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Journal

Molecular Neurobiology, 52(1)

ISSN

0893-7648

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Publication Date

2015-08-25

DOI

10.1007/s12035-014-8881-8

Peer reviewed

Robust Endoplasmic Reticulum-Associated Degradation of Rhodopsin Precedes Retinal Degeneration

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Received: 22 July 2014 / Accepted: 25 August 2014
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Abstract Rhodopsin is a G protein-coupled receptor essential for vision and rod photoreceptor viability. Disease-associated rhodopsin mutations, such as P23H rhodopsin, cause rhodopsin protein misfolding and trigger endoplasmic reticulum (ER) stress, activating the unfolded protein response (UPR). The pathophysiologic effects of ER stress and UPR activation on photoreceptors are unclear. Here, by examining P23H rhodopsin knock-in mice, we found that the UPR inositol-requiring enzyme 1 (IRE1) signaling pathway is strongly activated in misfolded rhodopsin-expressing photoreceptors. IRE1 significantly upregulated ER-associated protein degradation (ERAD), triggering pronounced P23H rhodopsin degradation. Rhodopsin protein loss occurred as soon as photoreceptors developed, preceding photoreceptor cell death. By contrast, IRE1 activation did not affect JNK signaling or *rhodopsin* mRNA levels. Interestingly, pro-apoptotic signaling from the PERK UPR pathway was also

not induced. Our findings reveal that an early and significant pathophysiologic effect of ER stress in photoreceptors is the highly efficient elimination of misfolded rhodopsin protein. We propose that early disruption of rhodopsin protein homeostasis in photoreceptors could contribute to retinal degeneration.

Keywords Endoplasmic reticulum-associated degradation · Rhodopsin · Retinal degeneration · Unfolded protein response · ER stress · Rod photoreceptor

Abbreviations

ERAD	ER-associated protein degradation
IRE1	Inositol-requiring enzyme 1
ONL	Outer nuclear layer
PERK	PKR-like endoplasmic reticulum kinase
qPCR	Quantitative real-time PCR
RP	Retinitis pigmentosa
RIS	Rod inner segment
ROS	Rod outer segment
UPR	Unfolded protein response
XBP-1	X-box-binding protein 1

Electronic supplementary material The online version of this article (doi:10.1007/s12035-014-8881-8) contains supplementary material, which is available to authorized users.

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Introduction

The endoplasmic reticulum (ER) is essential for secretory and membrane protein folding and assembly. Impairment of ER protein folding creates ER stress, activating the unfolded protein response (UPR) intracellular signal transduction network [1]. The UPR regulates transcriptional, translational, and post-translational intracellular programs such as ER-associated protein degradation (ERAD) whereby ER chaperones, protein folding enzymes, and lectins identify malformed proteins in the ER and retrotranslocate them into the cytosol for

degradation by proteasomes [2]. The ERAD program therefore prevents misfolded protein accumulation, and disruption of ERAD components often leads to early embryonic lethality [3–7].

Inositol-requiring enzyme 1 (IRE1) is a key UPR regulator [1]. IRE1 encodes an ER-resident transmembrane protein with an ER-luminal domain coupled to cytosolic kinase and endoribonuclease (RNase) domains [8–10]. In response to ER stress, IRE1's RNase removes an inhibitory intron from *Xbp-1* mRNA [11, 12] to generate the potent XBP1s transcription factor whose transcriptional targets include numerous ERAD genes [13, 14]. By this mechanism, IRE1 signaling regulates the ERAD program. In addition to ERAD, IRE1 signaling can also activate the c-Jun kinase signaling module and non-specifically degrade mRNAs through regulated IRE1-dependent decay (RIDD) in response to strong ER stress [1]. PKR-like ER kinase (PERK) mediates a separate UPR signaling pathway [1]. In response to ER stress, PERK inhibits the ternary translation initiation complex formation on mRNAs [15] and therefore attenuates protein synthesis in the cell. PERK signal transduction also regulates a cell death program through its upregulation of pro-apoptotic transcriptional activators: ATF4 and CHOP [16, 17].

Rod opsin mutations are common causes of heritable retinitis pigmentosa, a blinding disease arising from photoreceptor cell death [18]. Rod opsin is a G protein-coupled receptor that forms the visual photopigment, rhodopsin, when coupled with 11-*cis*-retinal [19]. Rhodopsin is essential for photoreceptor function and survival, and rod opsin knockout mice (*Rho*^{-/-}) develop early retinal degeneration [20, 21]. In cell culture, many rod opsin mutants misfold and aggregate to trigger ER stress and activate the UPR [22–26]. But, the effects and role of UPR activation in photoreceptor cell death remain obscure.

Here, we analyzed UPR signaling in our recently developed P23H rod opsin knock-in mouse model of retinitis pigmentosa [27] that recapitulated patients' gene dosage and spatiotemporal retinal degeneration [27]. By contrast, transgenic models of retinal degeneration produced retinal degeneration regardless of whether misfolded or wild-type rhodopsin was expressed [28]. Here, we found an unexpected bias in UPR signaling in photoreceptors, with strong IRE1 but little PERK signaling. The pathophysiologic consequence of IRE1 activation was robust ERAD elimination of almost all P23H rhodopsin from photoreceptors early after their formation to create a *Rho*^{-/-}-like retinal degenerative phenotype.

Materials and Methods

Animals

Homozygous breeding pairs of P23H rod opsin knock-in mice [27] were maintained to produce *Rho*^{P23H/P23H} homozygotes

or crossed with wild-type C57BL/6J (*Rho*^{+/+}) mice to produce *Rho*^{P23H/+} heterozygotes. C57BL/6J mice were used as wild-type controls (*Rho*^{+/+}). *ERAI*^{+/-} mice were provided by Takao Iwawaki (RIKEN, Saitama, Japan) and Peter Walter (UCSF). *Chop*^{-/-} mice were obtained from Jackson Laboratory. Mice were maintained in a barrier animal facility in a 12:12 light cycle at in-cage irradiance of less than 125 lx and provided standard mouse chow (UCSF and UCSD) or in a 12-h light (127–255 lx)/12-h dark (<10 lx) cyclic environment (Case Western Reserve University). Retinal tissues were collected from animals of either sex at indicated ages. Mice were checked daily by the UCSF Animal Care staff. In addition, laboratory personnel checked births in the morning and evening during the light phase of the light cycle to determine exact birthdates because early eye developmental and degenerative events occurred rapidly once they commenced. To define phenotypes of each of the three genotypes from P4 to P15, we examined two litters of mice of either sex with exact birthdates from each line, examining one animal of either sex from each litter on successive dates, beginning on P4 (Table S1). All animal studies followed the guidelines of the institutional animal care committees at UCSF, UCSD, or Case Western Reserve University, and were conducted in accordance with the recommendations of the American Veterinary Medical Association Panel on Euthanasia and the Association of Research for Vision and Ophthalmology.

Tissue Harvesting and Eye Histology

Immediately after euthanizing mice, an enucleated eye of each mouse was immersed in a fixative of mixed aldehydes for histology [29]. The other eye also was removed, and the retina was dissected by the Winkling procedure [30] and rapidly frozen in dry ice for biochemical and molecular analyses. For histological analysis, eyes were bisected, post-fixed with osmium tetroxide, and embedded in an epoxy resin as previously described [29]. One-micrometer-thick sections were cut through the optic nerve head along the vertical meridian and stained with toluidine blue. Beginning at P8, the outer nuclear layer (ONL) thickness, a surrogate for photoreceptor number [31], was quantified by measuring three positions in nine 250- μ m adjacent microscopic fields on each side of the optic nerve head that produced 54 measurements of each retina [32]. Means of these measurements allowed quantitative comparisons of multiple animals at different ages and of different genotypes in the two hemispheres of the eye, and the resulting values could then be plotted in a “retinal spidergram” to illustrate hemispheric differences across the retina [33]. All quantitative measurements were from a single retina from five to seven different mice of each experimental cohort.

The young retinas, particularly at P4–P8, changed so rapidly in formation of the ONL from the outer neuroblastic layer in a strong central-to-peripheral gradient (which all features

follow in the developing eye) that they were compared based on the following four phenotypic features: (1) length of rod inner + outer segments (if present) measured with an eyepiece micrometer in several regions in the central retina; (2) development of the outer plexiform layer (OPL) based on the percentage of the length in which it was present in each hemisphere (measured from the posterior pole); (3) incidence of ectopic photoreceptor nuclei still in the inner nuclear layer [34] measured as a percentage of the length in each hemisphere in which they still comprised a continuous row (measured from the ora serrata); and (4) incidence of pyknotic nuclei in the ONL, as seen in most rodent retinal degenerations [35, 36], per 430- μ m microscopic field.

Immunofluorescence and Confocal Microscopy

Eyes were enucleated and fixed by immersion in 4 % paraformaldehyde in PBS buffer for 1 h at room temperature. After overnight incubation with 30 % sucrose at 4 °C, eyes were frozen in O.C.T. (Tissue-Tek). Eight-micrometer-thick sections were cut through the optic nerve head and stained with indicated antibodies. Primary antibodies used included 1D4 anti-rhodopsin 1:500 dilution (Santa Cruz Biotechnologies, CA), anti-calnexin 1:250 (GeneTex, Irvine, CA), and anti-GFP 1:250 (Invitrogen, Carlsbad, CA). Secondary antibodies included Alexa 546 goat anti-mouse (Molecular Probes, Invitrogen) and Alexa 488 goat anti-rabbit (Molecular Probes, Invitrogen) used at 1:500. Images were collected with an Olympus FluoView-1000 confocal microscope and processed with Olympus FluoView Ver.2.0a Viewer software.

Transmission Electron Microscopy

Four mice from each genotype (*Rho*^{+/+}, *Rho*^{P23H/+}, and *Rho*^{P23H/P23H}) were analyzed by transmission electron microscopy at P14. Fixation, sectioning, and staining were performed according to Sakami et al. 2013 [44]. Low magnification ($\times 2,000$) TEM images of photoreceptor cell inner segments were captured by a JEOL 1200 EX EM at 80 kV (JEOL, Musashino, Tokyo, Japan). Eight images from each genotype (two images from each mouse) were analyzed.

Molecular Biology

Retinas were lysed and total RNA was collected with a RNeasy mini kit (Qiagen, Germany). mRNA was reverse-transcribed with the iScript cDNA Synthesis Kit (Bio-Rad, Hercules, CA). For quantitative PCR analyses, primers included the following: mouse *rhodopsin* mRNA, 5'-TTCACCAC CACCCTCTACACATCAC-3' and 5'-CGGAAGTTGCTC ATCGGCTTG-3'; mouse *GNAT1* mRNA, 5'-GAGGATGC TGAGAAGGATGC-3' and 5'-TGAATGTTGAGCGTGG TCAT-3'; mouse *Xbp-1s* mRNA, 5'-GAGTCCGCAGCAGG

TG-3' and 5'-GTGTCAGAGTCCATGGGA-3'; mouse *Derl1* mRNA, 5'-CGCGATTAAAGGCCTGTTAC-3' and 5'-GGTA GCCAGCGGTACAAAAA-3'; mouse *VCP* mRNA, 5'-AAGTCCCCAGTTGCCAAGGATG-3' and 5'-AGCCGA TGGATTTGTCTGCCTC-3'; mouse *ERdj4*, 5'-TAAAAG CCCTGATGCTGAAGC -3' and 5'-TCCGACTATTGGCA TCCGA -3'; mouse *Grp78/BiP*: 5'-CCTGCGTCGGTGTG TTCAAG-3' and 5'-AAGGGTCATTCCAAGTGCG-3'; mouse *Rpl19*: 5'-ATGCCAACTCCCGTCAGCAG- 3' and 5'-TCATCCTTCTCATCCAGGTCACC-3'; and mouse *Chop*: 5'-ACGGAAACAGAGTGGTCAGTGC-3' and 5'-CAGGAGGTGATGCCCACTGTTC-3'. For *rhodopsin* mRNA levels, *GNAT1* mRNA levels, a transcript only expressed in rod cells, served as the normalization control. For all other quantitative real-time PCR (qPCR) analysis, *Rpl19* mRNA levels, a transcript with levels unaltered by ER stress, served as internal normalization standards. qPCR conditions were 95 °C for 5 min, 95 °C for 10 s, 60 °C for 10 s, 72 °C for 10 s, with 50 cycles of amplification.

Cell Culture

Cells were maintained at 37 °C, 5 % CO₂ in Dulbecco's modified Eagle medium (Mediatech, Manassas, VA) supplemented with 10 % fetal calf serum (Invitrogen) and 1 % penicillin/streptomycin (Invitrogen). Tetracycline-inducible P23H rhodopsin HEK293 cell line was previously described [25]. To induce the expression of P23H rhodopsin, 1 μ g/ml of doxycycline is added to the cell culture media for 20 h.

Immunoprecipitation

Rhodopsin proteins were immunoprecipitated for mass spectrometry (MS) analysis or immunoblotting analysis. Retinas from three to five mice per experiment were collected from *Rho*^{+/+}, *Rho*^{P23H/+}, and *Rho*^{P23H/P23H} mice. MEF in one 10 cm dish were used as controls. To lyse the retinas or MEF cells, 1 ml of PBS-D (PBS containing 1.0 % *n*-dodecyl- β -D-maltoside and protease and phosphatase inhibitor mixture) was added to MEF cells or retinas. After sonication, the suspension was rotated for 1 h at 4 °C. The supernatants were collected, and the protein concentrations were determined with a BCA assay kit (Pierce). Equal amounts of total protein were added to Dynabeads Protein G (Invitrogen) preincubated with 1D4 anti-rhodopsin antibody (Santa Cruz Biotechnologies) and incubated overnight at 4 °C. To elute rhodopsin protein for immunoblotting analysis, beads were heated with SDS sample buffer at 80 °C. For MS analysis, rhodopsin was eluted twice from the beads by heating the beads with elution buffer (PBS containing 8 M urea and 0.5 % *n*-dodecyl- β -D-maltoside) for 10 min at 80 °C. Total proteins from the two eluates were combined and subjected to methanol chloroform extraction to remove detergent.

Liquid Chromatography–Tandem Mass Spectrometry

Protein pellets were resuspended in 8 M urea. The samples then were reduced by a 20-min incubation with 5 mM tris(2-carboxyethyl)phosphine (TCEP) at room temperature and alkylated in the dark by treatment with 10 mM iodoacetamide for 20 additional min. Proteins were digested overnight at 37 °C with Sequencing Grade Modified Trypsin (Promega, Madison, WI, USA) and the reaction was stopped by acidification. A 100- μ m i.d. capillary with a 5- μ m pulled tip was packed with 10 cm of 5- μ m Aqua C18 material (Phenomenex). This desalting column then was equilibrated for 30 min with buffer A (5 % acetonitrile/0.1 % formic acid), and the protein digest was loaded onto it under pressure. The column was placed in line with an Agilent 1200 quaternary HPLC and analyzed after elution and separation. As peptides eluted from the microcapillary column, they were electrosprayed directly into an LTQ-Orbitrap mass spectrometer from Thermo Finnigan (Waltham, MA) with the application of a distal 2.4-kV spray voltage.

Protein identification and quantification and analyses were done with Integrated Proteomics Pipeline (IP2, Integrated Proteomics Applications, Inc., www.integratedproteomics.com/) using ProLuCID and DTASelect2. Spectrum raw files were extracted into ms1 and ms2 files from raw files with RawExtract 1.9.9 (<http://fields.scripps.edu/downloads.php>) [37], and the tandem mass spectra were searched against European Bioinformatic Institute protein databases. To estimate peptide probabilities and FDRs accurately, we used a target/decoy database containing the reversed sequences of all the proteins appended to the target database [38]. Tandem mass spectra were matched to sequences by using the ProLuCID [39] algorithm with 50 ppm peptide mass tolerance for precursor ions and 400 ppm for fragment ions.

Immunoblotting Analysis

Mouse retinas were lysed in 300 μ l lysis buffer (PBS, 0.5 g/ml *n*-dodecyl- β -D-maltoside (Calbiochem EMD Bioscience), protease inhibitors (Sigma-Aldrich), and phosphatase inhibitor (Thermo Scientific, Rockford, IL)). MEF cells and HEK293 cells with indicated drug treatments were lysed in SDS lysis buffer (2 % SDS, 62.5 mM Tris-HCl, pH 6.8, containing protease inhibitors and phosphatase inhibitor). The following antibodies and dilutions were used: anti-G α t1 and 1D4 anti-rod opsin at 1:1,000 dilution (Santa Cruz Biotechnologies); B630N anti-rhodopsin at 1:1,000 (gift of W.C. Smith, Gainesville, FL); anti-calnexin at 1:1,000 at anti-VCP at 1:1,000 at anti-HSP90 at 1:5,000 at and anti- β -tubulin at 1:5,000 (GeneTex); anti-P-JNK at 1:1,000 and anti-JNK at 1:1,000 (Cell Signaling); anti-flag at 1:5,000 (Sigma-Aldrich); and anti-ubiquitin at 1:1,000 (Dako). After overnight incubation with primary antibody, membranes were washed

followed by incubation with a horseradish peroxidase-coupled secondary antibody (Cell signaling). Immunoreactivity was detected with the SuperSignal West chemiluminescent substrate (Pierce).

Statistical Analyses

All results are presented as means \pm standard deviations from at least three mice per experimental condition as indicated. Student two-tailed *t* tests (for paired samples) were performed to determine *p* values. A *p* value of ≤ 0.05 was considered statistically significant.

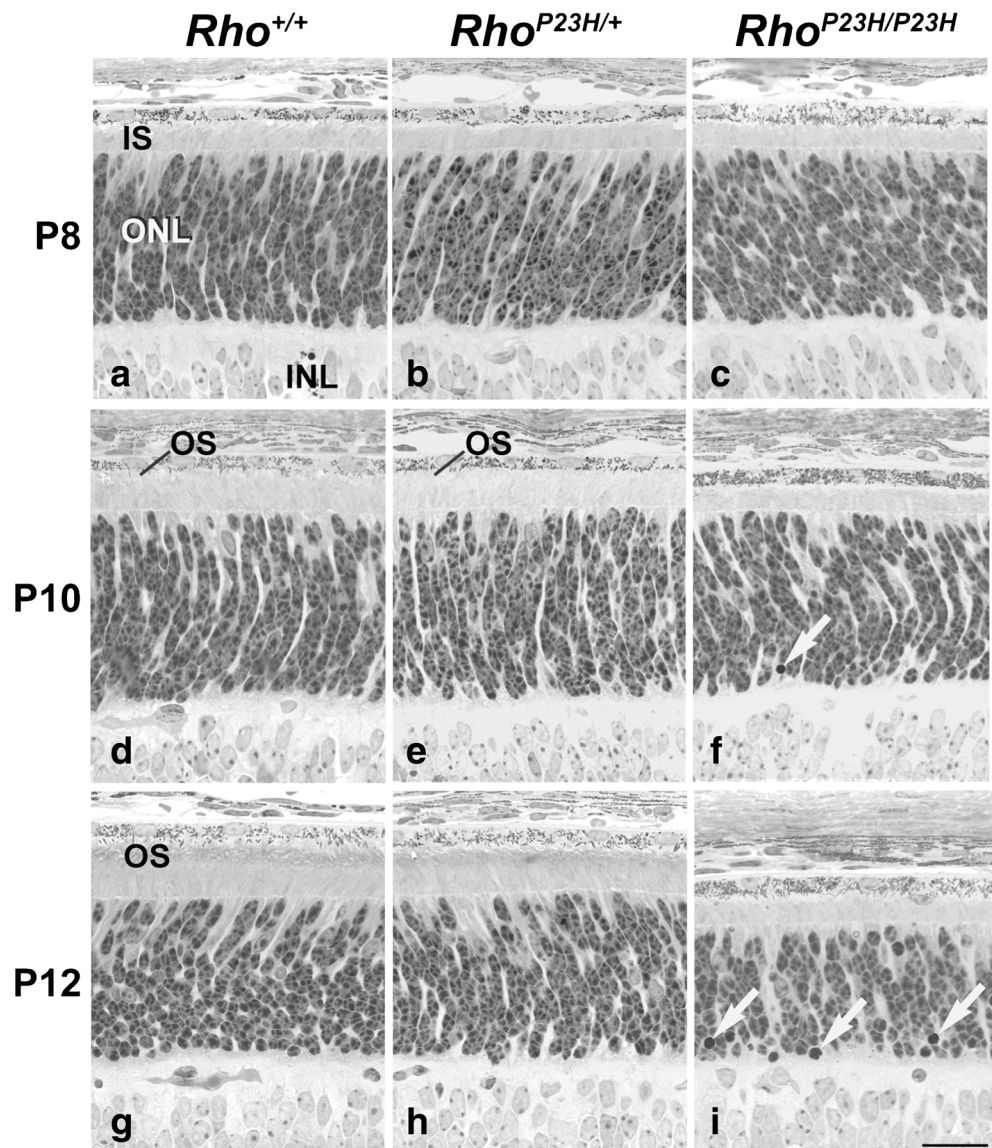
Results

Disruption of Photoreceptor Compartment Morphogenesis in P23H Rhodopsin Knock-in Mice During Early Retinal Degeneration

In wild-type mice (*Rho*^{+/+}), photoreceptor development and histogenesis follow well-described, stereotyped, temporal programs [40–42]. Photoreceptor neurogenesis starts in the embryo and lasts through approximately the first week of postnatal life (Fig. 1a). Post-mitotic photoreceptor cell bodies accumulate in the outermost neural lamina of the retina, termed the ONL (Fig. 1a). Newly formed rod photoreceptors cells develop morphologically and functionally distinct cellular compartments including the rod inner segment (RIS) and the more distal rod outer segment (ROS). Biosynthetic organelles, such as the ER, Golgi, and mitochondria, are found in the RIS (Fig. 3a), and the RIS cellular compartment can be readily distinguished in *Rho*^{+/+} retinas by postnatal day (P) 8 (Fig. 1a). The ROS is a specialized cilium that houses hundreds of membranous discs, embedded with rhodopsin protein for phototransduction [43]. In *Rho*^{+/+} mice, nascent ROS were clearly visible at P10 (Fig. 1d) and scattered ROS disc membranes were also seen as early as P7–P8 by electron microscopy [40]. The ROS elongated to about 2/3 of their adult lengths by P15 (Fig. 2a) and reached their maximum adult lengths shortly afterwards as shown at P30 and P60 (Fig. 2d, g).

In heterozygous *Rho*^{P23H/+} mice, the ONL thickness was indistinguishable from that of wild-type *Rho*^{+/+} mice from P4 through P12 (Fig. 1b, e, h, Table S1). By P15, scattered pyknotic nuclei could be identified (Fig. 2b, Table S1), accompanied by a slight reduction in the mean ONL layer thickness ($p < 0.01$), (Fig. 4a), indicating the earliest onset of photoreceptor cell loss and retinal degeneration. The lengths of the photoreceptor RIS and ROS compartments in *Rho*^{P23H/+} mice were also indistinguishable from those of wild-type mice until P15 (Fig. 1b, e, h, Table S1), when both RIS and ROS

Fig. 1 Light micrographs of wild-type and P23H mouse retinas at postnatal day (P) 8, P10, and P12. Degeneration of photoreceptor morphology in $Rho^{P23H/P23H}$ mice began by postnatal day 10. **a–c** At P8, the ROS had not formed, and the thicknesses of the rod inner segment (IS) layer and outer nuclear layer (ONL) were identical between $Rho^{+/+}$, $Rho^{P23H/+}$, and $Rho^{P23H/P23H}$ mice. **d–f** Also at P10, nascent ROS were identified in $Rho^{+/+}$ and $Rho^{P23H/+}$ mice but were absent in $Rho^{P23H/P23H}$ mice. By P10, the rod IS layer was thinner in $Rho^{P23H/P23H}$ retinas compared to those in $Rho^{+/+}$ and $Rho^{P23H/+}$ mice. By P10, pyknotic nuclei (white arrow) were found in $Rho^{P23H/P23H}$ mice at a significantly greater prevalence than in $Rho^{+/+}$ or $Rho^{P23H/+}$ mice. **g–i** By P12, ROS were present in $Rho^{+/+}$ and $Rho^{P23H/+}$ mice but are absent in $Rho^{P23H/P23H}$ mice. The rod IS layer and ONL are appreciably thinner in $Rho^{P23H/P23H}$ mice compared to $Rho^{+/+}$ and $Rho^{P23H/+}$ mice, and many pyknotic nuclei (white arrows) were identified in the inner aspect of the ONL in $Rho^{P23H/P23H}$ mice. Scale bar, 20 μ m



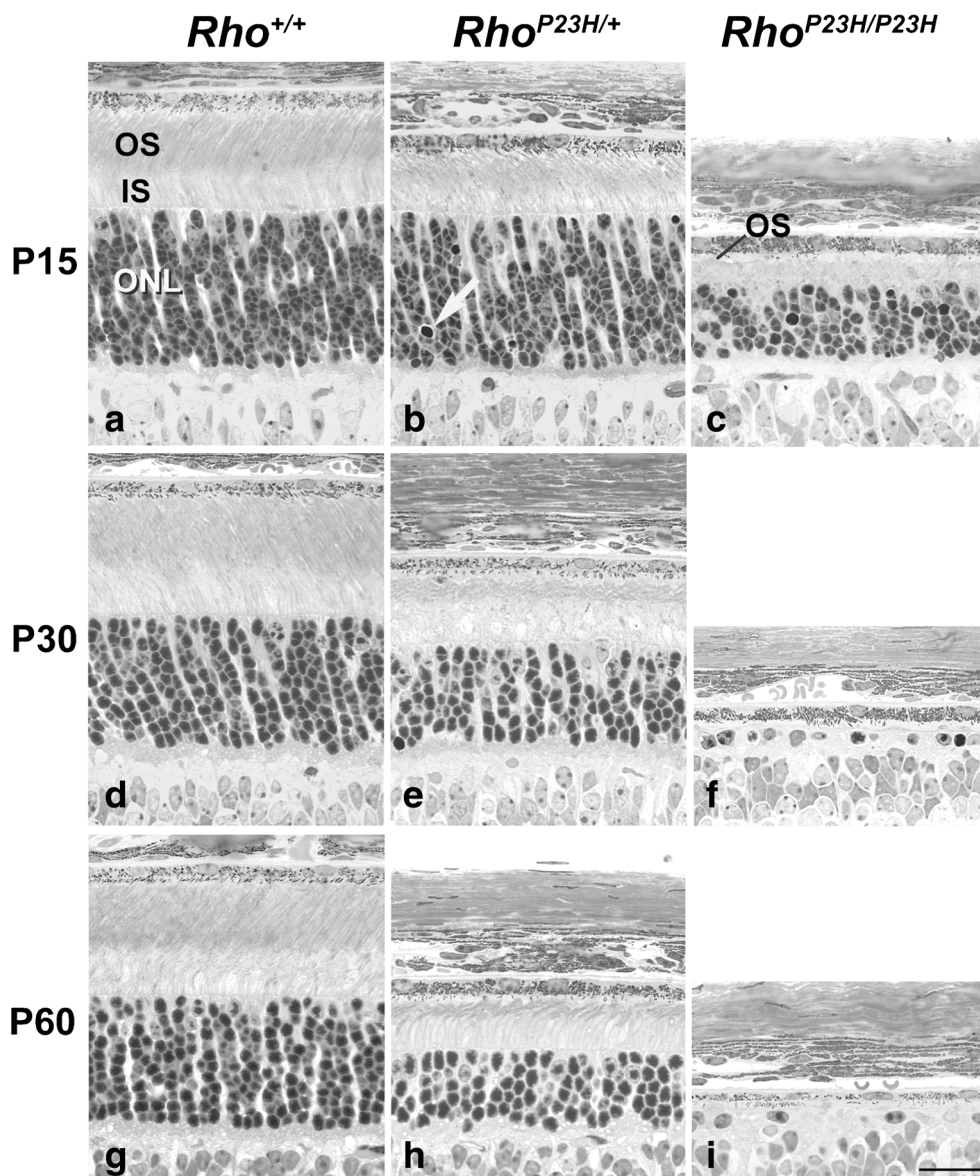
compartments were clearly shorter compared to their wild-type photoreceptor counterparts (compare Fig. 2a, b). Later, at P30 (Fig. 2e) and P60 (Fig. 2h), the retinas from heterozygous in $Rho^{P23H/+}$ mice showed progressive shortening of ROS, RIS, and loss of photoreceptor nuclei in the ONL (Fig. 2e, h) as previously described [27].

In homozygous $Rho^{P23H/P23H}$ retinas, thicknesses of the ONL and RIS were indistinguishable from those of wild-type mice from birth up to P10 (Fig. 1a, c, Table S1). By P10, rare pyknotic photoreceptor nuclei were evident in the ONL (Fig. 1f, Table S1), and photoreceptor RIS lengths were shorter in homozygous $Rho^{P23H/P23H}$ retinas compared to wild-type (Fig. 1d, f, Table S1). Little ROS was seen in homozygous $Rho^{P23H/P23H}$ retinas compared to wild-type at this age (Fig. 1d, f, Table S1). By P12, the prevalence of pyknotic photoreceptor nuclei increased significantly (Fig. 1i, Table S1), accompanied by a measurable reduction

in ONL thickness ($p < 0.001$) (Fig. 4a). Also, by P12, photoreceptor morphology was further compromised with markedly stunted ROS (Fig. 1i) bearing abnormally formed discs visualized by electron microscopy in ciliary protrusions [44]. By P14–15, vacuolization of IS became pronounced (Fig. 3c). The RIS remained short, and few visible ciliary protrusions were present (Figs. 2c, 3c). Forty to 50 % of the ONL thickness was lost (Fig. 2c, Table S1), and the nuclei of many of the remaining photoreceptors were clearly pyknotic (Fig. 2c). By P30 (Fig. 2f) and P60 (Fig. 2i), less than a full row of photoreceptor nuclei remained in the ONL, and some attenuation of the retinal pigmented epithelium was evident at P60 (Fig. 2i).

When the mean ONL values across the entire retina were plotted, sharp differences in the kinetics of photoreceptor loss and amount of retinal degeneration were evident between the heterozygous and homozygous P23H genotypes,

Fig. 2 Light micrographs of wild-type and P23H mouse retinas at P15, P30, and P60. Degeneration of photoreceptor morphology in $Rho^{P23H/+}$ mice began by P15. **a–c** At P15, the rod OS were shorter and more disorganized in $Rho^{P23H/+}$ mice (**b**) compared to $Rho^{+/+}$ mice (**a**), and only short fragments of ROS disc membranes were present in $Rho^{P23H/P23H}$ mice (**c**). At P15, the rod inner segments (IS) were shorter in $Rho^{P23H/+}$ mice than in $Rho^{+/+}$ mice and significantly shortened in $Rho^{P23H/P23H}$ mice. Pyknotic nuclei (white arrow) were evident in $Rho^{P23H/+}$ mice. The ONL was significantly thinner in $Rho^{P23H/P23H}$ mice (**c**). **d–f** By P30, the OS, IS, and ONL were significantly thinner in $Rho^{P23H/+}$ mice (**e**) compared to $Rho^{+/+}$ mice (**d**). Extensive photoreceptor cell loss was present in $Rho^{P23H/P23H}$ mice, with less than one full row of nuclei present in the ONL (**f**). **g–i** By P60, the OS, IS, and ONL had progressively thinned in $Rho^{P23H/+}$ mice (**h**) compared to $Rho^{+/+}$ mice (**g**), and even further loss of photoreceptors was evident in the ONL of $Rho^{P23H/P23H}$ mice (**i**). Scale bar, 20 μm



with an early and rapid loss of ONL thickness in the $Rho^{P23H/P23H}$ retinas significantly preceding the onset of loss seen in the heterozygous $Rho^{P23H/+}$ retinas (Fig. 4a). The ONL thicknesses at P30 and later in the homozygous $Rho^{P23H/P23H}$ and heterozygous $Rho^{P23H/+}$ retinas were similar to our prior findings [27], with the loss of most photoreceptors in the $Rho^{P23H/P23H}$ mice by P30. By contrast, the ONL thickness stabilized and retinal degeneration plateaued in $Rho^{P23H/+}$ retinas by P30 (Fig. 4a). Previously, we reported a greater loss of photoreceptors in the inferior hemisphere of the eye in heterozygous $Rho^{P23H/+}$ retinas than in the superior hemisphere by P112 [27]. Our retinal “spidergram” measurements of ONL thickness at different ages in the heterozygote $Rho^{P23H/+}$ retinas (Fig. 4b) revealed that this hemispheric asymmetrical degeneration was not present at P20, but present at all older ages

examined (Fig. 4b, P45 and P90 not shown). In the homozygous $Rho^{P23H/P23H}$ retinas, however, the hemispheric asymmetry in retinal degeneration was reversed, with the superior hemisphere more severely affected in all mice from P12 to P60 (Fig. 4c; P30 and P45 not shown), but not at ages older than P60. The difference in the least marked asymmetry at P12 was significant, however ($p < 0.005$).

Our analyses of early retinal development revealed an unexpected early onset of photoreceptor pathology in P23H mice. Although photoreceptor neurogenesis and retinal patterning appeared histologically normal in both heterozygous and homozygous P23H mice, severe defects in photoreceptor compartment morphogenesis, especially the proper generation and elongation of ROS and to a lesser extent RIS, appeared in $Rho^{P23H/P23H}$ mice as soon as P10,

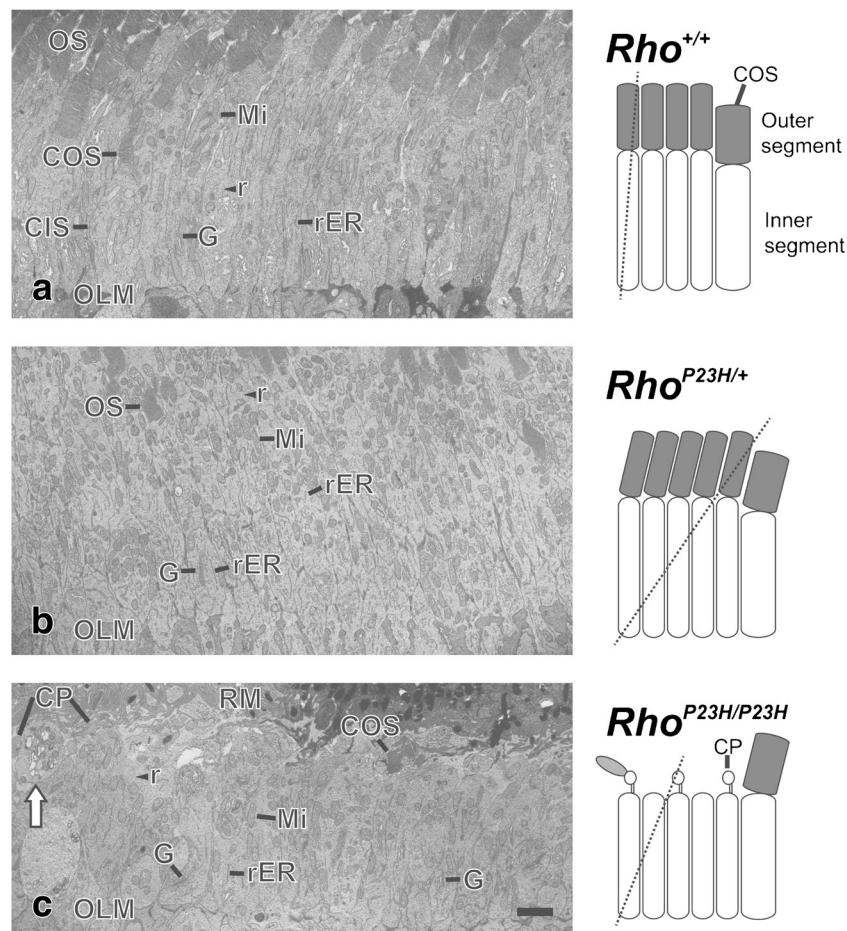


Fig. 3 Transmission electron microscopy images of photoreceptor cell inner segments at postnatal day 14. **a–c** Mitochondria (*Mi*), rough endoplasmic reticulum (*rER*), A Golgi apparatus (*G*), and ribosomes (*r*) were abundant in the inner segment. Golgi apparatus was observed at the proximal part of most inner segments. **a** Image of longitudinal-cut inner segments from *Rho*^{+/+} mouse retinas. Column-shaped inner segments were observed between photoreceptor cell outer segments (*OS*) and outer limiting membranes (*OLM*). In rare cases, the entire cone photoreceptor outer segment (*COS*) and inner segment (*CIS*) of a single cell was observed. **b** Image of tilted-cut inner segments from *Rho*^{P23H/+} mouse retinas. Two to four rows of oval or column-shaped inner segments were observed between the *OS* and *OLM*. Structures of the *rER* and Golgi apparatus in the inner segments of *Rho*^{P23H/+} mice were almost identical

to those in *Rho*^{+/+} mouse retinas especially compared to tilted-cut inner segment images from *Rho*^{+/+} mouse retinas. **c** Image of slightly tilted-cut inner segments from *Rho*^{P23H/P23H} mouse retinas. One to three rows of oval or column-shaped inner segments were observed between retinal pigmented epithelium microvilli (*RM*) and *OLM*. Ciliary protrusions (*CP*) and cone outer segments (*COS*) were noted in the space between *RM* and the distal ends of inner segments. Not all, but some inner segments showed multiple vacuolization (white arrow). Such vacuolization was seen in all four analyzed *Rho*^{P23H/P23H} mice, but not in TEM images of *Rho*^{P23H/+} and *Rho*^{+/+} mice (four mice were analyzed for each genotype). Hypothetical diagrams of photoreceptor cells, indicate the cutting planes with dashed lines. Scale bar, 2 μ m

coinciding with the onset and progressive death of nearly all photoreceptors by P30. Defects in photoreceptor ROS and RIS morphogenesis also appeared in heterozygous *Rho*^{P23H/+} mice beginning at a slightly later date, ~P15, and these also coincided with the onset of retinal degeneration. However, by contrast to homozygous mice, far fewer photoreceptors were lost in heterozygous *Rho*^{P23H/+} retinas such that the ONL thickness stabilized by ~P30 (Fig. 4a). These studies reveal a critical period within the first postnatal month of life when photoreceptors in both heterozygous and homozygous P23H mice undergo degeneration. We next examined UPR activity during this early period of photoreceptor cell death in P23H rhodopsin mice.

The IRE1 Signaling Pathway of the UPR Is Strongly Induced in Photoreceptors Expressing P23H Rhodopsin In Vivo

To investigate the role of the IRE1 signaling UPR pathway in native photoreceptors, we crossed *Rho*^{P23H/P23H} mice with *ERAI* reporter mice [45]. *ERAI* mice carry a modified FLAG-tagged truncated *Xbp-1* that lacks the DNA-binding and transcriptional activator domains but retains the native *Xbp-1* intron fused upstream of *Venus*, a green fluorescent protein (GFP) variant. Activated IRE1 splices out the *Xbp-1* intron, leading to the production of FLAG-tagged XBP1-Venus fusion protein [45]. Thus, the *ERAI* reporter mouse provides a highly specific monitor of the status of IRE1

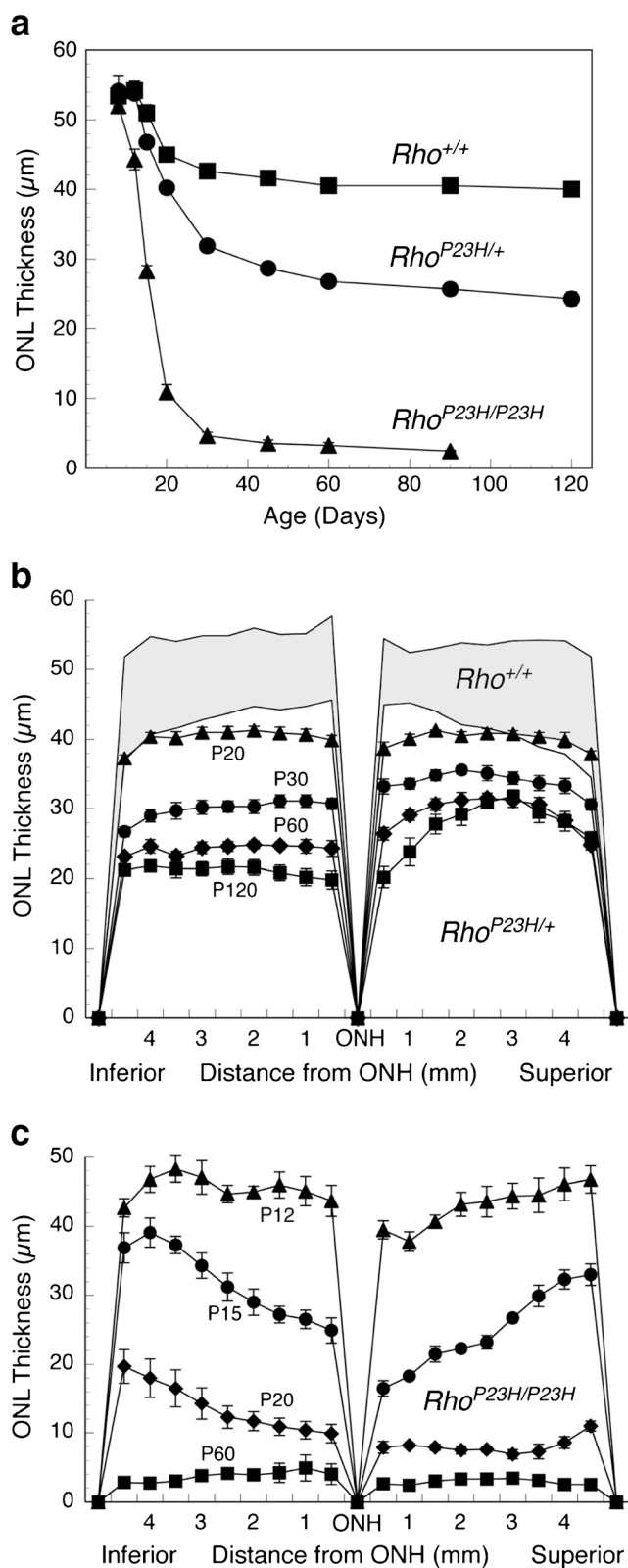


Fig. 4 Anatomic distribution of retinal degeneration in *Rho*^{+/+}, *Rho*^{P23H/+}, and *Rho*^{P23H/P23H} mice. **a** Mean ONL thickness of *Rho*^{+/+}, *Rho*^{P23H/+}, and *Rho*^{P23H/P23H} retinas. Each value is the mean ± SEM of five to seven retinas, one from each mouse; the mean for each retina was determined from 54 measurements around the eye from the inferior to superior poles. If error bars are not shown, they fall within the symbol. **b** Retinal “spidergram” showing ONL thickness from the optic nerve head (ONH) to the anterior optic margin (ora serrata) in the inferior and superior hemispheres of *Rho*^{+/+} retinas (at P12 and P120) and *Rho*^{P23H/+} retinas (at P20, P30, P60, and P120). The top border of the shaded area represents the *Rho*^{+/+} ONL thickness at P12, and the bottom border shows the *Rho*^{+/+} ONL thickness at P120. Thinning of the retina ONL in *Rho*^{+/+} mice between P12 and P120 is due to growth of the eye, not retinal degeneration. The ONL thickness progressively diminished in *Rho*^{P23H/+} retinas from P20 to P120 with more pronounced retinal degeneration in the inferior hemisphere. **c** Retinal spidergram showing a rapid decrease in ONL thickness in *Rho*^{P23H/P23H} retinas from P12 to P60. In *Rho*^{P23H/P23H} retinas, the superior hemisphere shows greater loss of the ONL, the opposite of that seen in *Rho*^{P23H/+} retinas

In *Rho*^{P23H/+}*ERAI*^{+/-} mice, we detected strong expression of XBP1-Venus protein in retinas compared to *Rho*^{+/+}*ERAI*^{+/-} littermates at P30 (Fig. 5a–h). Strong XBP1-Venus expression was localized to rod photoreceptor cells in the outer nuclear layer (Fig. 5d) with strong staining seen in the RIS and perinuclear region of all rod photoreceptors (Fig. 5f, h). XBP1-Venus expression was excluded from photoreceptor nuclei consistent with the deletion of the DNA-binding domain of XBP1 in the reporter construct (Fig. 5f, h and [45]). By contrast, in wild-type retinas, only rare XBP1-Venus staining was observed in photoreceptors in the ONL (Fig. 5c, e, g). At P120, we also observed strong XBP1-Venus staining in the *Rho*^{P23H/+}*ERAI*^{+/-} mice, but not the *Rho*^{+/+}*ERAI*^{+/-} littermates (Fig. 5i–l). In other retinal cells of *Rho*^{P23H/+}*ERAI*^{+/-} mice, XBP1-Venus staining was also rarely seen in cone photoreceptors (indicated by a white arrow in Fig. 5l), the inner nuclear layer, ganglion cell layer, and retinal pigment epithelium (data not shown). Whole retinal lysates also showed significantly increased FLAG-tagged XBP1-Venus protein expression in *Rho*^{P23H/+}*ERAI*^{+/-} mice compared to controls (Fig. 5m), consistent with the increased XBP1-Venus protein by imaging. These findings demonstrate that the ERAI reporter is efficiently and specifically activated in rod photoreceptors expressing P23H rhodopsin.

To confirm the activation status of the endogenous IRE1 signal transduction pathway in retina, we examined the splicing status of native *Xbp-1* mRNA in *Rho*^{P23H/+}*ERAI*^{+/-} mice. We found increased levels of spliced *Xbp-1* mRNA in animals expressing P23H rhodopsin (Fig. 6a). Spliced *Xbp-1* produces a potent transcription factor, XBP1s, with many transcriptional targets that are shared between diverse mammalian cells [14, 13, 48, 49]. In both *Rho*^{P23H/+} and *Rho*^{P23H/P23H} mice, we found significantly elevated mRNA levels of *Erdj4*, *VCP*, *Der11*, and *Grp78* (Fig. 6b–e) at P30. Furthermore, we found a P23H rhodopsin dose-dependent increase in the upregulation of *Xbp-1* splicing and XBP1s target genes, with

signaling at the cellular, tissue, and organ level. In the eye, the ERAI reporter mouse offers a useful way to distinguish IRE1 signaling status between the different laminar layers of the retina [46, 47].

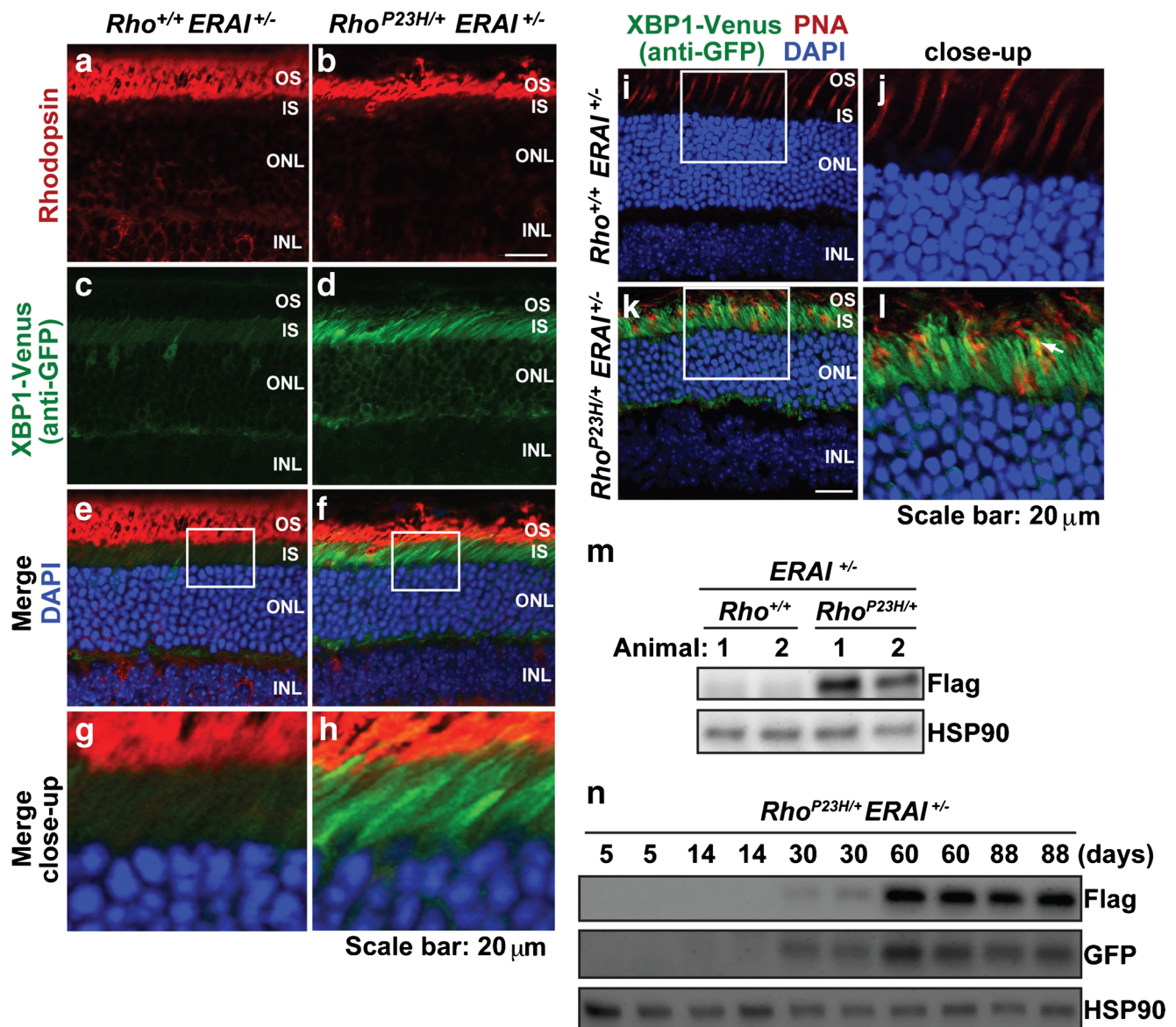


Fig. 5 The IRE1 signaling pathway of the UPR is strongly activated in photoreceptors of *Rho*^{P23H/+}*ERAI*^{+/-} mice. **a–h** Rhodopsin (red) and XBP1-Venus (by anti-GFP, green) were visualized in eyes from *Rho*^{P23H/+}*ERAI*^{+/-} mice and *Rho*^{+/+}*ERAI*^{+/-} mice at P30. DAPI (blue) highlights photoreceptor nuclei. Strong XBP1-Venus staining is visualized in the IS and perinuclear region of ONL photoreceptors expressing P23H rhodopsin. **i–l** Peanut agglutinin (PNA) staining of cones (red) and XBP1-Venus (by anti-GFP, green) were visualized in eyes from

Rho^{P23H/+}*ERAI*^{+/-} mice and *Rho*^{+/+}*ERAI*^{+/-} mice at P120. DAPI (blue) highlights photoreceptor nuclei. White arrow in **l** indicated a cone cell overlapping with the XBP1-Venus signal. **m** XBP1-Venus (by anti-flag) protein levels were detected in retinal lysates from *Rho*^{P23H/+}*ERAI*^{+/-} mice and littermate control *Rho*^{+/+}*ERAI*^{+/-} mice at P30. HSP90 served as a protein loading control. **n** XBP1-Venus protein levels were detected by anti-flag or anti-GFP in retinal lysates from *Rho*^{P23H/+}*ERAI*^{+/-} mice at indicated ages

significantly higher levels of spliced *Xbp-1*, *Erdj4*, *VCP*, *Der11*, and *Grp78* in retinas from *Rho*^{P23H/P23H} mice expressing two copies of P23H rod opsin gene compared to the *Rho*^{P23H/+} mice expressing only one copy (Fig. 6a–e). From these findings, we conclude that P23H rhodopsin induces ER stress and strongly activates the IRE1 pathway in photoreceptors in vivo.

Next, we examined the onset and duration of IRE1 signaling in *Rho*^{P23H/+}*ERAI*^{+/-} mice. We detected no expression of

FLAG-tagged XBP1-Venus protein by anti-Flag or anti-GFP antibodies at P5 (Fig. 5n). By P14, we saw faint expression of FLAG-tagged XBP1-Venus protein by anti-GFP immunoblotting, indicating that the IRE1 pathway was becoming activated in these retinas. The XBP1-Venus protein levels detected by both anti-GFP and anti-Flag progressively increased by P30, and these levels along with mRNA levels of downstream transcriptional targets of XBP1s, such as *Erdj4*, remained strongly elevated at older ages (P60 and P90) (Fig. 6f, g).

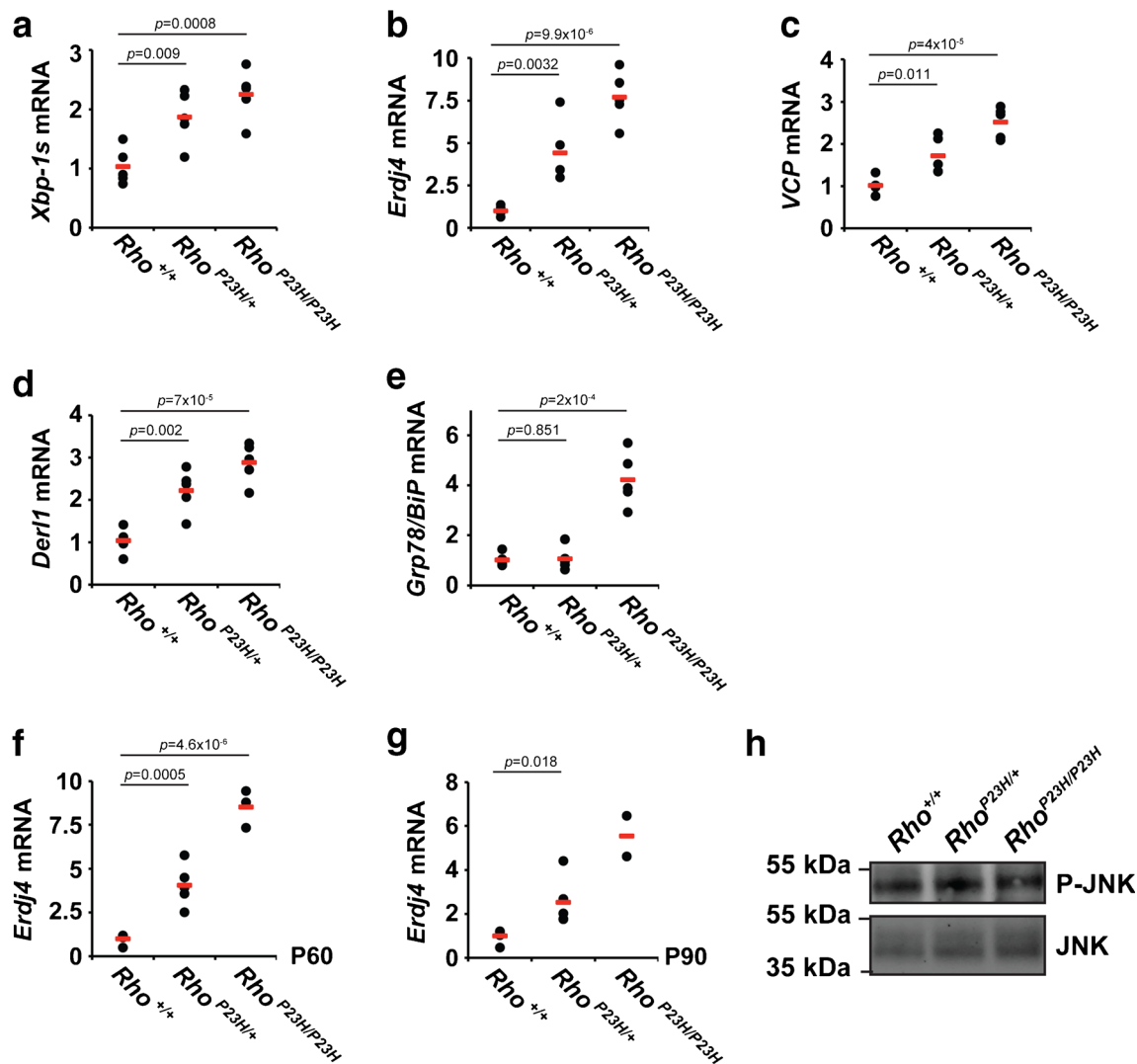


Fig. 6 The IRE1 signaling pathway of the UPR is strongly activated in photoreceptors expressing P23H rhodopsin. **a–e** *Xbp-1s* mRNA level (**a**) and XBP1s transcriptional target genes, *Erdj4* (**b**), *VCP* (**c**), *Derl1* (**d**), and *Grp78/BiP* (**e**), were measured in *Rho*^{+/+}, *Rho*^{P23H/+}, and *Rho*^{P23H/P23H} mice at P30 by quantitative PCR. **f, g** Upregulation of *Erdj4* was measured in *Rho*^{+/+}, *Rho*^{P23H/+}, and *Rho*^{P23H/P23H} mice at

P60 (**f**) or P90 (**g**) by quantitative PCR. **h** Retinas from three *Rho*^{+/+}, *Rho*^{P23H/+}, or *Rho*^{P23H/P23H} mice were collected at P15, pooled, and lysed. Forty micrograms of total retinal lysates was loaded onto the SDS-PAGE. Phosphorylated JNK protein levels were detected by immunoblotting. Total JNK protein levels served as a loading control

These findings demonstrated that photoreceptors expressing P23H rhodopsin activated the IRE1 signal transduction pathway by ~P14, and IRE1 signaling remained strongly activated thereafter. Interestingly, the onset of IRE1 signaling closely mirrored the onset of photoreceptor degeneration seen in *Rho*^{P23H/+} mice (Fig. 2b). Furthermore, IRE1 signaling remained strongly activated in photoreceptors even after retinal degeneration had plateaued in heterozygous *Rho*^{P23H/+} mice (Fig. 4a). IRE1 signaling can also activate the JNK pathway independent of its induction of XBP1s in vitro [50]. However, we saw no increases in activated phosphorylated JNK levels in P23H animals compared to controls (Fig. 6h). We subsequently focused our studies on the consequences of IRE1's induction of XBP1s in the retina.

ERAD Induction Leads to Ubiquitination and Rapid Degradation of P23H Rhodopsin In Vivo

Previously, we found that artificial activation of IRE1 signaling to selectively induce XBP1s targeted P23H rhodopsin for protein degradation in vitro [25]. The genes we found transcriptionally upregulated in retinas expressing P23H rhodopsin all play well-defined roles in ERAD (*p97/VCP* and *Derlin1*) and/or ER protein folding (*Erdj4* and *Grp78/BiP*) (Fig. 6b–e, and [51]). To determine if ERAD components and ER chaperones targeted P23H rhodopsin in photoreceptors, we performed mass spectrometry analysis on pure P23H rhodopsin immunoprecipitated from retinas of homozygous *Rho*^{P23H/P23H} mice and wild-type rhodopsin immunoprecipitated from *Rho*^{+/+} mice. We

consistently detected increased interaction of P23H rhodopsin with ER chaperones and ERAD components in triplicate experimental mass spectrometry analyses from retinal lysates compared to analogous studies with wild-type rhodopsin (Fig. 7a). We also confirmed some of the P23H rhodopsin interactions identified by mass spectrometry in retinal co-immunoprecipitation studies where we found increased levels of p97/VCP and calnexin after rhodopsin pull-down in photoreceptors from *Rho*^{P23H/P23H} mice compared to wild-type mice (Fig. 7b). The transcriptional induction of ERAD genes and

increased association between ERAD proteins and P23H rhodopsin protein suggested that photoreceptors of *Rho*^{P23H/P23H} mice target misfolded rhodopsin for ERAD. Consistent with this notion, we found a substantial increase in the amount of ubiquitination on the purified P23H rhodopsin (Fig. 7c). In heterozygous *Rho*^{P23H/+} mice that express both WT and mutant P23H rhodopsin protein, we found a significant increase in the amount of ubiquitinated rhodopsin protein after immunoprecipitation with an antibody that recognizes both WT and P23H rhodopsin proteins (Fig. 7c, lower panel). In homozygous

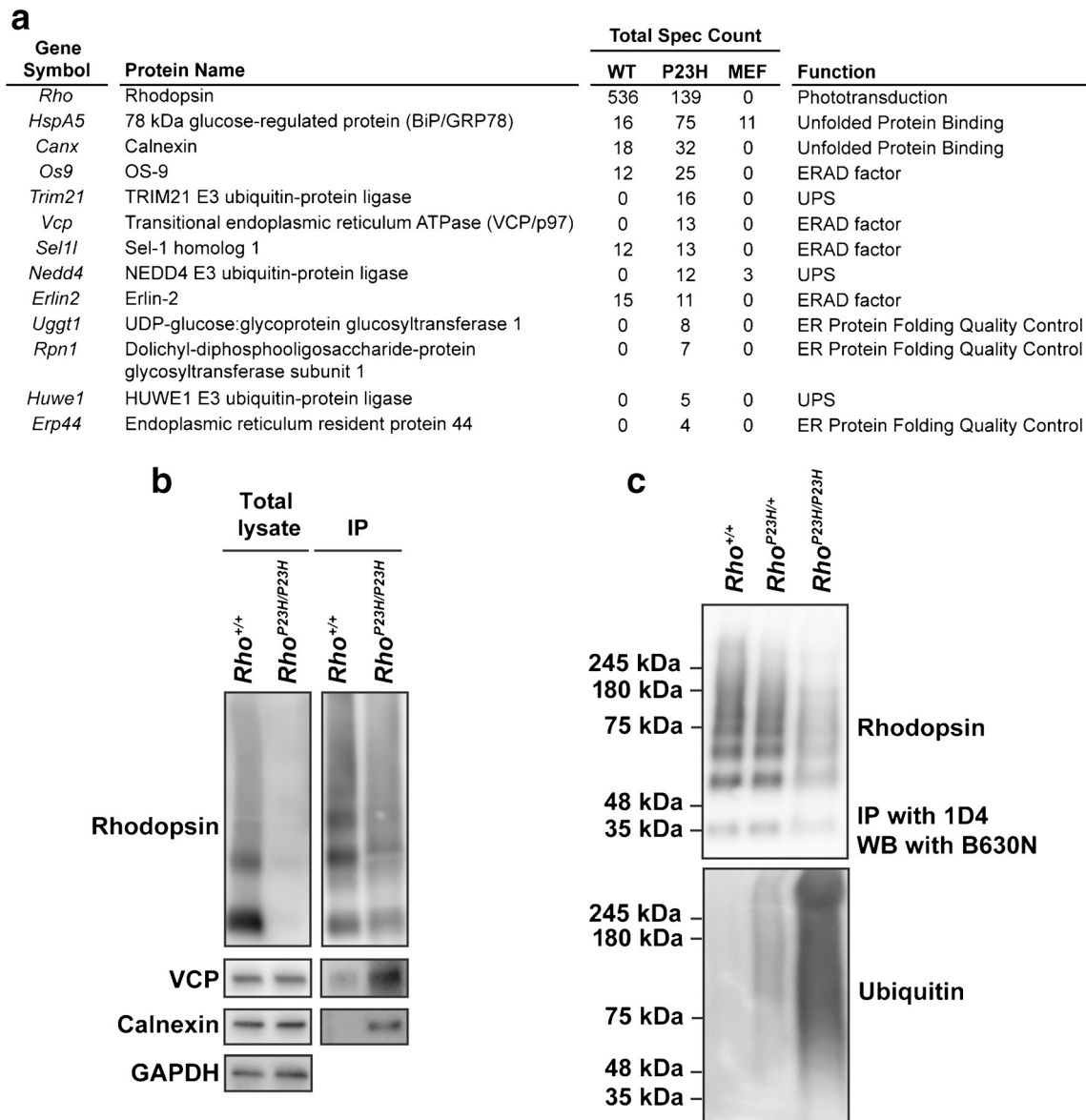


Fig. 7 P23H rhodopsin protein co-immunoprecipitates with ERAD components and is strongly ubiquitinated in photoreceptor cells. (a) ERAD-related proteins that co-immunoprecipitated with P23H rhodopsin were identified by mass spectrometry. Total spectra counts of the proteins co-immunoprecipitated with WT or P23H rhodopsin from three experimental mass spectrometry analyses are shown. Immunoprecipitation of MEF cell samples served as an experimental control. (b) Rhodopsin was

immunoprecipitated from retinal lysates of *Rho*^{+/+} and *Rho*^{P23H/P23H} mice at P15. The elutions were concentrated by chloroform/methanol precipitation. Rhodopsin, VCP, and calnexin proteins were detected by immunoblotting. (c) Rhodopsin was immunoprecipitated from retinas of *Rho*^{+/+}, *Rho*^{P23H/+} and *Rho*^{P23H/P23H} mice at P15, and rhodopsin (top panel) and ubiquitin (bottom panel) levels were detected by immunoblotting

$Rho^{P23H/P23H}$ that expresses only mutant P23H rhodopsin protein, we found a massive amount of ubiquitination on the immunoprecipitated P23H rhodopsin protein (Fig. 7c, lower panel). These data demonstrate that induction of ERAD by IRE1 signaling leads to profound ubiquitination of P23H rhodopsin in photoreceptors, indicating P23H rhodopsin is targeted *in vivo* for degradation. Indeed, when we examined steady-state levels of rhodopsin from wild-type, heterozygous $Rho^{P23H/+}$, and homozygous $Rho^{P23H/P23H}$ retinas, we found substantial loss of total rhodopsin protein in P23H rhodopsin-expressing mice with almost complete loss of rhodopsin protein from photoreceptors in homozygous $Rho^{P23H/P23H}$ mice (Fig. 8a), despite equivalent levels of *rod opsin* mRNA gene expression in wild-type and P23H rhodopsin-expressing mice at P14 (Fig. 8b). Immunofluorescence analysis confirmed the pronounced loss of rhodopsin protein in homozygous $Rho^{P23H/P23H}$ photoreceptor cells (Fig. 8c). In stark contrast to the pronounced retention of P23H rhodopsin in the ER when P23H rhodopsin is expressed in heterologous cell types, we detected no ER-retained P23H rhodopsin in native photoreceptors under non-saturating imaging conditions (Fig. 8c, note the absence of rhodopsin staining of the IS and perinuclear regions of the ONL as exhibited by calnexin expression). However, under these imaging conditions, P23H rhodopsin expression in $Rho^{P23H/P23H}$ photoreceptors was found outside of the ER at the base of the stunted OS (Fig. 8c), corresponding to ciliary protrusion and abnormal discs where P23H predominantly localized as we reported previously by immuno-electron microscopy [44].

Loss of rhodopsin protein in $Rho^{P23H/P23H}$ photoreceptors was apparent as early as P5 (Fig. 8d). Loss of P23H rhodopsin protein was not due to photoreceptor cell death, as we observed no frank degeneration of photoreceptor compartment morphology, pyknotic nuclei, or any ONL thinning at this early age (Fig. 1 and Table S1). Furthermore, we found normal protein levels of rod photoreceptor-specific transducin α -subunit (GNAT1) at P5 [52], indicating that photoreceptors still generated other rod-specific proteins (Fig. 8d). We conclude that loss of rhodopsin protein occurs shortly after photoreceptors are formed. Furthermore, the loss of rhodopsin protein precedes the onset of morphologic defects arising in photoreceptor compartments, obvious photoreceptor cell death, and retinal degeneration. ERAD was directly involved in rhodopsin protein degradation, as we prevented the degradation of P23H rhodopsin upon application of ERAD inhibitors, kifunensine, an inhibitor of ER mannosidase I [53, 54], or eeyarastatin I, an inhibitor of p97/VCP ATPase activity [55, 56], *in vitro* (Fig. 8e, f).

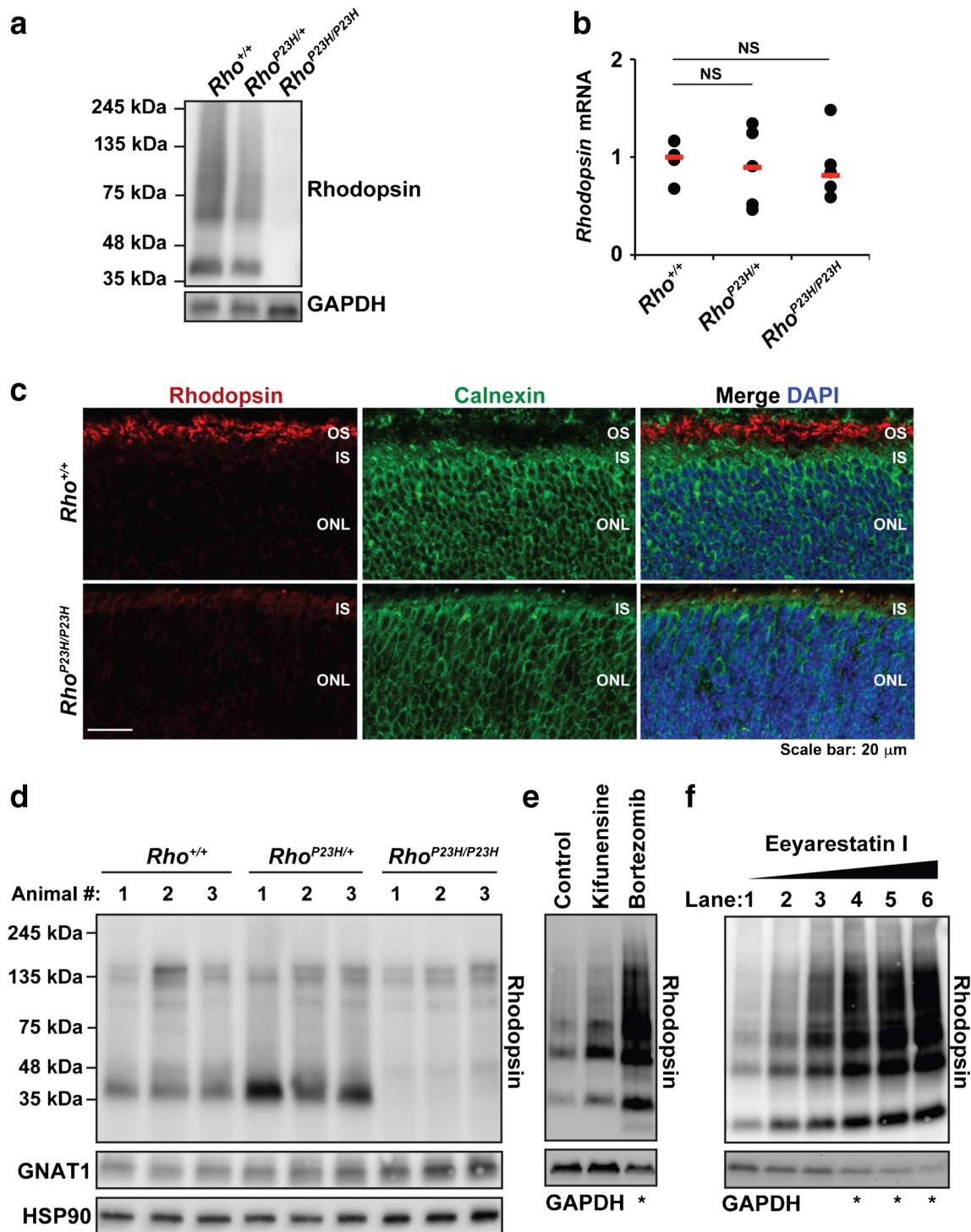
Chronic ER Stress-Induced Pro-apoptotic *Chop* Does Not Contribute to P23H Rhodopsin-Induced Retinal Degeneration

Uncorrectable ER stress ultimately triggers cell death, most notably by activating the PERK signaling pathway of the UPR

Fig. 8 Rhodopsin protein, but not mRNA, levels are significantly diminished in P23H mice. **a** Retinal protein lysates were collected from $Rho^{+/+}$, $Rho^{P23H/+}$, and $Rho^{P23H/P23H}$ mice at P15. Rhodopsin was detected by immunoblotting. GAPDH protein levels served as a loading control. **b** Rhodopsin mRNA levels from $Rho^{+/+}$, $Rho^{P23H/+}$, and $Rho^{P23H/P23H}$ mice at P12 were measured by quantitative PCR. Filled circles represent rhodopsin mRNA levels in individual mice. Horizontal red bars represent mean rhodopsin mRNA levels from each experimental cohort of mice. NS, statistically not significant. **c** Retinas from $Rho^{+/+}$ and $Rho^{P23H/P23H}$ mice at P10 were sectioned and rhodopsin (red) and calnexin (green) were visualized by confocal microscopy. DAPI counterstain highlighted nuclei. Under equivalent confocal microscopy detection settings, strong rhodopsin protein staining was evident in the developing OS of $Rho^{+/+}$ photoreceptors but only a scant amount of rhodopsin protein staining was visualized in the presumptive OS layer in $Rho^{P23H/P23H}$ photoreceptors. Rhodopsin protein did not colocalize with calnexin in either $Rho^{+/+}$ or $Rho^{P23H/P23H}$ photoreceptors. **d** Retinal protein lysates were collected from $Rho^{+/+}$, $Rho^{P23H/+}$, and $Rho^{P23H/P23H}$ mice at P5. Rhodopsin and GNAT1 were detected by immunoblotting. HSP90 protein levels served as a loading control. **e** P23H rhodopsin was expressed in HEK293 cells. The cells were treated with 10 μ M kifunensine for 20 h. Cells were also treated with bortezomib 10 μ M as a control for proteasomal-mediated degradation. **f** HEK293 cells expressing P23H rhodopsin were treated with 1.25, 2.5, 3.75, 5, or 7.5 μ M of eeyarastatin I for 20 h (lanes 2 to 6). **e, f** Rhodopsin protein was detected by immunoblotting. GAPDH protein levels served as a loading control. Significant cell toxicity was observed with some drug treatment conditions (indicated with asterisk)

which phosphorylates eIF2 α leading to strong induction of ATF4 transcriptional activator and its downstream pro-apoptotic *Chop* gene [17, 1]. ER stress-inducing toxins, tunicamycin and thapsigargin, strongly induced *Chop*, and overexpression of P23H rhodopsin in heterologous cell types or atop native wild-type rhodopsin in transgenic mice also upregulated *Chop* [26, 57]. Furthermore, *Chop*^{-/-} mice were resistant to cell death and damage induced by ER protein misfolding [58–60]. These findings support a model whereby P23H rhodopsin expression in photoreceptors leads to strong *Chop* induction that drives photoreceptor cell death *in vivo*. However, our analysis of $Rho^{P23H/P23H}$ retinas revealed low steady-state levels of P23H rhodopsin in photoreceptors (Fig. 8a, d) and no obvious aggregation or retention of P23H rhodopsin within the ER (the RIS/perinuclear compartment) of photoreceptors (Fig. 8c). Furthermore, ER dilation and fragmentation, ultrastructural abnormalities seen with irreparable ER stress, were not observed in the RIS of $Rho^{P23H/P23H}$ photoreceptors, (Fig. 3c). These findings raise an alternative scenario whereby the robust degradation of P23H rhodopsin by induction of ERAD in photoreceptors effectively eliminated misfolded rhodopsin from the ER. In this model, ER stress is alleviated, and therefore, *Chop* would not be induced nor causally contribute to photoreceptor cell death.

To address these competing models of *Chop*'s role in the photoreceptor cell death seen in P23H rhodopsin mice, we measured *Chop* mRNA levels at time points spanning the



entire period of retinal degeneration in heterozygote *Rho*^{P23H/+} and homozygous *Rho*^{P23H/P23H} mice. In both *Rho*^{P23H/+} and *Rho*^{P23H/P23H} mice, we saw no significant increase in *Chop* mRNA levels compared to age-matched wild-type mice at all ages (Fig. 9a, b). We also found no increase in ATF4 protein in either *Rho*^{P23H/+} or *Rho*^{P23H/P23H} mice and only a mild elevation in phosphorylated eIF2 α protein levels (Fig. 9c). By contrast, all of these markers were strongly induced by a

strong ER toxin, thapsigargin (Tg), in vitro (Fig. 9c). These findings indicate that PERK signaling is weakly activated in vivo in photoreceptors expressing P23H rhodopsin, and PERK signaling does not rise sufficiently to trigger its downstream pro-apoptotic ATF4-CHOP circuit. Consistent with this interpretation, when we examined retinal degeneration in *Rho*^{P23H/+}*Chop*^{-/-} mice, we found no significant change in ONL thickness, a proxy for photoreceptor cell numbers,

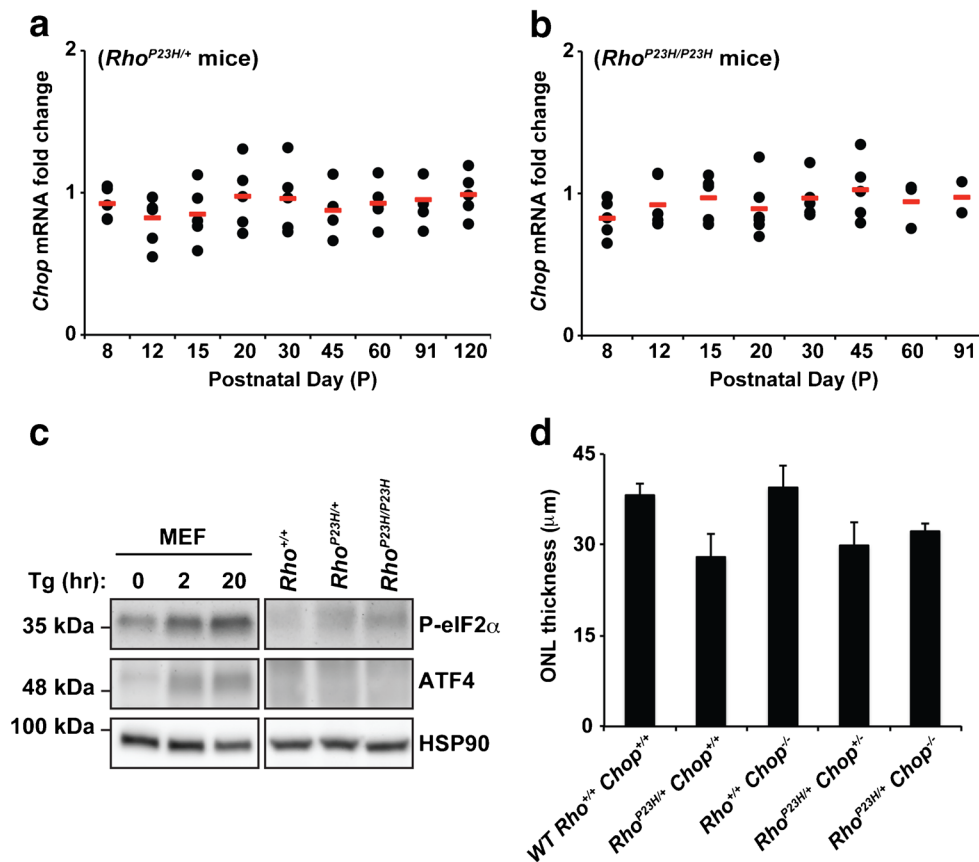


Fig. 9 *Chop* is not induced in retinas of P23H rhodopsin mice, and loss of *Chop* does not affect P23H-induced retinal degeneration. **a**, **b** *Chop* mRNA levels were quantified in the retinas of *Rho*^{P23H/+} and *Rho*^{P23H/P23H} mice and are shown relative to levels in age-matched wild-type mice at indicated ages. **c** PERK downstream targets, phospho-eIF2 α and ATF4, were analyzed by immunoblotting. Retinas from three *Rho*^{+/+}, *Rho*^{P23H/+}, or *Rho*^{P23H/P23H} mice were collected at P15, pooled, and lysed. MEF cells were treated with thapsigargin (500 nM) for indicated

times. Then, 10 μ g of MEF total lysates or 40 μ g of total retinal lysates are loaded onto the SDS-PAGE. HSP90 protein levels served as a loading control. **d** Mean ONL thicknesses of wild-type *Rho*^{+/+} *Chop*^{+/+}, *Rho*^{P23H/+} *Chop*^{+/+}, *Rho*^{+/+} *Chop*^{-/-}, *Rho*^{P23H/+} *Chop*^{+/-}, and *Rho*^{P23H/+} *Chop*^{-/-} from perfused eye enucleations were quantified at P95. Error bars represent SDs based on measurements from at least three mice of each genotype. For *Rho*^{P23H/+} *Chop*^{-/-}, only two mice were analyzed

compared to *Rho*^{P23H/+} *Chop*^{+/+} mice (Fig. 9d). We conclude that photoreceptor cell death induced by P23H rhodopsin in vivo occurs independently of CHOP.

Discussion

Our studies reveal an unexpected asymmetry in the use of the UPR by photoreceptor cells expressing P23H rhodopsin where we found strong IRE1 signaling but minimal PERK signaling. How does P23H rhodopsin trigger IRE1 so well with lesser effect on PERK? Direct interaction between IRE1's ER-luminal domain and misfolded protein is an important step in activation of IRE1 [61]. One possibility could be that misfolded P23H rhodopsin protein preferentially interacts with IRE1 but not PERK in the ER lumen of photoreceptors, and therefore the IRE1 pathway is more activated. This appears unlikely given the strong homology between the

luminal domains of IRE1 and PERK [62]. A second possibility is that PERK is actively repressed in photoreceptors through inhibition of its cytosolic kinase by binding of p58IPK or Nck proteins [63, 64]. A third possibility is that photoreceptors eliminate misfolded P23H rhodopsin so efficiently that the ambient levels of ER stress never rise to the threshold needed to trigger strong PERK signaling and its induction of the ATF4-CHOP pro-apoptotic circuit. The ER of photoreceptors is highly specialized toward the production of rhodopsin because rhodopsin is expressed in massive quantities only by rod photoreceptors [65, 43]. To cope with continuous rhodopsin protein production throughout life, the ER of photoreceptor cells must be exquisitely optimized toward folding rhodopsin as well as recognizing and removing any damaged/misfolded rhodopsin. In support of this model, we observed no obvious aggregates or retention of misfolded rhodopsin in the ER of photoreceptors and no ultrastructural alterations in ER morphology in P23H rhodopsin photoreceptors. By contrast, mutant rhodopsin aggregates are

consistently found in the ER of heterologous cell types possibly because the ER of these cells is not as well equipped to process rhodopsin as the ER of native photoreceptor cells.

Heterozygous *Rho*^{P23H/+} mice closely model the genetic dosage and slow retinal degeneration seen in patients with the P23H mutation, in whom decades typically pass before vision loss becomes detrimental [27]. In *Rho*^{P23H/+} mice, IRE1 and ERAD are also activated in photoreceptors as in *Rho*^{P23H/P23H} mice. However, after an early wave of photoreceptor cell death during the first postnatal month, photoreceptor numbers stabilize at ~50 % the number of photoreceptors found in wild-type mice (Figs. 2, 4a). Our analysis of *Rho*^{P23H/+}*ERAI*^{+/-} mice (Fig. 5) revealed coincident peak induction of the IRE1 reporter during this period as well. These findings suggest that IRE1 signaling is perpetually activated in *Rho*^{P23H/+} photoreceptor cells throughout life and cowl maintain a fine proteostatic balance within the cell whereby P23H rhodopsin (synthesized from the mutant allele) is removed, whereas wild-type rhodopsin (produced by the remaining normal rhodopsin allele) is properly folded, exported to the ROS, and vision is retained. Here, *Rho*^{P23H/+} mimics the rhodopsin haplo-insufficiency seen in *Rho*^{+/-} mice. Mild ROS and RIS photoreceptor compartment disorganization is seen in *Rho*^{+/-} mice [21]. However, rhodopsin haplo-insufficiency does not cause photoreceptor cell death or vision loss in *Rho*^{+/-} mice or people [21, 20, 66, 67]. So, why are heterozygous *Rho*^{P23H/+} photoreceptors prone to cell death, albeit it at a slow protracted pace?

One possibility is that the small amount of P23H rhodopsin that exits the ER and escapes ERAD exerts a cytotoxic effect in photoreceptor cells, not seen in cell lines. This residual P23H rhodopsin concentrates in misshapen discs within stunted rod outer segments in *Rho*^{P23H/P23H} mice whereas wild-type rhodopsin is found in normally stacked discs in the rod outer segments of *Rho*^{+/+} mice [44]. It is unclear if subcellular mislocalization of P23H rhodopsin is causally linked to the disorganization of discs or dysgenesis of the ROS compartment. However, disruption of rod outer segment structure or function would be deleterious for photoreceptor viability. Another factor affecting why heterozygous *Rho*^{P23H/+} photoreceptors are more prone to death as they age is that exogenous insults that trigger ER stress in the retina could exacerbate ER stress already induced by P23H rhodopsin. Numerous environmental factors including phototoxicity, nutrient imbalances, infections, inflammation, and toxins (e.g., tobacco) have been shown to trigger ER stress in the eye [68–71]. Addition of environmentally induced ER stressors on top of P23H rhodopsin-induced ER stress could upset the proteostatic balance maintained by the UPR signaling network leading to photoreceptor cell death.

Our analysis of misfolded rhodopsin reveals how its preferential activation of different programs and modules of the UPR, in particular ERAD and IRE1, may drive the pathophysiology of photoreceptor cell death and retinal degeneration

arising from misfolded rhodopsin-induced ER stress. Other neurodegenerative diseases associated with misfolded proteins also cause ER stress, but the UPR signaling pathways and transcriptional, translational, or post-translational UPR programs most relevant to each disease may also vary depending on the specific misfolded protein, morphologies, and functions found in the targeted neuronal cell types.

Acknowledgements We thank J. Han, N. Hiramatsu, C. Sigurdson, W. C. Smith, S. Tsang, and L. Wiseman for helpful suggestions and reagents. These studies were supported by NIH grants EY001919, P30EY002162, and EY020846, Foundation Fighting Blindness, UCSD Neuroscience Microscopy Shared Facility P30 NS047101, and VA Merit award BX002284. W.-C. Chiang received postdoctoral support from the Fight-for-Sight Foundation.

Conflict of Interest The authors declare no conflict of interest.

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Table S1. Early P23H Rod Opsin Knock-in Phenotypes*

Phenotype	Age	WT (+/+)	HET (P23H/+)	HZ (P23H/P23H)
ROS	P4	None	None	None
	P5	None	None	None
	P6	None	None	None
	P8	None	None	None
	P10	Definite short ROS	= WT	None
	P12	> P10	= WT	None**
	P15	~66% adult length	< WT	Short Nubs
RIS	P4	Normal	= WT	= WT
	P5	Normal (> P4)	= WT	= WT
	P6	Normal (> P5)	= WT	= WT
	P8	Normal (> P6)	= WT	= WT
	P10	Normal (> P8)	= WT	= or < WT
	P12	Normal (> P10)	= WT	< WT & HET
	P15	Normal (> P12)	< WT	< WT & HET
ONL Thickness/PN***	P4	Normal/0-2	= WT	= WT
	P5	Normal/2-4	= WT	= WT
	P6	Normal/1-3	= WT	= WT
	P8	Normal/0-2	= WT	= WT
	P10	Normal/0-2	= WT	= or < WT/3-12†
	P12	Normal/0-3	= WT	< WT & HET/25-34†
	P15	Normal/0-1	< WT/17-19	~50-60% WT/35+

ONL, outer nuclear layer; RIS, rod inner segments; ROS, rod outer segments (including ~3% cone RIS and ROS).

*As viewed by light microscopy of 1 μ m thick sections of epoxy resin-embedded retinas; posterior retinas scored.

Exact birthdates in 2 litters of each genotype were determined by checking for births in the morning and evening; one mouse was taken at each age from each litter.

**Electron microscopy revealed mostly ciliary protrusions with a very few ROS discs[44] not readily visible by light microscopy.

***PN, number of pyknotic nuclei per 430- μ m field; 3-4 fields were counted in the posterior retina.

†These retinas demonstrated strong central-to-peripheral gradients, as the far peripheral ONL had no PN/430- μ m field.