of total band signal) and a consequent increase of beta fraction (78.6% of total band signal), indicating that, even though the LDL-C values are low but among the reference levels (0–150 mg/dL), most of the circulating cholesterol was in beta migrating particles (LDL). The esterified cholesterol levels were very low: 6% of total cholesterol (reference value 60–80%). In addition, two of her three brothers have had recurrently low levels of HDL-C throughout their lives (down to 15–16 mg/dL).

The 6 exons in the *LCAT* gene of the proband were analyzed using the Sanger method to identify the mutations responsible for the pathology. Two missense variants were found: c.368G>C (p.Arg123Pro) classified as pathogenic (criteria: PM2, PM5, PP3, PP2, PP5 according to the American College of Medical Genetics and Genomics, ACMG)¹³ and c.803G>A (p.Arg268His) classified as pathogenic (criteria: PS3, PP3, PM2, PM5, PP5, according to the ACMG) located in exons 3 and 6 of the *LCAT* gene, respectively (Figure 3). Once the mutations were confirmed, carrier analysis was performed on the parents and three siblings, which determined that the proband inherited the variant Arg123Pro from her mother and the mutation Arg268His from her father. In addition to the patient, two of her male siblings are carriers of the variant located in exon 6 (Arg268His) (Figure 3).

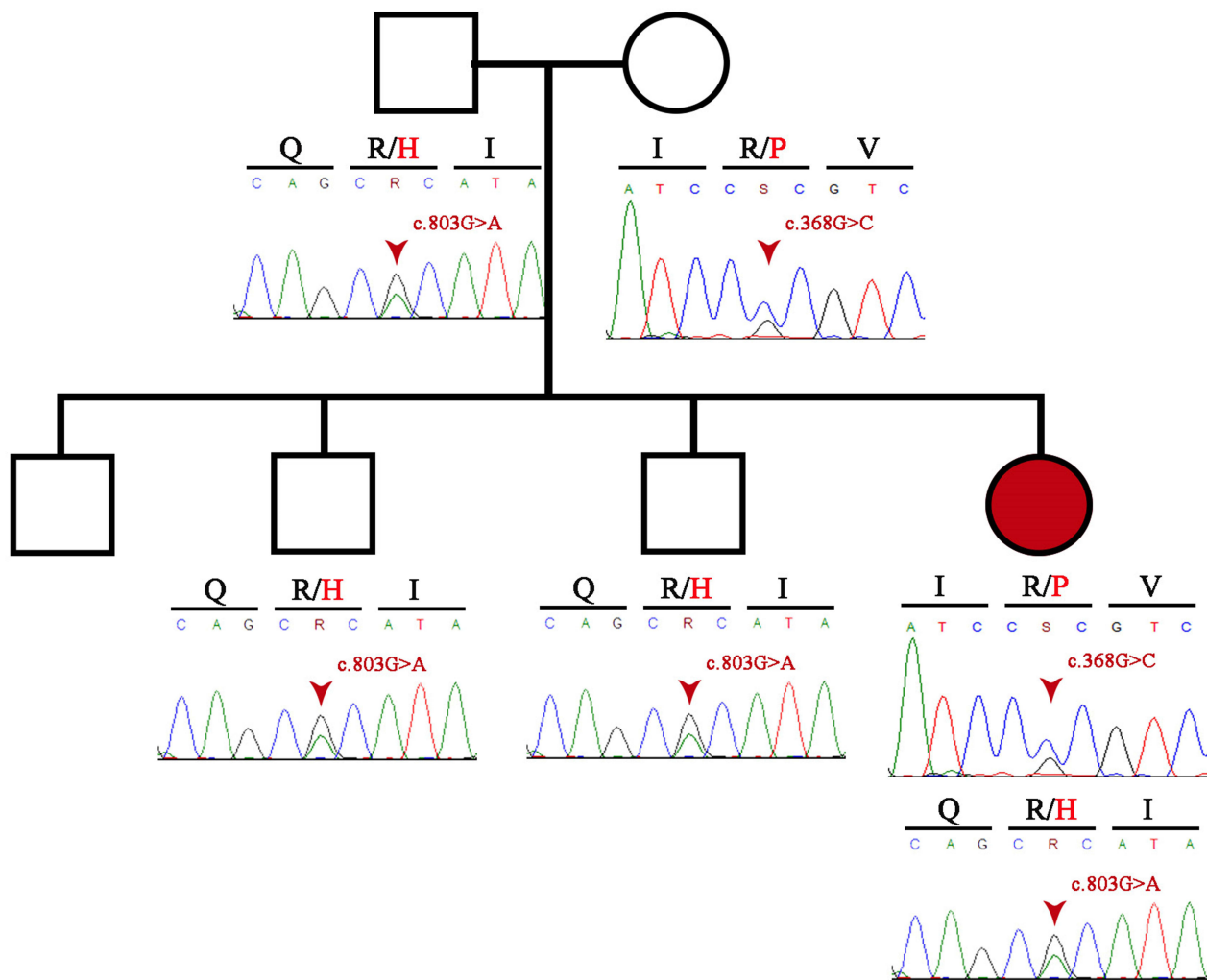


Figure 3 Pedigree information of the patient and Sanger sequencing electropherogram of both patient and her family.

Analysis of c.368G>C (p.Arg123Pro)

This mutation was classified as pathogenic according to the criteria of the ACMG. This variant has been described in the literature in a homozygous 44-year-old Spanish woman,¹⁴ although there is no functional information on its impact on the function of the protein. This variant is classified in this category because of other known mutations in the same codon and its Genome Aggregation Database (gnomAD) allelic frequency of 0.00000398. Variant rs199717050 (Arg123His), the mutation in the same codon reported in our patient, was associated with decreased ($\beta = -0.72$) HDL-C levels (P discovery = 5.9×10^{-10} , P conditional = 2.5×10^{-12}) in the Finnish population.¹⁵

Using in silico prediction tools it was possible to determine that the region of the mutation corresponds to an *LCAT* conserved region (Figure 4) and SIFT predicts that the residue change has a deleterious effect on the function of the protein (0.018). This variant has a score of 1.0 (maximum score) on the *PolyPhen-2* predictor and according to *I-mutant 2.0* this change decreases, in silico, the stability of the new mutated protein ($\Delta G = -1.60$). The mutation also obtained a high score in the *CADD* predictor (26.5). Finally, according to *HOPE*, the amino acid change corresponds to a smaller one with a neutral charge, possibly altering the physicochemical characteristics of the protein.

Analysis of c.803G>A (p.Arg268His)

The effects of this variant are quite similar. The mutation was analyzed with the *PolyPhen-2* and *SIFT* predictors that gave a score of 1.0 and 0.0 respectively, being the maximum values for predictions of deleterious effects of a mutation.

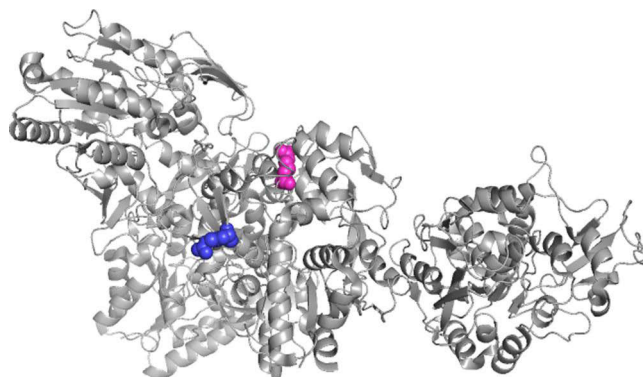


Figure 4 Visualization through PyMOL of the crystal structure of the human Lecithin-Cholesterol Acyltransferase (4X96 Protein Data Bank) reported by Glukhova et al 2015.¹⁶ Protein Data Bank accession number 4X96. Arginine residue 123 is visualized in blue and the purple residue corresponds to Arginine 268.

I-mutant 2.0 classified it as a variant that decreases protein stability in silico ($\Delta G = -1.05$) and it obtained a *CADD* score of 28.5. *HOPE* predicts physicochemical changes in the protein and loss of internal and external interactions, since the amino acid change differs in size and charge with the new one (Figure 4). The allelic frequency of this variant in gnomAD is 0.000036 and according to the ACMG classification the mutation corresponds to a pathogenic variant with known cases of this mutation related to *LCAT* deficiency.^{17–19}

Discussion

This is the first clinically and genetically diagnosed case of FLD in Colombia and the first report of a compound heterozygous patient with the *LCAT* variants Arg123Pro and Arg268His. Mutations in codon 123 of the *LCAT* gene have been described previously. Blanco-vaca et al described a Spanish woman homozygous for the Arg123Cys variant who was diagnosed with FED based on the appearance of corneal opacities at the age of approximately 54 years and vanishingly low plasma concentrations of cholesteryl esters.²⁰ However, our patient's corneal opacity and progressive deterioration of night vision started since the age of 10 years, a rather soon manifestation when compared to the initiation of ocular symptoms in most of FED cases. In addition, Bérard et al described a compound heterozygous case with the same mutation as the Spanish patient.²¹ This compound heterozygous case, harboring one of our patient's mutations, sheds some light on the possibility of our patient being a FLD case. The patient's plasma *LCAT* concentrations and alpha *LCAT* enzymatic activity were significantly reduced, indicating a virtual absence of *LCAT*. Furthermore, Arg123 is a conserved residue in the human, rabbit, rodent, *C. albicans* and yeast *LCAT* genes, suggesting a role in *LCAT* function or stability.²² Although there is no information about the specific alpha or beta *LCAT* enzymatic activity of mutation Arg123Pro, *LCAT* enzymatic activity was 15.2 nmol/mL/hour (reference range, 81 ± 12 mL/min/hour) in a homozygous patient.²⁰

Splenomegaly, anemia, and renal failure are some differential conditions that most FLD patients share, in contrast to most FED cases. Although important contrasting symptoms, the appearance and progression of renal failure are variable among FLD cases, and it is likely related to the biochemical phenotype rather than to the inherited mutation.²³ Clinical and biochemical heterogeneity is a challenging characteristic regarding the diagnosis of *LCAT* phenotypes. However, on the basis of the clinical and biochemical features of our case, the anemia episodes, and that both of our patient's mutations and variants in the same codons have been described in FLD patients,^{14,19–22} we diagnosed our compound heterozygous patient as an FLD case who had not developed renal failure, proteinuria, or splenomegaly, pointing to the possibility that additional genetic or environmental factors may have contributed to the apparently benign course of the patient's disease.

Likely due to the complexity of the *LCAT* biochemical reaction and despite the availability of a 3D model enzyme, it is impossible to predict the phenotype associated with the mutations.²⁴ The dyslipidemia profile is indistinguishable between subjects classified as FLD or FED. The differential diagnosis between these two phenotypes is limited to alternatives that are not available in clinical laboratories in Colombia, such as the measurement of the ability of individual plasma to esterify cholesterol in endogenous lipoproteins (alpha *LCAT* with beta *LCAT* enzymatic activity) and in a standardized exogenous HDL (alpha *LCAT* activity only), both of which are null in FLD cases but low or normal

in FED.^{3,24} Other useful alternatives to distinguish between these conditions are the CE/TC ratio in plasma, which is always reduced in FLD but not in FED; and through the expression of *LCAT* mutants in cultured cells, and subsequent measurement of *LCAT* concentration and activities in cell media.³ Nevertheless, we were unable to calculate this ratio as well as performing these experiments.

An association with increased risk of cardiovascular disease has been described in *LCAT* phenotypes by preserving the esterification of cholesterol in atherogenic lipoproteins such as LDL. Nevertheless, at the time of the patient's approach, no calcified atherosclerotic plaques were documented by coronary computed tomography (calcium score 0 Agatston units). Cardiovascular disease in *LCAT* diseases has been described with a median age at presentation of 56 years, meaning that the patient is still at risk of developing such condition.^{3,25}

Despite FLD being a recessive disorder, the two siblings of the proband are carriers of the Arg268His mutation who have HDL-C levels that are persistently below normal. Authors have described the behavior of lipoproteins in carriers of *LCAT* gene mutations who don't have clinical manifestations as severe as a homozygous or compound heterozygous patient, but whose HDL-C levels are persistently below the values of non-carrier subjects, meaning they express an intermediate phenotype.^{4,26}

Of the compound heterozygous mutations in the patient, Arg123Pro was found in a 44-year-old woman in Spain.¹⁴ The patient was homozygous for this *LCAT* gene variant and had corneal dystrophy, anemia and an altered lipid profile resembling our patient: HDL-C levels down to 6.6 mg/dL, triglycerides up to 173 mg/dL, LDL-C 131 mg/dL, VLDL 45 mg/dL and Apo-A1 62 mg/dL. However, unlike our patient, the Spanish *LCAT* patient had proteinuria and the histology study from the renal biopsy confirmed segmental hyaline lesions, irregular mesangial enlargement, and parietal thickening of the glomerular capillary walls. Despite having the same mutation, our patient didn't have any sign of renal compromise or proteinuria.

According to the in silico predictors, this variant is classified as a destabilizing and deleterious mutation, and according to the ACMG criteria it is classified as pathogenic. The Arg123Pro mutation is predicted to generate changes in the stability and functionality of the protein because of the difference in size and charge from the original residue as it is smaller and has a neutral charge, as well as being more hydrophobic than arginine.

The mutated residue is found in the membrane binding region, a region enriched with hydrophobic residues responsible for anchoring HDL to membranes to initiate the cholesterol esterification process (Figure 4).²⁷ Variations in the physicochemical properties of the new residue affect the formation of chemical bonds with other residues and could cause a loss of external interactions, which interferes in the function of the protein.¹²

The Arg268His variant was previously reported as a cause of *LCAT* enzyme activity deficiency, specifically of FLD in compound heterozygous patients.¹² This mutation is found in the Cap domain of the protein, specifically a part of the lid region (residues 257–271), which opens and closes the access through a hydrophobic tunnel that leads to the catalytic site of the enzyme. This tunnel is made up of hydrophobic residues whose function is also to protect these residues from interaction with water.^{16,28} In addition, this residue forms salt bridges with Asp 359 and Glu 265, which keep the lid in a closed conformation; but when they are broken the lid changes to an open conformation.²⁹ This variant corresponds to a substitution of a polar amino acid for a smaller one with a neutral charge, unlike arginine, which has a positive charge that allows it to form salt and hydrogen bonds with two leucines at positions 247 and 309, interactions that will be affected by this mutation.¹²

It is believed that the Cap domain interacts with Apo-AI (major HDL apolipoprotein) and is essential for the activation of the *LCAT* enzyme, the interaction involves a conformational change in the lid to an open state for a better binding with the substrate.^{30,31} In this case, the Arg268His mutation could interfere with the interaction of HDL with Apo-AI and the activation and the conformational change of the lid, causing poor cholesterol esterification specifically in HDL. In addition, Holleboom et al concluded that this residue is important for the expression and function of the enzyme based on the severe reduction of its expression in an in vivo study in the case of a mutation in the same codon (Arg268Cys).³²

Our study has some limitations. We were unable to obtain specific data of the siblings' corneal photographs, renal function, and specific blood data. In addition, limited to our research and clinical resources, we could not perform any of the assays that would clearly differentiate FLD from FED, such as the measurement of alpha and beta *LCAT* enzymatic activity. Also, we were not able to contact the patient for further ophthalmologic information such as intraocular pressure. Finally, we were not able to perform a molecular characterization of the variants, which would have required expression of the variants in cell systems and evaluation of their ability to promote alpha and beta *LCAT* enzymatic activity.

Conclusion

This is the first clinically and genetically confirmed report of FLD in Colombia. Moreover, this is the first case of a compound heterozygous patient with the *LCAT* (NM_000229.2):c.803G>A (p.Arg268His) and *LCAT* (NM_000229.2):c.368G>C (p.Arg123Pro) mutations. The in silico analysis of the mutations determined the affect of the physicochemical properties of the protein, mainly by altering the interaction of residues in their own domains or external interactions with other proteins, possibly causing instability and decreased function of the *LCAT* enzyme. This case highlights the clinical heterogeneity caused by *LCAT* mutations, demonstrating the possibility of other factors that may contribute to these phenotypes. The high risk of developing accelerated atherosclerotic disease as previously described may be an indicator for pharmacological (statin) and non-pharmacological interventions for its prevention in this patient. The description of these two variants will allow a better characterization of FLD and *LCAT* phenotype patients and support the identification of other individuals in heterozygosity and their appropriate clinical approach, as well as highlight the necessity to further evaluate the prevalence and clinical presentation of *LCAT* deficiency syndromes in Latin American countries.

Abbreviations

FED, Fish eye disease; *LCAT*, lecithin-cholesterol acyltransferase; UC, unesterified cholesterol; HDL, high-density lipoproteins; EC, esterified cholesterol; FLD, familial *LCAT* deficiency; LDL, low-density lipoproteins; VLDL, very low-density lipoproteins; HDL-C, high-density lipoprotein-cholesterol; LDL-C, low-density lipoprotein-cholesterol; TG, triglycerides; ACMG, American College of Medical Genetics; gnomAD, Genome Aggregation Database; RFLP, restriction fragment length polymorphism.

Data Sharing Statement

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Ethics Approval

All research was conducted according to the Declaration of Helsinki and the research protocol was registered with the number 1504 upon approval from the IRB Biomedical Research Ethics Committee of the Fundacion Valle del Lili. The patient and patient's family provided their written informed consent to participate in this study and the publication of the case details and accompanying images.

Consent for Publication

The patient and patient's family provided their written informed consent authorizing to perform genetic tests, use case details, pictures and publish the case along with the accompanying images.

Acknowledgments

The authors would like to thank the patient for agreeing to the publication of this report. We also thank the people who have contributed to the development and execution of this study.

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

Funding

This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Disclosure

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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