A genome-wide association study for myopia and refractive error identifies a susceptibility locus at 15q25

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Myopia and hyperopia are at opposite ends of the continuum of refraction, the measure of the eye’s ability to focus light, which is an important cause of visual impairment (when aberrant) and is a highly heritable trait. We conducted a genome-wide association study for refractive error in 4,270 individuals from the TwinsUK cohort. We identified SNPs on 15q25 associated with refractive error (rs8027411, $P = 7.91 \times 10^{-8}$). We replicated this association in six adult cohorts of European ancestry with a combined 13,414 individuals (combined $P = 2.07 \times 10^{-9}$). This locus overlaps the transcription initiation site of RASGRF1, which is highly expressed in neurons and retina and has previously been implicated in retinal function and memory consolidation. Rasgrf1−/− mice show a heavier average crystalline lens ($P = 0.001$). The identification of a susceptibility locus for refractive error on 15q25 will be important in characterizing the molecular mechanism responsible for the most common cause of visual impairment.

Myopia is the most common eye disorder. It is part of the spectrum of refractive error, measured as spherical equivalent, which is the quantitative strength of spectacle lens required to enable images to be focused on the retina of the eye. Clinically relevant myopia or nearsightedness (more severe than −1 diopter) affects 33% of the adult population in the United States1. Its prevalence is increasing in more recent birth cohorts across both developed and less developed countries, particularly in east Asia, where over 80% of Chinese young adults in Singapore and Taiwan are now affected2,3. Apart from the huge economic cost (estimated at $5.5 billion annually spent by adults in the United States alone⁴), severe refractive error is a major cause of blindness⁵–⁷, particularly in the working-age population⁸. Vision loss is greatest in the 2%–3% of the population with ocular structural changes associated with high myopia (termed ‘pathological myopia’). The World Health Organization has identified the prevention and treatment of myopia as a priority within the

Figure 1 Association scatter plot for SNPs in the RASGRF1 promoter region in the TwinsUK discovery cohort. Negative logarithms of the $P$ values for the association of each SNP with spherical equivalence are plotted. The lead SNP is plotted in blue, with the meta-analysis $P$ value for that SNP indicated. Genotyped SNPs are plotted as diamonds and imputed SNPs as circles, with the color indicating the degree of pairwise LD between the lead and neighboring SNPs. Red indicates strong pairwise LD, with $r^2 \geq 0.8$; orange indicates moderate LD, with $0.5 < r^2 < 0.8$ yellow indicates weak LD, with $0.2 < r^2 < 0.5$; and white indicates no LD, with $r^2 < 0.2$. The associated SNPs lie in a 120-kb region of high LD, delineated by two local high-recombination spots at approximately 77,150 kb and 77,270 kb. The recombination rates are shown as light blue line and the locations of the genes with respect to these SNPs are shown as dark green arrows.

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current global initiative to eliminate avoidable blindness7. There are currently no practical therapies to progress the elimination of myopia, even though the condition is easily diagnosed in affected children and adolescents.

The mechanism for the development of myopia is not fully understood. Although environment (in particular, education, urbanization, outdoor activity and close work) clearly plays a role in myopia9,10, numerous cross-sectional studies suggest that genetic heritability might be as high as 80%6,11. No individual genes have been consistently associated with myopia to date, although several candidate loci have been identified through linkage studies.

In order to further explore potential genetic influences of refractive error development in the general population, we conducted a genome-wide association study (GWAS) using the British TwinsUK adult twin registry cohort (see Online Methods for a detailed description of the population and clinical characteristics).

We analyzed a set of 4,270 individuals with complete information on a panel of 2.2 million SNPs, of which 286,481 directly genotyped in all individuals. All of these SNPs passed stringent quality control criteria (Hardy-Weinberg equilibrium P > 0.001, minor allele frequency of at least 0.04 and a genotyping success rate for the SNP of at least 95%). Homogeneity of the population was confirmed by comparison of sample set similarities with each other and with three reference panels from the HapMap Phase 2 data (Supplementary Fig. 1).

The genomic inflation factor (λ) was 1.04 (Supplementary Fig. 2). Multiple SNPs, both directly genotyped and with imputed values, showed genome-wide significance. Association was maximal for rs8027411 (P = 7.91 × 10⁻⁸), although association was consistently high across a region of approximately 120 kb on chromosome 15 (Fig. 1 and Table 1). This association was consistent with an additive model, and we found no evidence of dominance for any of these SNPs. In the TwinsUK cohort, each additional copy of the susceptibility allele for the top SNP lowered the refraction by an average of 0.35 diopters (Supplementary Table 1). Individuals carrying a genotype homozygous for the T allele on the rs8027411 locus were significantly more likely than those homozygous for the non-susceptibility allele to have myopia than to have no refractive error (odds ratio (OR) = 1.16, 95% CI 1.02–1.28, P = 0.02; Fig. 2, see Online Methods for definitions).

We performed a replication study in 13,414 adults from six independent population-based cohorts of European ancestry: four Dutch cohorts (three within the Rotterdam Study and the Erasmus Rucphen Family Study (ERF), a family-based study in a genetically isolated population in the southwest of The Netherlands), one British cohort (1958 British Birth Cohort) and one Australian cohort (Australian Twin Study). The replication analysis was performed using both observed and imputed genotypes. We observed an association for SNPs at chromosome 15q25 in the Dutch and British replication studies. Although the effect size was higher in the discovery panel (Fig. 3), the effect sizes and directions of the effect were consistent among all the cohorts, with the exception of the Australian cohort. The strongest evidence of association in the meta-analysis was found for rs939658 and rs8027411 (combined P = 1.85 × 10⁻⁹ and P = 2.07 × 10⁻⁹, respectively), with each copy of the myopia susceptibility allele (G and T for the two SNPs, respectively) altering the spherical equivalent by 0.15 diopters (s.e.m. = 0.026) (Table 1). It is of note that the Australian twin group was relatively underpowered (1,086 individuals from 525 families) and substantially younger than the other cohorts. It has been shown that there is evidence of a rapidly increasing prevalence of myopia and, therefore, a cohort effect is possible12.

All of the identified SNPs associated with the spherical equivalent measures were in high linkage disequilibrium (LD) with one another (Supplementary Fig. 3). This associated region spans about 120 kb on chromosome 15q25 and overlaps the transcription initiation site of RASGRF1 (encoding Ras protein-specific guanine nucleotide-releasing factor 1). This is a large gene spanning 130 kb with 28 known exons and multiple alternative mRNA transcripts. Possible gains or losses in transcription factor–binding sites, because of the DNA polymorphisms within the region of association, are predicted in silico (Supplementary Table 2). The relevance of individual SNPs that potentially influence transcription factor binding and expression of this gene is difficult to assess due to the high LD across the region.

RASGRF1 encodes a protein that is expressed at high levels in neurons and retina in mice13 and that activates Ras14. Our reverse transcription experiments identified high levels of expression of this protein in human retinas (data not shown). In mice, the protein is heavily imprinted in the early developmental phases15,16, but this imprinting fades with time16. RASGRF1 expression is upregulated by stimulation of muscarinic receptors17 and retinoic acid18. Muscarinic

| Table 1 Selection of SNPs associated with spherical equivalent measures in six cohorts of European ancestry at 15q25 |
|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|
| SNP | Discovery Cohort | Replication Cohorts | Meta-analysis |
|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|
|       | Allele | Effect | s.e.m. | P | Effect | s.e.m. | P | Effect | s.e.m. | P | Effect | s.e.m. | P | Effect | s.e.m. | P |
|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|
| rs17175798 | C | -0.13 | 0.05 | 0.01 | 0.02 | 0.08 | 0.9 | 0.30 | 0.08 | 0.00 | 0.16 | 0.07 | 0.03 | 0.19 | 0.04 | 0.06 | 0.04 | 0.03 | 0.15 | 0.026 | 1.99 | 10⁻⁹ |
| rs8027411 | T | -0.12 | 0.05 | 0.01 | 0.02 | 0.08 | 0.00 | 0.16 | 0.07 | 0.03 | 0.19 | 0.04 | 0.06 | 0.04 | 0.03 | 0.15 | 0.026 | 2.07 | 10⁻⁹ |
| rs8033963 | T | -0.12 | 0.05 | 0.01 | 0.02 | 0.08 | 0.00 | 0.16 | 0.07 | 0.03 | 0.19 | 0.04 | 0.06 | 0.04 | 0.03 | 0.15 | 0.026 | 1.86 | 10⁻⁹ |
| rs939658 | G | -0.12 | 0.05 | 0.01 | 0.02 | 0.08 | 0.00 | 0.16 | 0.07 | 0.03 | 0.19 | 0.04 | 0.06 | 0.04 | 0.03 | 0.15 | 0.026 | 1.85 | 10⁻⁹ |
| rs939661 | A | -0.11 | 0.05 | 0.01 | 0.02 | 0.08 | 0.00 | 0.16 | 0.07 | 0.03 | 0.19 | 0.04 | 0.06 | 0.04 | 0.03 | 0.15 | 0.026 | 1.16 | 10⁻⁹ |

TwinsUK, the Twin cohort recruited in the UK that served as a discovery GWAS (N = 4,270); RS-I, the first Rotterdam Study cohort (N = 5,328); RS-II, the second Rotterdam Study cohort (N = 2,157); RS-III, the third Rotterdam Study cohort (N = 2,082); ERF, the Erasmus Rucphen Family (N = 2,032); 1958BC, the 1958 British Birth Cohort (N = 1,000); Aus, the Australian Twin Study (N = 1,086). The combined column includes results from the meta-analysis of the discovery and replication cohorts. Effect, β coefficient of linear regression. The effect sizes denote changes in diopters per each additional copy of the reference allele.
inhibitors such as atropine inhibit the development of myopia in animal models and human intervention studies\textsuperscript{19,20}, and choroidal retinoic acid synthesis is altered in animal models of myopia\textsuperscript{23}. \textit{RASGRF1} knockdown mouse models have normal brain structure; however, they perform worse than wild-type mice in exercises involving long-term memory\textsuperscript{22}.

Lack of expression of \textit{RASGRF1} in knockdown models results in a morphologically normal retina but causes pronounced deficiencies in photoreception and visual sensory processes without any other obvious sensory impairments\textsuperscript{23}. Furthermore, lack of expression of \textit{RASGRF1} causes changes in the downstream expression of many genes\textsuperscript{23}, including some genes whose mutations have been reported to cause severe Mendelian vision disorders. Thus, \textit{RASGRF1} appears to be involved in the maintenance of normal function of the retina and possibly in the signaling pathways determining myopia.

We further studied the effect of \textit{Rasgrf1} on mouse eye parameters. For this purpose, 19 wild-type mice were compared to 16 \textit{Rasgrf1} knockdown mice of the same age group (between 3- and 4-months-old, when eye growth has stabilized). We found that the average crystalline lens weight was significantly heavier in the knockdown mice (\(P = 0.001\)). This was independent of the weight of the whole eye, as the ratio of crystalline weight to the total eye weight remained significantly higher in the knockdown mice (\(P = 0.003\); \textit{Supplementary Table 3}). Axial length was not significantly different between the groups; however, it is known that the measurement of this trait in mice is technically difficult and inaccurate\textsuperscript{24}.

In humans, refractive error is mostly due to axial length variations, but refractive error is highly correlated with lens thickness in mice\textsuperscript{25}.

In an accompanying study reported in this issue, C. Klaver and colleagues report a genome-wide association study for refractive error in a Dutch population-based cohort\textsuperscript{26}. Researchers from that study identified a susceptibility locus at genome-wide significant levels on chromosome 15q14. \textit{GJD2}, also known as \textit{Connexin 36}, is the gene nearest to the associated chromosome 15q14 locus. In our study, the chromosome 15q14 locus showed suggestive association (\(P = 6 \times 10^{-7}\) in the TwinsUK Cohort GWAS), but this locus was not taken forward for inclusion in our replication study (\textit{Supplementary Fig. 4}).

Both \textit{RASGRF1} and \textit{GJD2} are involved in synaptic transmission and play important roles in the transmission and processing of visual signals\textsuperscript{23,27}. Animal models of myopia demonstrate that the signal that continues or halts eye growth originates within the retina, and these signals are transmitted through the choroid to stimulate or inhibit scleral remodeling\textsuperscript{28}.

We report, to our knowledge, the first genome-wide association for common myopia and refractive error in multiple European populations, identifying an associated locus upstream of the transcription site of \textit{RASGRF1}. \textit{Rasgrf1} knockdown mice have larger crystalline lenses than wild-type mice, which are likely to result in increased refractive power. \textit{RASGRF1} is involved in learning, visual processing and muscarinic signaling pathways, all of which have been associated or implicated with myopia. We believe that identification of this susceptibility locus and the suggestive evidence for involvement of the candidate gene \textit{RASGRF1} will greatly improve our understanding of the pathogenesis of myopia.

\textbf{METHODS}

Methods and any associated references are available in the online version of the paper at http://www.nature.com/naturegenetics/.

\textit{Note: Supplementary information is available on the Nature Genetics website.}

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Outdoor activity reduces the prevalence of myopia in children.

The development of the refractive status and ocular growth in children is influenced by various factors, including lifestyle factors such as outdoor activity. This is supported by studies showing a decrease in myopia prevalence among children who engage in outdoor activities compared to those who do not.

Additionally, the article mentions the importance of the 1958 British Birth Cohort study, which was funded for biomedical assessment (Medical Research Council). This study has contributed significantly to our understanding of eye development and the prevalence of eye diseases over time.

Competing financial interests

The authors declare no competing financial interests.

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Genotyping was carried out using three genotyping platforms from Illumina: the HumanHap300k Duo for part of the UK Twin Cohort and the HumanHap610-Quad array for the rest of the UK Twin Cohort. The RS-I, RS-II and RS-III studies used the Infinium II HumanHap550 chip v3.0 array to conduct the genotyping, with the exception of 133 of 2,516 subjects in RS-II who were genotyped using the HumanHap610-Quad array. The ERF cohort was genotyped with the Infinium HumanHap300-2 318K array. The Australian cohorts were genotyped on the HumanHap610-Quad array, with part of the sample typed alongside the UK Twin Cohort and the remainder of the sample typed as part of a separate contract with deCODE Genetics. Illumina’s Human1M-Duo chip was used for genotyping of the 1958 British Birth Cohort. Imputation was calculated with reference to HapMap release 22 CEU population data using IMPUTE (see URLs) version 2 for the UK Twin Cohort and the 1958 Birth Cohort. MACH (see URLs) was used for imputation in the Dutch and Australian cohorts.

Statistical analyses. Association analysis was performed using MERLIN in the UK and Australian Twin Cohorts and using PLINK (see URLs) in the Rotterdam cohorts. Ancestry for these individuals was determined initially through self-reporting and was verified through principal component decomposition of their ancestry with and without comparison with HapMap phase 2 standard populations. Meta-analysis was carried out using METAL (see URLs).

Laboratory animal handling and eye measurements. In an effort to minimize the number of animals used and their suffering, all procedures with laboratory mice were carried out in accordance with European Union guidelines (directives 86/609/EEC and 2003/65/EC), current Spanish legislation (RD 1201/2005. BOE 252/34367-91, 2005) and the US National Institutes of Health guidelines for the use and care of animals in research. Rasgrf1-null mice were obtained as described previously29. Mice were bred in a pathogen-free facility with water and food ad libitum.

For eye measuring, 16 3–4-month-old Rasgrf1-null mice and 19 wild-type control mice29 were anesthetized with chloral hydrate (5% wt/vol) in saline, administered intraperitoneally, at 10 ml/g of body weight) and killed by cervical dislocation. After measuring the body weight of each mouse in a precision balance (Sartorius), its eyes were immediately extracted using 7-inch curved tweezers (Lawton), eliminating all rests of the optical nerve, and all vessels and surrounding tissues were cut away. Eye weight was immediately measured in an AX-120 Analytical Weight Balance (Cobos Precision, S.L.). The precision of the balance was 0.001 g. After being weighed, the eye balls were positioned on a glass slide under a microscope (Olympus Provis AX70 photomicroscope, Olympus Optical Co., Ltd.). The coronal and longitudinal axes were captured using an Olympus DP70 digital camera and measured at x40 magnification using Olympus’ image capture software DP controller, version 1.2.1.108 (Olympus Optical Co., Ltd.).

The lens from each eye was then manually isolated using 7-inch tweezers (Lawton) under a binocular lens (Zeiss, Stemi 2000-C, Carl Zeiss) and measured in an M-220 analytical balance (precision = 0.1 mg) (Denver Instruments GmbH).
