Photoreceptors in a mouse model of Leigh syndrome are capable of normal light-evoked signaling

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Mitochondrial dysfunction is an important cause of heritable vision loss. Mutations affecting mitochondrial bioenergetics may lead to isolated vision loss or life-threatening systemic disease, depending on a mutation’s severity. Primary optic nerve atrophy resulting from death of retinal ganglion cells is the most prominent ocular manifestation of mitochondrial disease. However, dysfunction of other retinal cell types has also been described, sometimes leading to a loss of photoreceptors and retinal pigment epithelium that manifests clinically as pigmentary retinopathy. A popular model of mitochondrial disease that lacks NADH:ubiquinone oxidoreductase subunit S4 (NDUFS4), a subunit of mitochondrial complex I, phenocopies many traits of the human disease Leigh syndrome, including the development of optic atrophy. It has also been reported that ndufs4−/− mice display diminished light responses at the level of photoreceptors or bipolar cells. By conducting electretroretinography (ERG) recordings in live ndufs4−/− mice, we now demonstrate that this defect occurs at the level of retinal photoreceptors. We found that this deficit does not arise from retinal developmental anomalies, photoreceptor degeneration, or impaired regeneration of visual pigment. Strikingly, the impairment of ndufs4−/− photoreceptor function was not observed in ex vivo ERG recordings from isolated retinas, indicating that photoreceptors with complex I deficiency are intrinsically capable of normal signaling. The difference in electrophysiologial phenotypes in vivo and ex vivo suggests that the energy deprivation associated with severe mitochondrial impairment in the outer retina renders ndufs4−/− photoreceptors unable to maintain the homeostatic conditions required to operate at their normal capacity.

Mitochondria house the enzymatic machinery for oxidative phosphorylation (OX-PHOS)3 and serve as the major source of cellular ATP production in most eukaryotic cells. Although mitochondrial DNA (mtDNA) encodes a minority of the subunits of OX-PHOS protein complexes, most subunits are encoded by the cell’s nuclear DNA and trafficked post-translationally to the mitochondria (1). Disruption of the enzymatic reactions of OX-PHOS not only reduces the efficiency of ATP production but also leads to an increase in reactive oxygen species, which destabilize the mitochondrial genome and damage proteins and lipids throughout the cell, potentially leading to apoptosis (2). Retinal neurons are highly sensitive to mitochondrial dysfunction. Inherited deficiencies in mitochondrial OX-PHOS enzymes may result in vision loss, often due to death of retinal ganglion cells (RGCs) and subsequent atrophy of the optic nerve, which comprises RGC axons (3). Disease of the outer retina in the form of pigmentary retinopathy has also been described in some forms of mitochondrial disease, but whether this reflects a primary pathologic of the retinal photoreceptors or of supporting retinal pigment epithelium (RPE) cells is uncertain (4, 5).

The severity of human disease related to mitochondrial dysfunction is highly variable. On one end of the spectrum is Leber hereditary optic neuropathy (LHON), which is among the most common human diseases with a mitochondrial inheritance pattern (6). Although it causes severe vision loss from optic atrophy, systemic manifestations are rare, and patients can typically expect to have a normal lifespan (5). Most LHON cases result from hypomorphic mutations in mtDNA encoding three of the 14 core subunits of complex I of OX-PHOS (ND1, ND4, and ND6) and reducing complex I catalytic activity to varying degrees (7, 8). At the other end of the mitochondrial disease spectrum is Leigh syndrome, a rare neurometabolic disease manifesting as hypotonia and ataxia in early childhood, with progressive cardiac and respiratory failure often leading to death within a few years of life (9). Visual impairment secondary to optic atrophy and pigmentary retinopathy is commonly noted during infancy (10). Some mutations associated with

3 The abbreviations used are: OX-PHOS, oxidative phosphorylation; NDUFS4, NADH:ubiquinone oxidoreductase subunit S4; ERG, electroretinography; LHON, Leber hereditary optic neuropathy; mtDNA, mitochondrial DNA; RPE, retinal pigment epithelium; RGC, retinal ganglion cell; cd, candelas( ); P, postnatal day; PKC, protein kinase C; CNG, cyclic nucleotide–gated channel; ChAT, choline acetyltransferase; Rmax, maximal response amplitude; 0.5, half-saturating light intensity.
Leigh syndrome affect subunits of complex I, and they likely produce more profound dysfunction of the enzyme complex than those mutations associated with LHON (11).

Genetic mouse models of complex I dysfunction have produced phenotypes similar to their human counterparts, also demonstrating a dependence on the severity of OX-PHOS impairment. For instance, a mouse line carrying a hypomorphic P25L mutation in the ND6 subunit of complex I resulting in ~30% reduction in complex I activity had no reduction in lifespan but demonstrated a delayed RGC degeneration phenotype manifesting by 24 months of age (12). In contrast, a mouse line with a deletion of the accessory complex I subunit encoded by the nuclear gene ndufs4 demonstrated severe systemic pathology due to profound instability of complex I as a whole (13). Similar to Leigh syndrome patients harboring ndufs4 mutations, homozygous ndufs4−/− mice developed a rapidly progressive myocencephalopathy, with animal death commencing around postnatal day 50 (P50). Although these mice have normal RGC counts initially, RGC degeneration was found to begin prior to death of the mice (14). Interestingly, both of these mutant mouse models were reported to demonstrate abnormal electroretinography (ERG) responses, suggesting pathology of preganglionic retinal neurons. However, histologic analysis of the outer retina has not yet been described, and it is unclear whether photoreceptor degeneration is a feature of these mouse models of complex I deficiency. We now demonstrate that the retinal signaling anomalies observed in ndufs4−/− mice can be completely rescued by altering the retinal extracellular environment and, therefore, are not due to irreversible photoreceptor dysfunction but rather to abnormal homeostatic conditions in the outer retina.

Results

ndufs4−/− mice produce abnormal in vivo ERG responses

Retinas from ndufs4−/− mice have previously been shown to have >80% reduction in complex I activity and a corresponding severe reduction in oxidative metabolism (14). The impact of this metabolic deficiency on retinal signaling has been assessed by analyzing ERG b-waves, which were profoundly reduced (13, 14). As the recordings had been performed as early as P21, this raised the question of whether lack of NDUF54 resulted in aberrant retinal development or an early degenerative process. To explore these possibilities, we performed comprehensive in vivo ERG analysis of ndufs4−/− mice and their WT littermates at multiple ages (Fig. 1). Light responses were recorded from dark-adapted mice (Fig. 1A) and in the presence of 30 cd/m² illumination saturating rod responses (Fig. 1B). The data were fit using a double-hyperbolic function as in Herrmann et al. (15) (Table 1). In contrast to the findings of Kruse et al. (13), we observed robust ERG b-waves in P22 ndufs4−/− mice under both scotopic and photopic conditions. The b-wave amplitudes were modestly reduced compared with WT, although this difference did not reach statistical significance at most flash intensities (Fig. 1C). Similarly, modest reductions in the amplitude of the ERG a-wave and in the responses to flashes in the presence of rod-saturating light were observed in ndufs4−/− mice at P22. These amplitude reductions were more pronounced at P36 (Fig. 1D) and then appeared to plateau, showing a similar reduction at age P47, the last time point tested within the lifespan of the mice (Fig. 1E). The average maximal ERG b-wave and a-wave responses at P36 and P47 in ndufs4−/− mice were all roughly half of the maximal responses in the WT littermates (Table 1).

ndufs4−/− mice display normal outer retina morphology without degeneration

Because ERG a-waves originate from photoreceptors and b-waves reflect primarily downstream responses of ON-bipolar cells, the proportionate reduction of ERG a- and b-wave amplitudes observed in ndufs4−/− mice of all ages points to a signal-amplification abnormality at the level of photoreceptors. This could either be due to abnormal phototransduction or to a reduction of the total number of photoreceptors. Morphological analysis of ndufs4−/− retinas by light and EM revealed healthy appearing photoreceptors with well-ordered outer segments at all ages, indicating no deficiency in outer segment morphogenesis or maintenance (Fig. 2, A and B). Furthermore, expression of several representative outer segment proteins participating in or regulating rod phototransduction was unchanged in ndufs4−/− retinas compared with WT (Fig. 2C). We also confirmed proper compartmentalization of rod photoreceptors, as rhodopsin and the β1-subunit of the cyclic nucleotide–gated channel (used as representative proteins of outer segment discs and plasma membrane, respectively) retained their normal localization in ndufs4−/− retinas, whereas the inner segment marker Na⁺/K⁺-ATPase remained excluded from outer segments (Fig. 2D).

Consistent with the lack of gross morphological abnormalities, there was no evidence of photoreceptor degeneration, as the number of photoreceptor nuclei within the outer nuclear layer in ndufs4−/− mice was unchanged compared with WT littermates, even at P47 (Fig. 3A). Furthermore, there was no significant difference in mean outer segment length at this late time point (p = 0.60; Fig. 3B). It has been reported that rod bipolar cells are the first retinal neurons to degenerate in ndufs4−/− mice (16), which could theoretically explain the reduction in ERG b-wave amplitudes, at least under scotopic conditions. However, in contrast to this previous report, when we immunolabeled retinal cross-sections at P49 with the rod bipolar cell marker PKC-α, no difference was found in rod bipolar cell count between ndufs4−/− and WT retinas (p = 0.71; Fig. 3C). This lack of detectable rod bipolar cell pathology further supports our conclusion that the ERG defect in ndufs4−/− mice originates primarily from photoreceptors. We confirmed a previously reported reduction in the number of choline acetyltransferase–positive starburst amacrine cells in ndufs4−/− retinas (p < 0.01; Fig. 3D) (14). However, prior ERG analysis of mouse retinas in which starburst amacrine cells were ablated revealed no effect on a-wave amplitude and minimal effect on b-wave amplitude (≤10% reduction and only at the highest stimulus intensities) (17). Thus, loss of starburst amacrine cells would seem to contribute little to the decreased ERG amplitudes of ndufs4−/− mice.

Because degeneration of preganglionic retinal neurons did not appear to explain the electrophysiology phenotype of ndufs4−/− mice, we asked whether these findings could be...
Retinal signaling in NDUFS4-deficient mice

A P47  
dark-adapted

B P47  
30 cd/m² background

C P22

D P36

E P47

scotopic b-wave amplitude (µV)

flash intensity (log cd·s/m²)

photopic b-wave (µV)

flash intensity (log cd·s/m²)

wt

ndufs4⁻/⁻
related to pathology within the RPE. The RPE plays important roles in supporting photoreceptors, including delivery of metabolic substrates from the choroidal circulation and regeneration of the 11-cis-retinal chromophore. We therefore assessed morphologic signs of RPE pathology by EM. No difference in RPE cell thickness was found, and although the posterior segments of some ndufs4−/− eyes demonstrated occasional RPE cells with large cytoplasmic vacuoles (Fig. 4A), these were not observed in every mouse, whereas diminished ERG amplitudes were observed in every ndufs4−/− mouse studied. No overt differences in RPE mitochondrial morphology or organization of cristae were observed. However, ndufs4−/− RPE cells displayed a higher density of mitochondria than WT (1.01 ± 0.06 versus 0.65 ± 0.01 mitochondria μm−2, respectively; p < 0.001), as observed previously in other cell types of humans (18) and mice (12) with complex I dysfunction.

We next determined whether the ERG abnormalities reflected an overall decrease in photoreceptor responses due to reduced regeneration of 11-cis-retinal by RPE by measuring the retinal content of rhodopsin using difference spectroscopy. However, in concordance with our Western blot analysis (Fig. 2C), no difference in rhodopsin content was noted between P47 ndufs4−/− and WT retinas after 6 h of dark adaptation, the same duration of dark adaptation given to animals prior to ERG testing (p = 0.21; Fig. 4B).

**The ERG defect in ndufs4−/− mice can be rescued by modulating the extracellular environment**

The lack of any overt pathology in ndufs4−/− photoreceptors may suggest that ERG defects observed in these mice derive from insufficient energy production within photoreceptors or from dysfunction of neighboring cell types adversely affecting their extracellular environment. One possibility, previously described for MCT-3 lactate transporter-deficient mice (19), is decreased uptake of photoreceptor-derived lactic acid by the RPE. We explored this hypothesis by measuring the lactate content in isotonic washes of freshly dissected retinas. However, we observed no difference in the amount of lactate recovered from ndufs4−/− retinas compared with WT controls after washes of 2 (p = 0.32) or 10 min (p = 0.83) (Fig. 4C), suggesting that pathological acidification of the interphotoreceptor matrix from accumulation of lactic acid was unlikely to occur in the mutant mice.

To further explore the role of the extracellular environment in the ERG phenotype of ndufs4−/− mice, we performed ex vivo ERG recordings from isolated P41 retinas equilibrated in Ames’ medium. Both retinas from five male ndufs4−/− mice and four male WT mice were recorded. To examine whether the deletion of NDUF54 affects the rod phototransduction cascade, we recorded families of transretinal responses to test flashes of increasing light intensities (Fig. 5). The presence of postsynaptic inhibitors in the perfusion solution blocked contributions of higher-order response components (such as ERG b-waves driven by ON-bipolar cells), allowing isolation of the pure rod photoreceptor response (20).

Strikingly, in contrast to the ERG results obtained in vivo, retinas of dark-adapted control and NDUF54-deficient mice produced rod light responses of comparable amplitudes (Fig. 5, A and B). The dim-flash response kinetics for ndufs4−/− and control rods were also comparable, with average time-to-peak values of 128 and 130 ms, respectively. The activation kinetics of rod phototransduction, measured from the rising phase of the dim-flash response, was unaffected by NDUF54 deletion (Fig. 5C). The initial phase of response inactivation was also unaltered in ndufs4−/− rods, whereas the very late phase of the recovery revealed a slight acceleration. Surprisingly, the recovery following saturating flashes was moderately slower in the mutant mice, as evident from the average kinetics of maximal rod responses (Fig. 5D). The molecular mechanisms for these subtle changes in kinetics are unclear, although they might be related to small, long-term compensatory changes in expression and/or function of phototransduction proteins due to the altered metabolic state of the retina. Notably, the average rod photosensitivity (defined as a half-saturating light intensity (I0.5)) in mutant mice was not statistically significantly different from that in WT animals (23 ± 3 versus 29 ± 5 photons μm−2, respectively; p > 0.05) (Fig. 5, E and F).

Taken together, these results indicate that the lack of the NDUF54 subunit in mouse rods impairs their signaling in vivo but not ex vivo. Thus, the intrinsic capacity of mutant photoreceptors to respond to light is preserved and can be restored under ex vivo conditions when the retinal tissue is supplemented with proper nutrients to maintain its function. This further suggests that mutant photoreceptors do not undergo major irreversible changes. Rather, the energy deprivation in the outer retina either restricts their energy supply or modifies the molecular or ionic composition of the interphotoreceptor matrix, such that rods and cones cannot operate at their normal capacity.

**Discussion**

Our electrophysiological characterization of ndufs4−/− mice in vivo has confirmed the presence of abnormal retinal signaling and further revealed that ERG a- and b-waves are proportionally diminished in these mice, implicating photoreceptors as the primary origin of this visual impairment. Our data suggest that this phenomenon cannot be explained by an irreversible process, such as aberrant photoreceptor development or photoreceptor degeneration. Instead, ndufs4−/− photoreceptor function can be completely restored by bathing isolated retinas in standardized recording media, suggesting that photoreceptors with diminished capacity for oxidative metabolism are

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**Figure 1. Reduced rod and cone light responses in ndufs4−/− mice in vivo.** A, representative traces from ERG recordings of dark-adapted WT and ndufs4−/− mice at P47 with stimulus intensities ranging from 0.0001 to 500 cDs/m2. Black arrowhead denotes the application of the light stimulus. B, representative photopic light responses in P47 WT and ndufs4−/− mice, recorded at the three indicated stimulus intensities. C–E, ERG responses of dark-adapted WT and ndufs4−/− mice at P22, P36, and P47 are plotted as a function of flash intensity and fit using a double- or single-hyperbolic function. The left panels show scotopic b-wave amplitudes, middle panels show scotopic a-wave amplitudes, and right panels show photopic b-wave amplitudes. Four eyes were analyzed for each genotype at P22 and P36, and six eyes were analyzed per genotype at P47; Error bars represent S.E. Results of two-tailed t tests between responses from WT and ndufs4−/− mice at each flash intensity are presented as follows: *, p < 0.05; **, p < 0.01; ***, p < 0.001. See Table 1 for fitting parameters.
Table 1

Fitting parameters for in vivo ERG recordings and statistical analysis of the differences among selected groups

| Group | P22 KO | WT KO | P22 WT | WT KO |
|-------|--------|-------|--------|-------|
| Rmax | 0.590 | 0.860 | 0.570 | 0.860 |
| I0.5 | 0.090 | 0.070 | 0.150 | 0.170 |
| n    | 0.700 | 0.800 | 0.040 | 0.080 |
| p    | 0.0001 | 0.0001 | 0.0001 | 0.0001 |

There is precedence for mice with morphologically normal photoreceptors producing abnormal ERG responses in vivo. Examples of such observations include mice with deletions of MCT-3 (19), phospholipase C β4 (21), and carbonic anhydrase XIV (22). Notably, the first two cases also demonstrated normal ex vivo light-evoked electrical responses (suction electrode recordings of isolated rods in both cases), whereas the carbonic anhydrase–knockout mouse was not further characterized in this context. None of these mouse lines suffers from a severe systemic phenotype, in contrast to the ndufs4<sup>−/−</sup> mice used in the present study, and they continue to demonstrate normal photoreceptor morphology well into adulthood. In the cases of MCT-3 and carbonic anhydrase XIV, neither protein is expressed by photoreceptors but instead by RPE (MCT-3) and/or Müller glia (both proteins). Loss of MCT-3 results in accumulation of extracellular retinal lactate (19), likely due to the inability of excess lactate to traverse the basolateral membrane of the RPE into the choroidal circulation. Although not directly tested, reduced efflux of lactic acid would be predicted to result in acidification of the interphotoreceptor matrix. Similarly, carbonic anhydrase has been shown to buffer the pH of the interphotoreceptor matrix, with its inhibition resulting in acidification of this compartment (23). Interphotoreceptor matrix acidification might explain the abnormal in vivo ERG responses in both mutations, as low pH is known to decrease the activity of the rod outer segment Na<sup>+</sup>/Ca<sup>2+</sup>/K<sup>+</sup> exchanger NCKX1 (24), which maintains Ca<sup>2+</sup> homeostasis in the outer segment and is critical in supporting normal rod sensitivity and response amplitude (25). The mechanism behind reduced ERG amplitudes in phospholipase C β4–knockout retinas is less clear but may also be related to altered Ca<sup>2+</sup> homeostasis.

In this context, what could explain why ndufs4<sup>−/−</sup> photoreceptors produce diminished light responses in vivo but normal responses ex vivo? One possibility is that the mutant photoreceptors themselves are incapable of generating enough energy to support electrochemical gradients of ions involved in producing electrical light responses. This could be compensated in the recording medium by 1) providing photoreceptors with an essentially unlimited supply of glucose and 2) providing constant extracellular ion conditions, thereby reducing the burden of establishing ion gradients across the cell membrane. Indeed, an emerging understanding of metabolic flux within vertebrate retina indicates that photoreceptors rely on aerobic glycolysis rather than OX-PHOS as their primary energy source (26, 27), suggesting that an unlimited glucose supply may compensate for reduced mitochondrial function.

Another, not mutually exclusive, possibility is that abnormal energy metabolism within other neighboring cell types contributes to in vivo pathology. One plausible hypothesis is related to the role of the RPE in supporting the flux of glucose and lactate between the choroid and the retina. RPE cells serve as a conduit for glucose entry to the retina by expressing the facilitative glucose transporter GLUT1 at both the basolateral and apical membranes (28) and for efflux of retinal lactate by expressing the facilitative lactate transporters MCT-1 and MCT-3 on the apical and basolateral membranes, respectively (29). Impor-
tantly, in addition to providing a route for retinal lactate to exit into the choroidal circulation, RPE cells also have a high capacity to use photoreceptor-derived lactate as a substrate for oxidative metabolism (26). In fact, metabolizing lactate allows the RPE to spare glucose passing from the choroid to the interphotoreceptor matrix (26). Once there, the glucose can then be taken up by photoreceptors via GLUT1 in the inner segments (30). Impaired OX-PHOS within the

\textit{ndufs4}\textsuperscript{-/-} RPE could lead to a reduced ability to metabolize lactate and a greater reliance on glycolysis, thereby limiting glucose delivery to photoreceptors. Bathing the \textit{ndufs4}\textsuperscript{-/-} retinas in recording media eliminates their reliance on RPE for glucose supply, thus potentially explaining the restoration of normal photoreceptor signaling \textit{ex vivo}. Testing this hypothesis would require generation of transgenic mice with cell-specific deletion of \textit{ndufs4} in photoreceptors or RPE to determine which cell type is primarily responsible for the \textit{in vivo} ERG phenotype.

It is instructive to compare our results with those addressing the retinal phenotype of the \textit{agc1}\textsuperscript{-/-}aralar\textsuperscript{-/-} mouse also suffering from impaired OX-PHOS (31). AGC1/Aralar is the neuronal isoform of the mitochondrial aspartate–glutamate carrier, and its absence makes affected cells unable to efficiently regenerate NAD\textsuperscript{+} via oxidative metabolism due to impairment of the malate–aspartate shuttle on the mitochondrial membrane (32). Because cytosolic NAD\textsuperscript{+} is required for additional rounds of glycolysis, pyruvate must be reduced to lactate in the cytosol of these cells to regenerate NAD\textsuperscript{+}, rather than entering mitochondria as a substrate for the Krebs cycle. Similar to \textit{ndufs4}\textsuperscript{-/-} mice, \textit{agc1}\textsuperscript{-/-}aralar\textsuperscript{-/-} mice develop both systemic (32) and retinal (31) phenotypes. These mice also display a marked 40–50% reduction in a- and b-wave ERG amplitudes recorded from dark-adapted animals despite no evidence of retinal morphologic abnormalities. Their ERG responses were partially improved after brief exposure to background illumination. Because light adaptation is thought to reduce the metabolic demand of photoreceptors (33), this result suggests that the mutant photoreceptors function at the limit of their metabolic capacity. It would be interesting to determine whether the \textit{agc1}\textsuperscript{-/-}aralar\textsuperscript{-/-} mouse exhibits restoration of normal signaling in isolated retinas and whether this gene is expressed in both photoreceptors and RPE.

Finally, with regard to the applicability of our findings to human patients, there is scant literature available on ERG assessments of outer retinal function in patients with Leigh syndrome and LHON. Given that pigmentary retinopathy is known to develop in some patients with Leigh syndrome, abnormal ERG responses consistent with retinal dystrophy would be expected, and one study reported this finding in 23% of children with Leigh syndrome (10). However, the patients in this report had mutations in genes other than \textit{ndufs4}, so the
prevalence of ERG abnormalities in this particular genotype remains to be determined. Among rare reports of outer retinal functional assessments in LHON, an asymptomatic mother carrying the 11778 mtDNA mutation in the ND4 complex I subunit and her affected adolescent son both demonstrated full-field ERG abnormalities suggestive of cone dysfunction (34). Another study reported one of two patients with the 11778 mutation demonstrating an abnormal ERG suggestive of rod–cone dystrophy (35). A separate report on an extensive Brazilian pedigree of the 11778 mutation reported that multifocal ERG abnormalities were commonly noted among asymptomatic carriers, although it was not specified whether this reflected photoreceptor dysfunction in addition to inner retinal pathology (36). It would be important to gain further insight into the degree of visual impairment in these patients, such as decreased ability to see under dim-light conditions and to recognize low-contrast objects.

Experimental procedures

Animals

Mice heterozygous for a deletion of exon 2 of ndufs4 on a C57BL/6 genetic background (13) were purchased from The Jackson Laboratory (stock number 027058) and were free of RD1 and RD8 mutations. Heterozygous mice were crossed to generate homozygous ndufs4−/− animals and WT littermates as controls. Animals were reared under a normal day/night cycle and handled according to a protocol approved by the Institutional Animal Care and Use Committee of Duke University. All experiments were performed during the day.
indicate mean (1:1000; Santa Cruz Biotechnology, sc-47778, lot number C10C10 (1:1000; Ref. 37), goat polyclonal anti-CNG-

clonal anti-NDUFS4 1-E-4 (1:200; Santa Cruz Biotechnology, used for Western blot analysis of retinal lysates: mouse mono-

tler, University of Alabama, Birmingham, AL), mouse mono-

dopsin kinase (GRK1) G-8 (1:500; Santa Cruz Biotechnology, sc-8143, lot number L1205), and mouse monoclonal anti-rho-

gated cation channel (CNG)-Abcam, ab98887), rabbit polyclonal anti-cyclic nucleotide-

orescence experiments; 1:500 dilution) or Alexa Fluor 680 appropriate species conjugated to Alexa Fluor 488 (immunoflu-

gated retinas. In separate experiments, WT and

cgenotype. Scale bar

Figure 4. Retinal pigment epithelium morphology and function in ndufs4−/− mice. A, electron micrographs of RPE cells at P47. Although most ndufs4−/−: RPE (middle panel) had a morphology similar to WT (left), occas-

Figure 4. Retinal pigment epithelium morphology and function in ndufs4−/− mice. A, electron micrographs of RPE cells at P47. Although most ndufs4−/−: RPE (middle panel) had a morphology similar to WT (left), occas-

cellular vacuoles. Scale bar, 2 μm. B, rhodopsin content of isolated WT and ndufs4−/− retinas at P47, as measured by difference spectroscopy. n = 6 retinas per genotype. C, quantification of lactate liberated from isotoxic washes of iso-

Antibodies

The following antibodies were used for immunofluorescence experiments: mouse monoclonal 4D2 against rhodopsin (1:100; Abcam, ab98887), rabbit polyclonal anti-cyclic nucleotidediated cation channel (CNG)-β1 (1:200; a gift from Steven Pit-

tler, University of Alabama, Birmingham, AL), mouse mono-

clonal anti-Na+/K+-ATPase (1:500; Santa Cruz Biotechnology, sc-58628, lot number I2413), mouse anti-PKC-α H-7 (1:100; Santa Cruz Biotechnology, sc-8393, lot number I1306), and goat polyclonal anti-choline acetyltransferase (1:100; Millipore, Santa Cruz Biotechnology, sc-8004, lot number A289). Secondary antibodies against the appropriate species conjugated to Alexa Fluor 488 (immunofluo-

Ex vivo ERG recordings were performed on isolated mouse retinas. After dark adaptation overnight, mice were sacrificed by CO2 asphyxiation. Each retina was removed from its eyecup under IR illumination and stored in oxygenated Ames’ medium (Sigma-Aldrich) at room temperature. The retina was mounted on filter paper with the photoreceptor side up and placed in a perfusion chamber (39) between two electrodes connected to a differential amplifier. The sample was perfused with bicarbon-
ate-buffered Ames’ medium (Sigma-Aldrich) supplemented with 40 μM L-2-amino-4-phosphonobutyric acid to block postsynaptic components of the photoresponse (40) and with 100 μM BaCl₂ to suppress the slow glial PIII component (41). The perfusion solution was continuously bubbled with a 95% O₂, 5% CO₂ mixture, and its temperature was maintained at 36–37 °C with a heater.

The retina was stimulated with 20-ms test flashes of calibrated 505-nm light-emitting diode (LED) light. The light intensity was controlled by neutral density filters and a computer in 0.5 log-unit steps. Intensity–response relationships were fitted with the following Naka–Rushton hyperbolic functions (Equation 3 for raw data and Equation 4 for normalized data).

\[
R = \frac{R_{\text{max}} \cdot I}{I + I_{0.5}} \tag{Eq. 3}
\]

\[
\frac{R}{R_{\text{max}}} = \frac{I}{I + I_{0.5}} \tag{Eq. 4}
\]

Photoresponses were amplified by a differential amplifier (DP-311, Warner Instruments), low-pass-filtered at 300 Hz (eight-pole Bessel), and digitized at 1 kHz. Data were analyzed with Clampfit 10.4 and Origin 8.5 software. All data were expressed as mean ± S.E. and analyzed with the independent two-tailed Student’s t-test (using an accepted significance level of \( p < 0.05 \)).

Figure 5. Normal rod light responses in isolated ndufs4⁻/⁻ retinas. A, representative family of rod responses from isolated WT mouse retina. Test flashes of 505-nm light with intensities of 0.5, 1.4, 4.8, 14, 33, 114, 392, and 1188 photons μm⁻² were delivered at time 0. B, representative family of rod responses from isolated ndufs4⁻/⁻ mouse retina. Test flashes of 505-nm light had the same intensities as in A. C, kinetics of rod phototransduction activation and inactivation in control and ndufs4⁻/⁻ mice. Population-averaged dim-flash responses to test stimuli of 4.8 photons μm⁻² (n = 8 for controls and n = 10 for mutants) were normalized to \( R_{\text{max}} \) of respective retinas. D, comparison of saturated rod responses from WT and ndufs4⁻/⁻ mice. Population-averaged responses to test stimuli of 1188 photons μm⁻² (n = 8 for controls and n = 10 for mutants) were normalized to \( R_{\text{max}} \) of respective retinas. E, averaged rod intensity–response functions for WT (n = 8) and ndufs4⁻/⁻ (n = 10) retinas. Points were fitted with hyperbolic Naka–Rushton functions as described under “Experimental procedures.” F, normalized averaged rod intensity–response relationships for control (n = 8) and ndufs4⁻/⁻ (n = 10) retinas. Naka–Rushton fits yielded \( I_{0.5} \) values of 29 and 23 photons μm⁻² for WT and ndufs4⁻/⁻ mice, respectively. The statistical significance of the data in E and F is represented as follows: *, \( p < 0.05 \); **, \( p < 0.01 \); ***, \( p < 0.001 \). All recordings were from P41 control and ndufs4⁻/⁻ mice. Error bars represent S.E. in all panels.
Retinal signaling in NDUFS4-deficient mice

Histological techniques

The eyes of euthanized mice were enucleated, and posterior eyecups were dissected and fixed for 1 h in 4% paraformaldehyde and rinsed in PBS. Agarose-embedded retinal cross-sections were prepared as described (42). 100-μm-thick cross-sections from the central portion of the retina were collected with a vibratome (Leica VT1200S) in 24-well plates, and floating sections were then incubated for 4 h with the blocking solution containing 5% goat serum and 0.03% Triton X-100 in PBS buffer on an orbital shaker. Sections were further incubated overnight with the corresponding primary antibodies, washed three times with PBS, incubated for 2 h with appropriate secondary antibody conjugated to Alexa Fluor 488 (1:500), and washed three times in PBS. Alternatively, retinal cryosections were prepared by cryoprotecting fixed eyecups in 30% sucrose and embedding them in optimal cutting temperature (OCT) medium (Tissue-Tek, Sakura Finetek). Retinal 20-μm cross-sections were collected using a cryostat microtome (Microm HM 550, Thermo Fisher Scientific). Sections were rehydrated with PBS, blocked with 5% goat or donkey serum in PBS with 0.03% Triton X-100 for 1 h, incubated in primary antibody in the same blocking solution overnight, washed three times, incubated in secondary antibodies in blocking solution, and then finally washed three more times. Both agarose sections and cryosections were mounted with Vectashield (Vector Laboratories) under glass coverslips. Images were acquired using a Nikon Eclipse Ti2 inverted confocal microscope, CFI Plan Fluor 20× or 60× (oil) objectives, and an A1 confocal scanner controlled by NIS-Elements software (Nikon). In the case of PKC-α– and choline acetyltransferase (ChAT)-stained cryosections, 45,000-μm² images spaced 500 μm on either side of the optic nerve head were obtained on three sections for each retina. The number of positive nuclei/μm² of inner nuclear layer or ganglion cell layer was averaged for each retina.

For experiments involving analysis of mouse retinas in plastic sections, the superior limbus of each eye was marked with cautery to facilitate proper orientation prior to enucleation and fixation in 2% glutaraldehyde and 2% paraformaldehyde. 1-μm plastic-embedded cross-sections of the mouse retina were prepared as described (43) and stained with toluidine blue for light microscopy. Nuclear counts in 100-μm segments of the outer nuclear layer were performed in sections cut through the optic nerve at 500-μm steps from the optic nerve head. The same specimens were also processed for transmission EM. Thin sections of 60–80 nm were collected on copper grids, counterstained with uranyl acetate and Sato’s lead, and examined using an electron microscope (JEM-1400, JEOL) at 60 kV. Images were collected using a charge-coupled device camera (Orius, Gatan).

Rhodopsin quantification

Rhodopsin concentration was determined by difference spectroscopy using the molar extinction coefficient 40,500 cm⁻¹ M⁻¹ (44, 45). Individual retinas were extracted under dim-red illumination and sonicated in 250 μl of deionized water. A 100-μl aliquot was mixed with 20 μl of 200 mM hydroxylamine, pH 7.5, containing 10% n-octyl β-D-glucopyranoside. The sample was centrifuged in a tabletop microcentrifuge, and rhodopsin concentration in the supernatant was determined from the absorbance at 500 nm before and after complete bleaching of the sample.

Retinal washes and extracts for lactate quantification and Western blotting

Measurements of extracellular lactate in WT and ndufs4−/− mouse retinas were performed, adapted from a previously described protocol (19). Briefly, dark-adapted mice were euthanized in the light and immediately enucleated. Posterior eyecups were dissected, and each retina was removed using forceps. Retinas were then placed in a microcentrifuge tube containing 50 μl of PBS and incubated for 1 or 9 min. The samples were centrifuged at 14,000 × g for 1 min, and the supernatant was removed for a lactate assay, performed in duplicates using a colorimetric lactate assay (Cell Biolabs). The pelleted retinas were then solubilized in 100 μl of lysis buffer (25 mM HEPES buffer, pH 7.4, 150 mM NaCl, 5 mM MgCl₂, and protease inhibitors (Complete Mini, Roche Applied Science) containing 1% Triton X-100), and the protein concentration of each lysate was determined with a colorimetric assay (Bio-Rad). The lactate concentration in the retinal wash was normalized to detergent-soluble protein concentration for each retina.

The detergent-soluble retinal lysates were used for Western blot analysis. After mixing with SDS-PAGE sample buffer, retinal lysates from WT and ndufs4−/− mice were separated on 4–20% SDS-PAGE gels (25 μg of protein per lane), transferred onto polyvinylidene fluoride (PVDF) membranes and blotted with the indicated primary and secondary antibodies. The blots were imaged using an Odyssey imaging system (LI-COR Biosciences).

Experimental design and statistical analysis

All histological experiments and the in vivo electrophysiological experiments were performed on ndufs4−/− mice and littermate controls with both sexes represented. The in vivo ERG recordings were performed at P22 (four eyes of two ndufs4−/− mice and four eyes of two WT mice), at P36 (four eyes of two ndufs4−/− mice and four eyes of two WT mice), and P47 (six eyes of three ndufs4−/− mice and six eyes of three WT mice). In the case of ex vivo ERGs, all recordings were performed on retinas from P41 male ndufs4−/− mice (10 retinas from five mice) and age-matched male C57BL/6 controls (eight retinas from four mice). Male mice were selected for this experiment because of the more severe visual dysfunction often seen in male human patients with complex I dysfunction.

To compare sensitivities of different experimental groups, a-wave and b-wave amplitudes recorded at each flash intensity were compared using a two-tailed t test in GraphPad Prism version 7.00 for Windows (GraphPad Software). No statistical methods were used to predetermine sample sizes, but sample sizes are similar to those used in previously published studies. No randomization was used, and animal genotypes were not masked.
Retinal signaling in NDUF54-deficient mice

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