Complement factors C1q, C3 and C5b–9 in the posterior sclera of guinea pigs with negative lens–defocused myopia

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Abstract

• AIM: To investigate the expression of complement factors in the posterior scleral fibroblasts of guinea pigs with negative lens–defocused myopia.

• METHODS: Eighteen guinea pigs were assigned randomly to two groups: the negative lens–defocused group (NLD group, n=9) and the normal control without treatment group (NC group, n=9). The effect of myopic induction was compared in three subgroups: eyes treated with a −10.00 D negative lens in the NLD group (NL group), eyes treated with a plano (0 D) lens in the NLD group (PL group), and untreated right eyes in the NC group (NC group). The following analyses were conducted at four weeks: examination of the refractive error via retinoscopy, assessment of complement C5b–9 expression in the posterior scleral fibroblasts using immunohistochemistry, and measurements of complement C1q and C3 protein levels in the posterior sclera by Western blot.

• RESULTS: After an induction period of four weeks, a significant myopic shift was detected in the eyes of the NL group, relative to that of the PL and NC groups (P<0.05). Data analysis showed a significant increase in the percentage of C5b–9 immunopositive fibroblasts in the posterior sclera of the NL group eyes, compared to the PL group (q=11.50, P<0.001). Significantly higher levels of C1q (q=4.94, P=0.01) and C3 (q=4.07, P=0.03) protein were detected in the posterior sclera of NL group eyes, compared to the PL group. There were no significant difference between the PL and NC groups for C5b–9 (q=2.44, P=0.10), C1q (q=1.55, P=0.53) and C3 (q=0.98, P=0.77) in the posterior sclera.

• CONCLUSION: The data from present study provide evidence of the up-regulation of C5b–9, C1q and C3 in the posterior scleral fibroblasts in a NLD myopic animal model. The results suggest that the complement system may be involved in the development of myopia.

• KEYWORDS: experimental myopia; complement factors; sclera; inflammation

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INTRODUCTION

It is evident that genetic and environmental factors are involved in the development of myopia, however, the exact mechanisms are not yet understood[1-3]. Animal models have been valuable in investigating the mechanisms involved in the development of myopia. We have previously used negative lens-defocus to successfully induce axial myopia in newborn guinea pigs [4,5]. Numerous studies have shown that degradation of the visual input using form deprivation or lens defocus can induce axial myopia during the early postnatal period[6-8]. During this process, the retina is thought to initiate signal transduction triggered by blurred vision; however, the sclera ultimately facilitates ocular elongation and myopia[4-8]. Theoretically, the factors that affect the function of the sclera, more specifically the posterior sclera, may modulate the induction of myopia[9,10].

We have previously reported that serum concentrations of high sensitivity C-reactive protein (hs-CRP) and complement C3 and CH50 were significantly higher in pathological myopia (PM) patients than in age- and gender-matched normal controls. This suggests that myopia may share features similar to those seen in autoimmune diseases and may represent a state of low-grade systemic or local inflammation[11]. Based on these findings, we hypothesize that the complement system may also contribute to the transformation of the sclera during the development of myopia.

Activation of the complement cascade occurs by three major pathways: classical (triggered by C1q activation), lectin and...
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alternative. This leads to the production of complement C3b, a central component in the complement system, that then joins other complement factors to form C5b, which along with C6, C7, C8 and C9 (referred to as C5b-9), forms the membrane attack complex (MAC), eventually triggering cell lysis or a sublytic attack[12-14].

To our knowledge, the relationship between the above complement factors and experimental myopia has not been previously reported. Therefore, the aim of our study is to investigate the expression of complement factors in the posterior sclera of experimentally-induced myopia. The structural organization of the sclera depends largely on the cellular activity of fibroblasts[15,16]. To test our hypothesis, we examined the expression of complement factors C1q, C3 and C5b-9 in posterior scleral fibroblasts.

MATERIALS AND METHODS

Animals Eighteen pigmented guinea pigs (Cavia porcellus, approximately 3 days old, weighing 70-90 g) were purchased from the animal center at the Chinese Academy of Medical Sciences (Beijing, China). All animals were treated according to the Association for Research in Vision and Ophthalmology (ARVO) statement for the Use of Animals in Ophthalmic and Vision Research.

Materials Custom poly (methyl methacrylate) (PMMA) concave lenses were purchased from Beijing Jingdejiarun Contact Lenses Co. (Beijing, China). The -10.00 diopter (D) lens (confirmed by lensometer) parameters were as follows: overall diameter (including the rim of the lens), 17.00 mm; optical diameter: 11.5 mm; inside optical curve radius, 7.50 mm; outside optical curve radius, 8.927 mm. The plano (0 D) lenses (confirmed by lensometer) parameters were as follows: overall diameter, 17.00 mm; optical diameter: 11.5 mm; inside optical curve radius, 7.50 mm; outside optical curve radius, 7.642 mm. There were 6 holes in the rim of each lens enabling suturing to the skin around the eye of the guinea pigs.

Myopic Induction and Ocular Measurement The guinea pigs were assigned randomly to two groups: the negative lens-defocused group (NLD group, n=9) and the normal control group (NC group, n=9). Animals in the NLD group were treated for four weeks with -10.00 D lenses fitted over one eye and plano (0 D) lenses fitted over the contralateral eye, while the animals in the NC group received no treatment. The effect of myopic induction was compared in three subgroups: the eyes treated with a negative lens in the NLD group (NL group), the eyes treated with a plano (0 D) lens in the NLD group (PL group), and the right eyes in the NC group (NC group). During the experimental period, measurements were taken to prevent form deprivation, such as keeping the lens clean at all times. All animals underwent cycloplegic ocular refraction measurement using streak retinoscopy (Suzhou Medical Equipment Factory, Suzhou, Jiangsu Province, China), prior to the experiment and four weeks after induction.

Immunohistochemistry At 4wk, guinea pigs were sacrificed by intraperitoneal injection of a lethal dose of thiopental. Both eyes of the guinea pigs in the NLD group (n=6) and right eyes of the guinea pigs in the NC group (n=6) were enucleated, and the remaining muscle and connective tissue were carefully removed. The eyeballs were fixed in 10% formalin at room temperature for 4h, and paraffin sections (3 µm) were cut on a microtome. After deparaffinization and rehydration, antigen retrieval was performed using the microwave oven method with a working solution from BD Pharmingen. Nonspecific binding was prevented by incubating the section with 1% goat serum in PBS for 1h at 37℃, followed by incubation with the primary antibody C5b-9 (1:100 dilution; Calbiochem/204903) overnight at 4℃. After washing, the sections were incubated with the following agents: biotinylated secondary antibodies for 30min at 37℃, Avidin-Biotin Complex reagent for 30min at 37℃, and DAB substrate for 10min. Counterstaining was performed with hematoxylin for 2min. Slides were then dehydrated in gradient alcohol washes, cleared with Xylene, and mounted for analysis. Sections without incubation with the primary antibody were used as negative control. Staining was examined by microscopy (Zeiss) equipped with a digital camera (Nikon). Data analysis of C5b-9 expression was performed by counting the number of immunopositive posterior scleral fibroblasts on each slide using a high powered lens (LM ×400). The data for each eye were calculated by averaging three non-overlapping visual fields in the posterior sclera area. All images were captured using the same light filter settings.

Western Blot At 4wk, the eyeballs were obtained as mentioned above. After removing the surrounding connective tissue, the eyeballs were sectioned with a razor blade and the anterior half discarded. The vitreous body, retina, pigment epithelium and choroid were peeled off on ice. The posterior scleral tissue was snap frozen in liquid nitrogen, followed by homogenization. Lysis buffer was added and the samples were centrifuged for 20min at 12 000 rpm at 4℃ in a microcentrifuge. The tubes were removed from the centrifuge and placed on ice. The supernatant was transferred into a fresh tube on ice and the pellet was discarded. The protein concentration was determined using the bicinchoninic acid method (Pierce, Rockford, IL, USA). Fifty micrograms of total protein per sample were denatured in SDS with a reducing agent. The samples were loaded into 15% SDS-polyacrylamide gels and subsequently electrophoretically transferred to nitrocellulose membranes in blotting buffer using an 80 V current applied for 60min. Blots were washed for 15min in PBS containing 0.05% Tween 20 (PBST), pre-incubated with blocking solution (5% Tween20 (PBST), pre-incubated with blocking solution (5%
Figure 1 C5b–9 immunostaining in the posterior sclera. Positive cells showed brown staining. Immunopositive staining of C5b-9 was increased in the posterior sclera of NL group eyes, compared to those of the PL and NC groups; no C5b-9 staining was observed when the primary antibody was omitted (negative control).

non-fat milk powder in PBST), washed three times with PBST, and subsequently incubated with primary antibodies, anti-C1q antibody (1:50 dilution; Abcam/ab71940) or anti-C3 antibody (1:50 dilution; Abcam/ab11887) at 4°C overnight. After washing, membranes were incubated with horseradish peroxidase-conjugated goat anti mouse secondary antibody for 1h with agitation and then washed four times for 5min in PBST. Antibody binding was visualized using enhanced chemoluminescence reagents (Amersham Biosciences), exposed on Kodak X-OMAT-R films (Rochester, NY, USA) and quantified using the Image J 2.1.4.7 software (developed by Wayne Rasband, National Institutes of Health, Bethesda, MD, USA). Protein loading was assessed by probing the blot for glyceraldehyde phosphate dehydrogenase (GAPDH) at 1:2000 dilution (Proteintech). The densitometry showed no significant differences in loading between the various lanes. The molecular mass of specific bands was determined using the BenchMark pre-stained protein ladder (Invitrogen) electrophoresed alongside the experimental samples.

Statistical Analysis Data were analyzed using SPSS 11.5 for Windows statistical software (SPSS, Chicago, IL, USA) and GraphPad Prism 5.0 (GraphPad Software, La Jolla, CA, USA). Between-group analyses of immunostaining data were carried out utilizing one-way analysis of variance (ANOVA) and Student-Newman-Keuls-q test (SNK-q). Numerical variables were expressed as average and standard deviation (mean±SD). A P<0.05 was considered statistically significant.

RESULTS

Myopic Shift in Negative Lens–defocused Eye of the Negative Lens–defocused Group No significant differences were observed for the refractive power between eyes in the NL, PL and NC groups before treatment (one-way ANOVA, F=1.50, P>0.05). After four weeks of induction, the refractive error in the eyes of the NL, PL and NC groups were (-1.61±0.49) D, (4.36±1.08) D and (4.78±0.98) D, respectively. A significant myopic shift was detected in eyes of the NL group, relative to that of the PL group (one-way ANOVA, q=20.16, P<0.001). There was no significant difference in refractive error between eyes of the PL and NC groups(one way ANOVA, q=1.41, P=0.59). This is consistent with the findings of our previous studies [8,9].

Increased C5b–9 Immunostaining in the Posterior Scleral Fibroblasts of Negative Lens–defocused Eye C5b-9 positive cells showed DAB positive brown staining, while negative cells and negative controls were stained with the hematoxylin counterstain only. Immunostaining for C5b-9 revealed little cellular labeling in the posterior sclera of the PL and NC group eyes. However, in the posterior sclera of the NL group eyes, Immunopositive staining of C5b-9 was present in many scleral fibroblasts (Figure 1). Data analysis showed a significantly increased percentage of C5b-9 immunopositive fibroblasts in the posterior sclera of NL group eyes, compared to those of the PL group (71.11%±14.10% vs 39.44%±11.62%, one-way ANOVA, q=11.50, P<0.001). The percentage of C5b-9 immunopositive fibroblasts revealed no significant difference between the PL and NC groups (39.44%±11.62% vs 32.78%±8.26%, one-way ANOVA, q=2.44, P=0.10; Figure 2).

Elevated C1q and C3 Expression in the Posterior Sclera of Negative Lens–defocused Eye The target bands for C1q and C3 were 26 KDa and 186 KDa, respectively. At four weeks, the densitometric analysis of Western blots showed significantly higher levels of C1q and C3 protein in the posterior sclera of NL group eyes, compared to those of the PL group (C1q:1.65±0.39 vs 1.15±0.15, one-way ANOVA, q=4.94, P=0.01; C3:0.45±0.06 vs 0.29±0.11, one-way ANOVA, q=4.07, P=0.03). No statistical difference in C1q
and C3 protein expression was shown between the PL and NC groups in the posterior sclera (C1q: 1.15 ± 0.15 vs. 0.99 ± 0.10, one-way ANOVA, \(P = 0.53\); C3: 0.29 ± 0.11 vs. 0.25 ± 0.10, one-way ANOVA, \(P = 0.77\)). A strong band at 37 kDa for GADPH indicated equal amounts of protein in each lane (Figures 3, 4).

**DISCUSSION**

The purpose of this study was to investigate the expression of complement factors in the posterior scleral fibroblasts of guinea pigs with negative lens-defocused myopia. Our results showed that C5b-9 immunostaining was elevated in the posterior scleral fibroblasts of the NL group eyes. This was accompanied by an increase in the expression of proteins C1q and C3 in the posterior sclera of NL group eyes, compared to the PL and NC groups. This corroborates our previous hypothesis that the complement system may be involved in the deformation of the sclera during the development of myopia.

The complement system is an important component of innate immunity and consists of approximately 30 fluid-phase cell-membrane proteins. It plays a key role in driving the immune system and triggering inflammatory responses\(^{[17,18]}\). Some evidence has suggested a possible role of the complement system in the pathogenesis of myopia. There is a broad consensus that the development of myopia is associated with active remodeling of the scleral extracellular matrix (ECM), evident by the thinning of the posterior sclera\(^{[19,20]}\). Different aspects have been studied in the search for factors involved in scleral ECM remodeling. Matrix metalloproteinases (MMPs) and tissue inhibitors of metalloproteinases (TIMPs) are considered to be the most important enzymes involved in the degradation of the ECM and have been proven to regulate the remodeling of the sclera\(^{[5,21,22]}\). Besides MMPs, some growth factors, such as transforming growth factor \(\beta\) (TGF-\(\beta\)) and basic fibroblast growth factor (bFGF), have also been reported to play important roles in the pathogenesis of myopia by modulating the function of either MMPs or scleral fibroblasts\(^{[23,24]}\). Studies have shown that the complement factors can affect the expression of MMPs, TIMPs, TGF-\(\beta\) and bFGF, which subsequently act on ECM remodeling\(^{[25,26]}\). Furthermore, a number of studies have shown associations between the complement system and ECM-related systemic and ocular disorders, such as coronary artery disease, systemic lupus erythematosus (SLE), age-related macular degeneration.
Although the elevated expression of C5b-9 and C3 in the posterior sclera of induced myopic eyes suggests an association between the complement system and scleral ECM remodeling, the precise mechanisms involved are not clearly defined. To address this issue, we investigated C1q, an initiating component of the classical complement cascade that belongs to a family of proteins called defense collagens [12,28]. It has been shown to serve as a bridging molecule to enhance the phagocytosis of antigen-antibody complexes and apoptotic cells through its collagen-like domain [39,40]. In addition, C1q plays an important role in mediating the adhesion of fibroblasts to ECM proteins such as collagen and fibronectin [41,42]. Evidence indicates that patients with high myopia have elevated serum levels of circulating immune complexes and autoantibodies to collagen [43,44]. This may explain our finding of increased C1q expression in the posterior sclera of myopic induced eyes, suggesting that the classical pathway may, at least in part, contribute to scleral ECM remodeling.

In summary, the results from our in vivo studies provide preliminary evidence showing an increase in complement in the sclera of myopic eyes. The data suggest that the complement system may be involved in the development of myopia and would be further confirmed by enlarging the sample size in a future study. As a powerful defense mechanism of the immune system, complement activation is a double-edged sword. Not only can it damage host tissue by causing cell lysis or sublytic attack, it can also promote the clearance of antigen-antibody complexes and enhance cell survival. Our observations support and encourage further investigation into the dynamic changes within the complement system during the various stages of myopic induction and suggest using specific complement inhibitors to elucidate the relationship between complement factors and negative lens-defocused myopia. A thorough understanding of these changes could lead to potential complement-targeted treatments in clinical myopia to ensure a balance between the helpful and harmful activities of the complement system.

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