Genetic Deletion of the Adenosine A2A Receptor Confers Postnatal Development of Relative Myopia in Mice

Xiangtian Zhou, Qinzhu Huang, Jianbong An, Runxia Lu, Xiaoyi Qin, Liqin Jiang, Yuan Li, Jianhua Wang, Jiangfan Chen, and Jia Qu

PURPOSE. To critically evaluate whether the adenosine A2A receptor (A2AR) plays a role in postnatal refractive development in mice.

METHODS. Custom-built biometric systems specifically designed for mice were used to assess the development of relative myopia by examining refraction and biometrics in A2AR knockout (KO) mice and wild-type (WT) littermates between postnatal days (P)28 and P56. Ocular dimensions were measured by customized optical coherence tomography (OCT), refractive state by eccentric infrared photorefraction (EIR), and corneal radius of curvature by modified keratometry. Scleral collagen diameter and density were examined by electron microscopy on P35. The effect of A2AR activation on collagen mRNA expression and on soluble collagen production was examined in cultured human scleral fibroblasts by real-time RT-PCR and a collagen assay kit.

RESULTS. Compared with WT littermates, the A2AR KO mice displayed relative myopia (average difference, 5.1 D between P28 and P35) and associated increases in VC depth and axial length from P28 to P56. Furthermore, the myopic shift in A2AR KO mice was associated with ultrastructural changes in the sclera: Electron microscopy revealed denser collagen fibrils with reduced diameter in A2AR KO compared with WT. Last, A2AR activation induced expression of mRNAs for collagens I, III, and V and increased production of soluble collagen in cultured human scleral fibroblasts.

CONCLUSIONS. Genetic deletion of the A2AR promotes development of relative myopia with increased axial length and altered scleral collagen fiber structure during postnatal development in mice. Thus, the A2AR may be important in normal refractive development in mice. We hypothesized that the adenosine A2A receptor (A2AR) plays an important role in postnatal refractive development in mice by controlling collagen synthesis in scleral fibroblasts. The extracellular adenosine level in mammalian retina is regulated by light/dark conditions, an important component of the visual signal that contributes to eye growth control and possibly to development of myopia. A2ARs are expressed in ocular tissue, including in the ciliary processes, retina, retinal pigment epithelium, and scleral fibroblasts. Of interest, A2AR activity can modulate collagen synthesis and extracellular matrix production in various cell types and tissues. For example, the A2AR agonist CGS21680 dose dependently increases collagen production in cultured human hepatic stellate cells and promotes tissue repair, wound healing, and matrix production in skin in vivo. Conversely, genetic deletion or pharmacologic blockade of A2ARs attenuates the fibrogenic process in liver and skin. Together, these studies raise the interesting possibility that A2AR activity influences the development of myopia by modulating collagen synthesis in the sclera.

In this study, we used A2AR KO mice and custom-built biometric systems specifically designed for mice to critically evaluate the role of A2AR in development of relative myopia at refractive and biometric levels, and we used electron microscopy to examine the ultrastructure. We detected a greater myopic shift, increased vitreous chamber depth and axial...
length, and reduced scleral collagen fibril diameters in A2AR KO mice than in their wild-type littermates. In addition, we found that A2AR activation increases expression of mRNAs for collagens I, III, and V and soluble collagen production in cultured human fibroblasts. Together, these results provide the first evidence that the A2AR plays a role in controlling postnatal development of myopia in mice.

MATERIALS AND METHODS

Animals

The treatment and care of animals was conducted according to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research, and the protocol for handling animals was approved by the Animal Care and Ethics Committee at Wenzhou Medical College (Wenzhou, China). A2AR KO mice were generated by Chen et al., originally in the mixed C57BL/6 background and more recently in a congenic C57BL/6 background which significantly reduces the potential confounding effect of genetic background. In this study, heterozygous female mice were mated with heterozygous males so that both A2AR KO and WT littermates were generated from the same breeding pairs. The genotypes of the mice were determined by PCR analysis of tail DNA, as described previously.

Biometric Measurements

Biometric measurements, including corneal radius of curvature, refractive state, and ocular dimensions, were taken once per week from postnatal day (P)28 to P56. These measurements were performed by a research optometrist who was blind to the genotypes of the mice.

Keratometry. Corneal curvature was measured with a keratometer (OM-4; Topcon, Tokyo, Japan) which was modified by mounting a +20.0-D aspherical lens. A group of stainless-steel ball bearings with diameters ranging from 2.0 to 3.98 mm were used for calibration. The corneal radius of curvature measured in the mice was then deduced from the readings on the balls with known radii. Each eye was measured three times to obtain a mean value.

Refraction and Pupil Size. Refractive state and pupil size were measured in a darkened room with an eccentric infrared photorefractor (EIR). Briefly, the mouse was gently restrained by grabbing its tail and positioning it on a small stage in front of the photoretinoscope. The position of the stage was swiftly adjusted until a clear first Purkinje image was seen in the center of the pupil, meaning an on-axis measurement. The data were then recorded by using software designed by Schaeffel et al. Measurements were repeated at least three times for each eye.

Ocular Dimensions. Ocular dimensions, including anterior chamber (AC) depth, lens thickness, vitreous chamber (VC) depth, and axial length (AL), were measured by real-time optical coherence tomography (a custom-built OCT). AC depth was defined as the distance from the posterior surface of the cornea to the anterior surface of the lens. AL was defined as the distance between the anterior surface of the cornea and the vitreous-retina interface. The configuration of the OCT was described in detail elsewhere, as were the principle and the measurement process. After being anesthetized, each mouse was placed in a cylindrical holder mounted on a positioning stage in front of the optical scanning probe. A video viewing system for observing the eye was used to monitor orientation and positioning. In addition, the optical axis of the mouse eye was aligned with the axis of the probe by detecting the specular reflex on the corneal apex and on the back apex of the lens in two-dimensional OCT images. Ocular dimensions were determined by moving the focal plane with the stepper motor and recording the distance traveled between two ocular interfaces. The recorded optical path length was then converted into a geometric path length by using the appropriate refractive indexes for each component of the eye. Each eye was scanned three times.

Development of Myopia in A2AR Deficient Mice

Mice were killed by an overdose of pentobarbitone sodium at P35. Three mice from each genotype group were evaluated. The anterior segment of the eye (including the cornea, iris, and crystalline lens) was cut away from the anterior scleral rim, and the vitreous body and retina were also dissected and discarded, leaving only the sclera. A 1.5-mm diameter trephine was used to punch out a tissue button from the sclera adjacent to the temporal margin of the optic disc. The tissue button was immediately fixed in 2.5% glutaraldehyde with phosphate buffer (pH 7.4) for approximately 2 to 3 hours. The scleral tissue was processed further and sectioned for electron microscopic examination, as described elsewhere. For transmission electron microscopy (model 7650; Hitachi, Tokyo, Japan), the scleral tissue button was equally divided into outer, middle, and inner layers (from the episclera), and five to six electron micrographs were taken from the middle layer of the sclera for each eye at 30,000.

Analysis of fibril diameter and density was performed by a technician who was blind to the genotype. The diameter of fibrils was measured with image analysis software (Phototshop 10.0; Adobe Systems, San Jose, CA). Micrographs from nonoverlapping regions of the central portion of the sclera were taken from cross sections at 30,000. For statistical analysis of fibril diameters, we adapted the method described by Ezura et al. and Chakravarti et al., in which the median of 100 to 300 diameter measurements from each micrograph was calculated and treated as one sample (WT: n = 14 micrographs, KO: n = 20 micrographs). Since the normality assumption was supported by the data (P > 0.05, K-S test), parametric methods were used in this analysis. To eliminate the difference in fibril number that may have affected histogram shape, we applied a percentage frequency histogram in the analysis (WT: n = 1220, KO: n = 1897).

The mean density of the scleral fibrils was estimated by determining the number of fibril cross sections from a randomly selected area (0.4 μm²) in each micrograph. Fibrils showing only partial cross sections at boundaries of a counting area were excluded from analysis.

Cell Biology

Chemical Preparation. CGS21680 was first dissolved with DMSO to a concentration of 10 mM and then serially diluted with FBS-free DMEM, to obtain different concentrations. SCH58261 was dissolved in DMSO to a concentration of 1 mM, which was then diluted with FBS-free DMEM to 1 μM. Dissolving and diluting were performed in the darkroom under red lights.

Cell Culture. Normal human scleral fibroblasts (HSFs) were isolated from donor eyeballs. The donor eyes were obtained and managed in accordance with the guidelines in the Declaration of Helsinki for research involving human tissue. The HSFs were maintained in DMEM supplemented with 10% fetal bovine serum, 100 U/mL penicillin, and 100 μ/mL streptomycin. For treatment, culture medium was replaced by serum-free DMEM, and the cells were treated for 24 hours with or without CGS21680 (1 mM to 1 μM). For the group treated with both CGS21680 (1 μM) and SCH58261 (1 μM), the cells were incubated with 1 μM SCH58261 for 15 minutes before addition of CGS21680. After treatment, the medium was harvested, and total soluble collagen concentration was determined by collagen assay (Sircol; Biocolor, Ltd., Carrickfergus, UK). The cells were harvested for determination of collagen mRNA expression.

Immunohistochemistry of A2ARs in HSFs. Untreated HSFs were used for immunohistochemistry of A2ARs. The primary antibody was a monoclonal mouse anti-A2AR (1:400; Upstate Biotechnology, Lake Placid, NY), and the secondary antibody was Cy3-conjugated goat anti-mouse (1:1500; Jackson Immunochemicals, West Grove, PA). Fluorescent images were then captured (Axiovision software; Carl Zeiss Meditec, Dublin, CA).

Real-Time RT-PCR. Total RNA was extracted from HSF cultures (Trizol reagent; Invitrogen-Gibco, Grand Island, NY) and 0.2 μg of RNA from each sample was reverse transcribed with M-MLV reverse tran
scriptase according to the manufacturer’s instructions (Promega, Madison, WI). The primers were designed on computer (Primer Express 3.0 software; Applied Biosystems, Inc. [ABI], Foster City, CA), and the sequences and product length are provided in Table 1. PCRs were performed in a real-time PCR system (model 7500; ABI), with 2× SYBR green PCR master mix (ABI). The results were normalized to the housekeeping gene GAPDH.

Sircol Collagen Assay. The total soluble collagen concentration in the cell supernatants was measured with the collagen assay kit (Sircol; Biocolor, Ltd.). In brief, a 200-μL aliquot of supernatant was added to 1 mL of the dye from the kit, mixed in a mechanical shaker at room temperature for 30 minutes, and centrifuged to precipitate the collagen-dye complex at the bottom of the tube. The pellet was dissolved in 0.5 M NaOH, to release the collagen-dye complex, and the concentration of collagen was determined by absorbance at 540 nm in a spectrophotometer.

Statistics

To determine significance in our comparisons of biometric measurements, we used commercial software (Prism 4; GraphPad, San Diego, CA) for two-way, repeated-measures ANOVA, with one factor, to determine the main effects (genotype and postnatal day) and their interaction. Differences between genotypes at specific postnatal days were analyzed by the Bonferroni post hoc test. For collagen fiber diameter, among all AL components, only VC length declined during postnatal development. VC in both genotypes declined between P28 and P56. The most rapid increase took place between P28 and P42, followed by a small but significant increase during the remaining time. The AL in A2AR KO mice was significantly longer than in WT littermates at P28 (Fig. 1c; two-way ANOVA for repeated measurements; main effect: $F_{(1,125)} = 20.94$, $P < 0.001$; interaction: $F_{(4,125)} = 0.432$, $P = 0.785$; post hoc Bonferroni test indicated a significant difference between WT and A2AR KO groups at postnatal day 28 ($P < 0.05$).

Among all AL components, only VC length declined during postnatal development. VC in both genotypes gradually decreased between P28 and P56. However, the VC was greater in the A2AR KO mice than in the WT group (two-way ANOVA for repeated measurements; main effect: $F_{(1,150)} = 65.83$, $P < 0.0001$; interaction: $F_{(4,150)} = 1.48$, $P = 0.211$). There were significant differences between the genotypes at P28 ($P < 0.001$; two-way ANOVA), followed by post hoc Bonferroni test, P42 ($P < 0.001$), P49 ($P < 0.05$), and P56 ($P < 0.01$; Fig. 1b, Table 2).

Results

Relative Myopia and Associated Increase in AL and VC Length from P28 to P56

Consistent with previous reports, the WT mice developed hyperopia over the period between P28 and P56, with greatest hyperopia at P56 (4.5 ± 1.1 D, mean ± SE; Fig. 1a). A2AR KO mice also developed hyperopia during this period, with greatest levels at P56 (2.4 ± 1.6 D). A2AR KO mice were more myopic than were the WT mice (WT: $n = 18$, KO: $n = 13$, main effect: $F_{(1,115)} = 15.25$, $P < 0.001$; interaction: $F_{(4,115)} = 0.54$, $P = 0.709$, two-way ANOVA for repeated measurements).

Between P28 and P35, the A2AR KO was an average of 5.1 D more myopic than were the WT littermates. Post hoc comparison revealed a significant difference between genotypes at P28 ($P < 0.05$, two-way ANOVA, followed by post hoc Bonferroni test).

To further characterize the ocular growth in A2AR KO mice, we used our custom-built biometric equipment, which was specifically designed for mice, to determine the AL and other biometric parameters. AL in both genotypes increased significantly from P28 to P56. The most rapid increase took place between P28 and P42, followed by a small but significant increase during the remaining time. The AL in A2AR KO mice was significantly longer than in WT littermates at P28 (Fig. 1c; two-way ANOVA for repeated measurements; main effect: $F_{(1,125)} = 20.94$, $P < 0.001$; interaction: $F_{(4,125)} = 0.432$, $P = 0.785$; post hoc Bonferroni test indicated a significant difference between WT and A2AR KO groups at postnatal day 28 ($P < 0.05$).

Normal Growth in the Anterior Segment in A2AR KO Mice

Corneal radius of curvature in both genotypes increased rapidly during postnatal development, but was not significantly different between the two genotypes (main effect: $F_{(1,150)} = 1.35$, $P = 0.25$, Fig. 2a). Corneal radius of curvature in A2AR KO mice increased from 1.37 ± 0.006 mm at P28 to 1.49 ± 0.005 mm at P56, comparable to the increase from 1.36 ± 0.008 to 1.48 ± 0.008 mm seen for WT littermates. Similarly, AC depth in both genotypes increased during postnatal development. The AC in WT mice increased from

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### Table 1. Sequences for Primers and RT-PCR Product Length

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward Primer (5′−3′)</th>
<th>Reverse Primer (5′−3′)</th>
<th>Length (bp)</th>
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<tr>
<td>Collagen I</td>
<td>CGACCCTGTGCTGACAGGTC</td>
<td>CTGACCACGTTCACCAGGCG</td>
<td>158 bp</td>
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<tr>
<td>Collagen III</td>
<td>ATCTGAAATGCTGTTAAATGGAC</td>
<td>TATTTTGCCAGTGTCGTTGTC</td>
<td>120 bp</td>
</tr>
<tr>
<td>Collagen V</td>
<td>GAATTGACCGCAGGACACCA</td>
<td>TTTTTACCCGTTGGGACCTGGA</td>
<td>173 bp</td>
</tr>
<tr>
<td>GAPDH</td>
<td>GCCCTCTGCTGCTCCCTGTTTC</td>
<td>GACTCAGAACCTTCACCTTTCC</td>
<td>100 bp</td>
</tr>
</tbody>
</table>

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FIGURE 1. A2AR KO mice exhibited a greater myopic shift and greater VC depth and AL than in the WT littermates during postnatal development. Comparison of refractive status (a), VC depth (b), and AL (c) between A2AR KO mice and WT littermates at the indicated postnatal time points. *$P < 0.05$; two-way ANOVA for repeated measurements post hoc Bonferroni test, WT: $n = 18$, KO: $n = 13$. 

FIGURE 2. A2AR KO mice exhibited a greater myopic shift and greater VC depth and AL than in the WT littermates during postnatal development. Comparison of refractive status (a), VC depth (b), and AL (c) between A2AR KO mice and WT littermates at the indicated postnatal time points. *$P < 0.05$; two-way ANOVA for repeated measurements post hoc Bonferroni test, WT: $n = 18$, KO: $n = 13$. 

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309 ± 4 μm (mean ± SE) at P28 to 369 ± 3 μm at P56. AC in A2AR KO mice also increased from 299 ± 6 μm at P28 to 371 ± 6 μm at P56. There was no significant difference between the two genotypes (main effect: \( F_{(1,150)} = 2.53, P = 0.114 \), Fig. 2b). Lens thickness increased comparably in both genotypes during postnatal development (main effect: \( F_{(1,150)} = 2.47, P = 0.118 \), Fig. 2c). At P28, lens thickness (LT) in WT mice was 1544 ± 8 μm (mean ± SE), comparable to 1553 ± 8 μm (mean ± SE) in A2AR KO mice. At P56, LT was 1734 ± 7 μm in WT and 1722 ± 7 μm in A2AR KO.

To evaluate whether the development of myopia is confounded by pupil size or body weight in A2AR KO mice, we examined both during postnatal development. Pupil size increased in both genotypes with no significant difference between them (main effect: \( F_{(1,145)} = 0.123, P = 0.727 \), Fig. 2d). Similarly, body weight increased steadily in both genotypes during postnatal development (Table 2), comparable with results in a previous report,\(^{45}\) with no significant difference between the two genotypes at any time point (main effect: \( F_{(1,155)} = 2.650, P = 0.105 \), Table 2).

**Myopia Associated with Thinner but Denser Collagen Fibrils in the Posterior Sclera**

To investigate the possible ultrastructural basis for the development of myopia in the absence of A2ARs, we evaluated the fibril diameter in sclera of A2AR KO mice and WT littermates at P35 by electron microscopy. Ultrastructural analysis revealed that the collagen fibrils from posterior scleral regions of WT mice had wider mean cross-sectional diameters than those in A2AR KO mice (Figs. 3a, 3c). Quantitative analysis showed that this difference in diameters was significant (WT: 69.78 ± 11.87 nm, KO: 59.32 ± 11.87 nm, \( P = 0.009 \), independent \( t \)-test; Figs. 3b, 3d). Consistent with the smaller diameter, the density of the scleral fibrils per square millimeter in A2AR KO mice (173.31 ± 42.26/mm²) was significantly higher than in WT littermates (141.72 ± 34.19/mm²; \( P = 0.02 \), independent \( t \)-test).

**Effect of Activation of the A2A R on Expression of mRNAs for Collagens I, III, and V and Production of Total Soluble Collagen in Cultured HSFs**

To further investigate the biochemical basis for reduced scleral fibril diameter, we examined the effect of adenosine agonists and antagonists on expression of the mRNAs for collagen I, III, and V and production of total soluble collagen in cultured HSFs.

Further exploration of regulation of collagen synthesis by A2AR, we used a collagen assay kit (Sircol; Biocolor, Ltd.) to determine the effect of A2AR agonists and antagonists on total soluble collagen production in HSFs.\(^ {45}\) At 1 μM, the A2AR agonist CGS21680 significantly increased the level of collagen I, III, and V mRNAs compared with that in vehicle-treated cells (\( P < 0.01 \), one-way ANOVA, post hoc Bonferroni test). Conversely, SCH58261 (1 μM) significantly decreased collagen production.

Furthermore, co-treatment with the A2AR antagonist SCH58261 also abrogated the effect of CGS21680 on collagen mRNA expression (\( P < 0.01 \), one-way ANOVA, post hoc Bonferroni test).

To further explore regulation of collagen synthesis by A2AR, we used a collagen assay kit (Sircol; Biocolor, Ltd.) to determine the effect of A2AR agonists and antagonists on total soluble collagen production in HSFs.\(^ {45}\)

**Table 2. Various Biometric Parameters Comparison between WT and KO Mice between Days 28 and 56**

<table>
<thead>
<tr>
<th>Age (d)</th>
<th>Group</th>
<th>Refraction (D)(^{a})</th>
<th>Pupil Size (mm)</th>
<th>AC Depth (μm)</th>
<th>Lens Thickness (μm)</th>
<th>VC Depth (μm)(^{b})</th>
<th>AL (μm)(^{b})</th>
<th>Body Weight (g)</th>
<th>Corneal Curvature (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>P28</td>
<td>WT</td>
<td>0.0 ± 1.5</td>
<td>2.18 ± 0.02</td>
<td>309 ± 4</td>
<td>1544 ± 8</td>
<td>697 ± 7</td>
<td>2943 ± 17</td>
<td>12.31 ± 0.58</td>
<td>1.37 ± 0.006</td>
</tr>
<tr>
<td>P28</td>
<td>KO</td>
<td>-5.4 ± 1.1†</td>
<td>2.17 ± 0.02</td>
<td>299 ± 6</td>
<td>1535 ± 8</td>
<td>758 ± 7</td>
<td>3002 ± 14†</td>
<td>11.28 ± 0.69</td>
<td>1.36 ± 0.008</td>
</tr>
<tr>
<td>P35</td>
<td>WT</td>
<td>2.6 ± 1.5</td>
<td>2.28 ± 0.02</td>
<td>355 ± 4</td>
<td>1609 ± 6</td>
<td>695 ± 7</td>
<td>3052 ± 13</td>
<td>17.03 ± 0.47</td>
<td>1.42 ± 0.005</td>
</tr>
<tr>
<td>P35</td>
<td>KO</td>
<td>-2.2 ± 1.7</td>
<td>2.30 ± 0.02</td>
<td>354 ± 4</td>
<td>1606 ± 7</td>
<td>658 ± 7</td>
<td>3087 ± 16</td>
<td>16.54 ± 0.62</td>
<td>1.42 ± 0.007</td>
</tr>
<tr>
<td>P42</td>
<td>WT</td>
<td>3.4 ± 1.3</td>
<td>2.37 ± 0.02</td>
<td>354 ± 4</td>
<td>1667 ± 6</td>
<td>711 ± 10†</td>
<td>3158 ± 14</td>
<td>18.70 ± 0.51</td>
<td>1.43 ± 0.05</td>
</tr>
<tr>
<td>P42</td>
<td>KO</td>
<td>0.3 ± 1.3</td>
<td>2.34 ± 0.02</td>
<td>343 ± 4</td>
<td>1659 ± 6</td>
<td>711 ± 10†</td>
<td>3158 ± 14</td>
<td>18.70 ± 0.51</td>
<td>1.43 ± 0.05</td>
</tr>
<tr>
<td>P49</td>
<td>WT</td>
<td>4.1 ± 1.4</td>
<td>2.36 ± 0.02</td>
<td>355 ± 4</td>
<td>1695 ± 8</td>
<td>655 ± 8</td>
<td>3164 ± 12</td>
<td>20.23 ± 0.54</td>
<td>1.47 ± 0.005</td>
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<tr>
<td>P49</td>
<td>KO</td>
<td>1.8 ± 1.5</td>
<td>2.36 ± 0.02</td>
<td>350 ± 3</td>
<td>1689 ± 7</td>
<td>688 ± 10†</td>
<td>3191 ± 16</td>
<td>19.33 ± 0.55</td>
<td>1.46 ± 0.008</td>
</tr>
<tr>
<td>P56</td>
<td>WT</td>
<td>4.5 ± 1.1</td>
<td>2.42 ± 0.01</td>
<td>360 ± 3</td>
<td>1734 ± 7</td>
<td>630 ± 6</td>
<td>3191 ± 11†</td>
<td>22.12 ± 0.47</td>
<td>1.49 ± 0.005</td>
</tr>
<tr>
<td>P56</td>
<td>KO</td>
<td>2.4 ± 1.6</td>
<td>2.43 ± 0.01</td>
<td>371 ± 6</td>
<td>1722 ± 7</td>
<td>609 ± 11†</td>
<td>3235 ± 11</td>
<td>21.38 ± 0.64</td>
<td>1.48 ± 0.008</td>
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</tbody>
</table>

Data are expressed as the mean ± SE (WT: \( n = 18 \), KO: \( n = 15 \)). Significance was determined by two-way ANOVA for repeated measurements for the main effects (genotype and postnatal day) and their interaction, followed by Bonferroni post hoc comparison to determine the difference between the genotypes at specific postnatal days.

\(^{a}\) Significant main effect (i.e. genotype) by two-way ANOVA for repeated measurements. For refraction: \( F_{(1,155)} = 15.24, P < 0.01 \); for vitreous chamber depth: \( F_{(1,150)} = 64.83, P < 0.01 \); for axial length: \( F_{(1,125)} = 20.94, P < 0.01 \).

\(^{†}\) Significant difference \((P < 0.05)\) between the genotypes (Bonferroni post hoc comparisons).
DISCUSSION

Recently, three studies have shown that relative myopia can be induced in mice by targeted deletion of genes, including Egr-1, Nyx, and a double deletion of lumican and fibromodulin. However, because of the technical limitations of biometric measurements in mice, refractive status, and important biometric parameters of myopia, such as VC depth and crystalline lens thickness, were not evaluated in these studies. Our custom-built OCT and the EIR designed by Schaeffel et al. allowed us to overcome these limitations and more accurately measure refractive status and other important biometric parameters in mice. The WT mice in our present study displayed similar trends within the time frame of the study. Of note, A2AR KO mice were on average 5.1 D more myopic than were WT mice at P28 and P35. Detailed measurements of VC length, AC depth, and corneal curvature demonstrate that development of relative myopia in A2AR KO mice is associated with longer AL, particularly in the VC, but not with altered corneal curvature or AC depth. These biometric changes in A2AR KO mice closely resemble those seen in human myopia which features higher myopic diopters, longer VC depth and AL, but no change in corneal radius of curvature and lens thickness. In contrast to the Egr-1 mutant mice, where the result was complicated by the potential confounding effects of reduced body weight, indicating general growth impairment, the effect we see in A2AR KO mice cannot be attributed to gross growth changes, since body weight is indistinguishable between A2AR KO mice.
and WT littermates. An earlier report stated that a 0.1-mm increase in pupil size could contribute to 0.8-D hyperopia when measured with EIR.** Our measurements showed that pupil size was not a factor in the difference we saw between A2AR KO and WT littermates. Furthermore, the normal development of corneal curvature, AC depth, and lens thickness strongly argue that the development of relative myopia in A2AR KO mice is not the result of any general difference in eye growth. The refractive shift in A2AR KO mice is also unlikely to be due to alterations in other organs and systems, because few physiological changes (with the exception of heart rate and metabolic rate) have been described during postnatal development in naive animals of this line of A2AR KO mice.53 Most studies, including ours, have uncovered different phenotypes in these animals after pharmacologic insults or stress, or in response to particular pharmacologic treatments, consistent with an important pathophysiological role of the A2AR.

The magnitude of the change in refractive status in A2AR KO animals was comparable to that in Egr-1 KO mice. However, Egr-1 knockout mice exhibited a persistent refractive change, but only a temporary change in AL, whereas A2AR KO mice displayed a transient refractive change but a persistent change in AL. It was speculated that additional changes in lens thickness, curvature, and power may account for the difference between refractive and AL changes in Egr-1 KO mice.46 Our finding that WT and A2AR KO mice have identical lens thickness suggests that this parameter is not responsible for the return to normal refractive status that we see in A2AR KO mice. It is possible that the retarded emmetropization process may in part account for the myopic shift observed in the A2AR KO mice. Studies of myopia induced by form deprivation or lens defocus are needed, to clarify whether the myopic phenotype in A2AR KO mice is specifically associated with visual signals.

According to the schematic mouse eye,42,54 1 D of change corresponds to only a 4- to 5-μm change in AL. However, we found that at P28, when there is significant refractive difference between A2AR KO and WT mice, the AL difference between the genotypes was 59 nm, significantly larger than the values predicted by the model, which predicts changes of 20 to 25 μm. The reason for this discrepancy is not clear. However, it should be noted that, to date, no published study has reported AL changes and refractive errors that agree with the predictions of the mouse model eye.46-55 A possible explanation for this is that the mouse model eye was based on in vitro measurements,42 whereas most of the published studies, including ours, present in vivo measurements. Consistent with this explanation, in two reports published by the same group, the AL reported in frozen sections42 was larger than that measured in vivo in mice of the same age.55

Differences between A2AR KO mice and WT littermates in refraction and AL were apparent only in young animals. By P56, neither refraction nor AL in A2AR KO was significantly different from those in WT. There are several possible explanations for this transient effect. It may reflect developmental changes in A2AR expression. Although there is no information available on developmental expression of A2AR in eye, A2AR mRNA in the brain increases during the first 3 weeks of postnatal development and then declines slightly to adult levels.56-59 Thus, it is possible that A2AR activity is at its highest level in the eye.

### Figure 4

Activation of the A2AR increases the expression of mRNAs for collagen I, III, and V and the total soluble collagen production in cultured HSFs. (a) Immunofluorescent analysis of A2AR distribution in HSFs. Cultures were immunostained with antibody specific for A2AR and with DAPI, to visualize the nuclei. Left: to right: anti-A2AR staining (red); 5× enlargement of the marked region; DAPI (blue) staining of the same enlarged region; and the merged image. Bar, 20 μm. (b) Collagen mRNA levels. HSFs were cultured and treated with different concentrations of CGS21680 and SCH58261. RQ (relative quantity) value is the mean change ratio in expression of the target gene. *P < 0.05, CGS21680 versus vehicle. **P < 0.05, SCH58261 versus vehicle. ▲P < 0.05, CGS21680 versus CGS21680 and SCH58261 (n = 6, one-way ANOVA, post hoc Bonferroni test). (c) Soluble collagen production. HSFs were cultured and treated with different concentrations of CGS21680 and SCH58261. **P < 0.05, CGS21680 versus vehicle. ▲P < 0.05, CGS21680 versus CGS21680 + SCH58261 (n = 6, one-way ANOVA, post hoc Bonferroni test).
around the third postnatal week, resulting in the refractive change seen around this particular postnatal stage. Another possibility is that A2AR activity exerts the same effect on refractive status throughout development, but other factors contribute in later development, counteracting or minimizing the effect of the A2AR.

In contrast to the transient change in refractive status and AL, VC depth remained significantly greater in A2AR KO than WT littermates throughout postnatal development. This difference in duration of effect is consistent with our recent finding that in hyperopic-defocus-induced myopia in the guinea pig, the defocused eyes rapidly developed toward hyperopia after removal of the defocusing lens, whereas the VC length in the defocused eyes returned toward normal much more slowly. Changes in lens curvature, index, and power may compensate for the longer VC depth later in development, resulting in return to normal refraction without a change in VC depth.

Current understanding of refractive development suggests that the sclera plays a pivotal role in the control of eye size in the development of myopia. Thus, our finding of longer AL, particularly in the VC, led us to investigate possible alterations in scleral structure in A2AR KO mice. The association we observed between postnatal development of relative myopia in A2AR KO mice and ultrastructural changes in collagen fibers in the sclera is consistent with previous studies. Studies in human myopia and in animal models of myopia have demonstrated that scleral thinning, tissue loss, and reduced collagen fibril diameter are associated with ocular enlargement and development of myopia. Scleral thinning is greatest at the posterior pole of the eye. Consistent with this notion, Chakravarti et al. found that mice with a double mutation of lumican and fibromodulin (important regulators of scleral fibril synthesis) show certain features of high myopia including increased AL, thin sclera, and retinal detachment. Our electron microscopic examination (which provides more sensitive measurements than the thickness and dry weight used in many previous studies) revealed that collagen fibril diameter is reduced and the density of collagen fibrils is increased in the sclera of A2AR KO mice. This finding is consistent with the reduced collagen fibril diameter observed in myopic tree shrews and highly myopic human eyes, as well as with the finding of increased numbers of fibril bundles across the sclera thickness in myopic tree shrews.

There is little information about changes in scleral collagen fibril density during postnatal development of myopia. In the fibromodulin-lumican double mutants just described, collagen fibrils with both abnormally large and abnormally small diameters are found in the sclera, whereas mice with fibromodulin mutation alone have only the abnormally small diameter collagen fibrils. However, refraction was not evaluated in these animals. In one study of form deprivation in tree shrews, McBrien et al. reported an increased number of small diameter collagen fibrils in the sclera after long-term form deprivation, which is consistent with findings in humans and is likely to contribute to the weakened biomechanical properties of the sclera that have previously been reported. This conclusion is in apparent agreement with our finding of increased density of collagen bundles in myopic A2AR KO mice.

Our ultrastructural finding in A2AR KO mice suggested a possible histologic basis for the development of relative myopia in these animals and prompted us to investigate A2AR modulation of collagen synthesis in cultured HSFs (Fig. 4a). Consistent with the ultrastructural changes, treatment with the A2AR agonist CGS21680 dose dependently increased collagen I, III, and V production in HSF in vitro, an effect that was blocked by the selective A2AR antagonist SCH58261. This finding suggests that A2AR activity directly affects collagen synthesis in scleral fibroblasts.

The modulation of scleral collagen by the A2AR is also in agreement with results reported in human hepatic stellate cells and human dermal fibroblasts. In contrast with our results, an early report showed that 7-methylxanthine (7-mx), a metabolite of the nonselective adenosine receptor antagonist caffeine, increases collagen concentration and the diameter of collagen fibrils in the posterior sclera in rabbit. More recently, 7-mx also was shown to reduce axial eye elongation and myopia progression in both guinea pigs and patients (Cui et al. IOVS 2008;49:ARVO E-Abstract 1736). However, it was not clear which adenosine receptor was responsible for the effects seen in those studies, since there are four receptor subtypes and 7-mx is nonselective. One explanation for the discrepancy between our study and the 7-mx studies is that different adenosine receptors exert different effects on eye development. The 7-mx may block several receptor subtypes in the sclera, whereas the A2AR KO affects only the A2AR. Caffeine and its methylxanthine metabolites block both A1R and A2AR with similar low affinities; thus, it is possible that the activity of A1R receptors also modulates the development of myopia, possibly by antagonizing the A2AR effect, since A1R is mainly coupled to Gs and inhibits adenylyl cyclase, whereas A2AR usually stimulates adenylyl cyclase through coupling to Gs proteins. Another possibility is that the effects are species specific, as has been reported for other factors associated with myopia. For example, retinoic acid levels change in the opposite direction in the choroids of marmoset and chicken during induction of myopia.

Our biometric, histologic, and biochemical findings suggest that activation of the A2AR is essential for normal growth of refraction, AL, and VC in mice during postnatal development and possibly through control of collagen synthesis in the sclera. Our findings raise the possibility that therapies targeting the A2AR can modify the development of myopia. In addition, these findings invite genetic studies of A2AR polymorphism and epidemiologic studies of human caffeine consumption as they relate to the development of myopia.

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References