Induction of Ocular Complement Activation by Inflammatory Stimuli and Intraocular Inhibition of Complement Factor D in Animal Models

Maura A. Crowley, Omar Delgado, Adrian Will-Orrego, Natasha M. Buchanan, Karen Anderson, Bruce D. Jaffee, Thaddeus P. Dryja, and Sha-Mei Liao

Novartis Institutes for Biomedical Research, Cambridge, Massachusetts, United States

Correspondence: Sha-Mei Liao, Department of Ophthalmology, Novartis Institutes for Biomedical Research, 22 Windsor Street, Cambridge, MA 02139, USA; sha-mei.liao@novartis.com.

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Purpose. Genome-wide association studies suggest a role for the complement system in age-related macular degeneration (AMD). We characterized ocular complement activation and evaluated a complement factor D (FD) neutralizing antibody.

Methods. Mice were treated with toll-like receptor (TLR) ligands, intravitreal injection (IVT), or corneal debridement. Levels of complement proteins and mRNA were measured. A FD neutralizing antibody was administered IVT into eyes of rabbits that were challenged with LPS (lipopolysaccharide) administered intravenously.

Results. Levels of C3 and factor B (FB) mRNA and protein in the eye were increased following intraperitoneal injection of TLR4 ligand LPS. Increased levels of C3 and FB breakdown products were observed in both eye tissues and plasma. Complement activation products were markedly reduced in C3−/− and Cfb−/− mice challenged with LPS. Ocular complement levels were also markedly reduced in eyes treated systemically with TLR2 and 3 ligands, injured by IVT injection or corneal debridement, or even in normal aging. IVT administration of a complement FD neutralizing antibody in rabbits inhibited LPS-induced complement activation in the posterior segment of the eye, but not in the anterior segment of the eye or in plasma.

Conclusions. Systemic TLR stimulation and eye tissue injury induced time-dependent alternative complement pathway activation in the eye. Ocular complement levels were gradually elevated during aging. An anti-FD antibody IVT potently inhibited LPS-induced complement activation in the posterior segment of the eye. This study provides insights into the dynamic profile of ocular complement activation, which is valuable for complement research in eye diseases and for developing complement therapeutics for AMD.

Keywords: alternative complement pathway, animal model, complement factor D, toll-like receptor

Age-related macular degeneration (AMD) is the leading cause of central vision loss in the aging population. Increasing evidence implicates the involvement of the immune system, specifically abnormalities in the alternative complement pathway (AP), in the pathogenesis of AMD. 1-7 Genetic linkage and genome-wide association studies have identified AMD susceptibility alleles in multiple complement genes (CFH, CFI, C2/CFB, C3, C9, CFHR1/3). 8-20 Complement proteins are frequently detected in drusen, the deposits in the macular area that are a common early sign of AMD. 11,21-23 Moderate elevation of complement components and activation products has been reported in plasma and eye samples from AMD patients. 24-31 Multiple animal models of AMD have been utilized, such as laser-induced choroidal neovascularization, light damage-induced retinal degeneration, and cigarette smoke exposure, to assess the role of complement in the eye. 25-38 Several complement inhibitors are currently being tested in the clinic for the treatment of AMD. 39-42 However, the precise mechanism of how complement activation influences AMD disease pathology and progression and even the basic understanding of complement function in the eye are still under investigation.

The AP is a part of the host defense system of innate immunity. It opsonizes and destroys invading pathogens and also responds to endogenous danger signals such as injured or apoptotic cells. Because increased complement activation is associated with AMD pathology, 24-31 identification of factors that activate complement in the eye could have implications for the pathogenic mechanisms of AMD. We characterized the changes in complement levels from eye tissues under various AMD-like pathophysiological conditions in mice. We also evaluated the ability of an intravitreally injected anti-FD antibody to block ocular complement activation in rabbits.

Materials and Methods

Animals and Induction of Acute Ocular Inflammation

All animal experiments adhered to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research and were approved by the Novartis Institutes for Biomedical Research Animal Care and Use Committee. Female C57BL/6 mice were
purchased from Taconic Biosciences (Rensselaer, NY, USA), Charles River Laboratories (Wilmington, MA, USA), and Jackson Laboratory (Bar Harbor, ME, USA). C57BL/6N mice from either Taconic Biosciences or Charles River Laboratories were used in all the mouse studies, except for one aging study with C57BL/6J mice from Jackson Laboratory. C57BL/6N mice from both Taconic Biosciences and Charles River Laboratories are B6N. The rd8 mutation in the Crb2 gene is present in C57BL/6N mice from both vendors as reported.43,45 Animals were housed in a pathogen-free facility on a 12-hour light and dark cycle and were fed standard laboratory chow and sterile water ad libitum. C57+/− (Jackson Laboratory) and C57−/− (University of Texas-Houston, Houston, TX, USA) mice and their wild-type (WT) littermates in C57BL/6N background strain were generated by heterozygous breeding.44,45 Acute ocular inflammation in mice was induced by intraperitoneal (IP) injection of either Salmonella typhimurium lipopolysaccharide (LPS) (Sigma-Aldrich Corp., St. Louis, MO, USA), Pam3CSK4 (Invivogen, San Diego, CA, USA), or poly I:C (polyinosinic-polycytidylid acid, InvivoGen) at the indicated doses formulated in sterile phosphate-buffered saline (PBS). Control animals received an IP injection of sterile PBS. On average, six to eight mice were used per time point in each treatment group. In IVT injection studies in mice, 1 μl sterile PBS was injected per mouse eye. Corneal epithelial debridement was performed as described previously.46 Male Dutch Belted rabbits at 8 weeks of age were purchased from Covance (Princeton, NJ, USA). Acute ocular inflammation in rabbits was induced by an intravenous injection of 20 μg/kg LPS. Twenty-four hours prior to LPS challenge, either anti-FD antibody (R&D Systems, Minneapolis, MN, USA) or a control goat IgG antibody (R&D Systems) was intravitreally administered bilaterally at 200 μg in 50 μl per eye. On average, five to seven rabbits were used per treatment group.

Tissue Collection and Preparation of Eye RNA and Protein Lysates

For mice, at indicated times post stimulation injection, plasma was collected. Eyes with lens removed were collected and frozen quickly on dry ice. While quickly collecting eye tissues, care was taken to avoid as much ocular muscle as possible. For rabbits, plasma was collected via ear vein 18 hours after LPS injection. Eye tissues were collected step by step on ice-cold petri dishes. First, aqueous humor was collected via ciliary body (CB) were collected. The lens was removed and vitreous humor was collected using a 1-ml pipette tip. After all the vitreous was carefully removed, the eye cup consisting of retina and retinal pigment epithelium (RPE) and choroid was collected with a number 21 scalpel. Each tissue was immediately placed on dry ice right after collection. All disposable tools were used once; other tools were cleaned between each tissue dissection to avoid cross-contamination. Total RNA was isolated from homogenized eye tissues using RNeasy micro columns (Qiagen, Venlo, Netherlands). Cell lysis buffer from Cell Signaling Technology (Danvers, MA, USA) was used to prepare eye protein lysates. Mouse eye tissues were dissociated in 200 μl lysis buffer. Rabbit iris/CB and retina/RPE/choroid were dissociated in 250 and 500 μl, respectively. Eye tissues were lysed using a TissueLyser II (Qiagen). The Bradford protein assay (ThermoFisher, Waltham, MA, USA) was used to determine protein concentrations of respective samples.

TaqMan

One microgram DNase I-treated total RNA from each eye sample was reverse-transcribed using the High Capacity cDNA Reverse Transcription kits (Applied Biosystems, Waltham, MA, USA). Reverse-transcribed cDNA was used for TaqMan quantitative real-time PCR reactions. Expression of the gene of interest was determined on the ABI Prism 7900HT (Applied Biosystems) with TaqMan mouse GAPDH endogenous control reagents. Primers used for real-time PCR purchased from Invitrogen (Waltham, MA, USA) were mouse C3 (Mm01252779_m1) and mouse C5b (Mm00435909_m1). The ΔCt value was calculated by subtracting the GAPDH Ct of the gene of interest Ct. The relative expression of the gene of interest = 2−ΔΔCt divided by the average of PBS control 2−ΔΔCt.

Protein Gel and Western Blot

Equal amounts of eye protein extract (40–60 μg protein) or 0.5 μl plasma from each sample was resolved on SDS gels (Novex NuPAGE 4-12% Bis-Tris; Invitrogen), and then transferred onto nitrocellulose membranes (iBlot Gel Transfer Stacks Nitrocellulose, Invitrogen). After incubation with Odyssey blocking buffer (LI-COR, Lincoln, NE, USA), the nitrocellulose membranes were probed with primary antibodies: either anti-mouse C3d (R&D Systems) or anti-factor B (FB) antibody (Quidel, San Diego, CA, USA). Secondary antibody was infrared fluorescently conjugated anti-goat antibody (IRDye800CW donkey anti-Goat IgG; Rockland, Limerick, PA, USA). Infrared fluorescent signals, which are directly proportional to the amount of the antigen on the Western blots, were detected with the Odyssey imager (LI-COR) and quantified using Odyssey software.

C3a and C5a MSD

Standard MSD plates (Meso Scale Discovery, Rockville, MD, USA) were coated with rat anti-mouse C3a or C5a (BD Pharmingen 558250 or 558027; East Rutherford, NJ, USA). Diluted mouse plasma and eye lysates that had been normalized to 5 mg protein/ml lysis buffer were added to MSD plates. Mouse C3a and C5a proteins (BD Pharmingen 558618; R&D Systems 2150-C5) were used to generate standard curves. Secondary biotinylated anti-mouse C3a or C5a antibody (BD Pharmingen 558251; R&D Systems BAF2150) and detection antibody (streptavidin-sulfotag, Meso Scale Discovery) were used and plates were read on a MSD plate reader.

Complement Alternative Pathway Functional Assay

Black MaxiSorp (Thermo Scientific, Waltham, MA, USA) plates were coated with Zymosan A (Sigma-Aldrich Corp.) diluted to 1 mg/ml in carbonate buffer pH 9.5 (Thermo Scientific) and washed three times with TBS-Tween 20 (TBS-T) buffer. A 7-point serial dilution of anti-human FD goat polyclonal antibody (AF1824, R&D Systems) starting at a concentration of 130 nM (final concentration in reaction) was prepared in saline. Normal human serum (Quidel) or normal rabbit serum (BioreclamationIVT, New Cassel, NY, USA) was diluted to 6% in ice-cold GVB-MG buffer (Boston Bioproducts, Ashland, MA, USA) supplemented to 2 mM MgCl2 final concentration for alternative pathway conditions. Diluted serum was first added to a polypropylene plate, and then diluted antibody was added. The plate was incubated for 15 minutes at room temperature, followed by transfer of 25 μl of the mixture to the coated reaction plate. The reaction plate was sealed with a plastic
adhesive film and incubated at 37°C for 30 minutes. The reaction was terminated by washing the plate three times with TBS-T. Membrane-attack complex (MAC) was detected with an alkaline phosphatase-conjugated mouse anti-human C9 neo-epitope monoclonal antibody (clone AE11; Diacel, Oslo, Norway) at 0.25 μg/ml. The plate was incubated for 1 hour at room temperature. After washing three times with TBS-T, substrate 4-MUP (Fisher, Waltham, MA, USA) at 0.18 mg/ml in 0.1 M Tris-HCl pH 9.0 containing 2 mM MgCl2 was added and the plate was incubated for 30 minutes at room temperature protected from light. Plates were read at 355-nm excitation/460-nm emission on a Fluoroskan plate reader (Thermo Scientific). Percentage inhibition of MAC deposition was calculated relative to the positive control (serum with assay buffer) after subtracting the background signal (serum with 25 mM EDTA added). These data were used to generate the IC50 value using nonlinear regression with GraphPad Prism software (GraphPad Prism, LaJolla, CA, USA).

Statistical Data Analysis

Data are presented as mean ± SEM. Mean differences were compared and statistical differences were determined by 2-tailed unpaired t-test. Differences were considered statistically significant when P < 0.05.

For FD antibody studies in rabbits, data were analyzed by subtracting the baseline values of the Ba levels (control IgG and PBS-treated group) from all values for each LPS-treated group. Using the baseline-subtracted values, the percent inhibition was calculated as % inhibition = (1 – [value of Ba with anti-FD Ab and LPS/average value of Ba in control IgG and LPS group]) × 100.

RESULTS

Systemic Administration of LPS Induces Transient Complement Synthesis and Activation in Mouse Eyes

Toll-like receptor (TLR) activation has been proposed to contribute to AMD pathology. Endotoxin/LPS-induced uveitis (EU) has been used as an acute model to evaluate anti-inflammatory reagents, such as anti-TNFα. To assess the effect of acute systemic TLR4 activation on complement activation in the eye, LPS was administered via IP injection in C57BL/6N mice. Plasma and eye tissues were collected at various time points from 3 hours to 14 days after a single LPS treatment. Levels of ocular C3 and C5b mRNA measured by TaqMan peaked 16 hours post LPS, and increased 20- to 50-fold, compared to the level of PBS-treated controls (Figs. 1A, 1B). Systemic LPS therefore induces local complement synthesis in eye tissues.

Western blots were used to evaluate the effect of LPS on AP proteins in mouse plasma and eye tissues. Representative Western blot images are presented in Figure 1C and 1D. Full-length C3 (~120 kDa) and FB (~90 kDa) protein levels were increased in eye tissues, peaking at 24 to 48 hours post LPS challenge at levels 2- to 3-fold higher than in PBS-treated control animals (Figs. 1A, 1B). The 10- to 20-fold difference between induced mRNA and protein levels is likely due to LPS-induced complement activation that results in breakdown of full-length C3 and FB proteins. Similar induction of full-length C3 and FB proteins was observed in plasma (data not shown).

The alternative pathway of the complement system is activated by a cascade of proteolytic steps. The FD protease cleaves FB into Bb and Ba. Bb serves as the catalytic subunit of C3 convertase (C3bBb) and splits C3 alpha into C3a and C3b. A majority of C3b is rapidly cleaved into iC3b, which is further degraded into C3d. A small portion of C3b binds to C3bBb to form the C5 convertase, which breaks down C5 into C5a and C5b. We evaluated these breakdown products to determine optimal readouts for complement activation in vivo. In naïve or PBS-treated control mice, Ba, C3d, and iC3b were barely detectable in eye tissues while a low level of each was present in plasma due to the continuous tick-over process of C3 cleavage by C3(H2O)FB. Systemic injection of LPS induced a time-dependent increase of Ba, C3d, and iC3b levels in both plasma and eye tissues (Figs. 1E, 1F). Since the C3b level is usually very low, the sum of iC3b and C3d levels was used to represent total C3 breakdown products. Levels of C3 and FB breakdown products increased gradually, starting at 3 hours post LPS challenge, reaching the highest levels at 24 to 48 hours, and thereafter gradually declining. Maximum induction of Ba or C3d+iC3b by LPS was approximately 4- to 6-fold when compared to PBS-treated controls. In plasma, the level of iC3b was approximately 5-fold lower than the C3d level (Fig. 1G).

However, in eye tissues, C3d and iC3b levels were comparable, indicating that iC3b may be more stable in the eye than in circulation. C3a and C5a levels, measured by MSD assays, also increased in plasma in a time-dependent manner, but were largely undetectable in eye tissue (Figs. 1H, 1I). C3a and C5a are known to be short-lived in vivo. Therefore, C3 and FB breakdown products Ba, C3d, and iC3b were suitable to monitor both ocular and systemic complement activation from tissue and plasma samples.

LPS induces C3 mRNA synthesis (Fig. 1A) and C3 protein breakdown (Fig. 1E). C3 is the convergence point of all three complement pathways (classical, alternative, and lectin). Increased Cfb mRNA levels (Fig. 1B) and increased complement breakdown product Ba (Fig. 1E) indicate that LPS-induced complement activation involves the alternative pathway and the amplification loop in the common pathway downstream of C3 activation. The role of the complement system in LPS-induced complement activation was further investigated by the use of C3-deficient and FB-deficient mice. Both strains are in C57BL/6N background. Ba levels are undetectable in C3−/− mouse eyes and plasma even following LPS challenge (Figs. 2A, 2B), indicating that C3 is essential for initiation of FB breakdown. C3b or C3H4O in C3FB complex was considered to be essential for FD to recognize FB and to cleave FB in in vitro studies. Using FB breakdown product Ba to monitor FB activation in C3-deficient mice, our data provided in vivo evidence that C3 is required for FB activation. Cfb−/− mice exhibited a significant (P < 0.01) but not complete reduction in C3 breakdown products C3d and iC3b in eyes and plasma comparing between LPS-challenged Cfb−/− and WT animals (Figs. 2C, 2D), indicating that the alternative pathway contributes to LPS-induced ocular complement activation.

TLR2 and TLR3 Ligands Also Induce Complement Activation in the Eye

LPS, the major ligand for TLR4, not only induces complement activation but also induces complement synthesis through the TLR4 signaling pathway. To address whether other TLR signaling pathways could also activate complement in the eye, Pam3CSK4 (a synthetic triacylated lipopeptide) and poly I:C (polynosinic-polycytidylic acid) were evaluated. Pam3CSK4 binds TLR2 and poly I:C signals through TLR3 and other intracellular RNA binding factors. Time-course studies in 7-week-old C57BL/6N mice challenged with IP injection of either Pam3CSK4 or poly I:C exhibited similar patterns of complement synthesis and activation, peaking in both eye and plasma at 8 to 24 hours (Figs. 3A, 3B), although at lower levels of induction compared to LPS challenge. Cfb
FIGURE 1. Time-dependent induction of complement mRNA and protein following LPS IP injection (2.5 mg/kg) in 7-week-old C57BL/6N mice. The dose of LPS was chosen based on both literature83 and dose-response studies for the optimal inflammatory responses without producing septic shock (data not shown). (A, B) C3 and Cfb mRNA levels in eye tissues were graphed relative to PBS-treated controls. Both C3 and Cfb mRNA levels peaked at 16 to 24 hours post LPS IP (P < 0.001). Full-length protein levels of complement components C3 and FB were graphed relative to the average value of PBS-treated controls. C3 and FB proteins peaked in eye tissues 48 hours (P < 0.01) and 24 hours (P < 0.001), respectively, post LPS. (C, D) Western blot analysis of eye lysates probed for C3 and FB proteins, both full-length proteins and breakdown products. The C3d antibody recognizes any mouse C3d containing
FIGURE 2. LPS-induced complement activation is dependent on the presence of C3 and FB. In order to run all samples on one gel (total 16 wells) for direct comparison between WT and complement-deficient mice in Western blots, seven PBS-treated samples of WT mice were pooled and loaded in 1 well. LPS-treated WT samples were run individually in seven wells. C3+/C0 or Cfb+/ mice were run in the same manner. The same amounts of protein lysate were loaded in each well. C3 and FB breakdown products C3d, iC3b, and Ba were measured in Western blots and were graphed relative to the average value of PBS-treated controls. (A, B) Comparing to LPS-treated WT controls, C3−/− mice exhibited a significant reduction (P < 0.01) in plasma and ocular Ba levels 24 hours after LPS IP injection (2.5 mg/kg). This is a representative of four independent studies. (C, D) Comparing to LPS-treated WT controls, Cfb−/− mice exhibited a significant reduction (P < 0.01) in plasma and ocular C3d and iC3b levels 24 hours after LPS IP injection. This is a representative of three independent studies.
mRNA was generated in the eye at earlier time points and at a higher level compared to C3 mRNA (Figs. 3C, 3D, 1B), suggesting that TLR signaling may preferentially activate the AP in the eye. Taken together, systemic and ocular complement activation can occur via systemic induction of TLR2, -3, or -4 signaling pathways.

Eye Tissue Damage Induces Rapid and Transient Local Complement Activation

Complement is known to be activated by trauma and in ischemia–reperfusion injury, presumably in response to endogenous danger signals. Intravitreal injection (IVT) is a common procedure to deliver anti-VEGF biologics for neovascular AMD. It involves a needle breaking through normal intact eye surface and retina to reach the vitreous cavity. To address whether ocular complement activation can occur via systemic induction of TLR2, -3, or -4 signaling pathways.

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**Corneal epithelial debridement is a model of inflammatory corneal neovascularization.** This model is initiated by mechanically removing the corneal epithelium of C57BL/6N mice. Eye samples were collected at various time points post corneal epithelial debridement, and complement levels were assessed. Similar to IVT, complement breakdown products C3d and iC3b were quickly elevated 2 hours post debridement and returned to baseline by 4 hours (Fig. 4B).

These results demonstrate that eye tissue damage causes rapid and transient local complement activation. Systemic complement levels were not affected by either challenge (data not shown).

**Aging Increases Baseline Complement Levels While Decreasing Response to LPS**

Aging is the greatest risk factor for AMD and other age-related diseases. We evaluated complement activation during aging in
naive C57BL/6N mice at 7, 21, and 48 weeks of age. Baseline levels of complement C3 and FB breakdown products gradually increased by 50% to 100% in eye tissues and 50% in plasma from 7 to 48 weeks old, indicating an increase of ocular and systemic complement activation with age (Figs. 5A, 5B). Aged C57BL/6N mice were reported to have increased accumulation of subretinal microglia/macrophages likely due to the presence of the rd8 mutation in the Crb1 gene.57 To address if the rd8 mutation plays a role in complement activation, we evaluated complement activation during aging in naive C57BL/6J mice, which do not carry the rd8 mutation, at 7, 22, and 54 weeks of age. Baseline levels of complement C3 and FB breakdown products gradually increased by 50% in both eye lysates and plasma from 7 to 54 weeks old, confirming an increase of complement activation with age that is independent of the rd8 mutation (Figs. 5C, 5D). Additionally, the ability to mount a response to LPS was reduced in older C57BL/6N mice. Complement breakdown products in eye tissues following LPS challenge were increased 5- to 8-fold in 7-week-old mice, but only 2- to 3-fold in 48-week-old mice (Fig. 5E). There was also a moderate reduction in LPS response in the plasma from 7 to 48 weeks old (Fig. 5F).

Factor D (FD) Neutralizing Antibody Inhibits Local Complement Activation in Rabbits

Human and rabbit FD proteins share 81% amino acid identity. A commercially available anti-human FD antibody from R&D Systems neutralized both human and rabbit FD in human and rabbit serum-based alternative pathway assays with similar potencies (Fig. 6A). However, possibly due to limited sequence homology between mouse and human FD proteins (66% identity), the anti-human FD antibody failed to inhibit AP complement activation in a similar assay with mouse serum (not shown). LPS-induced complement activation was adapted from mouse to rabbit to enable evaluation of the human FD neutralizing antibody via intravitreal delivery. The larger eye of rabbits is more suitable for preclinical testing of intraocular delivery of biologics with longer half-life. Since FB is the substrate for FD, the soluble FB breakdown product Ba was used as a direct readout for FD inhibition. Eighteen hours after LPS injection, the level of Ba in the posterior segment of the eye (retina/RPE/choroid) was 2- to 3-fold above PBS-treated controls. LPS-induced Ba generation in the posterior segment of the eye was inhibited by intravitreal administration of the anti-human FD antibody at 200 μg per eye 24 hours before LPS challenge. We assume that the concentration of FD in the eye should be lower than in serum (40–80 nM)58 and rabbit vitreous volume to be around 1.52 cm3.59 Intravitreal injection of 200 μg per eye would predict ~0.88 μM anti-FD, which is at least 11- to 22-fold higher than FD levels in the vitreous. LPS-induced alternative pathway activation in vitreous humor as measured by Ba generation was completely blocked (Fig. 6B) 42 hours after IVT of the anti-FD antibody. There was a 60% reduction of Ba levels in the retina/RPE/choroid (Fig. 6C). Since choroid is a highly vascular tissue, some of the Ba fragments detected in retina/RPE/choroid may have come from the blood. As expected, LPS-induced Ba generation in plasma was not affected by IVT anti-FD (Fig. 6D). LPS-induced Ba generation in the anterior segment of the eye (aqueous humor or iris/CB) was also unaltered by FD antibody pretreatment (Figs. 6E, 6F). This result demonstrates for the first time in rabbits that intravitreal administration of a complement inhibitor blocks local complement activation while complement levels in systemic circulation or even the adjacent tissue compartment remain unaffected.

**DISCUSSION**

Despite the strong genetic association with AMD and increasing evidence that upregulation of complement activation is associated with AMD pathology,24–31 little is known about how complement is activated in the eye. This is partly due to the lack of tools to monitor complement activity, particularly in eye tissue. Having established quantitative MSD immunoassays and Western blot assays, we were able to profile ocular and systemic alternative pathway complement activation under potential AMD-like pathophysiological conditions in mice. We demonstrated that multiple endogenous and exogenous danger signals that induce the innate immune system are capable of activating complement in the eye. Among them, aging is a well-known risk factor for AMD. TLR
FIGURE 5. Effect of aging on complement activation. Complement breakdown products Ba, C3d, and iC3b were assessed from naïve and LPS IP-injected C57BL/6N female mice at 7, 21, and 48 weeks of age and from naïve C57BL/6J female mice at 7, 22, and 54 weeks of age. LPS treatment was 24 hours. C3 and FB breakdown products were measured by Western blots and were graphed relative to the average value of 7-week-old samples. (A) Baseline levels of complement activation products in eye tissues of C57BL/6N mice increased with age from 7 to 48 weeks ($P < 0.001$ for C3d+iC3b and $P < 0.001$ for Ba). (B) Baseline levels of complement activation products in plasma of C57BL/6N mice also increased with age from 7 to 48 weeks ($P < 0.01$ for C3d+iC3b and $P < 0.01$ for Ba). (C) Baseline levels of complement activation products in eye tissues of C57BL/6J mice...
activation by pathogen products has also been implicated in AMD pathology.\textsuperscript{5,47} Furthermore, the stress response to eye tissue damage by the examples of IVT injection or corneal debridement could easily trigger complement activation. AMD is a complex and multifactorial disease, and complement activation could be a common pathogenic mechanism influenced by multiple AMD risk factors.

Most complement components are produced in the liver and circulate in the bloodstream as one of the main defense mechanisms against pathogens and endogenous danger signals. The role of complement activation in the eye is less well understood. Our results identify at least two types of local ocular complement activation induced by either eye tissue damage or systemic TLR activation. Local acute injury such as an IVT or corneal debridement breaks down the barrier between eye tissue and capillary blood vessels. This may cause a transient influx of complement proteins from blood, which, in combination with the presence of low levels of complement components in the eye, may lead to rapid ocular complement activation. Systemic challenge with TLR ligands may not directly activate ocular complement. Instead, TLR ligands upregulate complement gene expression in the eye, which leads to a slow induction of ocular complement activation, peaking 24 hours post challenge. Although some of the ocular complement activation by systemic TLRs might be due to residual blood from tissue collection, we observed similar levels of LPS-induced complement activation in eye tissue with or without removal of blood by perfusion (not shown). In addition, local complement inhibition would not be able to block complement breakdown products coming from circulation. The IVT FD antibody in this study inhibited complement activation in the eye, but not in the blood. This indicates that the ocular activation products are generated locally. The increased levels of C3 and FB mRNA and protein in the eye after inflammation could be produced by the infiltrating neutrophils and monocytes since leukocytes are good complement producers.\textsuperscript{60,61} They could be also synthesized by resident cells such as microglia, astrocytes, and RPE cells.\textsuperscript{62–66}

**FIGURE 6.** Intravitreal administration of an antibody against complement factor D inhibits LPS-induced Ba generation in rabbits in the posterior segment of the eye, but not systemically or in the anterior segment of the eye. (A) MAC deposition after Zymosan-induced alternative pathway activation in 6% human serum (solid line) or rabbit serum (dotted line) was blocked by a FD-neutralizing polyclonal antibody with IC\textsubscript{50} values of 8.1 and 4.9 nM, respectively. Twenty-four hours before LPS challenge, goat IgG or anti-FD was delivered IVT at 200 \(\mu\)g per eye to both eyes. Eighteen hours after intravenous injection of either PBS or LPS at 20 \(\mu\)g/kg, plasma (D) and eye tissue were collected. Eyes were dissected immediately into vitreous humor (B), retina/RPE/choroid (C), aqueous humor (E), and iris/ciliary (CB) (F). Complement Ba levels in indicated tissues were measured by Western blot and graphed relative to the IgG/PBS control group. LPS-induced Ba generation was inhibited 100% \((P < 0.01)\) in vitreous humor and was reduced by 60% \((P < 0.01)\) of PBS control levels in retina/RPE/choroid. The rabbit LPS study is a representative of two independent studies.
Ocular Complement Activation and Inhibition

The eye is considered an immunoprivileged site where complement is under tight control by intraocular complement regulators. However, we found that complement levels in the eye are easily altered in response to various inflammatory perturbations. Frequent exposure to systemic or local inflammatory stimuli, such as TLR ligands derived from pathogens, products or damaged tissue components, in combination with 50 to 60 years of aging could lead to a constant low level of complement activation in the eye, ultimately contributing to the development of AMD. This may be particularly relevant for those who are predisposed with high-AMD-risk genetic variants in complement genes. A notable difference we observed between systemic and ocular complement activation in mice is the higher level of iC3b relative to C3d level in eye tissues. The average ratio of ocular iC3b to C3d is 5-fold higher than in plasma. Plasma iC3b is barely detectable, while the ocular iC3b level is always comparable to the ocular C3d level. Increased deposition of iC3b along the glomerular basement membrane in complement factor H–deficient mice is responsible for the detrimental effect on renal function in the murine model of dense deposit renal disease. The interaction between iC3b and its receptor complement receptor 3 (CR3) is likely to play a similar role in the early synapse loss in Alzheimer mouse models. The presence of iC3b in drusen and on RPE cells has been proposed as one of the pathogenic mechanisms of AMD. Deposition of iC3b on the ocular surface could be recognized by infiltrating phagocytes or resident microglia that express CR3 or CR4, leading to macrophage activation and phagocytic damage to ocular cells such as RPE and photoreceptors. The presence of macrophages and subretinal microglia cells has been reported in AMD eyes.

Currently a number of complement therapies are in clinical trials for AMD. Lampalizumab, an anti-complement FD Fab (IVT), is in phase 3 clinical trials for the treatment of geographic atrophy (GA), an advanced form of AMD. However, Genentech recently announced that lampalizumab failed in one of the two phase 3 GA trials. Blocking complement FD, the first protease in the AP cascade, will prevent the generation of multiple inflammatory components in the alternative pathway and in the amplification loop. The effect of lampalizumab IVT on systemic complement activation was reported, but not its efficacy against complement activation in the eye. We demonstrated that complement activation in the vitreous was completely blocked 2 days after IVT of an IgG anti-human FD antibody in rabbits. However, in the retina/RPE/choroid fraction, which is considered to be the site of action for complement in the pathogenesis of AMD, only 60% inhibition was observed, possibly due to residual systemic complement products in choroidal and retinal vessels without PBS perfusion. Another possibility would be partial inhibition of local complement likely due to incomplete penetration of the antibody into the ocular tissues from vitreous. Thus the contribution of complement from choroidal and retinal vessels may account for only at most 40% of the total complement observed. Since only a modest increase in complement products is observed in AMD patients, it remains to be determined how much complement inhibition in eye and/or blood is required to achieve clinical efficacy in AMD trials. Complement model systems that we have developed, in combination with clinical results from complement trials, will be valuable to further understand the role of complement in AMD pathogenesis to design better complement therapeutics.

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References


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