Variations in \textit{LOXL1} associated with exfoliation glaucoma do not affect amine oxidase activity

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\textbf{Purpose:} Lysyl oxidase-like 1 (\textit{LOXL1}) is a copper-dependant amine oxidase that plays an essential role in elastogenesis. Two non-synonymous single-nucleotide polymorphisms of \textit{LOXL1}, R141L (rs1048661) and G153D (rs3825942), have been reported to significantly increase susceptibility to exfoliation glaucoma (XFG). To evaluate the impact of the R141L and G153D variations on the amine oxidase activity of \textit{LOXL1}, we generated four different haplotypes of \textit{LOXL1} with R141L and G153D and assessed the amine oxidase activity of the \textit{LOXL1} variant proteins.

\textbf{Methods:} The four different haplotype variants of \textit{LOXL1} were created by oligonucleotide-directed mutagenesis in an \textit{LOXL1} expression vector. Recombinant \textit{LOXL1} variant proteins were purified by nickel-affinity chromatography. The amine oxidase activities of the \textit{LOXL1} variant proteins were assessed using peroxidase-coupled fluorometric assays.

\textbf{Results:} All of the haplotype variants of \textit{LOXL1} (141R-153G, 141R-153D, 141L-153G, and 141L-153D) showed \(\beta\)-aminopropionitrile-inhibitable amine oxidase activity toward elastin, type I collagen, and cadaverine, indicating that each \textit{LOXL1} variant functions as an amine oxidase. However, there were no significant differences in amine oxidase activity between the \textit{LOXL1} haplotype variants toward the tested substrates.

\textbf{Conclusions:} The R141L and G153D variations in the NH\textsubscript{2}-terminal region of \textit{LOXL1} do not affect the amine oxidase activity of \textit{LOXL1}. This is consistent with recent genetic findings on the reversal of risk alleles of R141L and G153D in different ethnic backgrounds. Our results suggest that other unknown genetic factors or molecular mechanisms may be more relevant to the development of XFG.

Lysyl oxidase-like 1 (\textit{LOXL1}) was originally reported as a novel human gene with amino acid sequence homology to the COOH-terminus of lysyl oxidase (\textit{LOX}) [1,2]. \textit{LOX} is a copper-dependent amine oxidase responsible for the lysine-derived cross-links in collagen and elastin, modulating the essential step for the biogenesis of fibrillar extracellular matrix in most tissues [3]. The \textit{LOX} gene family consists of five paralogs (\textit{LOX},\textit{LOXL1},\textit{LOXL2},\textit{LOXL3}, and \textit{LOXL4}) in humans, each containing a copper-binding motif, lysyl-tyrosyl-quinone (LTQ) residues, and a cytokine receptor-like (CRL) domain in the highly conserved COOH-terminus. We have previously shown that each of the \textit{LOX}-like proteins functions as an amine oxidase [4-7], but functional differences of these \textit{LOX} family proteins have not been clearly determined. Recently, \textit{Loxl1} null mice were found to have significantly reduced levels of desmosine in various tissues, indicating that \textit{LOXL1} serves as the primary cross-linking enzyme for formation, maintenance, and remodeling of elastic fibers [8].

Exfoliation glaucoma (XFG) occurs in the context of the exfoliation syndrome (XFS), characterized by progressive accumulation of abnormal microfibrillar material throughout the anterior segment of the eye and various ocular tissues [9]. XFG is the most common identifiable cause of secondary open-angle glaucoma worldwide, causing rapid progression of glaucomatous optic neuropathy with a high resistance to medical treatments [9]. The abnormal microfibrillar deposits in XFG are composed of a crosslinked and insoluble glycoprotein-proteoglycan complex, predominantly with epitopes of elastic fibers [10], suggesting the relevance of \textit{LOXL1} to the formation of the exfoliative material. In fact, using direct mass spectrometry, \textit{LOXL1} was identified as a component of the exfoliative material surgically isolated from XFG patients [11]. Later, using atomic force microscopy-based imaging, \textit{LOX1} was found to be localized around the fibrous protein material on the lens capsule surface [12].

In a genome-wide search for genetic risk factors of XFG, two non-synonymous single-nucleotide polymorphisms (SNPs) in exon 1 of \textit{LOXL1}, rs1048661G/T (R141L) and rs3825942G/A (G153D), were identified with the G allele as a risk allele at both SNPs in Nordic populations [13]. Of four possible haplotypes involving the two SNPs, only three haplotypes (GG, GA, and TG) were detected, and the TA haplotype was not observed in either the XFG or control group. Relative to the GA haplotype, the GG and TG haplotypes showed odds ratios of 27.05 and 8.90, respectively, for the incidence of XFG [13]. Subsequent studies confirmed the association of these SNPs with XFG in...
Table 1. List of Primers Used for Oligonucleotide-Directed Mutagenesis.

| SNP     | Amino acid change | Sequence                                           | Tm (°C) |
|---------|-------------------|----------------------------------------------------|---------|
| rs1048661 | R→L               | F-5’- gggcatggcccGgcgcgccactecG -3’               | 76      |
|         |                   | R-5’- gggctgagggcAggcatgcce -3’                   | 76      |
| rs3825942 | G→D               | F-5’- agccgacgagggAcctcctcG -3’                   | 75      |
|         |                   | R-5’- caggagggagtgccgctcG -3’                     | 75      |

In the sequence, the bases mismatched with the template are indicated in capital letters.

different populations, including German, Italian, Caucasian, Australian, and United States populations [14-17].

To evaluate the effects of the R141L and G153D variations on the amine oxidase activity of LOXL1, we constructed the GG, GA, TG, and TA haplotype variants across rs1048661 and rs3825942 by oligonucleotide-directed mutagenesis, each corresponding to 141R-153G, 141R-153D, 141L-153G, and 141L-153D in the amino acid sequence of LOXL1, respectively. The LOXL1 variant proteins were expressed and purified as hexa-histidine tagged recombinant proteins. The amine oxidase activities of the purified LOXL1 variant proteins were accessed using in vitro peroxidase-coupled fluorometric assays. The four different haplotype variants of LOXL1 did not show significant differences in amine oxidase activity toward elastin, type I collagen, or cadaverine.

**METHODS**

Construction of expression plasmids for four different LOXL1 haplotypes: The expression plasmid for LOXL1 was generated as previously described [4]. A pET21a-derived expression plasmid containing the nucleotide sequence from codon 135D to the COOH-terminus was used as a template for oligonucleotide-directed mutagenesis. The mutagenesis was performed using the QuickChange II site-directed mutagenesis kit (Stratagene, La Jolla, CA) according to the manufacturer’s protocol. Thermocycling consisted of 16 cycles at 95 °C for 30 s, 58 °C for 1 min, and 68 °C for 7 min. The sequences of the oligonucleotide primers used for mutagenesis are listed in Table 1. The nucleotide sequences of all resulting constructs were confirmed by DNA-sequencing analysis using the BigDye™ Terminator Cycle Sequencing Ready Reaction Kit (Applied Biosystems, Foster City, CA) according to the manufacturer’s protocol.

Purification and refolding of the haplotype variant proteins of LOXL1: Purification and refolding of the four different haplotype proteins of LOXL1 were performed as we previously described for the LOX family proteins [4,7]. Briefly, the pET21a-derived expression constructs harboring the four different haplotype proteins of LOXL1 were introduced into the Escherichia coli strain BL21(DE3; Novagen, Darmstadt, Germany). After induction with 1 mM isopropyl-1-thio-β-D-galactopyranoside (IPTG) at 37 °C, the transformed bacterial cells were lysed in a buffer containing 50 mM Tris, pH 8.0, 1 mM EDTA, 100 mM NaCl, 1 mM PMSF, 1 mg/ml lysozyme, 1% Triton X-100, and 0.1 mg/ml DNase. After centrifugation, inclusion bodies were homogenized in a buffer containing 6 M urea, 10 mM K₂HPO₄, pH 8.2, and 3 mM β-mercaptoethanol. The LOXL1 variant proteins were then purified using Ni-NTA agarose resins (Qiagen, Valencia, CA) according to the manufacturer’s protocol. For refolding of the LOXL1 variant proteins, stepwise dialysis was performed in a buffer containing 10 mM K₂HPO₄, pH 9.6, 200 μM CuCl₂, and 2% sodium N-lauroylsarcosinate and then in a buffer containing 10 mM K₂HPO₄, pH 9.6, and 5 μM CuCl₂. The protein samples were further dialyzed in 10 mM K₂HPO₄, pH 9.6. The dialyzed proteins were lyophilized in the presence of 10 mM trehalose using a freeze dryer (Labconco, Kansas City, MO). All purification procedures were performed at 4 °C. The purity and sizes of the LOXL1 variant proteins were determined using sodium dodecyl sulfate-PAGE (SDS-PAGE).

Amine oxidase assays for the haplotype variant proteins of LOXL1: Amine oxidase activity of the haplotype variants of LOXL1 was assessed using a peroxidase-coupled fluorometric assay with the Amplex red hydrogen peroxide assay kit (Molecular Probes, Eugene, OR), as previously described [18]. Each reaction contained 10 μg of a purified LOXL1 variant protein and 20 pmol of substrate in a reaction volume of 200 μl. Parallel assays were performed in the absence or presence of 1 mM β-aminopropionitrile (BAPN) for 2 h at 37 °C. Bovine neck ligament elastin, calf skin type I collagen, and cadaverine (1,5-pentanediamine; Sigma-Aldrich, St. Louis, MO) were used as substrates for amine oxidase assays. Fluorescence was measured using a fluorescence microplate reader (BioTek, Winooski, VT) with excitation and emission wavelengths of 560 nm and 590 nm, respectively. Total amine oxidase activity was expressed as nM of H₂O₂ produced per μg of the LOXL1 variant protein, calculated by interpolation with fluorescence values from an H₂O₂ calibration curve.

**RESULTS**

We previously reported the expression and purification of an enzymatically active form of human LOXL1 as a recombinant protein (LOXL-p1) [4]. The NH₂-terminal site of the previously reported LOXL-p1 [4] was deduced from amino acids.
acid sequence homology with the bovine LOXL1 protein, of which potential NH$_2$-terminal proteolytic cleavage sites were suggested by incubation with bone morphogenetic protein (BMP)-1 [19]. Using the previously reported expression construct of LOXL-p1 (141R-153G haplotype) as a template, we generated three additional haplotype variants at R141L and G153D by a series of oligonucleotide-directed mutagenesis (Figure 1). All of the expected haplotype variants of LOXL1 (141R-153G, 141R-153D, 141L-153G, and 141L-153D) contain a copper-binding domain, a CRL-domain, and LTQ residues (477K and 511Y) in the COOH-terminal region, which are considered prerequisites for amine oxidase activity of the LOX family proteins. Upon induction with 1 mM IPTG in E. coli, all expression constructs of the four different LOXL1 variants showed high levels of expression. However, the LOXL1 variant proteins were within inclusion bodies. The insoluble fractions were solubilized with 6 M urea, and then the LOXL1 variant proteins were purified by nickel-chelating affinity chromatography using a hexa-histidine tag attached at the COOH-terminus of the variant proteins. To refold the LOXL1 variant proteins denatured by urea during purification, the protein samples were subjected to stepwise dialysis in the presence of N-lauroylsarcosinate and Cu$^{2+}$. The apparent sizes of the purified recombinant LOXL1 variant proteins were in good agreement with the deduced molecular mass, 56 kDa, and were over 95% pure on SDS–PAGE gels (Figure 2).

Amine oxidase activity of the four different haplotype variants of LOXL1 was evaluated using physiologic substrates of LOXL1, elastin and type I collagen, as well as cadaverine, an artificial substrate with a diamine structure. All four haplotype variants of LOXL1 showed significantly higher levels of amine oxidase activity in the absence of BAPN than in the presence of BAPN, a well known specific inhibitor of LOX and LOXL1, indicating that the purified LOXL1 variant proteins had BAPN-inhibitable amine oxidase activity toward the tested substrates (Figure 3). Regardless of the haplotype, the LOXL1 variant proteins showed higher activities toward cadaverine than elastin and type I collagen. However, there were no significant differences in amine oxidase activity between the different haplotype variants toward any of the tested substrates (Figure 3), suggesting that the 141R and 153G variations in the NH$_2$-terminal region do not affect the amine oxidase activity of LOXL1.

DISCUSSION
Since the first report on genetic association of rs1048661 and rs3825942 with XFG in Nordic populations [13], reversal of
the risk alleles have been reported in various ethnic backgrounds. For instance, the G allele of rs1048661 was identified as a risk allele in most populations studied, including Nordic, United States, German, Italian, Caucasian Australian, Finnish, and black South African populations [13-17,20], while the opposite T allele of rs1048661 was reported to increase the risk of XFG in Japanese and Chinese populations [21,22]. The G allele of rs3825942 has been reported as a risk allele in most ethnic backgrounds studied [13-17,21-26]; however, the A allele of rs3825942 was identified as a risk allele for XFG in black South Africans [20]. In addition, an association between rs1048661 and XFG was not observed in some populations, including Indian, Chinese, Latin American, and United States populations [23-26]. These inconsistent genetic findings suggest that the R141L and G153D variations may not be causative factors in the development of XFG. To investigate the effects of the R141L and G153D variations on amine oxidase activity of LOXL1, we generated four different haplotype variant proteins with R141L and G153D. In amine oxidase assays using the purified LOXL1 variant proteins, the four different haplotypes did not show significantly different amine oxidase activities toward elastin, type I collagen, or cadaverine. These assay results explain the discrepant genetic findings on the association of rs1048661 and rs3825942 with XFG in different ethnic backgrounds, suggesting that other unknown genetic factors or molecular mechanisms may be more critical in the development of XFG.

Compared to the highly conserved COOH-terminal domains of the LOX family proteins, LOXL1 shows little amino acid sequence homology with LOX in the NH2-terminal region. Both LOX and LOXL1 are secreted into the extracellular matrix and become enzymatically active after proteolytic cleavage of the NH2-terminal propeptide region by BMP-1 [19,27]. Using a recombinant protein approach, the NH2-terminal propeptide region (codons 1 to 303) of LOXL1 was shown to play a critical role in interaction with

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**Figure 2.** Expression and purification of the four different haplotype proteins of LOXL1. Each lane contains approximately 3 μg of purified recombinant LOXL1 protein of the 141L-153G, 141L-153D, 141R-153G, or 141R-153D haplotype. Lane M contains a molecular mass standard.

**Figure 3.** Amin oxidase activity of the four different haplotype variants of LOXL1 toward elastin, type I collagen, or cadaverine. The reactions without BAPN are in gray, and the reactions with BAPN are in white. The amin oxidase activity is expressed as nM of H2O2 produced per μg of the LOXL1 variant protein. Standard deviations are indicated in each graph, and the p-values between the reactions with and without BAPN are less than 0.05 in all cases tested.
tropoelastin, directing the LOXL1 protein onto elastic fibers [28]. Thus, the presence of the R141L and G153D variations in the propeptide region raises a possibility that these missense variations may be important in directing LOXL1 onto the sites of elastogenesis by influencing its interaction with tropoelastin. Alternatively, the missense variations of R141L and G153D may affect the interaction of LOXL1 with the scaffolding proteins required for elastogenesis. For instance, fibulin-5, a microfibril-associated protein, is critical for elastic fiber organization, functioning as a scaffold element for elastogenesis [29,30]. Further, fibulin-5 colocalizes and interacts with LOXL1, through the NH2-terminal region unique to LOXL1 [8]. These findings suggest that the R141L and G153D variations may affect the interaction of LOXL1 with fibulin-5, thus leading to disorganized formation and abnormal aggregation of elastic fibers. Further molecular studies on the effects of R141L and G153D on proteolytic processing of LOXL1 or interactions with tropoelastin and the microfibril-associated proteins will be necessary for resolving these intriguing questions.

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