An Analysis of Metabolic Changes in the Retina and Retinal Pigment Epithelium of Aging Mice

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Purpose. The purpose of this study was to present our hypothesis that aging alters metabolic function in ocular tissues. We tested the hypothesis by measuring metabolism in aged murine tissues alongside retinal responses to light.

Methods. Scotopic and photopic electroretinogram (ERG) responses in young (3–6 months) and aged (23–26 months) C57Bl/6J mice were recorded. Metabolic flux in retina and eyecup explants was quantified using U-13C-glucose or U-13C-glutamine with gas chromatography-mass spectrometry (GC-MS), O2 consumption rate (OCR) in a perifusion apparatus, and quantifying adenosine triphosphatase (ATP) with a bioluminescence assay.

Results. Scotopic and photopic ERG responses were reduced in aged mice. Glucose metabolism, glutamine metabolism, OCR, and ATP pools in retinal explants were mostly unaffected in aged mice. In eyecups, glutamine usage in the Krebs Cycle decreased while glucose metabolism, OCR, and ATP pools remained stable.

Conclusions. Our examination of metabolism showed negligible impact of age on retina and an impairment of glutamine anaplerosis in eyecups. The metabolic stability of these tissues ex vivo suggests age-related metabolic alterations may not be intrinsic. Future experiments should focus on determining whether external factors including nutrient supply, oxygen availability, or structural changes influence ocular metabolism in vivo.

Keywords: aging, retina, retinal pigment epithelium, metabolism, electroretinogram

Aging is associated with vision loss and physiological changes in ocular tissues.1–4 The retinal pigment epithelium (RPE) accumulates lipofuscin and drusen, Bruch’s membrane thickens, and cellular organization decreases.5–10 These alterations are also characteristic of age-related macular degeneration, a severe age-associated disease that is a leading cause of vision loss.2,11–14 With age, the neural retina thins,15–17 synaptic connections deteriorate,16,17 photoreceptor function declines,8,15 and their mitochondria are damaged.20–22

The amplitude of scotopic and photopic electroretinograms (ERGs) are reduced,18,23–26 and visual discrimination tasks18,27 suggest a reduction in spatial acuity and contrast sensitivity with aging. Retinal cells do not appear to be lost in significant number or are lost in selective areas of the retina,15,16,30 with age and rhodopsin levels remaining stable,31 suggesting the function of existing cells decline due to changes in other cellular processes, such as decreased electrical resistance.19,27 We hypothesized that a reduction in metabolic function in retinal cells may be an additional factor that contributes to diminished retinal responses to light in aging. Manipulating metabolism ex vivo directly influences ERG response.32 Previous studies suggest metabolic interplay exists between the highly glycolytic retina and the RPE33–35 and that metabolic changes in the RPE can affect the...
FIGURE 1. The study design is presented including the mouse groups, ages, and sex (A), the measures of visual function used (B), and the ex vivo approaches to characterizing metabolism (C) in the aging eye. The functional metabolic measurements are listed below the structure of the murine eye C, top left, from which we isolated two explants in this study C, center: the retina RPE-choroid complex that includes the RPE, choroid, and sclera that has been cleared of connective tissue (eyecup). The cellular composition of these explants C, right, is highlighted with retinal cells shown in shades of red, purple, orange, and yellow, whereas the eyecup is shaded in variations of gray.

retina. Increasing or disrupting glucose conduction in the RPE can cause retinal degeneration.\textsuperscript{36–38} Although steady-state metabolite levels differ in ocular tissues isolated from young and middle-aged mice,\textsuperscript{23} to our knowledge, the cross-talk between aging and central energy metabolism in the mammalian retinal ecosystem has not been examined in advanced age. Based on pilot studies in our laboratory, we hypothesized aging would have contrary effects on the retina and RPE and impact function. We expected decreased glycolytic and TCA flux in retina with increased oxidative metabolism, and anticipated the opposite in the RPE with increased glycolytic flux and reduced mitochondrial metabolism. To investigate the relationship between retinal response to light and energy metabolism in aging retina and RPE (study structure in Fig. 1), we measured retinal function in vivo by ERG and interrogated intrinsic alterations of metabolism by analyzing ex vivo tissue explants using stable isotope tracers\textsuperscript{39} and targeted metabolomics.

METHODS

Mouse Ages, Origin, and Housing

C57Bl/6J mice of both sexes were obtained from the National Institute of Aging aged rodent colony (Bethesda, MD, USA) and Jackson Laboratories (Bar Harbor, ME, USA; STOCK NO: 000664). Young (3–7 months) and aged (23–26 months) mice were housed in groups of 5 or less with ad libitum food (Rodent Diet 5053) and water. Mice were confirmed free of the Crb1 mutation found in Rd8 models of retinal degeneration.\textsuperscript{40,41} The light/dark cycle was 14/10 hours. Some aged male mice were treated with saline via osmotic minipumps or provided water by bottle for 8 weeks prior to euthanasia as controls for separate studies unrelated to the eyes. No differences were observed in ocular metabolism between untreated and control mice. Experiments complied with the policies of the Animal Care and Use Committee at the University of Washington and the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research.

Electroretinogram Set-Up

Mice were dark adapted overnight (approximately 17 hours), anesthetized with isoflurane, and the eyes were dilated with 2.5% phenylephrine (Akorn, Inc; NDC 174780201-15) and 1% tropicamide (Bauch + Lomb; NDC 24208-585-64). Gold electrodes were placed on each cornea. A reference and ground electrode were positioned on the back of the head. Mice were placed inside a UTAS Visual Diagnostic System with BigShot Ganzfeld with UBA-4200 amplifier (LKC Technologies, Gaithersburg, MD, USA). Readings were taken from both eyes, and the eye with the best response was considered. Scotopic assays were performed first, followed by photopic flicker or photopic single-flash experiments under a continuous 30 cd/m\textsuperscript{2} background light.

Single-Flash ERGs

Recordings were elicited using flashes of LED white light at increasing intensities with 2-minute pauses between flashes under scotopic (−50 to 50 dB) and photopic (0 to 100 dB flashes, 30 cd/m\textsuperscript{2} background light) conditions. Readings were calibrated such that 0 db = 2.5 cd*s/m\textsuperscript{2}. The a-wave amplitude was measured 8 ms after stimulus. The b-wave amplitude was measured as the magnitude from the a-wave minimum to the b-wave maximum.
Flicker-Fusion ERGs

Temporal resolution was measured under photopic conditions using a 5 dB flash of varied frequencies (20–50 hertz [Hz]) with 10 second pauses between frequencies and 9 replicates. Measurements were taken at 0.5 second intervals for a total of 512 points in 0.255 seconds (sampling frequency = 2003.91 samples/second). At each frequency, replicates were averaged. Figures 2F and 2G show examples of these raw values at 37 Hz. A waveform was generated and the magnitude was calculated using Fast Fourier Transform (Microsoft Excel Data Analysis ToolPak; sampling frequency = 2003.9 samples/second, step value = 3.91).

Isolation of Retinas and Eyecups

Animals were euthanized by awake cervical dislocation. Eyes were enucleated, cleared of excess tissue, and the retina separated from the RPE-choroid-sclera complex (eyecup) in Hanks’ Balanced Salt Solution (Gibco, Grand Island, NY, USA) within approximately 5 minutes. Tissues for ATP determination were snap frozen and stored in liquid nitrogen. Those for flux or O2 consumption rate (OCR) were used immediately.

Metabolic Flux and Metabolite Extraction

Metabolite standards and buffer components were purchased from Sigma-Aldrich (Millipore Sigma, St. Louis, MO, USA). Product numbers are included in Supplementary Table S2. Incubation medium was formulated as follows:

- Krebs’s Ringer Buffer (KRB): 98.5 mM NaCl, 4.9 mM KCl, 1.2 mM KH2PO4, 1.2 mM MgSO4, 20 mM HEPES, 2.6 mM CaCl2, and 25.9 mM NaHCO3
- Either 5 mM U-13C glucose alone or 5 mM unlabeled glucose and 2 mM U-13C glutamine

13C metabolic tracers (99% isotopic purity) included 13C labeled D-Glucose and L-Glutamine (Cambridge Isotope Laboratories, Inc., Tewksbury, MA, USA). Incubations, tissue extractions, and media extractions were performed as described with minimal changes. Some samples were split in half by volume before drying.

Gas Chromatography-Mass Spectrometry Analysis

Calibration curves were generated with metabolite standards (1.25–35 μM). Derivatization and selected ion monitoring (SIM) methods have been described previously with few changes. Incubation with N-tert-butyldimethylsilyl-N-methyltrifluoroacetamide (Millipore Sigma, St. Louis, MO, USA) was increased to 60 minutes. Samples were analyzed using a 49-minute gradient on an Agilent 7890/5975GC gas chromatography-mass spectrometry (GC-MS) system. For all derivatized metabolites, target ions were used for quantification and isotopologue distribution, and a qualifier ion for identity confirmation. Supplementary Table S2 describes SIM method details. Peak areas were obtained in MSD Chemstation (Agilent Technologies, Santa Clara, CA, USA) with manual verification of integrations. IsoCor version 2.42,43 corrected for natural 13C abundance and determined percent enrichment. In isotopologues, the number of incorporated 13C is represented shorthand by “Mx,” where x is the number of 13C (M0, M1, etc.).
Oxygen Consumption Rate

Two retinas or four eyecups per replicate were quartered and loaded into a perifusion system that assesses OCR as described\textsuperscript{44,45} with minimal changes. Tissue was perfused with KRB, 0.1 g/100 mL BSA, 1X antibiotic-antimycotic (Gibco, Grand Island, NY, USA) and 5 mM glucose. An artificial lung maintained 21% O\textsubscript{2}, 5% CO\textsubscript{2}, and 74% N\textsubscript{2}. Succinate (5 mM) was added when testing mitochondrial function. OCR was calculated as the product of flow rate times the difference in outflow and inflow oxygen. Data were reported as a change in OCR after subtracting the baseline OCR in 5 mM glucose alone.

ATP Measurements

ATP pools in retinas and eyecups extracted in boiling water\textsuperscript{46} were measured via luminescence with the Molecular Probes ATP Determination Kit, per the manufacturer's instructions.

Protein Concentration for Normalization

Flux and ATP pools were normalized to protein content. Protein pellets were solubilized in RIPA buffer (150 mM NaCl, 1% Triton X-100, 0.5% sodium deoxycholate, 0.1% SDS, 50 mM Tris pH 8.0, and 1X HALT protease/phosphatase inhibitor) and quantified using the Pierce BCA Protein Assay Kit, per the manufacturer's instructions.

Grouping and Statistics

Extractions, derivatization, and sample runs were processed in batches including ages, genders, matched retinas, eyecups, and/or their media, and random time points. To account for circadian contributions, the time of death for animals used in glucose flux, glutamine flux, and OCR are plotted in Supplementary Figure S1C. Sample sizes represent biological replicates from different animals. Normality was examined using the Shapiro-Wilk test. The majority of data was not normally distributed, thus nonparametric tests were used. Age differences were examined using Mann-Whitney tests. Changes associated with age and sex were considered using Kruskal-Wallis and Dunn’s multiple comparison tests. Significant results ($P < 0.05$) are marked with “*” and comparisons that did not reach statistical significance are marked “ns”.

RESULTS

Scotopic and Photopic ERG Responses are Diminished in Aging Mice

We recorded scotopic and photopic ERGs in young (4–5 months) and aged (26 months) male mice. The a-wave and b-wave amplitudes were extracted from raw traces of single flash scotopic ERGs (see Fig. 2A). The a-wave response measured 8 msec after the flash were significantly reduced between young and aged mice at −30 db and between 0 and 50 db (see Fig. 2C) and the b-wave amplitude was reduced from 0 to 50 db (see Fig. 2D). The scotopic a-wave and b-wave decreased proportionally with age. Cone-mediated retinal function was characterized using b-wave amplitudes in raw traces (see Fig. 2B) from single-flash photopic ERGs. The b-wave amplitude declined significantly in aged animals (see Fig. 2E). These findings correspond with previous studies where a- and b-waves were reduced in mice as early as 16 months\textsuperscript{33,34}.

Temporal resolution was considered with flicker-fusion ERG using 5 db flashing light (20–50 Hz). An overall loss in temporal resolution was seen in the averaged raw responses. Young (see Fig. 2F) and aged (see Fig. 2G) averages are shown at 37 Hz. After Fast Fourier Transformation, the magnitude declined significantly with age at all frequencies except 38.5 and 50 Hz (see Fig. 2H).

Metabolic Flux From Glucose Is Preserved in Retinal Explants Cultured Ex Vivo

We sought to determine if retina and RPE glucose usage is reduced at an age with reduced ERG responses (see Fig. 2). Retinas and eyecups were isolated from young (3–5 months) and aged male (26 months) and female (23–25 months) mice. Labeled metabolites generated from U-\textsuperscript{13}C-Glucose in tissues and incubation media were measured by GC-MS (Fig. 3A). Metabolites were quantified in terms of pmol or nmol metabolite per μg of protein. We examined metabolites in the tissue or exported by the tissue and considered the percent of the total pool labeled with \textsuperscript{13}C, and product:reactant ratios for particular steps in glycolysis and the TCA cycle.

Glycolytic intermediates (see Fig. 3B) in the explants were considered first. We found no consistent change in pool size, percent \textsuperscript{13}C-incorporation, and labeled isotopologues in either tissue (Supplementary Fig. S2). Export of M3 lactate from the retinas (see Fig. 3C) and eyecups (see Fig. 3D) into the incubation medium was stable. We examined product:reactant ratios at 30 minutes when the label incorporation had stabilized and found no substantial age-associated changes in glycolytic reactions in retinas (see Figs. 3E, 3G) or eyecups (see Figs. 3F, 3G). Glycolytic function appears preserved in both tissues.

Next, we considered how age impacts the Krebs cycle. As with glycolysis, we saw no consistent changes in retinas (Supplementary Fig. S3A–C) or eyecups (Supplementary Fig. S3D–F) in pool size, percent \textsuperscript{13}C incorporation, or labeled isotopologues. There were no significant age-related changes in product:reactant ratios in retinas (see Fig. 3H) or eyecups (see Fig. 3I) at the same 30-minute time point considered for glycolysis. Age did not significantly impact the Krebs cycle in retinas and eyecup explants.

Sexual dimorphism can influence outcomes in biological studies\textsuperscript{47} and differentially impact the visual system generally\textsuperscript{48} and in aging\textsuperscript{49,50}. Glucose processing in young and aged female mice was considered in parallel to males at 2 minutes. Total protein content in retinas was unchanged between males and females (see Supplementary Fig. S1A), however, in the eyecups (see Supplementary Fig. S1B) there was a change in aged males accounted for by normalizing all values to tissue protein content. There were no significant contributions of age or sex in pool size or isotopologues (Supplementary Figs. S4A, S4B) in the retina or eyecups (see Supplementary Figs. S4C, S4D). No sex-dependent differences in aging retinal or eyecup metabolism were found.

Glutamine Metabolism is Altered Ex Vivo in Aged Eyecups

Mitochondrial dysfunction is a hallmark of aging\textsuperscript{51}. To more directly investigate mitochondrial metabolism in aged ocular
FIGURE 3. Metabolic activity was examined by incubating retinal and RPE-choroid (eyecup) explants in U-\(^{13}\)C-glucose between 2 and 45 minutes. The tissue was washed and frozen, and aliquots of the incubation media were collected for analysis (A). Labeled intermediates downstream of glucose were quantified (B) in terms of percent \(^{13}\)C incorporation, pmol of \(^{13}\)C-labeled isotopologue per \(\mu\)g of protein in the retinal or eyecup explant, and product:reactant ratios for glycolytic reactions and those pathways that can be entered via pyruvate. Percent incorporation, pool sizes, and amount of isotopologue are shown in Supplementary Figures S3 and S4. To examine glycolytic activity, the amount of exported M3 lactate was measured in the incubation media of retinas (C) and eyecups (D). One aged eyecup at 30 minutes was found to be an outlier by Grubb’s test (alpha = 0.05, \(P < 0.05\)) and removed. The slopes of the lines were all non-zero in retina (\(p_{\text{young}} = 0.002\) and \(p_{\text{aged}} = 0.0003\)) and eyecups (\(p_{\text{young}} = 0.03\) and \(p_{\text{aged}} = 0.009\)), but showed no significant age-related change (regression slope: \(p_{\text{retina}} = 0.7\) and \(p_{\text{eyecup}} = 0.3\)). Within tissues, product:reactant ratios in glycolysis and common exit points to other pathways were plotted at 30 minutes because both tissues had reached a steady state. No significant age-related changes were seen in retina (E, G) or eyecup (F, G). Moving into the Krebs Cycle (B) at 30 minutes, no statistically significant age-related changes were observed in retina (H) or eyecup (I) explants. Normality of data was determined using the Shapiro-Wilk test and \(P\) values were calculated for age-related comparisons using Mann-Whitney tests (* \(P < 0.05\)). Error bars represent the standard deviation, except in panels C and D, which show the 95% confidence interval for the linear regression. GAP, glyceraldehyde 3-phosphate; DHAP, dihydroxyacetone phosphate; 3PG, 3-phosphoglycerate; PEP, phosphoenolpyruvate; Pyr, pyruvate; Ala, alanine; Cit, citrate; AKG, \(\alpha\)-ketoglutarate; Succ, succinate; Fum, fumarate; Mal, malate.

tissues we examined the usage of glutamine and succinate, which are oxidized by RPE mitochondria.\(^{52,53}\) Glutamine usage in eyecup and retina explants was measured with \(^{12}\)C-glucose and U-\(^{13}\)C-glutamine. Metabolite pool size was diminished in aged eyecups (Supplementary Fig. S5A) in glutamine, glutamate, and the downstream Krebs cycle metabolites \(\alpha\)-ketoglutarate, fumarate, malate, and aspartate. Because we did not observe this decline in eyecups provided only glucose (see Fig. 3), we hypothesized that aged eyecups may have a defect in mitochondrial glutamine metabolism. After incubation in \(^{13}\)C-labeled glutamine, M5 glutamine, glutamate, and \(\alpha\)-ketoglutarate.
FIGURE 4. Metabolic activity with glutamine – and in the mitochondria more broadly – was examined by incubating eyecup explants in U-13C-glutamine for 20 or 90 minutes. The M4 and M5 labeled isotopologues entering into and proceeding through the Krebs cycle (A) were quantified in terms of pmol of 13C-labeled isotopologue per μg of protein in eyecup explants (B–G). M5 glutamine B, glutamate C, and AKG D trended lower in the aged eyecups compared to young at both time points examined, although glutamine did not reach statistical significance at 90 minutes. In the downstream intermediates fumarate E, malate F, and aspartate G this decline was reproduced, but did not reach statistical significance. The percent incorporation of 13C remains essentially unchanged in aged eyecups at both time points in all intermediates examined (H–M). Young percent 13C incorporation rises past the aged in glutamate I and AKG J. However, in glutamine H, fumarate K, malate L, and aspartate M, the percent of 13C incorporation was consistently lower in young eyecups at 20 minutes, but had essentially matched the aged by 90 minutes. These changes can be related back to pool size and product:reactant ratios, which are included in Supplementary Figure S5. Normality of data was determined using the Shapiro-Wilk test and P values were calculated for age-related comparisons using Mann-Whitney tests (* = P < 0.05). Error bars represent the standard deviation.

(Fig. 4B–D) were significantly lower in aged eyecups. Levels of M4 fumarate, M4 malate, and M4 aspartate (see Fig. 4E–G) trended lower in aged eyecups compared to young. The percent labeled with 13C (see Fig. 4H–M) varied by metabolite. Notably, M5 glutamine, M4 fumarate, M4 malate, and M4 aspartate were lower in young eyecups at 20 minutes, but essentially matched the aged by 90 minutes. The product:reactant ratio of M5 glutamate/M5 glutamine (see Supplementary Fig. S5B) decreased with age. These data suggest glutamine anaplerosis is diminished in aged eyecups.

The same analysis was performed in retinas. In contrast, no significant age-related changes in pool size or 13C-labeled intermediates (see Supplementary Figs. S6A, S6C) were observed. Percent incorporation at 20 minutes was higher in later Krebs cycle intermediates of aged retinas, but younger retina label uptake nearly matched aged by 90 minutes (see Supplementary Fig. S6B), although it was still lower. No significant changes in product:reactant ratios (see Supplementary Fig. S6D) were observed.

We postulated the decline in eyecup glutamine metabolism was suggestive of reduced mitochondrial function, which we approximated by measuring OCR and ATP levels in young and old retinas and eyecups. Succinate is a mitochondrial respiratory substrate preferentially oxidized by the RPE.52 Succinate-driven OCR was calculated for each animal by taking the maximum steady-state OCR of tissue respiring on 5 mM glucose + 5 mM succinate and subtracting the 5 mM glucose baseline OCR (Supplementary Fig. S7A) and retinas (see Supplementary Fig. S7B). Steady-state OCR values are listed in Supplementary Table S1. As reported previously, succinate stimulates OCR in eyecups to a significantly greater degree compared to the retina.52 Neither retinas nor eyecups exhibited significant
age-related changes in succinate-stimulated \( \text{O}_2 \) consumption (see Supplementary Fig. S7C, S7D). Steady-state levels of ATP (see Supplementary Fig. S7E) also showed negligible changes.

**DISCUSSION**

Aging reduces retinal ERG response in humans\(^{25,26} \) and mice.\(^{18,19,21,24} \) We sought to examine responses in 26 month old C57BL/6j mice, at such time that about 25% of the population would have died\(^{54,55} \) and are roughly equivalent to 79 year old humans.\(^ {56} \) Previous trends were confirmed – photopic temporal resolution and both rod and cone electrical responses to light declined relative to young mice (see Fig. 2). Although there are confounding factors associated with the measurement of the ERG, other work utilizing optokinetic tracking and visual discrimination tasks found diminished visual acuity.\(^{27-29} \) We hypothesized this diminished response would be concurrent with reduced glucose metabolism and mitochondrial function in aged retina and eyecup explants.

Rapid aerobic glycolysis in the retina is required for visual function.\(^{32,51} \) Despite reduced ERG responses, retinal glucose and glutamine metabolism is stable ex vivo. Both glycolytic and Krebs cycle intermediates (see Fig. 3, Supplementary Figs. S2, S3) were essentially unchanged between young and aged retinas. Although glycolysis had reached a steady-state and thus we could not measure flux within tissues, the \( ^{13} \text{C} \) lactate exported into culture medium can approximate the rate of glycolysis. This complementary measurement was also unchanged with age. There was no observed effect of sex (Supplementary Fig. S4), and mitochondrial OCR and ATP levels were also unchanged. Glutamine metabolism is linked to synaptic transmission through the neurotransmitter glutamate\(^ {57} \) and additional experiments will be required to determine if the small differences with aging in retinas (see Supplementary Fig. S6) translate to changes in neurotransmission. We postulate that aging retina is robust and may retain metabolic function on par with young retinas despite significant physiological alterations in vivo.

Glucose metabolism appears to be preserved in aged eyecups (see Fig. 3, Supplementary Figs. S3, S4). However, glutamine metabolism in aged eyecups differs from young. Although our sample size \( (n = 4–5) \) leaves subtle differences challenging to identify, pool size (see Supplementary Fig. S5) and isotopologues decreased with age. However, the percent of \( ^{13} \text{C} \) incorporation into glutamine intermediates (see Fig. 4) remained stable in aging eyecups compared to the increasing percent in young eyecups. We hypothesize this indicates aged eyecups are deficient in glutamine metabolism and reach their maximum metabolic capacity faster than their younger counterparts. While the decreased M5 glutamate:M5 glutamine ratio could be indicative of decreased glutaminase activity in aging RPE, enzyme activity assays from glutaminase to fumarase would allow more targeted identification of the source of the defect in glutamine metabolism that we observed.

Finally, we considered that mitochondria have diverse functions outside cellular respiration.\(^ {58-60} \) Abundance of Krebs cycle intermediates provide a direct link to epigenetic changes.\(^ {61,62} \) Mitochondrial dysfunction is also involved in cellular signaling and aging having broad implications in an organ system.\(^ {51,59,63-66} \) Although the ocular tissues we examined – especially retina – may be resilient to age-related metabolic defects, future research will be critical to clarify how other functions of mitochondria in retina and RPE influence the broader ocular ecosystem in aging.

**Study Limitations and Considerations for Future Studies**

That glucose metabolism was relatively robust with age was unexpected compared with other aged tissues. Metabolic changes and mitochondrial dysfunction are hallmarks of aging.\(^ {51} \) Glycolytic capacity declines in the aged brain and effector T cells.\(^ {57-70} \) Young hearts subsist on fatty acid oxidation, but switch to glycolysis and ketone bodies in aging and heart failure.\(^ {71,72} \) Liver and skeletal muscle experience metabolic shifts.\(^ {73-78} \)

Our experimental design and technique influenced our findings. Subsequent experiments in our laboratory\(^ {52} \) have suggested glycolytic flux is best examined in earlier time points. Although we can approximate glycolytic flux in this study by examining lactate export in the culture medium (see Figs. 3C, 3D), a direct examination of time points of 2 minutes and below will be necessary. The explants in our analyses were exposed to physiological levels of glucose, but oxygen levels were nonphysiological and, unlike conditions within an eye, there were few barriers to transport. Blood vessel function around the retina declines in aging people\(^ {79-81} \) and \( \text{O}_2 \) extraction is lower in aged human retina,\(^ {82} \) which may indicate that nutrient and \( \text{O}_2 \) access is restricted in an aged mouse. Aged retina and eyecup appear to maintain most of their metabolic function ex vivo. However, in the eye of an aged animal, they may experience non-cell autonomous dysregulation of metabolism. Previous studies\(^ {83,84} \) using this ex vivo approach identified intrinsic, cell autonomous differences between the groups.

An in vivo analysis of metabolic flux will be necessary to establish whether metabolism is altered within the aging eye, and whether it would be due solely to intrinsic properties of the tissues or related to nutrient access. Direct infusion of \( ^{13} \text{C} \) labeled fuels\(^ {85} \) would be an effective way to test this. The analyses described in this report can be a starting point to a more physiological assessment of glycolysis, mitochondrial activities, and the access that retinas have in vivo to fuels. Our study focused on fuels of known importance in retina and RPE, but other metabolic pathways also need to be evaluated.\(^ {52,53,86-89} \)

Although the loss of capacity for metabolic flux appears to be less than the decrement in ERG responses, we do not have a way to unambiguously relate those observations. ERG a-wave and b-wave response amplitudes reflect not only the electrical activities of photoreceptors and downstream neurons, but also the overall electrical resistance between the cornea and the reference electrode placed in the body of the animal. Measurements of visual acuity using motion-tracking approaches are an effective way to evaluate visual function in the aging eye.

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