Localized Structural and Functional Deficits in a Nonhuman Primate Model of Outer Retinal Atrophy

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Purpose. Cell-based therapy development for geographic atrophy (GA) in age-related macular degeneration (AMD) is hampered by the paucity of models of localized photoreceptor and retinal pigment epithelium (RPE) degeneration. We aimed to characterize the structural and functional deficits in a laser-induced nonhuman primate model, including an analysis of the choroid.

Methods. Macular laser photocoagulation was applied in four macaques. Fundus photography, optical coherence tomography (OCT), dye angiography, and OCT-angiography were conducted over 4.5 months, with histological correlation. Longitudinal changes in spatially resolved macular dysfunction were measured using multifocal electroretinography (MFERG).

Results. Lesion features, depending on laser settings, included photoreceptor layer degeneration, inner retinal sparing, skip lesions, RPE elevation, and neovascularization. The intraretinal choroid was degenerated. The normalized mean MFERG amplitude within lesions was consistently lower than control regions (0.94 ± 0.35 vs. 1.10 ± 0.27, P = 0.032 at month 1, 0.67 ± 0.22 vs. 0.83 ± 0.15, P = 0.0002 at month 2, and 0.97 ± 0.31 vs. 1.20 ± 0.21, P < 0.001 at month 3.5). The intertest variation of mean MFERG amplitudes in rings 1 to 5 ranged from 13.0% to 20.0% in normal eyes.

Conclusions. Laser application in this model caused localized outer retinal, RPE, and choriocapillaris loss. Localized dysfunction was apparent by MFERG in the first month after lesion induction. Correlative structure-function testing may be useful for research on the functional effects of stem cell–based therapy for GA. MFERG amplitude data should be interpreted in the context of relatively high intertest variability of the rings that correspond to the central macula. Sustained choroidal insufficiency may limit long-term subretinal graft viability in this model.

Keywords: age-related macular degeneration, macular laser photocoagulation, nonhuman primate model, retinal atrophy, retinal transplantation

Geographic atrophy (GA), the hallmark retinal lesion of advanced nonexudative age-related macular degeneration (AMD), is characterized by localized degeneration of retinal pigment epithelium (RPE) and photoreceptor cells.1–5 The need for GA treatment is presently unmet.5 Stem cell–derived RPE transplantation is envisioned as a method of preserving vision, by regenerating the RPE layer at the border of GA lesions and protecting viable photoreceptor cells from decline.7–10 Additionally, photoreceptor transplantation is a potentially complementary strategy to regenerate photoreceptor cells and improve retinal function within GA lesions.17–33

A key challenge in stem cell therapy research for GA is the scarcity of naturally occurring or transgenic animal models with localized atrophy of RPE and photoreceptor cells.34–36 The term “complete RPE and outer retinal atrophy (cRORA)” was recently introduced by the Classification of Atrophy Meetings (CAM) expert panel, describing photoreceptor layer and RPE atrophy in patients with AMD based on optical coherence tomography (OCT) criteria.27 This underscores the widespread recognition of photoreceptor layer and RPE atrophy as a target of future clinical trials of cellular regeneration in AMD. Among genetic mouse models, small GA-like lesions occur in aged Ccl2–/–, Ccr2–/–, 28
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DICER1-deficient, Abr–/–, and Elovl4-mutant mice. Early AMD-like changes of RPE degeneration occur in the SOD2 knockdown model. Nonhuman primates with naturally occurring mild or intermediate nonexudative AMD-like lesions have been described, but none thus far with the cROA-like lesions of established GA. Chemically induced retinal degeneration in rodents produced localized RPE and photoreceptor degeneration. However, in large animals, this technique appears to cause selective photoreceptor damage while sparing the RPE cells—possibly, a dose-dependent effect—therefore, simulating the localized degenerative phenotype of GA only partially. Mechanical induction of retinal degeneration, by placing nonpermeable material in the subretinal space, also produces photoreceptor-only loss. Light-induced injury models including verteporfin photodynamic therapy application in mice appear to induce primarily choriocapillaris and/or RPE changes. All models other than the nonprimate lack a macula. Thus, they do not fully replicate the spatial localization of human GA lesions in the posterior pole of the ocular fundus. This feature is important for translational stem cell studies focused on optimizing stem cell delivery approaches to the subretinal space at the posterior pole. The normal outer retina lacks intrinsic vasculature. Therefore, the long-term viability of subretinally transplanted therapeutic cells depends on the adequacy of the choroidal vascular supply. Data suggest that the human choroid is structurally and functionally attenuated in GA. Histological and in vivo imaging data also implicate choroidal degeneration in another important cause of macular degeneration: Stargardt disease. Therefore, choroidal status may be an important determinant of stem cell therapy outcomes in at least two disease indications. Recent cell transplantation studies have shown surviving subretinal cells, but the impact of the choroid as a modulating factor remains poorly understood.

Greater understanding of relevant large animal models will enable further progress in cell therapy development for AMD. Here, we aimed to determine the extent to which precise lesion features could be optimized by modulating the laser intensity and pattern characteristics. We also aimed to understand structure-function correlation and characterize the temporal evolution of localized atrophy of RPE and photoreceptor cells in the lesioned nonhuman primate macula. In addition, we considered it a priority to analyze choroidal features to address this knowledge gap in the literature.

Materials and Methods

Animals

The study included four adult cynomolgus macaques (Macaca fascicularis), designated as subjects CM1 to CM4, that were aged 6 to 8 years (approximate human equivalent: 18–24 years) and weighed 4 to 7 kg. All animal procedures conformed to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research and the SingHealth Institutional Animal Care and Use Committee (IACUC, approval number 2015/S05/113/4) approved the study. All procedures were carried out in the SingHealth Experimental Medicine Centre, which is fully accredited by the Association for Assessment and Accreditation of Laboratory Animal Care International (AAALAC). The study protocol, including enrichment and caging system, was followed as per the National Advisory Committee for Laboratory Animal Research (NACLAR) guidelines. The temperature of the housing room was maintained at 25°C to 27°C and the relative humidity was kept between 45% to 55%. Equipment and toys were regularly changed. Food was provided twice daily, and water was available ad libitum. Before recruitment into the study, the animals were examined comprehensively to exclude ocular disease.

Laser Lesion Induction

Retinal atrophy was induced by laser photocoagulation on day 0 of the experiment in one eye each of four macaques using a 532 nm green frequency-doubled Nd:Crystal laser (PurePoint 532 nm Green Laser, Alcon Laboratories, Fort Worth, TX, USA) attached to a slit-lamp delivery system. 532 nm laser is primarily absorbed by melanin-containing RPE cells causing secondary photoreceptor damage. A handheld Mainster focal/grid laser contact lens was used for the laser procedures (High-power settings: spot size 50 μm, power 250 mW, duration 200 ms; Low-power settings: spot size 50 μm, power 150 mW, duration 200 ms). As a reference, prior studies for CNV induction used power, spot size, and duration settings that were approximately 700 mW, 50 μm, and 0.1 seconds. Thus, our power settings were chosen to be lower than those for CNV induction. For confluent application, the burns were placed with their edges touching but not overlapping. For nonconfluent application, burns were placed about one spot size apart. Single-shot mode was used to enable the precise control of spot location and separation. Laser application settings are shown in Supplementary Table S1.

Retinal Imaging

Indocyanine Green Angiography (ICGA). ICG (VERDYE, Diagnostic Green; dose 1 mg/kg) dissolved in water was administered via the intravenous route. The retinal and choroidal images were captured using scanning laser fundus imaging with high-resolution OCT Multimodal Imaging Platform (Spectralis, Heidelberg Engineering, Franklin, MA, USA). Images were obtained after dye injection at different phases as follows: early phase (0–2 minutes), midphase (4–7 minutes), and late phase (8–12 minutes onward).

Fluorescein Angiography. Ten percent sodium fluorescein (Curatis AG; dose 10 mg/kg) dye was given via the intravenous route. The retinal images were captured after dye injection at different phases as follows: early phase (0–3 minutes), midphase (4–8 minutes), and late phase (10–12 minutes onward) using scanning laser fundus imaging with high-resolution OCT Multimodal Imaging Platform Spectralis (Heidelberg Engineering, Heidelberg, Germany).

Optical Coherence Tomography. Retinal optical coherence tomography (OCT) imaging assay was performed using spectral-domain OCT (Spectralis, Heidelberg Engineering). The volume scan (20° × 20° scan, 61 B-scans per image) was captured at the laser area and nonlaser area of both eyes at baseline and months 1, 2, 3.5, and 4.5. Retinal layers were annotated based on published information on the use of this modality in NHPs. Manual correction was applied to ensure the accuracy of the layer segmentation.

OCT Angiography. High-resolution AngioScan OCT Angiography (OCT-A) was taken using Nidek RS3000 (Nidek CO. Ltd, Gamagori, Japan) at the laser area and nonlaser area.
of both eyes. Superficial vessel plexus consists of the large vessels and microvasculature within the retinal nerve fiber layer, and deep capillary plexus consists of the microvasculature within the inner plexiform layer and outer plexiform layer. Choriocapillaris was segmented from 31 to 40 μm underneath the retinal pigment epithelium. Manual correction was applied to ensure the accuracy of the layer segmentation. Choriocapillaris flow void density was calculated based on a reported method.66 The retinal perfusion density and choriocapillaris flow void density was evaluated from the 3×3 mm high magnification scans. The retinal OCTA images were thresholded by the mean grayscale value of the image, and the perfusion density was calculated as the ratio between the vascular regions per the total imaged area.

**Multifocal Electroretinography**

Multifocal electroretinography (MFERG) recordings were obtained using the RETImap system (Roland Consult, Brandenburg a.d. Havel, Germany). Jet electrodes were used to perform MFERG. The 61- and/or 103-segment stimulus protocols were used. Responses to the 61-segment test protocol were recorded at months 1, 2, and 3.5. Responses to the 103-segment test protocol were recorded only at month 3.5. The results were averages of eight cycles. The response amplitude at each locus was normalized to the baseline (average amplitude in control eyes from 1 month to 3 months) amplitude at the approximate same locus. The upper limit of normalized values was capped at 100%. Normalized amplitudes of lesion area and nonlesion area were longitudinally compared over time. The 61 segments in each eye were grouped into six rings from the center to the periphery per manufacturer’s readout. The amplitude ratio of each ring was compared between lesion and control eyes. We did not analyze implicit time values because available data do not support the correlation of this metric with photoreceptor loss in photic retinal injury.

**Histopathology and Immunohistochemistry**

The whole macaque eye was enucleated and fixed in 10% neutral buffered formalin solution (Leica Surgipath, Leica Biosystems Richmond, Inc., Richmond, IL, USA) for 24 hours. The eye was then dissected on the following day to remove the cornea and lens. It was then fixed again in 10% neutral buffered formalin solution (Leica Surgipath, Leica Biosystems Richmond, Inc.) for 24 hours, dehydrated in increasing concentrations of ethanol, cleared in xylene, and embedded in paraffin (Leica-Surgipath, Leica Biosystems Richmond, Inc.) for 24 hours, dehydrated in increasing concentrations of ethanol, cleared in xylene, and embedded in paraffin. The sections were dried in a 37°C oven for at least 24 hours. The sections were heated on a 60°C heat plate, deparaffinized in xylene, and rehydrated in decreasing concentrations of ethanol, cleared in xylene, and embedded in paraffin (Leica-Surgipath, Leica Biosystems Richmond, Inc.). Four-micron sections were cut using a rotary microtome (RM2255, Leica Biosystems Nussloch GmbH, Nussloch, Germany) and collected on POLYSINE microscope glass slides (Gerhard Menzel, Thermo Fisher Scientific, Newington, CT, USA). The sections were dried in a 37°C oven for at least 24 hours. The sections were heated on a 60°C heat plate, deparaffinized in xylene, and rehydrated in decreasing concentrations of ethanol. Hematoxylin and eosin (H&E) staining was performed according to a standard procedure. A light microscope (Axioplan 2; Carl Zeiss Meditec GmbH, Oberkochen, Germany) was used to examine the slides and capture the images.

In parallel, colocalization immunofluorescence staining was performed. Antigen retrieval was performed by incubating slides in sodium citrate buffer (10 mM sodium citrate, 0.05% Tween 20, pH 6.0) for 20 minutes at 95°C to 100°C and then cooling the slides down for 20 minutes at room temperature (RT). The sections were then washed three times for 5 minutes each with 1X PBS. Nonspecific sites were blocked with a 5% goat serum solution containing 2% bovine serum albumin (BSA) and 0.1% Triton X-100 in 1X PBS for 1 hour at RT in a humidified chamber. The primary antibodies shown in Supplementary Table S2 were diluted in 1X PBS and applied and incubated overnight at 4°C in a humid chamber. After washing the slides twice with 1X PBS and once with 1X PBS containing 0.1% Tween for 10 minutes each, corresponding secondary antibodies (conjugated with Alexa 594; Molecular Probes, Eugene, OR, USA) were applied at a concentration of 1:1000 in 1X PBS and incubated for 90 minutes at RT as shown in Supplementary Table S2. The slides were then washed twice with 1X PBS and once with 1X PBS containing 0.1% Tween for 5 minutes each. The slides were mounted with Prolong Diamond Antifade Reagent and DAPI (Invitrogen, Eugene, Oregon, USA) to visualize cell nuclei. For negative controls, the primary antibody was omitted. A fluorescence microscope (Axioplan 2; Carl Zeiss Meditec GmbH) was used to examine the slides and capture the images. Experiments were repeated in duplicate for antibody staining.

**Data Analysis**

MFERG amplitude data were analyzed using the generalized estimating equation (GEE) model, variance correction (jackknife variance estimator), considering the repeated measures for two eyes in the same animal and the multiple loci measured per time point per eye. Coefficients of variation were calculated by dividing the means of the repeated measurement by the standard deviation of those means.

**Results**

We modulated the overall laser intensity in terms of spot power (high versus low) and spot separation (confluent versus nonconfluent) (Supplementary Fig. S1). The appearance of the retina after low-power and high-power application is shown in Figure 1. Confluent application resulted in lesions that were immediately contiguous by biomicroscopic observation (Figs. 1a–b'). With nonconfluent application, we placed the laser spots approximately one spot size apart initially. However, by 15 minutes postapplication, the penumbra of individual burns had expanded (Figs. 1d' and d''). By 4.5 months, the pigmented lesions’ edges appeared indistinct (Figs. 1e' and e''). We observed pigmented changes on fundus photos at all laser intensity settings. Pigmentary reaction, including hypo- and hyperpigmentation, was especially marked with high-power laser applied at confluence (Fig. 1b′).

Lesions showed atrophic or neovascular characteristics, depending on laser intensity (Fig. 2). Confluent and nonconfluent application produced predominantly atrophic lesions as long as the power was kept low. Features indicating atrophy included reduced fluorescence due to hypoperfusion, fluorescence blockage due to intraretinal hyperpigmentation, and fluorescence transmission windows due to RPE atrophy or depigmentation (Figs. 2a’ and b’). Focal areas of fluorescein leakage indicated that choroidal neovascularization (CNV), when present, occupied a minor proportion of total lesion area (Fig. 2a”). High-power laser application produced significant fluorescein leakage by 4.5 months,
**FIGURE 1.** Evolution of pigmentary changes over 4.5 months. a. Images of the right fundus from 15 minutes to 3.5 months after laser application in a confluent pattern (subject CM1). b. Outcome image of the right fundus 4.5 months after confluent laser application in subject CM1. c. Control left fundus without laser application (subject CM1). d. Images of the fundus from 15 minutes to 3.5 months after laser application in a nonconfluent pattern in subject CM2. e. Outcome image of the right fundus 4.5 months after nonconfluent laser application in subject CM2. f. Control left fundus without laser application (subject CM2). Scale bar (2 mm) applies to a–f. 10× magnification of the white squares: 15-minute and 4.5-month images of low-power (a' and b') and high-power (a'' and b'') confluent laser application in CM1. 15-minute and 4.5-month images of low-power (d' and d') and high-power (d'' and d'') nonconfluent laser application in subject CM1. Foveal (f') and perifoveal (f') control images of CM2 showing normal macular pigmentation. Retinal blood vessels are indicated by white arrowheads. The perifoveal inner retinal light reflex is indicated by the black arrowhead.

**FIGURE 2.** Development of choroidal neovascularization (CNV) with intense laser. a. Late fluorescein angiogram of the lesioned right fundus of subject CM2 after nonconfluent laser application at low (a') and high power (a''). b. Late fluorescein angiogram of the lesioned right fundus of subject CM1 after confluent laser application at low (b') and high power (b''). c. Control left fundus of subject CM1. Scale bar (2 mm) applies to a–c. 5× magnification of white squares: Lower power, whether applied in a nonconfluent (a') or confluent (b') pattern, produced hypofluorescence (blue circles in a' and b'). Well-demarcated late hyperfluorescent areas (black arrowheads in a' and b') indicated fluorescence transmission windows due to RPE atrophy. Focal (yellow circles in a' and b'') or diffuse (a'') fluorescein leakage indicated the formation of active CNV as a result of using higher power. A control area of perifoveal retina without laser lesions is shown in c'. Retinal blood vessels are indicated by white arrowheads in a', b', and c'.
FIGURE 3. Photoreceptor layer and retinal pigment epithelium (RPE) changes. a. Serial macular optical coherence tomography (OCT) after confluent laser application, taken across areas of high- and low-power laser application in subject CM1 (F in top row indicates the location of the fovea). At 2 months, with low-power laser application, a sharp demarcation (arrowhead, third row) separated lesioned and nonlesion areas. The hyporeflective outer nuclear layer (ONL) band (white asterisk) was present at the nonlesion area. b. Serial macular OCT after nonconfluent laser application, taken across areas of high- and low-power laser application in subject CM2, through the fovea (arrowhead, top row). Nonconfluent laser results in skip areas between individual laser burns (arrows, second row). In the skip areas, ONL was relatively preserved (arrowheads, third row; images are not coregistered). The lesions continued to evolve through month 3.5 and showed incomplete ONL ablation by 4.5 months. c. High-magnification image of confluent high-power laser application outcome at 4.5 months. Intraretinal hyperreflective foci (empty arrowheads) were present, and the photoreceptor layer was ablated (red line). Black asterisk denotes RPE elevation/thickening. d. High-magnification image of confluent low-power laser application outcome at 4.5 months. The photoreceptor layer was selectively ablated (red line). e. High-magnification image of nonconfluent high-power laser application outcome at 4.5 months. The photoreceptor layer was selectively ablated (red line). f. High-magnification image of nonconfluent low-power laser application outcome at 4.5 months. The photoreceptor layer was incompletely ablated (between red lines). g. Control perifoveal retina, showing the physiological lamination pattern. 1: retinal nerve fiber layer, 2 (vertical white lines in c–g): ganglion cell layer and inner plexiform layer complex, 3: inner nuclear layer, 4: outer plexiform layer, 5: ONL, 6: hyperreflective bands corresponding to (from top to bottom) the ellipsoid zone (EZ), retinal pigment epithelium–Bruch’s membrane, and choriocapillaris complex. Blue line: external limiting membrane. Red lines, when separated,
indicating the presence of large and active CNV (Figs. 2a” and b”).

Retinal thickening occurred immediately after laser application (Fig. 3 and Supplementary Fig. S2). Subsequent retinal thinning stabilized between 2 and 4.5 months (Fig. 3). We did not quantify inner versus outer retinal thinning in this dataset due to the small number of animals. Confluent laser application produced broad areas of photoreceptor layer ablation, relatively sparing the inner retina. Skip lesions, that is, areas of photoreceptor ablation alternating with spared areas, occurred with nonconfluent applications. The presence of skip lesions correlated with incomplete photoreceptor layer ablation at 4.5 months (Figs. 3e–f). High-power laser application produced intraretinal hyperreflective foci that may correspond to intraretinal pigment-laden cells on histology, although more precise anatomical correlation is required to validate this assumption (Supplementary Fig. S3). Local RPE elevation and/or subjective thickening (Figs. 3a and c) further supported the presence of CNV with high-power confluent laser application. Low-power application caused less RPE thickening and/or elevation than high power. Immunohistology subsequently confirmed photoreceptor and RPE layer damage (Figs. 3h–i and Supplementary Fig. S4). The inner retina was relatively preserved despite laser-induced atrophy of the photoreceptor layer. As this is a small dataset, representative histology from all animals (CM1–CM4) are shown in Supplementary Figs. S3 and S4.

Laser-induced GA lesions in this model showed reduced choroidal vascular perfusion on indocyanine green (ICG) and OCT angiography (Figs. 4a–d). Choroidal neovascularization was faintly visible on ICG angiography (Fig. 4e) and more clearly seen on OCT angiography (Fig. 4i). The choroidal vascular loss involved the choriocapillaris in all animals and relatively spared the larger choroidal vessels (Fig. 4b shows CM3 and g–i show CM4). Retinal vasculature remained relatively intact. Retinal perfusion density and choriocapillaris flow void density quantification are shown in Supplementary Table S3. These in vivo imaging findings were supported by histology showing choriocapillaris attenuation and relative preservation of large choroidal vessels (Figs. 4i–j). We detected evidence of choriocapillaris damage at all laser intensity settings included in this study.

Multifocal electroretinography (MFERG) provided a quantitative readout of localized retinal function. Qualitatively, the spatial distribution of dysfunctional loci in the lesioned eyes showed a center-surround pattern in general, with central response reduction and peripheral preservation (Fig. 5). We saw intertest variability in lesion and control eyes. The 103-segment readout was noisier than the 61-segment readout in one subject, CM4.

We assigned responses as being inside or outside the lesion area by referring to the infrared reflectance images that showed the lesion boundary clearly (Fig. 6a). With the 61-segment protocol, the mean normalized response amplitude in the lesions was lower than responses outside the lesions in the same eye, at all time points tested (0.94 ± 0.35 vs. 1.10 ± 0.27, P = 0.032 at month 1, 0.67 ± 0.22 vs. 0.83 ± 0.15, P = 0.0002 at month 2, and 0.97 ± 0.31 vs. 1.20 ± 0.21, P < 0.0001 at month 3.5). With the 103-segment protocol, the mean normalized response amplitude in lesions was lower than the nonlesion area in the same eye 0.79 ± 0.27 vs. 0.94 ± 0.23, P = 0.032 at month 3.5) (Fig. 6b).

We also studied the feasibility and utility of an unblinded approach to MFERG analysis. In this approach, we did not designate responses as being located inside or outside lesions, but instead categorized them into concentric rings per the manufacturer’s software. This method could be useful for investigators when normalization is impossible, such as when both eyes are lesioned, or when prelaser readings are not available. We adapted a method that has been used to evaluate toxic maculopathy in patients taking hydroxychloroquine.67 This method is based on the pattern of hydroxychloroquine-induced retinal damage that selectively involves the perifoveal retina during the early stages, sparing the macular periphery. We normalized the mean ring 2 (R2) amplitude, reflecting central responses, to either ring 4 (R4) or ring 5 (R5), reflecting responses in the macular periphery. Thus, we derived the R2/R4 and R2/R5 ratios in each eye. R2/R4 and R2/R5 ratios were generally lower in lesioned eyes compared to the fellow control eyes almost (R2/R4: 1.16 ± 0.38 vs. 1.63 ± 0.46, P = 0.046 at month 1, 0.84 ± 0.19 vs. 1.43 ± 0.67, P = 0.14 at month 2, and 0.83 ± 0.21 vs. 1.74 ± 0.58, P = 0.014 at month 3.5 ; R2/R5: 1.09 ± 0.42 vs. 1.58 ± 0.45, P < 0.0001 at month 1, 0.78 ± 0.21 vs. 1.22 ± 0.48, P = 0.035 at month 2, and 0.79 ± 0.24 vs. 1.64 ± 0.40, P = 0.0056 at month 3.5). We also calculated MFERG ratios based on ring 1 amplitudes (instead of ring 2), that is, R1/R4 and R1/R5. Both R1/R4 and R1/R5 differed between lesion and control eyes at months 1 and 2 (Supplementary Fig. S5).

In this dataset, we had obtained MFERG results in normal eyes on five occasions (OD and OS prelaser, and OS at months 1, 2, and 3.5), using the 61-segment protocol. We used these data to estimate the intertest variation of the 61-segment MFERG assay in nonlesioned nonhuman primate eyes. The coefficients of variation (CV) of the mean amplitudes of R1, R2, R3, R4, and R5 in nonlaser eyes (n = 5) were 26.0%, 21.5%, 17.8%, 13.0%, and 13.8%, respectively, as shown in the Table.

**Discussion**

The data demonstrate that laser-induced GA-like lesions in nonhuman primates can be characterized in depth by serial in vivo retinal testing over time. We found that lesions evolved for several weeks before reaching a stable anatomical configuration. We validated the use of MFERG to measure localized functional deficits topographically, identifying the importance of mitigating intertest variation. We also present novel information regarding choriocapillaris loss in this GA model.
FIGURE 4. Choriocapillaris loss in the nonhuman primate model of geographic atrophy. 

a. Indocyanine green angiogram (ICGA) showed decreased cyanescent signal in the lasered region (boundaries denoted by *arrowheads*) in subject CM3. 
b. 4× magnification of *white square* in (a) showed the relative preservation of large choroidal vessels (*arrowheads*) at the laser site, compared to the relatively normal cyanescent signal in the adjacent retina (*asterisk*) of subject CM3. 
c. Control left eye of subject CM3, late phase. 
d. Early phase ICGA showing choroidal hypoperfusion (*arrowhead*) corresponding to the area of laser application that was visible biomicroscopically (*inset*). This ICGA was obtained 3.5 months after laser application in subject CM4. 
e. Late phase ICGA showed faint late hypercyanescence (*arrowhead*) within the lesion. 
f. Control left eye of the same animal, early phase ICGA. 
g-i. Optical coherence tomography angiogram (OCTA) showed choriocapillaris degeneration (*white asterisks*), relatively preserved large choroidal vessels, and choroidal neovascularization (surrounding the *red asterisk*) within the lesion area (*yellow box* in h is enlarged in i). 

j. Hematoxylin & eosin (H&E) stain of control retina. Inner retina (In), photoreceptor (P), and choroidal (C) layers are intact. The melanin-pigmented retinal pigment epithelium (RPE) layer, smaller choroidal vessels (*white arrowheads*), and retinal pigment epithelium are shown (*black arrowhead*). A large choroidal vessel is denoted with the *asterisk*. 
k. H&E–stained retina after laser application, showing total photoreceptor layer loss. The choroid (*white arrowhead*) is severely reduced in thickness and is hyperpigmented. The smaller choroidal vessels are lost and the larger choroidal vessels (*asterisk*) are relatively preserved. Pigment-
laden cells are globular and hyperpigmented, and show intraretinal migration (black arrowheads). The RPE layer was completely absent in parts (white arrows). Scale bar, 50 μm for f–g. Red circles denote the optic nerve head.

FIGURE 5. Spatial distribution of multifocal electroretinogram response amplitudes showing localized macular dysfunction in lesioned eyes. Color-coded heat maps showing normalized multifocal electroretinogram (MFERG) amplitudes at each test locus, for subjects CM3 and CM4, over serial testing. Right eyes (OD) were lesioned and left eyes (OS) are presented as nonlesion controls at each time point. Lesion boundaries in each right eye are denoted by yellow dashed lines and are laterally reflected into each left eye to indicate the contralateral lesion equivalent area.

It is well known that laser application outcomes in animal models, including CNV formation, can be unpredictable. In this experiment, confluent low-power application produced optimal results, including: 1. photoreceptor layer ablation sparing the inner retina, 2. absence of skip areas, 3. minimal CNV formation, and 4. acceptable interval to lesion stabilization. In a laser-induced C57BL/6J mouse model, 810 nm diode laser application (60-second application per spot) produced mild to severe lesions, depending on the power delivered in each case.68 In that study by Ibbett and colleagues, full-field ERG amplitudes were affected, presumably because the lesions occupied a large portion of the total retinal area in mice and thus reduced the summed retinal response. In this study, manual application of laser spots produced uneven spacing. In humans, AMD does not produce significant full-field ERG deficits,69 due to the small lesion size relative to total retina and the lack of panretinal photoreceptor dysfunction. Therefore, we did not evaluate full-field ERG responses in this study.

In GA lesions of nonexudative AMD, the inner nuclear and ganglion cell layers are relatively preserved despite advanced photoreceptor and RPE atrophy.4,70,71 We found it possible to selectively ablate the photoreceptor layer by carefully titrating laser intensity. Factors other than power are known to influence laser energy absorption and thermal cell damage. RPE and choroidal melanin pigmentation, and the orthogonality of the laser beam to the curvature of the posterior pole, varied considerably between animals in this study. Placing a test spot in a noncentral location can be a useful method of titrating the power before GA lesion induction. Caution should be exercised to avoid heavy laser application in a GA model. As we found in this study, excessive laser intensity is known to produce CNV,62 which is undesirable in a GA model.

In human GA, evidence suggests that the choriocapillaris is degenerated and hypoperfused to some extent.46–49,72–77 In macular atrophy secondary to Stargardt disease, the choriocapillaris appears to be severely attenuated.46–49,51–58,78 Therefore, the laser-induced nonhuman primate GA model may also be relevant for preclinical retinal stem cell research in Stargardt disease.79,80 It may be possible to avoid or reduce choriocapillaris damage in this model by changing...
FIGURE 6. Consistency of multifocal electroretinogram (MFERG) response reduction in GA lesions. a. Infrared reflectance reference images of subject CM4 of the prelaser right eye (OD), postlaser OD at month 3.5, and control left eye (OS) at month 3.5, with registered overlay of multifocal electroretinogram (MFERG) tracings. Red-shaded area in the post-laser OD image delineates the GA lesion boundaries and identifies traces designated as “lesion” (red boundary with shading) or “non-lesion” (outside red-shaded area, within grey line) in that eye. The equivalent control region (lateral mirror image, black line) in the OS was demarcated for normalization purposes. Inset shows postlaser OD retina at 3.5 months without lesion boundary markup. b. Mean normalized MFERG response amplitudes in the lesion and nonlesion areas at months 1, 2, and 3.5 (n = 4 animals, CM1–4). Data using the 61-segment MFERG stimulus were obtained at all three time points. The 103-segment stimulus protocol was used at month 3.5 only. c. MFERG ring amplitude ratios (mean ± standard deviation) over time, comparing lesion eyes to the fellow control eyes (n = 4 animals, CM1–4), without fellow-eye normalization. R2, R4, and R5 denote rings 2, 4, and 5, respectively.

certain laser parameters. For example, micropulse laser exhibits selective thermal localization.\textsuperscript{11,81–83} In the rabbit model of sodium iodate injection, photoreceptor and RPE loss was accompanied by dose-dependent choroidal
atrophy. \(^5^9\) Rat and rabbit choriocapillaris appear to regenerate spontaneously following injury,\(^{84,85}\) although it is unclear if this regenerative capacity exists in the primate choroid. MFERG is an objective assay of localized macular dysfunction. Its use in this context has been validated in a nonhuman primate model with naturally occurring bilateral macular dystrophy.\(^{25}\) MFERG has also been used to map retinal function in the visual streak of the pig model\(^{11}\) and in the wide-type primate macula.\(^{16}\) Notably, in our study, MFERG amplitudes in the lesions were not reduced to the same extent as is typically found in the absolute scotomas of established human GA. Full-field ERG, as a summed retinal response,\(^{23,34,86,87}\) is poorly suited to detect and quantify localized functional deficits. This study was not adequately powered to statistically compare MFERG amplitudes between laser settings; the data here may be used to design a larger study for this purpose.

The CVs of R1, R2, and R3 were above the typically acceptable limit of intertest variability, that is, 15%. In the 61-segment protocol, the number of test loci in R1, R2, R2, R4, and R5 are 1, 6, 12, 18, and 24, respectively. Therefore, there may be a higher tolerance for variation (due to inconsistent stimulus centration) in the peripheral rings (R4 and R5) than in the central rings (R1–R3). One strategy to minimize ocular drift during the MFERG is by administering a pericentral local anesthetic injection. The potential for inconsistent centration, and therefore the central CVs, may be even higher after laser application, by the absence of normal foveal landmarks. Researchers could consider determining the CV of control and test loci that is specific to their experimental conditions.

Human GA is associated with several molecular abnormalities, including complement activation and lipofuscin accumulation. Bruch’s membrane senescence,\(^{68,88}\) glial cell infiltration,\(^{30,90}\) and inner retinal remodeling are important facets of GA with relevance to cell therapy. In particular, the health of Bruch’s membrane may be a key determinant of cell transplantation efficacy in humans, as it influences transplanted RPE cell adherence and viability.\(^{52,93}\) We did not study those aspects in this relatively small project wherein our resources were focused on serial in vivo evaluation. This model is also limited in that it does not demonstrate the long-term progression of atrophy that is typical of human disease. Furthermore, photochemical or oxidative damage models may reflect the molecular pathophysiology of GA compared to the presumably predominantly photothermal effect of the model reported here.

### Conclusions

Laser-induced macular degeneration in this nonhuman primate model resulted in photoreceptor, RPE, and choriocapillaris loss. The use of OCT, OCTA, dye angiography, and MFERG enabled the spatiotemporal correlation of localized structural and functional deficits in vivo. The lesion features in this model are similar to those of cRORA as defined by the CAM expert panel. With further optimization, this model could provide a useful platform to validate stem cell–derived RPE and photoreceptor cell transplantation approaches for retinal regeneration in AMD. Given the high intertest variability of MFERG data, it is important to ascertain the magnitude of variability for the different experimental conditions specific to each laboratory. MFERG change data in the fovea and perifovea should be interpreted in the context of the high variability of the ring amplitudes that correspond to those loci. Choroidal insufficiency, if sustained, may limit the long-term viability and functionality of subretinal cellular grafts in this model.

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### References

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83. Song Q, Risco R, Latina M, Berthiaume F, Nahmias Y, Yarmush ML. Selective targeting of pigmented retinal pigment epithelial (RPE) cells by a single pulsed laser