A Genome-Wide Association Study of Vertical Cup-Disc Ratio in a Latino Population

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PURPOSE. Vertical cup-disc ratio (VCDR) is used as a clinical assessment measure to identify and monitor glaucomatous damage to the optic nerve. Previous genetic studies conducted in European and Asian populations have identified many loci associated with VCDR. The genetic factors in other ethnic populations, such as Latino, influencing VCDR remain to be determined. Here, we describe the first genome-wide association study (GWAS) on VCDR in Latino individuals.

METHODS. We conducted this GWAS on VCDR using 4537 Latino individuals who were genotyped by using either the Illumina OmniExpress BeadChip (~730K markers) or the Illumina Hispanic/SOL BeadChip (~2.5 million markers). Study subjects were 40 years of age and older. Linear regression, adjusting for age, sex, and principal components of genetic ancestry, was conducted to assess the associations between single nucleotide polymorphisms (SNPs) and VCDR. We imputed SNPs from the 1000 Genomes Project to integrate additional SNPs not directly genotyped.

RESULTS. We replicated two previously reported SNPs that reached GWAS significance, rs1900005 and rs7916697, in the ATOH7-PBLD region, as well as identified two suggestive associations in the CDC7-TGFBR3 region on chromosome 1p22.1 and in the ZNF770-DPH6 region on chromosome 15q14. We discovered a novel SNP, rs56238729 (P = 1.22 × 10⁻¹³), in the ATOH7-PBLD region that is significantly associated with VCDR in Latino individuals. We replicated eight previously reported regions, including COL8A1, CDKN2B-CDKN2BAS, BMP2, and CHEK2 (P < 2.17 × 10⁻³).

CONCLUSIONS. Our results discovered a novel SNP that is significantly associated with VCDR in Latino individuals and confirmed previously reported loci, providing further insight into the genetic architecture of VCDR.

Keywords: GWAS, VCDR, Latino Individuals

The optic disc, situated at the back of the eye, is the location where the retinal ganglion cell axons exit the eye to form the optic nerve. Inspection of the optic disc is conducted during routine eye examinations to assess the optic nerve. In particular, the ratio of the vertical diameter of the optic cup to the vertical diameter of the optic disc, called the vertical cup-disc ratio (VCDR), is often calculated to identify and monitor possible glaucomatous damage to the optic nerve. As an important endophenotype for primary open-angle glaucoma (POAG), VCDR characterizes the extent of cupping of the optic nerve, and subsequently, the loss of retinal ganglion cell axons.

In numerous epidemiologic studies investigating the prevalence of POAG, VCDR is often used as a clinical diagnostic criterion to identify either suspects or cases of POAG.¹⁻⁵ Due to the complex etiology of POAG, identifying the factors that influence individual quantitative traits, such as VCDR, may better elucidate the biological mechanisms underlying this disease.

Previous studies have shown substantial differences in VCDR among various ethnic populations. Overall, individuals of African descent reported having the highest VCDR and individuals of European descent exhibiting the lowest, with Chinese and Hispanic individuals displaying VCDR measurements between African and European individuals.⁶⁻⁷ Furthermore, heritability studies have provided additional evidence of a genetic component for VCDR determination. These studies estimate the heritability of this trait to be 48% to 66%.⁸⁻⁹ Even more, these results suggest genetic factors have a major role in VCDR determination and specific genetic variants may influence this trait in different ethnic populations.

There have been a limited number of genome-wide association studies (GWAS) conducted to identify genetic variants associated with VCDR. A previous meta-analysis of GWAS conducted in individuals of European descent identified nine loci associated with VCDR, including CDKN2B, SIX1, SCY11, CHEK2, ATOH7-PBLD, DCLK1, BCAS3, RERE, and ARID3A.¹⁰ Additionally, a meta-analysis of European and Asian individuals identified several novel loci associated with VCDR, such as CDC7-TGFBR3, TMTC2, RPAP3, and BMP2.¹¹ Although these previous studies have identified a number of loci...
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associated with VCDR, these studies were conducted in individuals of European or Asian descent. Because specific genetic variants may be more enriched in one ethnic population compared with another, identifying genetic factors that are both common and unique to different ethnic populations will aid in elucidating racial differences in VCDR, as well as potentially uncovering the pathogenesis of various ocular diseases affecting the optic nerve. To date, there have been no reported GWAS of VCDR in Latino individuals.

Latino individuals are traditionally an underrepresented population in genetic studies, despite being the largest minority group in the United States. We hypothesize conducting a GWAS of VCDR in Latino individuals will replicate previously reported loci that are common to various ethnic populations and, potentially, identify novel loci specific to Latino individuals. Moreover, identifying genetic variants associated with VCDR in Latino individuals may help to elucidate the observed racial differences for this ocular parameter. As such, the purpose of this study was to conduct a GWAS of VCDR in Latino individuals. This study represents the first GWAS of VCDR in a Latino population.

Materials and Methods

Ethics Statement

The institutional review boards at the University of Illinois at Chicago, the University of Southern California Health Sciences Campus, and the Los Angeles Biomedical Research Institute at Harbor-University of California, Los Angeles (UCLA) approved this research and all clinical investigation was conducted according to the principles outlined in the Declaration of Helsinki.

Study Sample

This study was conducted using 4537 samples collected from the Los Angeles Latino Eye Study (LALES), a population-based epidemiologic study examining the prevalence, incidence, and impact of visual impairment and ocular disease in 6357 Latino individuals living in the city of La Puente, Los Angeles County, California, United States.12 All study subjects were 40 years or older and written informed consent was obtained from all study participants.

Measurements of VCDR

Stereoscopic optic disc photography was conducted using the Topcon TRC 50EX Retinal Camera (Topcon Corp. of America, Paramus, NJ, USA) with Ektachrome 100 film (Kodak, Rochester, NY, USA) and was evaluated using a stereoscopic viewer (Asahi viewer; Pentax, Englewood, CO, USA). VCDR measurements were determined by a board-certified ophthalmologist. We used the average VCDR between the left and right eyes as the final VCDR measurement to convert these data to a parameter. To further analyze. Duplicates, study participants with sex inconsistency between reported and genetically inferred sex, and missing VCDR measurements were further excluded. Implementation of these exclusion criteria resulted in 4537 study participants remaining for downstream analysis, including 3596 unrelated individuals used as a discovery set (stage 1), and 941 first-degree relatives (417 families) used as a replication set (stage 2, independent of stage 1). Quality control of the genotyped data for the study participants was conducted using the program PLINK (v1.90; available in the public domain at https://www.cog-genomics.org/plink2).13 Overlapping SNPs between the two genotyping chips were retained for analysis and were further omitted if the minor allele frequency (MAF) was less than 1%, the call rate was less than 95%, or the Hardy-Weinberg equilibrium P values were less than 10−6. These exclusion criteria resulted in 576,798 SNPs remaining for downstream analysis. To facilitate the imputation process, SNPs were coded on the forward strand.

Genotype Imputation

To identify additional SNPs associated with VCDR that were not directly genotyped, we performed genotype imputation using Shapeit2,14 Minimac3 (available in the public domain at http://genome.sph.umich.edu/wiki/Minimac3#Download), and the 1000 Genomes Project (1KGP) reference panels. These haplotype reference panels contain approximately 39.7 million variants, substantially increasing the number of SNPs to be tested through the use of linkage disequilibrium (LD). We used the AMR+CEU+YRI reference panels (a combination of Mexican, Colombian, Puerto Rican, CEPH, and Yoruba haplotypes) for imputation because of the unique admixture of different ancestral populations in Latino individuals. This panel has previously been shown to result in the highest genotype imputation accuracy for Latino individuals.15 Using the phased genotypes from Shapeit2, genotype imputation on the phased data was conducted using Minimac3. Imputed genotypes were coded as allelic dosages (estimated counts ranging from 0–2). Low-quality imputed SNPs (i.e., Rsq < 0.80 and MAF < 1%) were filtered out. After applying these quality control parameters, 6,844,888 imputed SNPs were retained for downstream analysis.

Statistical Analysis

We inferred principal components of genetic ancestry using the program EIGENSOFT.16 To make comparisons with reference populations of known ancestry, we included reference panels of unrelated Northern Europeans (CEU, n = 87) and West Africans (YRI, n = 88) from the 1000 Genomes Project,17 and Native Americans (n = 105).18 We retained and included the first four principal components as covariates in downstream association analyses. To assess control of population stratification, the genomic control inflation factor19 was calculated and a quantile-quantile (Q-Q) plot was generated to visualize the distribution of the test statistics. Linear regression, adjusting for age, sex, and principal components of genetic ancestry, was conducted to assess the associations between SNPs and VCDR among study participants in the discovery set using PLINK (v1.90).15 Additionally, an additive genetic effects model was assumed. To account for
relatedness among individuals in the replication set, we used a linear mixed-effects model (Proc Mixed procedure of SAS v9.4; SAS Institute, Cary, NC, USA) to test the associations between SNPs and VCDR, adjusting for age, sex, and principal components of genetic ancestry. The empirical “sandwich” estimator and compound symmetry covariance structure were used during linear mixed-effects modeling. The software EMMAX (Efficient Mixed-Model Association Expedited; available in the public domain at http://csg.sph.umich.edu/kang/emmax/download/) was used to analyze genotyped and imputed SNPs for the full study sample, the discovery and replication sets combined, using linear mixed-effects modeling to account for population stratification and relatedness, adjusting for age, sex, kinship, and principal components of genetic ancestry. Allelic dosage was used to account for genotype imputation uncertainty for imputed SNPs in EMMAX. SNPs with a \( P < 1 \times 10^{-6} \) in the discovery set were retained and analyzed in the replication set. SNPs reaching the genome-wide significance threshold (\( P < 5 \times 10^{-8} \)) were declared significant and SNPs reaching \( P < 1 \times 10^{-6} \) were declared suggestive during the full study sample analysis. The genome-wide significance threshold (\( P < 5 \times 10^{-8} \)) was retained and analyzed in the replication set. Together, the average VCDR (SD) was 0.34 (0.18; range, 0.10–0.90), with the average of the discovery and replication sets as 0.34 (0.18; range, 0.10–0.90) and 0.35 (0.19; range, 0.10–0.90), respectively.

**Genome-Wide Association Results**

The genomic control inflation factor was moderate, \( \lambda = 1.03 \). Supplementary Figure S1 displays the Q-Q plot of the observed \( P \) values versus the expected \( P \) values. As seen in the plot, the observed \( P \) values do not deviate from the null, except at the extreme tail. Both the genomic control inflation factor and the Q-Q plot indicate proper control of population stratification in this sample of Latino individuals.

Figure 1 displays a Manhattan plot of the genome-wide \( P \) values from the discovery set association analysis. The results for the top SNPs (\( P < 1 \times 10^{-8} \)) are summarized in Table 2. One SNP, rs1900005 (\( P = 4.17 \times 10^{-8} \), GRCh37/hg19 position 69,998,055), on chromosome 10q21.3 reached the GWAS significance level \( P < 5 \times 10^{-8} \). The minor allele A (MAF = 0.36) was associated with a reduction in VCDR with \( \beta (SE) = -0.13 (0.02) \). This SNP is located 6.2 kb upstream of the ATOH7 (atoh bHLH transcription factor 7) gene and 44 kb downstream of the PBLD (phenazine biosynthesis-like protein domain containing) gene. The second most significant SNP, rs7916697 (\( P = 5.44 \times 10^{-8} \), GRCh37/hg19 position 69,991,853), is located 6.2 kb upstream of rs1900005 and was borderline GWAS significant. This SNP is situated in the 5′ untranslated region (5′UTR) of ATOH7. The minor allele A of rs7916697 (MAF = 0.38) is associated with a decrease in VCDR with \( \beta (SE) = -0.15 (0.02) \). The third most significant SNP is rs16960773 (\( P = 3.15 \times 10^{-7} \), GRCh37/hg19 position 35,604,502), located on chromosome 15q12 and situated 324 kb upstream of ZNF770 (zinc finger protein 770) and 53.2 kb downstream of DPH6 (diphthamide biosynthesis 6) . The minor allele G (MAF = 0.08) is associated with a reduction in VCDR with \( \beta (SE) = -0.25 (0.04) \). The last most significant SNP is rs1192419 (\( P = 4.85 \times 10^{-7} \), GRCh37/hg19 position 92,080,059), located on chromosome 1p22.1 and positioned 88.7 kb downstream of CDC7 (cell division cycle 7) and 65.8 kb downstream of TGFBR3 (transforming growth factor, beta receptor III). The minor allele A (MAF = 0.29) of rs1192419 is associated with an increase in VCDR with \( \beta (SE) = 0.15 (0.03) \).

We then analyzed these top SNPs in the replication set using linear mixed-effects models. As displayed in Table 2, the direction of effect for the top four SNPs are consistent with the directions observed in the discovery set. The associations for these SNPs (rs1192419, rs7916697, and rs1900005) were strengthened when the full study sample was analyzed. In addition to rs1900005 remaining significant, rs7916697 also became genome-wide significant (\( P = 1.97 \times 10^{-11} \)) and rs1192419 became borderline genome-wide significant (\( P = 9.56 \times 10^{-8} \)) after analyzing the discovery and replication sets together. The SNPs rs7916697 and rs1900005 have previously been associated with VCDR.
been reported to be associated with VCDR in European and Asian individuals and the results reported in this study confirm these associations in a Latino population.

**Results From Imputed SNPs**

To interrogate additional SNPs not directly genotyped, we performed genotype imputation on the full study sample. After retaining SNPs of high quality ($R^2 \geq 0.80$), no additional genomic regions reached GWAS significance. Within the **ATOH7-PBLD** region, numerous imputed SNPs reached genome-wide significance, including several SNPs more significant than those directly genotyped (Supplementary Table S1). The most significant SNP in this region is rs56238729, located 9.8 kb downstream of rs7916697. This SNP represents a novel association with VCDR. The regional SNP association plot for the **ATOH7-PBLD** region is presented in Figure 2A. Genotyped SNPs are plotted as squares, and imputed SNPs as circles.

**Conditional Analysis**

To determine whether additional SNPs contribute to the VCDR association, we conducted conditional analysis in the **ATOH7-PBLD** region. As shown in Figure 2B, conditioning on the most significant SNP, rs56238729, by including this SNP as a covariate into linear regression models resulted in all immediate SNP associations to be reduced toward the null with no other SNP remaining significant. These data suggest rs56238729 is the leading SNP of the VCDR association and further nullifies the associations of surrounding SNPs in the **ATOH7-PBLD** region, including SNPs that have previously been reported.

**Analysis of Previously Reported Loci for VCDR**

We investigated previously reported VCDR loci identified in populations of European and Asian descent to determine whether these associations are consistent in a Latino population. Table 3 summarizes these results. Of the previously reported SNPs, 25 SNPs exhibited high imputation quality ($R^2 \geq 0.80$) and when further analyzed, we observed 32% (8/25) having a $P < 0.05$ in our Latino population. Furthermore, we observed consistent directions of associations with all previous SNPs, except one (rs301801). To account for multiple testing and to avoid penalties associated with a traditional Bonferroni correction, we used *simpleM* to calculate the effective number of independent tests. This method identified 23 SNPs with $P < 1 \times 10^{-6}$ in the discovery set are included in the table and were analyzed in the replication set. Gene name is in boldface if the SNP is located inside the gene. Single nucleotide polymorphism positions are according to GRCh37/hg19. Chr, chromosome; A1/A2, allele 1/allele 2.
FIGURE 2. Regional SNP association plots for the \textit{ATOH7-PBLD} region. (A) The most significant directly genotyped SNP, rs1900005, using the entire study sample is plotted in purple. Genotyped and imputed SNPs (Rsq ≥ 0.80) are plotted as squares and circles, respectively. Genes are shown below the SNPs and the arrows indicate the strand orientation for each gene. The color coding in the plot represents the level of LD with rs1900005. After imputation, rs56238729 is the most significant SNP in the \textit{ATOH7-PBLD} region. (B) Regional SNP association plot conditioning on rs56238729, the most significant SNP in the \textit{ATOH7-PBLD} region.
# Table 3. Comparison With Previously Reported SNPs Associated With VCDR in Latino Individuals

| SNP ID   | Chr | Position | Genes Nearby | Effect | Allele | Freq | Reference | Consistency of Direction | A1/A2 | A1F1 | P Value | Imputed | Most Significant Hits ≤ 100 kb |
|----------|-----|----------|--------------|--------|--------|------|-----------|--------------------------|-------|-----|---------|---------|-------------------------------|
| rs301801 | 1   | 8495945  | RERE         |        | C/T    | 0.35 | 0.01      | 11                       | T/C   | 0.79 | 0.01    | 6.67 × 10⁻¹ | Y                     |
| rs12025126 | 1    | 8759554  | RERE         |        | C      | 0.28 | -0.01     | 10                       | T/C   | 0.59 | 0.00    | 8.85 × 10⁻¹ | Y                     |
| rs4658101 | 1    | 92077409 | CDCC7-FGFR3  |        | A/G    | 0.18 | 0.02      | 11                       | A/G   | 0.29 | 0.12    | 4.58 × 10⁻⁷ | Y                     |
| rs6253425 | 3    | 99131755 | COL6A1      |        | A/C    | 0.13 | 0.02      | 11                       | C/A   | 0.77 | -0.04   | 8.67 × 10⁻¹ | Y                     |
| rs17658299 | 5   | 172191052| DUSP1       |        | C/T    | 0.05 | -0.02     | 11                       | T/C   | 0.99 | 0.05    | 5.88 × 10⁻¹ | Y                     |
| rs17756172 | 6   | 625071   | EXOC2       |        | G/A    | 0.18 | 0.01      | 11                       | G/A   | 0.82 | -0.05   | 6.73 × 10⁻² | Y                     |
| rs680153 | 6    | 122389555| HSF2        |        | G/T    | 0.36 | -0.01     | 11                       | T/G   | 0.74 | 0.05    | 5.94 × 10⁻² | Y                     |
| rs10615192 | 9   | 22003567 | CDKN2B      |        | G      | 0.45 | -0.01     | 3            | G/A   | 0.19 | -0.09   | 9.86 × 10⁻⁴ | N                     |
| rs7865618 | 9    | 22031005 | CDKN2B      |        | G/A    | 0.43 | -0.01     | 11                       | G/A   | 0.19 | -0.09   | 7.25 × 10⁻⁴ | Y                     |
| rs1900005 | 10   | 69998055 | ATOH7       |        | A/C    | 0.23 | -0.02     | 11                       | A/C   | 0.36 | -0.15   | 6.41 × 10⁻¹²| N                    |
| rs1900004 | 10   | 70000881 | ATOH7-PBDL  |        | T      | 0.22 | 0.01      | 10                       | T/C   | 0.64 | 0.15    | 1.50 × 10⁻² | Y                     |
| rs7072574 | 10   | 98056306 | PLCE1       |        | A/G    | 0.33 | 0.01      | 11                       | G/A   | 0.72 | -0.02   | 4.87 × 10⁻¹ | Y                     |
| rs17146964 | 11   | 65249145 | SCYL1       |        | G      | 0.21 | -0.01     | 10                       | G/A   | 0.86 | 0.05    | 2.89 × 10⁻¹ | N                     |
| rs13146 | 11   | 65337251 | SSCA1       |        | T/A    | 0.19 | -0.01     | 11                       | A/T   | 0.86 | 0.04    | 2.30 × 10⁻¹ | Y                     |
| rs936099 | 11   | 130280725| ADAMTS8     |        | C/A    | 0.42 | 0.01      | 11                       | C/A   | 0.29 | -0.01   | 8.02 × 10⁻² | Y                     |
| rs11168187 | 12  | 48014011 | RPAP3       |        | G/A    | 0.16 | 0.01      | 11                       | G/A   | 0.87 | 0.01    | 8.12 × 10⁻¹ | N                     |
| rs10862688 | 12  | 83922012 | TMC2        |        | G/A    | 0.45 | 0.01      | 11                       | G/A   | 0.75 | 0.00    | 9.19 × 10⁻¹ | N                     |
| rs192630 | 13   | 36562617 | DCLK1       |        | C      | 0.24 | 0.01      | 10                       | T/C   | 0.69 | -0.02   | 3.72 × 10⁻² | Y                     |
| rs4901977 | 14   | 60789716 | SIX1-SIX6   |        | T/C    | 0.31 | 0.01      | 11                       | C/T   | 0.75 | -0.05   | 5.27 × 10⁻² | N                     |
| rs10483727 | 14  | 61072875 | SIX1        |        | T      | 0.4   | 0.01      | 11                       | T/C   | 0.35 | 0.03    | 1.33 × 10⁻¹ | N                     |
| rs1345467 | 16   | 51482321 | SALL1       |        | G/A    | 0.27 | 0.01      | 10                       | G/A   | 0.17 | 0.06    | 2.63 × 10⁻² | Y                     |
| rs8006952 | 17   | 59286446 | BCA5        |        | G/C    | 0.24 | -0.01     | 10                       | G/C   | 0.19 | -0.03   | 3.51 × 10⁻¹ | Y                     |
| rs1251928 | 19   | 9503840 | ADRD3A      |        | G      | 0.13 | -0.02     | 10                       | G/T   | -     | -       | -       | NA                 |
| rs6054571 | 20   | 6578596 | BMP2        |        | T/C    | 0.42 | -0.01     | 11                       | T/C   | 0.46 | 0.06    | 1.02 × 10⁻² | Y                     |
| rs1547014 | 22   | 29100771 | CHEK2       |        | T      | 0.29 | -0.01     | 10, 11 (T/C, Freq = 0.3, β = -0.013) | T/C   | 0.35 | -0.05   | 3.43 × 10⁻² | N                     |
| rs5756813 | 22   | 38175477 | CARD10      |        | G/T    | 0.39 | 0.01      | 11                       | G/T   | 0.41 | 0.01    | 5.74 × 10⁻¹ | Y                     |

Additional allele, frequency, and effect size information is given in parentheses for SNPs with multiple references. The frequency of allele 1 is given for our Latino sample and is modeled as the effect allele. To correct for multiple testing for correlated SNPs, the program simpleM was used, and identified 23 independent tests, resulting in a Bonferroni correction P value of 0.05/23 = 2.17 × 10⁻³. Shown in bold are P values meeting this threshold. Using directly genotyped SNPs, the most significant hit within ≤ 100 kb of previously reported SNPs are listed (P values < 2.17 × 10⁻³ are italicized). All SNPs, except rs2159128, had an Rsq ≥ 0.80 from Minimac3. Most SNPs exhibited consistent direction of effect, except for rs301801. Single nucleotide polymorphism positions are according to GRCh37/hg19. Freq. frequency.
independent tests, resulting in a Bonferroni correction significance level of $P = 2.17 \times 10^{-3}$. We were able to replicate five index SNPs in three regions (**CD7-C2**,**CD32B**,**CDKN2AS**,**ATOH7-PBLD**) after multiple testing correction (shown in boldface in Table 3). To replicate genomic regions associated with VCDR, we extracted the most significant directly genotyped SNP within ±100 kb of the previously reported SNPs. This approach identified an additional five SNPs in five regions surviving the multiple testing correction, replicating the associations between VCDR and the **COL8A1**, **HSF2**, **RPAP3**, **TMTC2**, and **BMP2** regions.

**Pathway Analysis**

To determine whether canonical pathways were enriched with genes associated with VCDR, we performed pathway analysis using IPA. After adjusting for multiple testing, the only pathway significantly associated with VCDR was the “pathogenesis of multiple sclerosis (MS)” pathway ($P = 7.41 \times 10^{-3}$). Of the nine genes comprising this pathway, five genes from our dataset overlapped with these genes, including **COL13**, **COL14**, **CXCL9**, **CXCL10**, and **CXCL11**.

**DISCUSSION**

This study represents the first GWAS conducted on VCDR in Latino individuals. We identified two genome-wide significant SNPs, rs1900005 and rs7916697, associated with VCDR, confirming the involvement of the **ATOH7-PBLD** region. We also identified suggestive associations in the **CD7-C2**,**CD32B**,**ZNF770-DPH6** regions. We discovered a novel SNP, rs65238729, in the **ATOH7-PBLD** region to be significantly associated with VCDR in Latino individuals after genotype imputation from the 1KGP reference panels. Moreover, we were able to replicate genomic regions previously associated with VCDR, including **COL8A1**, **HSF2**, **RPAP3**, **TMTC2**, and **BMP2**. Results from our pathway analysis identified one canonical pathway associated with VCDR. The most significant SNPs in our study reside in the **ATOH7-PBLD** region. Previous GWAS studies, including Ramdas et al. and Springelkamp et al., identified many SNPs in the **ATOH7-PBLD** region to be associated with VCDR. In addition to VCDR, earlier GWAS have also associated this region with optic disc area, cup area, and POAG. Both rs1900005 and rs7916697, the most and second most significant SNPs in our study, respectively, have previously been associated with VCDR. In particular, rs7916697 resides in the 5’UTR region of **ATOH7**, a single exon gene that plays a role in retinal ganglion cell development. Moreover, rs7916697 has been associated with a reduction in optic disc area and was identified to have a significant interactive effect with rs1063192 in an Afro-Caribbean population, resulting in a reduction in POAG risk. Taken together, our results are consistent with previous studies of these SNPs being strongly associated with glaucoma-related quantitative traits and may have a biological role in the pathogenesis of POAG.

The third and fourth most significant SNPs indicate suggestive associations in the **CD7-C2**,**CD32B** and **ZNF770-DPH6** regions, respectively. Similar to the previous region, the **CD7-C2**,**CD32B** region has been reported to be associated with VCDR, optic disc area, and POAG. Moreover, expression of both **CD7** and **TGFBR3** have been observed in numerous human ocular tissues, most notably the optic disc and optic nerve. The SNP rs1192419 has specifically been associated with VCDR and an increase in disc area. Additionally, common variants within the genomic region on chromosome 15q14, in which **ZNF770-DPH6** resides, has previously been associated with refractive error and myopia.

This study independently confirms the associations of rs1900005 and rs7916697 in the **ATOH7-PBLD** region with VCDR in a sample of Latino individuals and suggests additional loci in the **CD7-C2**,**CD32B** and **ZNF770-DPH6** regions. Although ophthalmologists routinely assess the VCDR to diagnose and monitor the progression of POAG, the cupping of the optic nerve may not solely be a result of glaucoma and may result from other conditions, such as optic neuritis. Our pathway results implicate an association between the pathogenesis of MS and VCDR. Multiple sclerosis is a demyelinating disease of the central nervous system that commonly affects vision. Patients with MS were found to have a higher VCDR compared with healthy controls, suggesting enlarging of the optic cup due to the thinning of the retinal nerve fiber layer may be explained by the predilection of the disease to affect the optic nerves. Moreover, several of the genes included in this pathway code for chemokines that were shown to be at higher concentrations in the aqueous humor of glaucomatous eyes compared with cataract controls. Collectively, our pathway results suggest the biological mechanisms influencing VCDR and MS may share common genetic constituents.

This is the first GWAS of VCDR in Latino individuals and several limitations exist. First, the Latino population is historically understudied. As far as we know, our dataset is currently the only Latino genetic dataset with ophthalmic phenotypes. Furthermore, the three-way admixture of Latino individuals makes it even more challenging in genetics research (Supplementary Fig. S2). Using STRUCTURE (available in the public domain at http://pritchardlab.stanford.edu/structure.html), we estimated that our Latino subjects on average had 3%, 53%, and 44% African, European, and Native American ancestries, respectively (Supplementary Fig. S3). We performed a fixed-effects meta-analysis for the discovery and replication sets on the top genotyped SNPs using METAL (available in the public domain at http://csg.sph.umich.edu/abecasis/Metal/download/) and obtained similar results as EMMAX (Supplementary Table S2). These results suggest population stratification and genetic relatedness were properly controlled for in our analysis, despite Latino individuals being a three-way admixed population. However, we emphasize the need for replication in an independent Latino cohort. Second, we did not conduct secondary analyses adjusting for disc area, an ocular parameter known to be correlated with VCDR. Unfortunately, at the time of the VCDR data collection, disc area was not collected. Given a previously reported reduction in significance for VCDR-associated variants after adjusting for disc area, a similar trend may be observed for our results.

In conclusion, in the first GWAS of VCDR in Latino individuals, we discovered a novel SNP that is significantly associated with VCDR in Latino individuals. In addition, two SNPs reached genome-wide significance, replicating associations in the **ATOH7-PBLD** region. We were also able to replicate associations with several previously reported genomic regions for VCDR in this population. Our pathway results identified a novel association between pathogenesis of MS and VCDR, suggesting potential shared genetic factors influencing both VCDR and MS. The findings from this study suggest that many genetic factors influencing VCDR are shared among ethnic populations.

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