Amelioration of Mouse Retinal Degeneration After Blue LED Exposure by Glycyrrhizic Acid-Mediated Inhibition of Inflammation

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Glycyrrhizic acid (GA) is a major component in the root and rhizomes of licorice (Glycyrrhiza glabra), which have been used as an herbal medicine, because of its anti-inflammatory activity. GA is known as an inhibitor of high-mobility group box 1 (HMGB1), which is involved in the pathogenesis of various inflammatory diseases including inner retinal neuropathy. In this study, we examined the effect of GA in a mouse model of retinal degeneration (RD), the leading cause of blindness. RD was induced by exposure to a blue light-emitting diode (LED). In functional assessment, electroretinography showed that the amplitudes of both a- and b-waves were reduced in RD mice, whereas they were significantly increased in GA-treated RD mice (P < 0.05), compared to those in non-treated RD animals. In histological assessment, GA treatment preserved the outer nuclear layer where photoreceptors reside and reduced photoreceptor cell death. GA-treated retinas showed significantly reduced expression of proinflammatory cytokines such as TNF-α, IL-6, IL-1β, CCL2 and 6, iNOS, and COX-2 (P < 0.05), compared to that in non-treated retinas. Immunohistochemistry showed that Iba-1 and GFAP expression was markedly reduced in GA-treated retinas, indicating decreased glial response and inflammation. Interestingly, HMGB1 expression was reduced in non-treated RD retinas whereas GA paradoxically increased its expression. These results demonstrate that GA preserves retinal structure and function by inhibiting inflammation in blue LED-induced RD, suggesting a potential application of GA as a medication for RD. In addition, we propose a potential retinal protective function of HMGB1 in the pathogenesis of RD.

Keywords: glycyrrhizic acid, HMGB1, inflammation, photoreceptor, retinal degeneration

INTRODUCTION

Retinal degeneration (RD) is a heterogeneous group of diseases characterized by the irreversible and progressive degeneration of photoreceptor cells in the retina, leading to blindness (Papermaster and Windle, 1995; Gregory-Evans and Bhattacharya, 1998; Kim G.H. et al., 2016). In the pathogenesis of a representative RD type of age-related macular degeneration (AMD) resulting from aging and long-term light exposure, inflammation is thought to be critical (Coleman et al., 2008; Ding et al., 2009; Karlstetter et al., 2010; Kim G.H. et al., 2016). Microglia and macrophages...
A total of 39 male albino BALB/c mice, 7 weeks of age, were
employed for the screening of GA effect and determination of an
appropriate concentration of GA; 18 mice (n = 6, each group)
were used for the main study. They were kept in a plastic cage
in a climate-controlled laboratory with a 12 h (7 a.m. to 7
p.m.) light/dark cycle. The animals were randomly assigned to
the treatment group. All mice-related experiments were handled
according to the regulations of the Catholic Ethics Committee
of the Catholic University of Korea, Seoul, which conform to
the National Institute of Health (NIH) guidelines for the Care
and Use of Laboratory Animals (NIH Publication No. 80-23), as
revised in 1996. Experimental procedures were approved by the
Institutional Animal Care and Use Committee at the College of
Medicine, The Catholic University of Korea (Approval Numbers:
CUMC 2016-0172-12 and 2017-0241-02).

Electroretinography (ERG)
Electroretinography (ERG) recordings followed procedures
described in our previous study (Kim G.H. et al., 2016). In brief,
the mice were kept in a completely dark room for 16 h before the
ERG recording. All animals were anesthetized intraperitoneally
with zolazepam (20 mg/kg) and xylazine (7.5 mg/kg). The
corneae were coated with hydroxypropyl methylcellulose gel
and covered with gold ring contact electrodes. A ground
electrode and reference electrode were placed subcutaneously
in the tail and ear, respectively. Stimuli were brief white
flashes delivered via a Ganzfeld stimulator (UTAS-3000; LKC
Technologies, Gaithersburg, MD, United States). Signals were
amplified and filtered through a digital band-pass filter ranging
from 5 to 300 Hz to yield a- and b-waves. Scotopic ERG, rod-
mediated responses were obtained at the following increasing
light intensities: 0.025 and 3.96 cd/s·m². Photopic, cone-mediated
responses were obtained following 5 min light adaptation on the
background light intensity. Recordings were obtained at the light
intensity of 6.28 cd/s·m². Each record was the average of three
responses obtained within a 15-seconds inter-stimulus interval.
The amplitude of the a-wave was measured from the baseline to
the maximum a-wave peak, and the b-wave was measured from
the maximum a-wave peak to the maximum b-wave peak.

Histological Analysis
At 3 days after blue LED exposure, the eye cup was enucleated
and fixed in 4% paraformaldehyde for 2 h. After fixation, the
eye cup was rinsed in 0.1 M phosphate buffer (PB; pH 7.4),
transferred to 30% sucrose, infiltrated overnight, and embedded
in a supporting medium for frozen tissue specimens (Tissue-
Tek O.C.T. compound; Sakura Finetek, Alphen aan den Rijn,
Netherlands). As described in our previous report (Kim G.H.
et al., 2016), retinal sections (7-µm thickness) were cut in the
sagittal plane, and stained with hematoxylin and eosin (H&E).
Quantitative analysis was performed in the stained sections;
outer nuclear layer (ONL) thicknesses were measured at 240-µm
intervals (superior to inferior) on vertical strips of the retina.
nuclei were counterstained with 4′,6-diamidino-2-phenylindole (DAPI; dilution, 1:1000; Invitrogen, Eugene, OR, United States). Light microscopic images were acquired on a Zeiss LSM 510 Meta confocal microscope (Carl Zeiss Co., Ltd., Oberkochen, Germany). The number of TUNEL-positive cells in the ONL was counted in two sections at a distance between 480 and 720 µm from the optic nerve in the superior area of the retina.

Immunohistochemistry

After washing with 0.01 M PBS, retinal sections were pre-incubated in 10% normal donkey serum for 1 h at room temperature. The sections were then incubated with rabbit polyclonal anti-ionized calcium binding adaptor molecule 1 (Iba-1) antibody (1:1000; Wako Pure Chemical Industries, Osaka, Japan), rabbit polyclonal anti-glia fibrillary acidic protein (GFAP) antibody (1:1000; Chemicon, Temecula, CA, United States), or rabbit polyclonal anti-HMGB1 antibody (dilution, 1:000; Abcam, Cambridge, MA, United States) diluted into PBS, for 1 d at 4°C. Sections were subsequently washed in PBS and incubated with Cy3-conjugated (dilution, 1:3000; Jackson ImmunoResearch, West Grove, PA, United States) or Alexa 488-conjugated donkey anti-rabbit IgG (1:3000; Molecular Probes, Eugene, OR, United States) for 2 h at room temperature. After rinsing several times in PBS, the cell nuclei were fluorescently stained with DAPI for 10 min and then mounted with anti-fading mounting media (Vector Laboratories, Burlingame, CA, United States). Images were obtained using a Zeiss LSM 510 Meta confocal microscope (Carl Zeiss Co., Ltd.). Quantitative image analysis was performed using Zen 2.3 software (Blue edition; Carl Zeiss). Region of interest was selected at a distance of 300 µm from the optic disk of each retinal section, and intensity of Iba-1 and GFAP immunoreactivities was automatically measured.

Quantitative Real-Time PCR (Real-Time qPCR)

Total RNA was purified with easy-Blue reagent (iNtRON Bio) according to the manufacturer's instructions. First-strand cDNA was synthesized with reverse transcriptase using the PrimeScript RT reagent kit (Takara Bio, Japan) in a total volume of 10 µL containing 0.5 µg of total RNA.

Real-time qPCR was performed in a final volume of 20 µL containing 10 µL of 2× SYBR Premix Ex Taq (Takara Bio, Japan), 1 µL each of 10 pmol/µL forward and reverse primers, and 2 µL cDNA template (1/100 dilution), using a commercial PCR detection system (LightCycler® 480, Roche, Mannheim, Germany) following the manufacturer's instructions. The annealing temperature was increased to 56°C for amplification. Melting curve analysis confirmed that each product was homogeneous and specific. Relative expression was calculated by comparison with a standard curve after normalization to the expression of the housekeeping gene GAPDH chosen as the control. The following primer sets were used: 5′-CCTTGCTACTCCCA GGTTC-3′ (forward) and 5′-AGGAGGTGACTTTCTCCTG-3′ (reverse) for TNF-α; 5′-TGTTCAAAAGAGGCC TGTGT-3′ (forward) and 5′-ATGTCCTGGATGAACTTT-3′ (reverse) for IL-1β; 5′-CCATCCAAATTCATCTGGAAA-3′ (forward) and 5′-GAGGATCTGGCAAACTGA-3′ (reverse) for IL-6; 5′-GCTAATCATTCCAGCAAG-3′ (forward) and 5′-TGAGCTTGGTGACAAACT-3′ (reverse) for CCL2; 5′-TGTTGTCGACTGAAAGAC-3′ (forward) and 5′-AGGGTCAGAATCAAGAACC-3′ (reverse) for CCL5; 5′-GCTGT-3′ (forward) and 5′-AGGAGGTGACTTTCTCCTG-3′ (reverse) for iNOS; 5′-AAAAATGCTGAGTATCAA-3′ (forward) and 5′-ACCCCTTTTGGTTGAGAT-3′ (reverse) for COX-2; 5′-TGATGTATGGAGACGCTG-3′ (forward) and 5′-TTGCCACACCTCTTGATGTC-3′ (reverse) for GAPDH. Real-time qPCR was performed three times for each group.

Western Blotting

Western blot analyses were performed on retinal extracts of the retina, which were homogenized in ice-cold lysis buffer (1% sodium dodecyl sulfate, 1.0 mM sodium orthovanadate, 10 mM Tris, pH 7.4). Aliquots of lysed tissue, each containing 50 µg of total protein were heated at 100°C for 10 min with an equivalent volume of 2× sample buffer and were loaded onto 10% polyacrylamide gels. Proteins were electrophoresed and subsequently blotted onto a polyvinylidene fluoride membrane. The membrane was blocked with 5% non-fat dry milk dissolved in 0.01 M PBS (pH 7.4) containing 0.05% Tween-20 for 1 h at room temperature. The membrane was then incubated for 15 h at 4°C with rabbit anti-HMGB1 polyclonal antibody (1:1000; Abcam) in blocking solution. The membrane was rinsed 3 times with PBS containing 0.05% Tween-20 (10 min per wash), and was then incubated with peroxidase-conjugated donkey anti-goat IgG antibody (1:1000; Jackson ImmunoResearch) for 2 h at room temperature. Blots were developed using the Enhanced Chemiluminescence Detection Kit (Amersham, Arlington Heights, IL, United States) and densitometry was performed using the Eagle Eye TMII Still Video System (Stratagen, La Jolla, CA, United States).

Statistical Analysis

Data are presented as mean ± SEM. All statistical analyses for ERG amplitude, histology image analysis, TUNEL-positive quantification, immunohistochemistry, real-time qPCR, and western blot analyses were conducted in Graphpad Prism 5.0 (GraphPad Software, San Diego, CA, United States) by one-way ANOVA with Bonferroni’s post hoc test comparing the mean of each group with the mean of every other group. For all tests, the differences were considered statistically significant at P < 0.05.

RESULTS

Determination of GA Dosage

First, we screened the inhibitory effect of GA on blue LED-induced RD and determined the appropriate dose of GA. For this purpose, we performed scotopic ERG on the mouse eye at 3 days after blue LED exposure and assessed the histology in retinal sections stained with H&E after ERG recording (n = 3,
FIGURE 1 | Dose-dependent protective effect of GA against blue LED-induced RD. (A) ERG analysis. The amplitudes of both a- and b-waves of the ERG responses increased in a dose-dependent manner. (B) H&E staining. Retinal structure was better preserved in the group treated with 10 mg/kg GA than in other groups treated with various concentrations of GA. Scale bar, 50 µm.

GA Preserves Retinal Function in Blue LED-Induced RD

We evaluated the functional effect of GA against RD with ERG in detail. Figure 2 shows the representative scotopic and photopic ERG recordings in age-matched normal control mice, DW-treated RD control, and GA-treated RD mice at 3 days after blue LED exposure, under different light intensities. The flash intensity of 0.025 cd.s/m² was found to be the lowest scotopic level to yield the a-wave, with a reliable b-wave (Figures 2A,B) and the optimal luminance to produce both ERG components with maximal amplitudes was about 3.96 cd.s/m² (Figures 2C,D). In this scotopic condition (Figures 2C,D), a- and b-wave amplitudes in GA-treated RD mice (n = 6, a-wave: 68.6 ± 6.6 µV, b-wave: 180.8 ± 12.2 µV) were almost comparable to those in the normal control mice (n = 6, a-wave: 104.2 ± 3.5 µV) and were significantly higher than those in RD mice (n = 6, 22.9 ± 0.7 µV, b-wave: 53.1 ± 3.2 µV) (P < 0.05). The differences in amplitudes of a- and b-waves between RD and GA-treated RD mice were 299% (22.9 ± 0.7 µV vs. 68.6 ± 6.6 µV) and 340% (53.1 ± 3.2 µV vs. 180.8 ± 12.2 µV), respectively. The photopic ERG recordings are represented in Figures 2E,F at a flash intensity of 6.28 cd.s/m², which evoked b-waves (Figures 2E,F) without a-waves. Under the photopic condition, ERG responses in mice groups were almost similar to the scotopic responses, which show reduction and enhancement of the amplitudes in RD mice and in GA-treated RD mice, respectively. Figure 2F shows that b-wave amplitudes in GA-treated RD mice are about 80% (n = 6, 36.5 ± 0.8 µV) of those in the normal control mice (n = 6, 45.6 ± 3.5 µV) and are significantly higher than those in RD mice (n = 6, 14.6 ± 2.5 µV) (P < 0.05).

GA Preserves Retinal Histology in Blue LED-Induced RD via Inhibition of Photoreceptor Cell Death

After ERG recordings, histological analysis was performed. Although laminar structure of the retina was generally preserved in RD control and GA-treated RD mice, two prominent changes were easily observed in the ONL where photoreceptors are located (Figure 3). The most prominent was the change in the ONL thickness. That is, at 480 µm from the optic disk in the inferior retina, the thickness of the ONL in the RD control and GA-treated RD mice corresponded to 54 and 77%, compared to that in normal control mice (Figures 3a–d). The other change was the arrangement of the ONL. In normal control mice, the two borderlines of the ONL appeared to be clean lines (Figure 3a), whereas those in RD mice were much wobbly (Figure 3b). In GA-treated RD mice, both lines appeared to be modest (Figure 3c).
Next, we evaluated the GA effects on photoreceptor cell death using the TUNEL assay (Figure 4). As previously reported (Kim G.H. et al., 2016), none or few TUNEL labeling was seen in the ONL (Figure 4a), while numerous TUNEL-positive photoreceptors were identified throughout the ONL in RD mice (Figure 4b). However, in GA-treated RD mice, TUNEL-positive photoreceptors were significantly reduced (Figure 4c). Quantification (n = 5 in each group) confirms and demonstrates a significant reduction in TUNEL-positive photoreceptors (P < 0.01) (Figure 4d). These results suggest that GA inhibits...
photoreceptor cell death and preserves the structure and function of the retina in blue LED-induced RD.

**GA Inhibits Inflammation in Blue LED-Induced RD**

We assessed inflammation in blue LED-induced RD and the anti-inflammatory effects of GA against RD. Analysis of gene expression by real-time qPCR showed the differential expression of several genes related to inflammation in normal control, RD control, and GA-treated RD mice (Figure 5). In the RNA isolated from the RD mice, the following seven genes of inflammatory mediators were upregulated, compared to the normal controls (Figure 5): three proinflammatory cytokines, TNF-α \( (P < 0.01) \), IL-1β \( (P < 0.01) \), and IL-6 \( (P < 0.01) \); two chemokines, CCL2 \( (P < 0.01) \) and CCL5 \( (P < 0.01) \); two other inflammatory mediators, iNOS \( (P < 0.05) \) and COX-2 \( (P < 0.05) \). In GA-treated RD mice, all genes were significantly downregulated compared to those in the RD controls \( ^* P < 0.05 \), \( ^{**} P < 0.01 \) in Figure 5), and were thus comparable to the normal controls (Figure 5).
Therefore, these results demonstrate that GA preserves the retina by inhibiting inflammation in LED-induced RD.

GA Inhibits Glial Responses in Blue LED-Induced RD

As previously reported (Chang et al., 2016; Kim G.H. et al., 2016), two retinal glial cells, Müller cells and microglia are activated in blue LED-induced RD. We assessed the effects of GA on glial responses using immunohistochemistry with anti-GFAP, a marker of activated Müller cells, and anti-Iba-1, a microglial marker (Figure 6). In normal control retina, GFAP was expressed in the endfeet and proximal processes of Müller cells in the ganglion cell layer (GCL) and the inner plexiform layer (IPL) (Figure 6a). The expression level of GFAP in control RD mice was higher than that in normal control mice; thus, GFAP immunoreactivity was frequently seen in the inner nuclear layer (INL), as well as in the GCL and the IPL (Figure 6b), and infrequently in the ONL. However, GFAP expression in GA-treated RD mice was significantly decreased ($P < 0.05$) and thus, appeared to be similar to that in normal control mice (Figures 6c,d).

Iba-1 expression patterns were quite similar to GFAP expression patterns. That is, Iba-1 was weakly expressed in microglia in the IPL in the normal control (Figure 6e), whereas it was strongly expressed in microglia in the ONL and in the subretinal space in RD (Figure 6f). In GA-treated RD mice, Iba-1 expression in microglia in the ONL and in subretinal space was significantly decreased ($P < 0.05$) and thus, rarely observed (Figures 6g,h). Taken together, GA might have strong inhibitory effects on retinal glial responses in blue LED-induced RD.

GA Increases HMGB1 Expression Level by Inhibition of HMGB1 Release From Photoreceptors During RD

As the main mechanism of GA against inflammation involves inhibition of HMGB1 (Mollica et al., 2007; Kim et al., 2012; Shen et al., 2015), we first examined the changes in HMGB1 levels by
FIGURE 6 | Immunohistochemistry and image analyses to evaluate glial responses of GA on blue LED-induced RD. (a–h) Confocal micrographs from vertical sections of normal (a,e), untreated RD control (b,f), and GA-treated RD retinas (c,g) processed for GFAP (green in a–c) and Iba-1 (red in e–g) immunoreactivities. Quantitative image analyses of the immunoreactivities for GFAP (d) and Iba-1 (h) were conducted. Both GFAP and Iba-1 immunoreactivities were significantly reduced in the retinas of the GA-treated group, compared with the untreated RD group. Scale bar, 50 µm. Data are shown as mean ± SEM; n = 5, **P < 0.01.

western blot analysis (Figures 7a,b). The analysis demonstrated that HMGB1 expression in the RD retinas was significantly decreased compared to that in normal controls (P < 0.05), whereas that in GA-treated RD retinas was significantly increased compared to that in RD controls (P < 0.05) and was thus similar to that in normal controls (Figures 7a,b).

We also performed immunohistochemistry with anti-HMGB1 to examine changes in HMGB1 expression (Figures 7c–e). In the retina of normal control mice, HMGB1 was expressed in all retinal neurons (Figure 7c). Consistent with a previous report (Hoppe et al., 2007), HMGB1 immunoreactivity was found homogeneously in the somata of the bipolar, amacrine, ganglion cells, and in the retinal pigment epithelium, whereas it was observed in periphery of somata in the photoreceptors. In the retina of RD mice, HMGB1 immunoreactivity in most photoreceptors that were localized in the ONL disappeared (Figure 7d). In GA-treated RD retinas, the pattern of HMGB1 immunoreactivity changed in the RD retinas and was restored to that in the normal control retinas (Figure 7e). These results indicate that HMGB1 expression is decreased in the retina in RD whereas GA attenuates HMGB1 release from the photoreceptors during RD.

DISCUSSION

There have been growing evidences that inflammation is an important event in the pathogenesis of RD. Recent clinical reports have shown that inflammatory reactions underlie AMD (Coleman et al., 2008; Nita et al., 2014; Kauppinen et al., 2016), retinitis pigmentosa (RP) (Yoshida et al., 2013; McMurtrey and Tso, 2018), and retinal detachment (Arimura et al., 2009; Murakami et al., 2013), and that anti-inflammatory agents can ameliorate them (Coleman et al., 2008; Becerra et al., 2011; Viringipurampe et al., 2013; Wubben et al., 2016; Bandello et al., 2017). Previously, we introduced the blue LED-induced RD mouse model and demonstrated its inflammatory characters: advent and activation of microglial cells in the ONL and increased expression of retinal inflammation markers, such as GFAP and osteopontin (Chang et al., 2016; Kim G.H. et al., 2016). In this study, we demonstrated that expression of proinflammatory cytokines, such as TNF-α, IL-1β, and IL-6, chemokines, such as CCL2 and CCL5, and other inflammatory mediators, such as iNOS and COX-2, was significantly increased. These inflammatory signs have been reduced by administration of GA, which has an anti-inflammatory effect. These results corroborate the concept that inflammation is a critical event in the pathogenesis of RD, and indicate that the blue LED-induced RD model is a useful model to study RD.

In RD, apoptosis has been considered the main mechanism of photoreceptor cell death (AMD (Xu et al., 1996; Dunaief et al., 2002), RP (Portera-Cailliau et al., 1994; Cottet and Schorderet, 2009), and retinal detachment (Cook et al., 1995; Arroyo et al., 2005), and thus, many pharmacological trials targeting anti-apoptotic molecules have been conducted. Unfortunately, the trials have not produced significant therapeutic effects (Donovan and Cotter, 2002; Murakami et al., 2013; Chinskey et al., 2014; Guadagni et al., 2015). A recently growing body of studies has shown that photoreceptor cell death in RD involves necrosis and autophagy as well as apoptosis (Sancho-Pelluz et al., 2008; Murakami et al., 2013; Chinskey et al., 2014; Guadagni et al., 2015; Lin and Xu, 2019). Previously, we have also demonstrated that the blue LED-induced RD model used in this study showed characters of necrotic photoreceptor cell death and that a necrosis inhibitor, NecroX-5 significantly prevented
Glycyrrhizic acid binds directly to HMGB1 by interacting with two shallow concave surfaces formed by the two arms of both HMG boxes (Sakamoto et al., 2015; Sakamoto et al., 2017). Further, binding of HMGB1 and GA has been shown to inhibit the chemoattractant, mitogenic, and cytokine-like activities of HMGB1, leading to anti-inflammatory effects (Mollica et al., 2007). These characters make HMGB1 a critical molecular target in a variety of human diseases, and thus, GA as a pharmacological strategy to inhibit HMGB1 has been used both in vivo and in vitro (Ming and Yin, 2013; Kang et al., 2014; Kao et al., 2014). In retinal diseases, it has been demonstrated that GA has inhibitory effects on inner retinal neuropathy, such as diabetic retinopathy (Chen et al., 2013; Abu El-Asrar et al., 2014), NMDA-induced injury (Sakamoto et al., 2015; Sakamoto et al., 2017), and ischemia-reperfusion injury (Dvorianchikova et al., 2011; Liu et al., 2017), in which amacrine and ganglion cells die. In addition, RD models with photoreceptor death are reported to show increased levels of HMGB1 in a rat retinal detachment model and in human eyes with retinal detachment (Viringiparampeer et al., 2013) as...
FIGURE 8 | Cartoon summary. The diagram illustrates the role of HMGB1 in the pathogenesis of blue-LED induced RD and amelioration of RD by GA. HMGB1 released from damaged photoreceptors by blue-LED exposure acts as a damp, which activates microglia. Activated microglia release proinflammatory cytokines such as TNF-α, IL-6, IL-1β, CCL2 and 6, INOS, and COX-2, which lead to the enhancement of inflammation or cell-death signaling in photoreceptors, such as caspase-3 and RIP1/RIP3 kinases. In this pathologic process, GA acts in two modes: (1) the classical mode is to inhibit retinal inflammation by binding to released HMGB1, thereby inhibiting the activation of microglia and proinflammatory cytokine release; (2) the photoreceptor-specific mode is to protect photoreceptors by binding to nuclear HMGB1 in photoreceptors, thus leading to the stabilization of the genome against injury or inhibition of the expression of death-related genes. ONL, outer nuclear layer; OPL, outer plexiform layer; SRS, subretinal space.

In the present study, the HMGB1 level was decreased in control RD mice, compared to that in normal control mice with restored HMGB1 in GA-treated RD mice. This result appears to be different from previous mentioned results showing increased HMGB1 in a rat retinal detachment model and in human eyes with retinal detachment (Arimura et al., 2009) as well as in dsRNA-induced RD model (Murakami et al., 2014). However, this discrepancy might be caused by the difference in the samples used for measuring the HMGB1 expression level, i.e., the two previous studies measured HMGB1 (released from photoreceptors and accumulated) in the vitreous (Arimura et al., 2009; Murakami et al., 2014), whereas the present study measured it directly in the retina. This explanation is corroborated by two previous reports, one mentioning that HMGB1 is downregulated in a light-induced RD model (Krishnan et al., 2008) and another that HMGB1 immunohistochemistry shows decreased/absent staining in the ONL in an AAG-dependent alkylation-induced RD model (Allocca et al., 2019). Our immunohistochemical results also clearly demonstrate decreased HMGB1 in photoreceptors in the blue LED-induced RD model.

Lastly, we suggest that HMGB1 is a key molecule for photoreceptor survival in RD. This proposal is also illustrated in Figure 8 and derived from the following basis. First, although HMGB1 as a representative DAMP is suggested to trigger inflammation and aggravate RD
there has been no direct evidence to prove this hypothesis. In addition, HMGBl injection into the vitreous induces ganglion cell loss in the ganglion cell layer, but little photoreceptor loss in the ONL (Sakamoto et al., 2015). Second, although almost all photoreceptors in control RD retinas lose HMGBl immunoreactivity, most of them look normal, not necrotic, based on the DAPI staining results in this study and the EM findings in our previous study (Kim G.H. et al., 2016). The same finding was reported in an AAG-dependent alkylation-induced RD model (Allocca et al., 2019). These findings do not match our general knowledge that HMGBl is passively released from necrotic cells (Scaffidi et al., 2002; Lotze and Tracey, 2005; Klune et al., 2008). Taken together, we argue that HMGBl may be a key survival factor for photoreceptors, as proposed in several pathological systems, including the heart (Funayama et al., 2013) and brain (Qi et al., 2007; Enokido et al., 2008). Considering that GA binds to HMGBl that is bound to DNA and inhibits HMGBl release from apoptotic chromatin (Mollica et al., 2007), and that HMGBl functions in nuclear structure to regulate gene expression and DNA repair (Lange et al., 2008; Kang et al., 2014), GA administered before RD induction may bind the photoreceptor DNA and stabilize the genome against injury to photoreceptors or inhibit the expression of death-related genes such as caspase-3 and RIP1/RIP3 kinases. To confirm this point, we might need to evaluate retinal function in an RD model with photoreceptor-specific HMGBl knockout mice in a future study.

CONCLUSION

The present study reveals that GA inhibits the expression of proinflammatory cytokines, chemokines, and other inflammatory mediators in the retina via released HMGBl inhibition, and additionally suppresses the expression of death-related genes in photoreceptors via binding to nuclear HMGBl, and thus, prevents RD progression and preserves retinal structure and function. These results suggest that HMGBl is a key molecule in RD pathogenesis and a potent molecular target for the inhibition of RD, and that GA is a potential medication for the prevention or treatment of RD.

DATA AVAILABILITY

The raw data supporting the conclusions of this manuscript will be made available by the authors, without undue reservation, to any qualified researcher.

ETHICS STATEMENT

All mice-related experiments were handled according to the regulations of the Catholic Ethics Committee of the Catholic University of Korea, Seoul, which conform to the National Institute of Health (NIH) guidelines for the Care and Use of Laboratory Animals (NIH Publication No. 80-23), as revised in 1996. Experimental procedures were approved by the Institutional Animal Care and Use Committee at the College of Medicine, The Catholic University of Korea (Approval Numbers: CUMC 2016-0172-12 and 2017-0241-02).

AUTHOR CONTRIBUTIONS

I-BK conceived and designed the experiments. GK and S-SP set up the RD mouse model, performed ERG, TUNEL staining, and immunohistochemistry. YP contributed to taking confocal images. HK performed western blotting and densitometry. GK, S-SP, and I-BK wrote the manuscript. All authors read and approved the final manuscript.

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