



Chromophore hydrolysis and release from photoactivated rhodopsin in native membranes

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For sustained vision, photoactivated rhodopsin (Rho*) must undergo hydrolysis and release of all-*trans*-retinal, producing substrate for the visual cycle and apo-opsin available for regeneration with 11-*cis*-retinal. The kinetics of this hydrolysis has yet to be described for rhodopsin in its native membrane environment. We developed a method consisting of simultaneous denaturation and chromophore trapping by isopropanol/borohydride, followed by exhaustive protein digestion, complete extraction, and liquid chromatography–mass spectrometry. Using our method, we tracked Rho* hydrolysis, the subsequent formation of *N*-retinylidene-phosphatidylethanolamine (*N*-ret-PE) adducts with the released all-*trans*-retinal, and the reduction of all-*trans*-retinal to all-*trans*-retinol. We found that hydrolysis occurred faster in native membranes than in detergent micelles typically used to study membrane proteins. The activation energy of the hydrolysis in native membranes was determined to be 17.7 ± 2.4 kcal/mol. Our data support the interpretation that metarhodopsin II, the signaling state of rhodopsin, is the primary species undergoing hydrolysis and release of its all-*trans*-retinal. In the absence of NADPH, free all-*trans*-retinal reacts with phosphatidylethanolamine (PE), forming a substantial amount of *N*-ret-PE (~40% of total all-*trans*-retinal at physiological pH), at a rate that is an order of magnitude faster than Rho* hydrolysis. However, *N*-ret-PE formation was highly attenuated by NADPH-dependent reduction of all-*trans*-retinal to all-*trans*-retinol. Neither *N*-ret-PE formation nor all-*trans*-retinal reduction affected the rate of hydrolysis of Rho*. Our study provides a comprehensive picture of the hydrolysis of Rho* and the release of all-*trans*-retinal and its reentry into the visual cycle, a process in which alteration can lead to severe retinopathies.

rhodopsin | chromophore | photoreceptors | retinoids | eye

Rod photoreceptors possess a unique cellular structure, including the rod outer segment (ROS), composed of a long stack of lipid discs packed with rhodopsin (Rho) to optimize photon capture (1). This dense packing allows for the high sensitivity to light required for scotopic vision. The responsiveness of Rho to light is afforded by the photochemistry of its Schiff base (SB)–bound 11-*cis*-retinal (11cRAL) chromophore, whose *cis-trans* isomerization induces a series of protein conformational changes, leading to the photoactivated Rho (Rho*) state (2, 3). The binding site of 11cRAL has been meticulously explored by multiple groups and finally determined in 1982 to be Lys²⁹⁶ by using sodium borohydride (NaBH₄) reduction to trap the retinylidene SB (RSB) in place, thereby preventing loss of chromophore to hydrolysis during proteolysis and chromatography (4).

Rho* generated after light exposure exists in an equilibrium of metarhodopsin states, of which metarhodopsin II (MII) greatly dominates and interacts with the G protein transducin to initiate phototransduction (5). Therefore, the *cis-trans* photoisomerization of retinal converts the native Rho ligand, which behaves as an inverse agonist, into a transient agonist that must be hydrolyzed from Rho* (6). Hydrolysis provides all-*trans*-retinal (aRAL) substrate for the visual cycle, where it is isomerized back to 11cRAL. The amine group of the Lys²⁹⁶ residue then reacts with 11cRAL to regenerate ground-state Rho. Thus, hydrolysis primes the photoreceptor for restoration of dark sensitivity (dark adaptation), which in humans takes up to 40 min for complete recovery of maximal light sensitivity (7). Notably, 11cRAL enters and aRAL exits the protein through different sites (8). The entry site does not bind aRAL and is highly selective for 11cRAL. The exit site solely allows aRAL to leave without reentry (8). Released aRAL is subsequently reduced to all-*trans*-retinol (aROL) catalyzed by NADPH-dependent retinol dehydrogenases (RDHs) (9).

To date, the crucial step of all-*trans*-RSB (aRSB) hydrolysis in the Rho photocycle has largely been studied using indirect spectrophotometric techniques that require the solubilization of Rho in detergent micelles. As such, these methods do not capture Rho

Significance

Vision starts when a photon isomerizes the 11-*cis*-retinylidene chromophore of rhodopsin to initiate phototransduction. The all-*trans*-retinylidene product is hydrolyzed and all-*trans*-retinal released, allowing rebinding of 11-*cis*-retinal to regenerate rhodopsin for sustained vision. Subsequently, all-*trans*-retinal is cleared, preventing aldehyde toxicity by reduction to all-*trans*-retinol. Defects in these metabolic processes could lead to severe retinopathies. Studying the biochemistry of key proteins responsible for these fundamental steps of vision in their native membrane environments has remained challenging. Using rapid quantitative chemical and analytical methods, we directly captured and quantified the kinetics and energetics of these essential biochemical processes occurring in native membranes. Our results shed light on the entire rhodopsin photocycle and chromophore regeneration and are broadly applicable to other retinylidene proteins.

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biochemistry in the native membrane environment (10). The broad overlapping absorbance bands of Rho, its metastates after light exposure, and released aRHAL present challenges in the study of Rho* aRSB hydrolysis using spectrophotometric techniques (11). The metastates differ in structure, including *anti*- versus *syn*-stereoisomerism and protonation state of the aRSB (12). Interestingly, aRSB protonation in Rho* behaves inversely with pH in the following manner: lower pH stabilizes the deprotonated aRSB, MII, whereas higher pH stabilizes the protonated aRSBs, MI and MIII (6, 13, 14). Among these photoproducts, MII is understood to be the signaling state of Rho and has also been observed to be most susceptible to aRSB hydrolysis (5, 15). However, the hydrolysis of SBs is facilitated via nucleophilic attack of water on a protonated SB-carbon rather than a deprotonated SB-carbon, as found in MII (16). Therefore, the mechanisms of the hydrolysis and the Rho intermediate most susceptible to hydrolysis require more investigation.

Another factor that complicates the study of aRSB hydrolysis with detergent-solubilized Rho is the loss of phospholipids, in particular phosphatidylethanolamine (PE), which can form adducts with the released aRHAL and potentially affect the rate of the hydrolysis reaction (17). Moreover, these *N*-retinylidene-PE (*N*-ret-PE) adducts have documented roles as substrates for the visual cycle transport protein ABCA4, as well as in photic regeneration pathways (18). As such, a consideration of their formation and clearance is critical for understanding alternative chromophore regeneration pathways, as well as the pathophysiology of ABCA4-associated retinal diseases, such as Stargardt disease-1 (19). Furthermore, the reduction of aRHAL by RDHs in the context of the retina, where it can form PE adducts, needs clarification.

In this study, the hydrolytic release of aRHAL from Rho* within native bovine ROS membranes was monitored by direct quantification of the opsin-bound RSBs in their different *cis-trans* configurations after light exposure of Rho in ROS membranes and of the aRHAL hydrolysis product. This was achieved by using simultaneous alcohol protein denaturation and NaBH₄ reduction to trap RSBs as nonhydrolyzable retinyl secondary amine adducts. After proteolysis, these retinyl signatures and their *cis-trans* isomeric configurations were detected by liquid chromatography–tandem mass spectrometry (LC-MS/MS) and used to monitor opsin RSBs over time. Concurrently, both free and PE-bound aRHAL were tracked in the alcohol-solubilized lipid fraction. The energetic landscape of Rho*-aRSB hydrolysis was further elucidated using quantum chemical methods.

Results

Proteolysis and LC-MS/MS Analysis of N^e-at-retinyl-peptide (bovine Rho 292–302). Stereospecific detection of the trapped RSB within opsin has not been achieved to date because of the high hydrophobicity of the resulting retinyl peptides and the lability of the retinyl moiety. Therefore, RSB isomerization and hydrolysis could not be studied, except indirectly by spectrophotometric techniques. To establish a direct LC-MS/MS detection method, we first tested a synthetic peptide consisting of residues 292 to 302 of the bovine Rho sequence (bRhoPep) (*SI Appendix, Fig. S1A*). NaBH₄ reduction trapped the aRHAL-SB adducted to the ε amine group of Lys²⁹⁶ of bRhoPep, forming a nonhydrolyzable retinyl secondary amine adduct (*SI Appendix, Fig. S1B*). The LC-MS peak of this retinyl peptide showed the characteristic ultraviolet-visible (UV-Vis) spectrum of the retinyl moiety ($\lambda_{\max} = 328$ nm), which greatly enhanced the hydrophobicity of the peptide, as reflected in the increased chromatographic

retention time from 5 to 31 min (*SI Appendix, Fig. S1 C and D*). MS/MS showed electrospray ionization (ESI)–source fragmentation of the N^e-retinyl-peptide analyte into a retinyl cation and a peptide ion, whose sequence was confirmed by collision-induced dissociation (CID) fragmentation to be that of the bRhoPep (*SI Appendix, Fig. S1 E–G*).

Because stereospecific MS detection of the whole N^e-retinyl-opsin protein is unfeasible, a proteolytic digestion of N^e-retinyl-opsin to N^e-retinyl-peptides would be necessary for detection and sequence determination by LC-MS/MS. This principle was tested by proteolysis of the N^e-retinyl-bRhoPep using proteinase K or the protease mixture pronase. The proteinase K digest yielded four unique N^e-retinyl-peptides, identified by MS/MS to be K^{ret}TS, K^{ret}T, AK^{ret}, and FAK^{ret}, each with a sequence containing Lys²⁹⁶ adducted to the retinyl moiety (*SI Appendix, Fig. S2*). Complete proteolysis of N^e-at-retinyl-bRhoPep down to N^e-at-retinyl-Lys by pronase at 40 °C yielded one prominent peak and two minor peaks, which were all identified to be N^e-retinyl-Lys (*SI Appendix, Fig. S3*). Based on the UV-Vis and MS spectra of each synthetic retinyl isomer standard of N^e-retinyl-Lys (*SI Appendix, Fig. S4*), the most prominent peak was determined to be the all-*trans* isomer, whereas the two minor peaks were determined to be the 9-*cis* and 13-*cis* isomers. These two minor isomers are known to form spontaneously by thermal isomerization of the all-*trans* isomer (20). Pronase was found to completely proteolyze N^e-retinyl-opsin at 10 °C, which minimized the thermal isomerization and retained the isomeric composition. The LC separation followed by UV-Vis and MS spectral identification of the four N^e-retinyl-Lys isomers (9-*cis*, 11-*cis*, 13-*cis*, and all *trans*) as standards (*SI Appendix, Fig. S4*) demonstrated the potential of this method for capturing the photochemistry of retinylidene proteins following trapping of the RSB by NaBH₄ and proteolysis by pronase.

Proteolysis and LC-MS/MS Analysis of N^e-retinyl-opsin. To study Rho biochemistry in native membranes, a saturated solution of NaBH₄ in isopropanol (*i*PrOH) was added to ROS to rapidly denature the protein and, simultaneously, reduce the RSB, thus trapping the retinal moiety bound to opsin. Remarkably, this procedure can be used to study Rho either in native membranes (Fig. 1*A*) or purified in detergent solutions (*SI Appendix, Fig. S5*). Pronase converted a suspension of the denatured proteins into a clarified solution that could be analyzed by LC-MS/MS. Prior to processing the samples, we used UV-Vis spectroscopy to determine the photostate of Rho in ROS membranes. The 500-nm absorbance maximum of Rho declines after light exposure, producing a new maximum at 380 nm, which is characteristic of MII (Fig. 1*B* and *C*). From the pronase digest, Rho yielded a dominant N^e-11-*cis*-retinyl-Lys peak, whereas Rho* yielded a dominant N^e-all-*trans*-retinyl-Lys peak, as confirmed by LC-MS/MS and comparison with our synthetic standards (Fig. 1*D–J*).

The N^e-retinyl-Lys peaks were confirmed to originate from Rho in ROS membranes by LC-MS/MS analysis of the N^e-retinyl-peptides generated from a proteinase K digestion of the ROS-protein precipitate. Proteinase K also clarified the suspension of protein precipitate into a solution of N^e-retinyl-peptides that were identified by LC-MS/MS to be similar to those from the proteinase K digest of N^e-retinyl-bRhoPep (Fig. 2). The proteinase K and pronase digests of ROS-protein precipitates from treatment of ROS membranes with sodium borodeuteride (NaBD₄) in *i*PrOH generated similar results, showing the same N^e-retinyl analytes with an increase in mass corresponding to

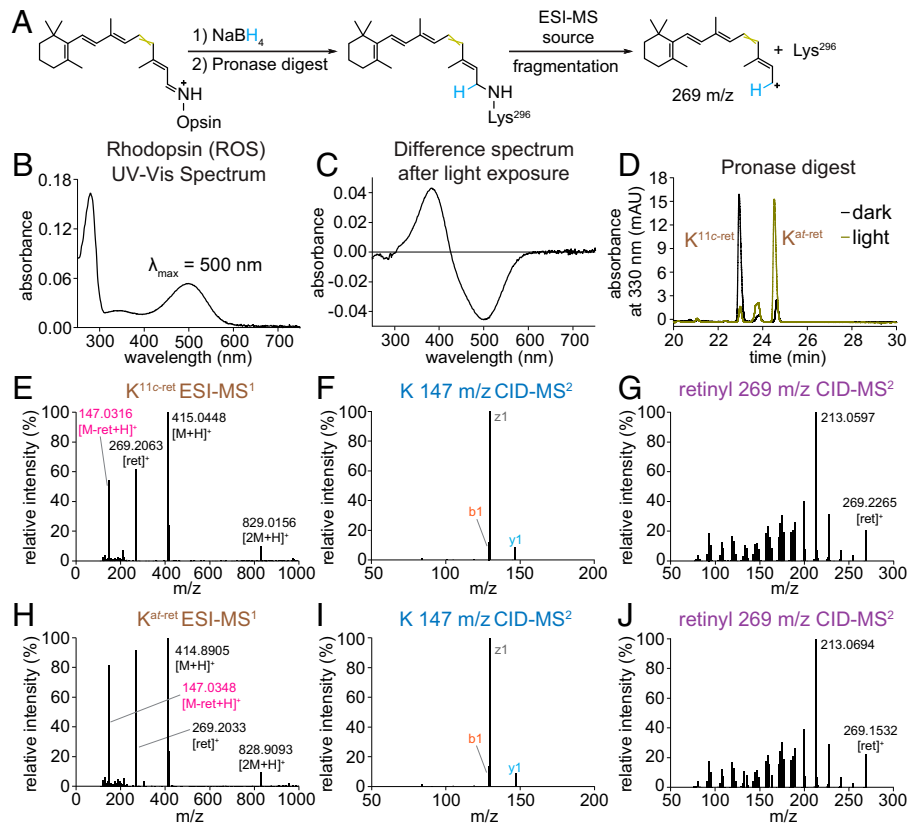


Fig. 1. LC-MS/MS analyses of pronase digests of Rho and Rho* from native membranes. (A) Schematic diagram of sample preparation and MS workflow. (B) UV-Vis spectrum of Rho from ROS membranes solubilized in DDM. (C) Difference absorbance spectrum of DDM-solubilized Rho versus Rho* in ROS membranes illuminated with 565-nm fiber light at 625 μ W for 10 s at 4°C. (D) Chromatographic separation of N^ε-retinyl-Lys peaks from the pronase digests of Rho and Rho* in ROS membranes treated with NaBH₄ in *i*PrOH. (E, H) ESI-MS spectrum showing characteristic cleavage of retinyl cation (269 m/z) from precursor N^ε-retinyl-Lys analyte (415 m/z), producing a product Lys ion (147 m/z). (F, I) CID-MS/MS spectrum of the product Lys ion exhibiting a characteristic fragmentation pattern of Lys. (G, J) CID-MS/MS spectrum of retinyl cation demonstrating a complex fragmentation pattern with a characteristic dominant 213-m/z signal (39, 40). mAU, milli absorbance unit.

deuteration and source fragmentation producing the C¹⁵-D¹-retinyl cation at 270 m/z (*SI Appendix*, Figs. S6 and S7). Our NaBD₄ results align with the NaBH₄ reduction of the RSB of Rho or Rho* and with the characteristic ESI-source fragmentation of the retinyl moiety.

Separation and Identification of *N*-ret-PEs. We expected that, concomitantly with the trapping of the bound aR_{AL} moiety to Rho* by NaBH₄, any *N*-ret-PEs formed by SB condensation of aR_{AL} with PE after Rho* hydrolysis would also be captured by NaBH₄ reduction as *N*-retinyl-PEs (Fig. 3A). Using LC-MS/MS and CID fragmentation of *N*-retinyl-PEs, we determined that the main fatty acyl chains in the sn-2 position of PE are docosahexaenoic acid (22:6), docosapentaenoic acid (22:5), and docosatetraenoic acid (22:4) (Fig. 3B), which are most abundant in ROS membranes (21) (*SI Appendix*, Fig. S8).

In the dark, ROS treated with NaBH₄ in *i*PrOH had a small signal of aR_{OL}, the NaBH₄-reduction product of aR_{AL}, with no other discernible signals of retinyl species (Fig. 3C). A small increase in aR_{OL} was observed in ROS briefly exposed to light. Incubation in the dark for 16 min after brief exposure to light led to a large increase in the aR_{OL} signal, as well as the appearance of six major peaks with distinct relative peak height/area ratios identified by LC-MS/MS as the *N*-retinyl-PEs described above. The addition of exogenous aR_{AL} to ROS membranes unexposed to light produced the same fingerprint of these six *N*-retinyl-PE peaks (Fig. 3D). Exposure to light followed by a 16-min incubation in the dark led to an increase in aR_{OL} and

N-retinyl-PE signals corresponding to the contribution from NaBH₄-reduced free and PE-bound aR_{AL} from Rho* hydrolysis (Fig. 3D). Therefore, addition of NaBH₄ in *i*PrOH to ROS membranes enabled continuous tracking of the aR_{AL} following its hydrolytic release from Rho*, even in the presence of exogenous aR_{AL}.

Kinetics of Rho* Hydrolysis and Subsequent *N*-ret-PE Formation.

Our trapping procedure enabled a detailed kinetic analysis of each step as it occurred in the native phospholipid bilayer (Fig. 4A). NaBH₄ in *i*PrOH was added to ROS at different time points after illumination to trap the remaining Rho*-bound aR_{AL}, the free aR_{AL}, and subsequently formed *N*-ret-PEs. Snapshots of the progressive hydrolysis of Rho* to form free aR_{AL} and the subsequent reaction of the aR_{AL} with the lipid membrane are shown by chromatograms of the NaBH₄-reduced opsin-bound RSBs over time in Fig. 4B and the NaBH₄-reduced free and PE-bound aR_{AL} over time in Fig. 4C. The decreasing N^ε-all-*trans*-retinyl-Lys signal and increasing aR_{OL} and *N*-retinyl-PE signals over time reflect the time course of Rho*-aR_{SB} hydrolysis and aR_{AL} release (Fig. 4B and C). The rates of hydrolysis followed pseudo first-order decay kinetics and increased with temperature (Fig. 4D).

The aR_{AL} released by hydrolysis of Rho* subsequently reacted reversibly with PE, reaching a steady state of ~40% aR_{AL} bound to PE as *N*-ret-PE (Fig. 4E). The kinetics of this *N*-ret-PE formation also followed pseudo first-order kinetics. The overall rate of *N*-ret-PE formation, starting from the hydrolysis of Rho*

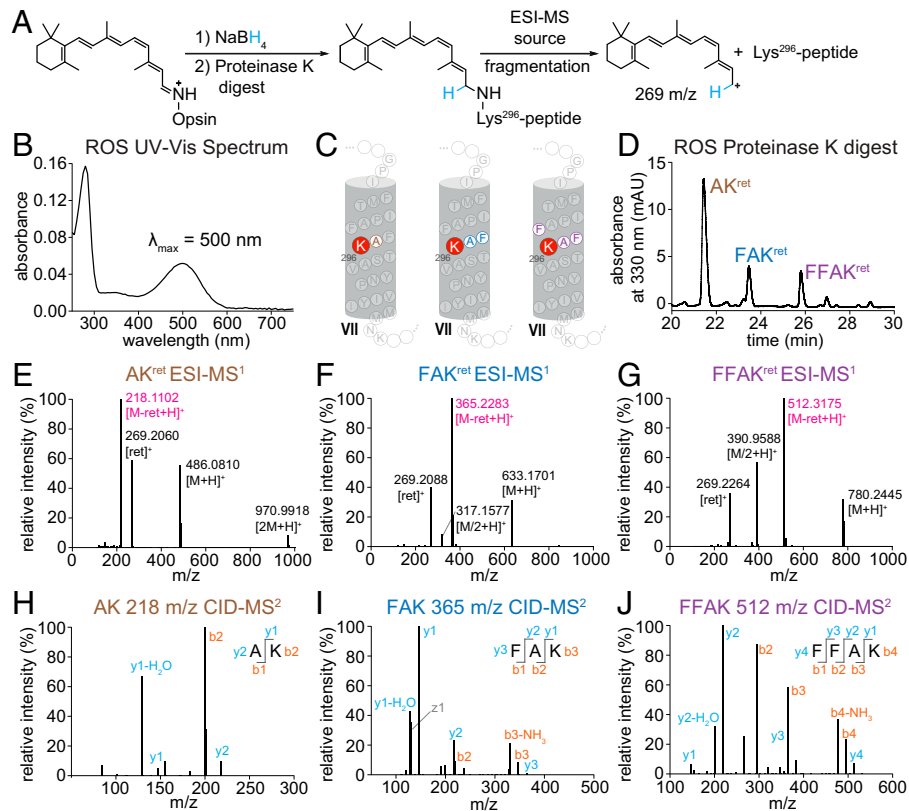


Fig. 2. LC-MS/MS analysis of proteinase K digest of Rho from native membranes. (A) Schematic diagram of sample preparation and MS workflow. (B) UV-Vis spectrum of Rho from ROS membranes solubilized in DDM. (C) Location of chromophore binding residue within helix VII of bovine Rho, with labeled N^{ϵ} -retinyl-peptide fragments detected from the proteinase K digest. (D) Chromatographic separation of N^{ϵ} -retinyl-peptides from proteinase K digestion of Rho treated with NaBH_4 in $i\text{PrOH}$. (E–G) ESI-MS spectrum showing characteristic cleavage of retinyl cation (269 m/z) from precursor N^{ϵ} -retinyl-peptide analyte, producing a product peptide ion. (H–J) MS/MS spectrum of CID fragmentation of the product peptide ion for sequence determination. mAU, milli absorbance unit.

to the release of $a\text{RAL}$ for SB condensation with PE, was found to increase with increased temperature; however, the steady-state level of N -ret-PE decreased with increased temperature (Fig. 4E). The proportion of $a\text{RAL}$ released via Rho^* hydrolysis and not bound to PE showed an inverse relationship with the formation of N -ret-PEs (Fig. 4F).

From the temperature dependence of the Rho^* hydrolysis reaction, an Arrhenius plot (Fig. 4G) was generated, yielding an activation energy (E_A) of 17.7 ± 2.4 kcal/mol. The rate of Rho^* hydrolysis was also measured in n-dodecyl- β -D-maltoside (DDM) micelles to mimic conditions of mild detergent solutions that are typically used for spectrophotometric measurements of Rho. The rate of hydrolysis in DDM was found to be $\sim 50\%$ slower than that in the ROS membranes (Fig. 4H).

pH Dependency of Rho^* Hydrolysis and N -ret-PE Formation.

The protonation state of the iminium nitrogen of Rho^* - $a\text{RSB}$ is known to behave inversely to changes in the external protic environment of Rho^* (22). This behavior was observed by UV-Vis spectroscopy, showing that higher pH shifts the metarhodopsin equilibrium from MII, the deprotonated state, toward protonated states, namely MI and MIII (Fig. 5A). Rho^* - $a\text{RSB}$ hydrolysis occurred to greater extents with decreasing pH, where MII is favored (Fig. 5B). However, lower pH appeared to slow down the overall rate of N -ret-PE formation (Fig. 5C). The amount of free $a\text{RAL}$ was inversely proportional to the amount of N -ret-PE (Fig. 5D). To isolate the reaction between $a\text{RAL}$ and PE and obtain the rate of N -ret-PE formation independently, exogenous $a\text{RAL}$ was added to ROS membranes and tracked by addition of NaBH_4 in $i\text{PrOH}$ at

different time points. The representative chromatograms of the SB condensation reaction of $a\text{RAL}$ with PE over time showed the decline in $a\text{RAL}$ starting material with the formation of N -ret-PEs (Fig. 5E). The rate of N -ret-PE formation exhibited pH dependency, with a faster rate of reaction as pH increased (Fig. 5F). In summary, Rho^* - $a\text{RSB}$ hydrolysis was faster at lower pH, which favors MII, whereas N -ret-PE formation was slower at lower pH.

Computational Analysis of Rho^* - $a\text{RSB}$ Hydrolysis. We explored the mechanistic basis for the pH dependency of Rho^* - $a\text{RSB}$ hydrolysis by computationally modeling the energy landscapes of $a\text{RSB}$ hydrolysis for each metarhodopsin species. We examined the influence of Glu¹¹³ and Glu¹⁸¹ on the energy levels of each metarhodopsin species and subsequent intermediates of hydrolysis, considering the location of these Glu residues in the chromophore binding pocket, their influence on spectral characteristics of the retinylidene chromophore, and their role in forming the water and H-bonding network that surrounds the chromophore and spans the Rho molecule (23, 24). Upon photoactivation to MII, Glu¹¹³ and Glu¹⁸¹ shift from their position in Rho, translocating so that both surround and interact with the $a\text{RSB}$ (23, 25). Plausibly, the protonation states of Glu¹¹³ and Glu¹⁸¹ could vary with pH and determine whether a charged or neutral carboxyl residue would exist near the $a\text{RSB}$ in Rho^* , which can exist in either a *syn*- or *anti*-configuration. In MI, the configuration of its protonated $a\text{RSB}$ is 15-*anti*, whereas in MIII, its protonated $a\text{RSB}$ is 15-*syn* (12, 26). The exact stereoisomerism of the deprotonated $a\text{RSB}$ of MII is unclear, but it could loosely adopt either a 15-*anti* or 15-*syn* configuration (12, 26). For either

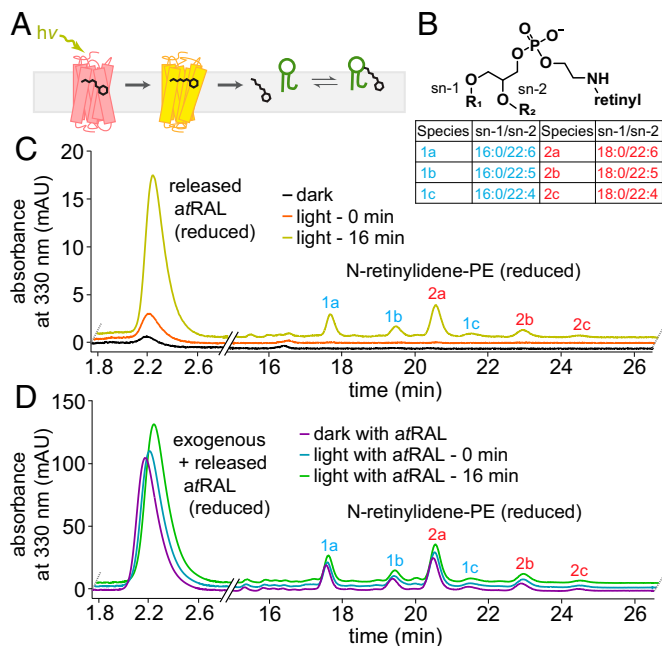


Fig. 3. Formation of *N*-ret-PE in native membranes after light exposure. (A) Diagram of *N*-ret-PE formation after illumination of ROS membranes. (B) Chemical structure of *N*-ret-PE, with a table of major species found in the ROS membranes. (C) Chromatogram of *aRtRAL* released and resultant *N*-ret-PE in ROS at 0 and 16 min after illumination for 10 s with 565-nm fiber light at 625 μ W. (D) Analogous experiment to that in C, but with exogenous *aRtRAL* added to the ROS membrane suspension prior to illumination. mAU, milli absorbance unit.

the protonated or deprotonated form of the *aRtRSB*, the *anti*-configuration is still dominant over *syn*, because the *anti-syn* interconversion requires additional light or thermal energy (12, 26). We carried out quantum chemical calculations to map the energetic landscape of hydrolysis as influenced by the *syn-anti* configuration and protonation state of the *aRtRSB*, as well the protonation states of Glu¹¹³ and Glu¹⁸¹. Our calculations were performed on a hydrogen-saturated cluster model of the Rho active site derived from the current highest-resolution structure of a metarhodopsin species to date, 3PQR, proposed to correspond to MII, with a deprotonated *anti-aRtRSB* structure (*SI Appendix*, Fig. S9).

For protonated *aRtRSB*, favored at high pH, our initial calculations showed preference for the deprotonated states of both Glu¹¹³ and Glu¹⁸¹. Notably, the protonated *syn-aRtRSB* was found to be 3.3 kcal/mol lower in energy (more stable) than the *anti*-configuration. However, water addition is only exothermic for the transition of the protonated *anti-aRtRSB* to an oxonium intermediate (Fig. 6 B, *Ib*) (−3.7 kcal/mol), followed by exothermic proton transfer (−2.0 kcal/mol) between the oxonium and nitrogen, as facilitated by the polar protic environment of the pocket (Fig. 6). The final C-N bond cleavage is considerably endothermic such that the final products (Fig. 6 B, *Id*) were 2.0 kcal/mol higher in energy than reactants (Fig. 6 B, *Ia*). The hydrolysis of the protonated *syn-aRtRSB* was also found to be an overall endothermic process (Fig. 6). In contrast, hydrolysis of either the *anti*- or *syn*-isomers of the deprotonated *aRtRSB* was overall exothermic (Fig. 6). Under low pH conditions favoring the deprotonated *aRtRSB*, our calculations showed a preference for the deprotonated state of Glu¹⁸¹ and the protonated state of Glu¹¹³. The deprotonated *anti-aRtRSB* was markedly lower in energy than the corresponding *syn*-isomer by 4.4 kcal/mol. Water addition to either the *anti*- or *syn*-isomers of deprotonated *aRtRSB* was considerably exothermic,

as was the proton transfer from Glu¹¹³ eventually to the nitrogen of the Lys²⁹⁶-carbinolamine intermediate, likely via water or the H-bonding network of the pocket (Fig. 6 C, *IId*). At low pH, Glu¹¹³ becomes protonated (Fig. 6 C, *IId*). The reprotonation of Glu¹¹³ followed by the C-N bond cleavage are both endothermic; however, the overall energy change in the process is exothermic by ~8.0 kcal/mol (Fig. 6 C, *IId*), in strong contrast to the endothermic reaction at higher pH. Assuming linear relationships between effective activation barriers and overall change in energy, we conclude that changes in pH affect the protonation patterns of Glu¹¹³, Glu¹⁸¹ carboxyl moieties, and the Lys²⁹⁶-*N*-atom, which in turn have a decisive effect on the energetics of hydrolysis of the different metarhodopsin species and release of *aRtRAL*.

Reduction of *aRtRAL* in Native Membranes. In the visual cycle, *aRtRAL* is reduced to *aRtROL* by RDHs (9). Our method using NaBD₄ in place of NaBH₄ allowed us to track this step of the visual cycle in ROS, thereby studying the fate of *aRtRAL* after its hydrolytic release from Rho* in the presence of the essential dinucleotide cofactor NADPH (Fig. 1A, *SI Appendix*, Fig. S10). We found that *aRtRAL* release from hydrolysis of Rho*-*aRtRSB* occurred to a similar extent in the presence or absence of NADPH (NADP was used as a control), suggesting that *aRtRAL* reduction by RDHs does not influence the biochemistry of Rho*-*aRtRSB* hydrolysis (Fig. 7B). However, nearly all of the released *aRtRAL* was reduced to *aRtROL* in the presence of NADPH at both 8 and 16 min postillumination (Fig. 7C). Notably, there were almost undetectable levels of *N*-ret-PEs in the presence of NADPH at both time points (Fig. 7C).

To determine whether *N*-ret-PEs play a role in mediating the delivery of *aRtRAL*, NADPH was added to ROS-containing *N*-ret-PE, as prepared by addition of exogenous *aRtRAL* followed by a 16-min incubation to reach the steady state of *N*-ret-PE formation at pH 7.4. The *aRtRAL* from *N*-ret-PEs appeared not to be a direct source of substrate for the RDHs, because the proportion of *N*-ret-PEs was unaffected at the instant of NADPH addition (Fig. 7D and E). Furthermore, the presence of NADPH did not accelerate the dissociation of *N*-ret-PEs as would have been observed if *N*-ret-PEs played an indispensable role in *aRtRAL*-substrate delivery to RDHs (Fig. 7D and E). Within the first 4 min, *N*-ret-PE levels were reduced to a small extent, but nearly 25% of free *aRtRAL* was reduced by the RDHs (Fig. 7E). Therefore, the RDHs exhibited their activity on free *aRtRAL* in native membranes rather than on PE-bound *aRtRAL*. By 32 min, nearly all of the *aRtRAL* had been reduced, with most *aRtRAL* dissociated from PE (Fig. 7E).

Discussion

A critical step in the visual cycle is the release of the spent chromophore, *aRtRAL*, which is ultimately transformed back to 11-*cRtRAL*. The steps immediately following retinal photoisomerization (i.e., hydrolytic release of *aRtRAL*, formation of *N*-ret-PE, and reduction of *aRtRAL* to *aRtROL*) have only been subjected to preliminary investigation because of technical issues, particularly in membranes. The problems, just to list a few, include the difficulty in isolating long hydrophobic peptides, heterogeneity of the digestion product from the transmembrane segments of Rho, complete decomposition of retinyl peptides by elimination of the retinyl carbocation moiety, and lack of quantitative preservation of the geometric isomers of the retinoids. Because of the importance of these fundamental biochemical processes in the physiology of mammalian vision, we focused on developing

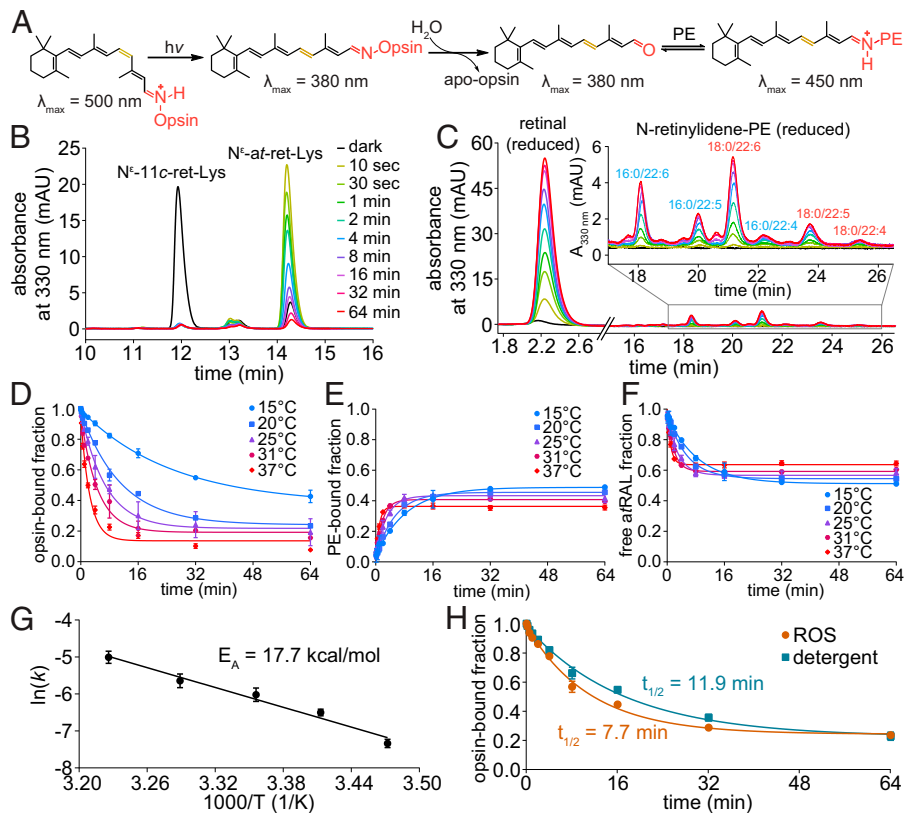


Fig. 4. Temperature dependence of Rho* hydrolysis and subsequent *N*-ret-PE formation. (A) Scheme of hydrolysis reaction following light exposure. (B) Chromatograms of pronase digests of Rho* protein pellets obtained from ROS membranes treated with NaBH₄ in *i*PrOH after brief light exposure (565-nm fiber light at 625 μW for 10 s) and subsequent incubation in the dark for different periods of time at 37°C. (C) Chromatograms of the *i*PrOH-solubilized lipid fractions from the corresponding samples from B for detection of NaBH₄-reduced free and PE-bound atRAL from Rho* hydrolysis at 37°C. (D) Rho* hydrolysis exhibited pseudo first-order decay kinetics at various temperatures. (E) The formation of *N*-ret-PEs at different temperatures from atRAL produced by Rho* hydrolysis followed pseudo first-order kinetics. (F) The proportion of the atRAL from Rho* hydrolysis that was unbound to PE tracked inversely to the proportion that formed PE adducts. (G) The Arrhenius plot of hydrolysis rate constants at different temperatures yielded an E_A of 17.7 kcal/mol. (H) Rho* hydrolysis was faster in ROS membranes than in DDM detergent micelles at 20°C. mAU, milli absorbance unit.

robust and direct methods to study them, thereby furthering our understanding of these biochemical reactions.

The highly robust proteases, proteinase K and pronase, were effective in the digestion of N^E-retinyl-opsin for subsequent MS analysis. Proteinase K unequivocally produced peptide profiles unique to Lys²⁹⁶, whereas pronase digested N^E-retinyl-opsin

completely, yielding the definitive retinyl-Lys²⁹⁶ adduct. The complete proteolysis of *i*PrOH/NaBH₄-treated Rho and Rho* by pronase and subsequent LC-MS/MS analyses enabled the determination of the *cis-trans* isomeric composition of the RSB. Thereby, Rho photochemistry was captured, with the observation of a dominant N^E-11-*cis*-retinyl-Lys from Rho and a

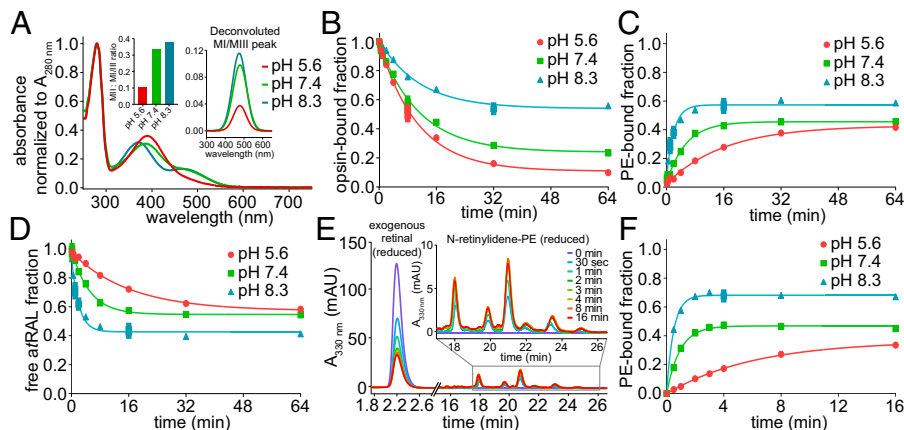
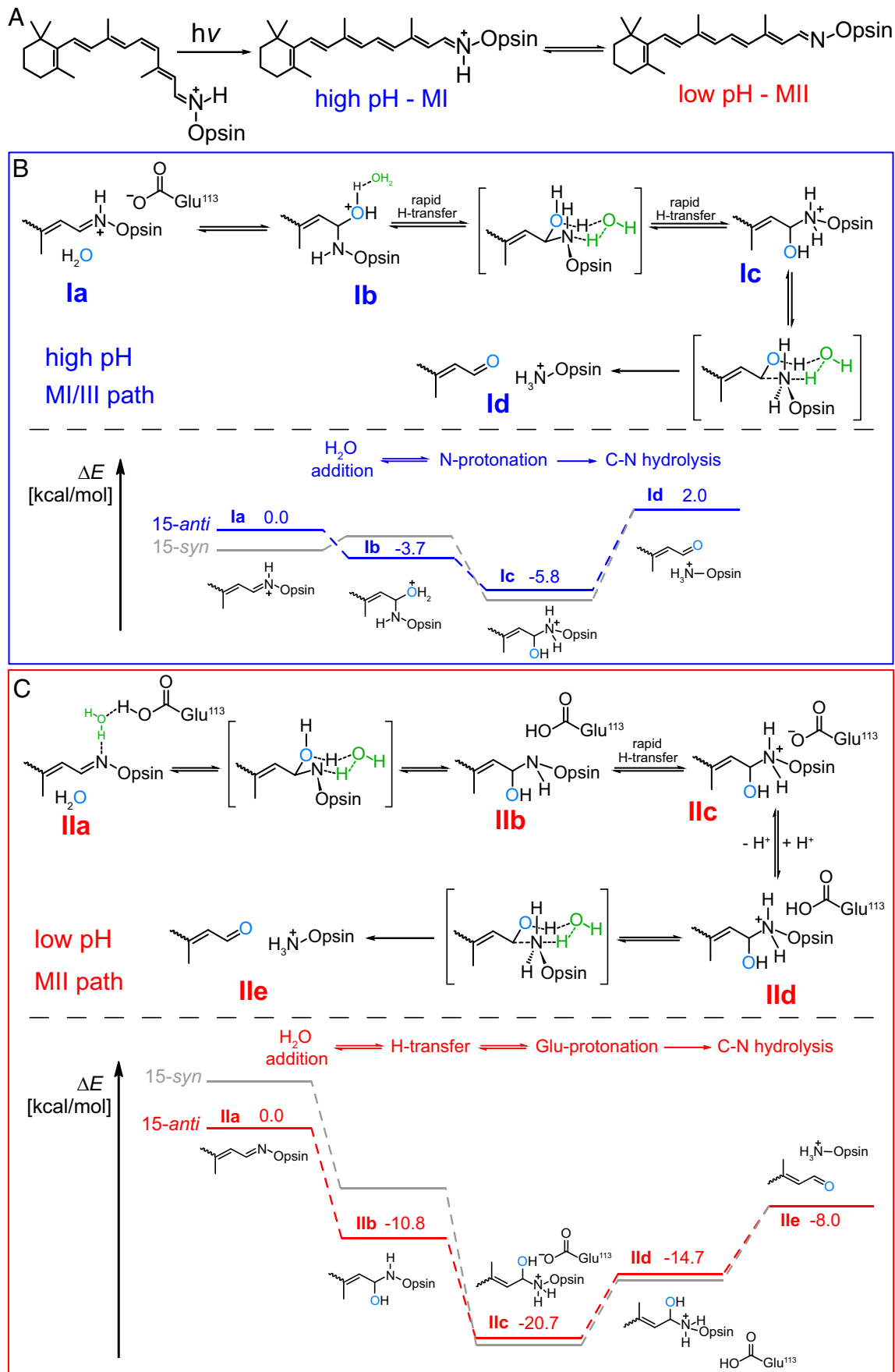


Fig. 5. pH-dependent Rho* hydrolysis and *N*-ret-PE formation in native membranes. (A) UV-Vis spectrum of Rho after illumination with a 565-nm fiber light at 625 μW for 10 s and incubation for 12 h at 4°C. Spectral analysis and deconvolution of the protonated retinylidene SB peak demonstrated inverse pH dependency, with a greater proportion of protonated retinylidene SB (MI/III) appearing with increasing pH. (B) Hydrolysis of atRAL from Rho* at 20°C occurred to the greatest extent at lower pH in which the MI-III equilibrium shifts toward MII. (C) The rate of *N*-ret-PE formation at 20°C exhibited pH dependency and was fastest at high pH. (D) The proportion of Rho* hydrolysis-derived atRAL that did not form PE adducts tracked inversely to the proportion that formed PE adducts. (E) Chromatographic traces of *N*-ret-PE formation over time at pH 8.3 and 20°C, after addition of exogenous atRAL to ROS membranes. (F) The mole fraction of *N*-ret-PE formed at 20°C from exogenous atRAL over time exhibited pseudo first-order kinetics, with faster rates and higher steady state as pH increased. mAU, milli absorbance unit.



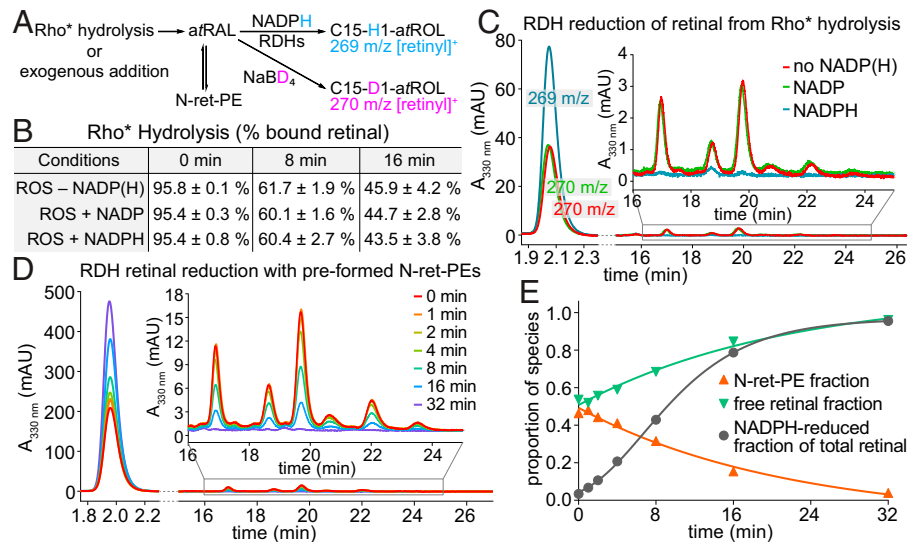


Fig. 7. Reduction of atRAL by RDHs in native membranes. (A) Simplified diagram from *SI Appendix, Fig. S10* depicting NaBD₄-based monitoring of RDH activity by tracking NADPH-reduced vs. NaBD₄-reduced retinal. (B) In ROS membranes, Rho* hydrolysis at pH 7.4 and 20 °C is unaffected by addition of dinucleotides, NADP, or NADPH. (C) At 16 min of Rho* hydrolysis, the released atRAL was reduced by RDHs in the presence of NADPH, substantially attenuating the formation of *N*-ret-PEs. (D) In the presence of both free atRAL and *N*-ret-PEs in equilibrium at pH 7.4 and 20 °C after addition of exogenous atRAL, RDH activity was initiated by addition of NADPH. The NADPH-reduced atRAL and NaBD₄-reduced free and PE-bound atRAL were monitored over time by LC-MS. (E) In samples corresponding to those of D, the fraction of NADPH-reduced atRAL followed sigmoidal growth kinetics, with the concurrent reduction of the initially available free atRAL and the atRAL gradually supplied from the equilibrium with *N*-ret-PE, as demonstrated by the slow decline of RDH activity over the time course. mAU, milli absorbance unit.

dominant N^e-all-*trans*-retinyl-Lys from Rho*. Thus, our methods captured the 11-*cis* to all-*trans* photoisomerization characteristic of Rho and the precise Lys²⁹⁶ residue that formed visual pigment via SB condensation with 11atRAL chromophore, demonstrating the ability to directly follow the course of Rho photochemistry.

MS analyses of N^e-retinyl-peptides and N^e-retinyl-Lys revealed the tendency of the retinyl group to cleave as a retinyl cation during source fragmentation, likely because of the resonance stabilization of the retinyl cation and the formation of a stable peptide or Lys leaving group upon protonation of the ε-amine by formic acid in the mobile phase. With other sites of protonation, residual precursor molecules can also be readily detected, but with retinyl esters or retinol, the retinyl cation is the predominant species observed on ESI-MS full scan (27–30). The partial source fragmentation of the precursor molecule consistently yields a product retinyl cation and a product peptide or Lys ion, which can be further characterized by CID fragmentation for sequence determination. We expect that our method can be extended to other molecules that possess a retinyl moiety as a fluorophore or a tag, together with a variety of protonatable functional groups.

After characterizing the photochemistry of Rho, we tracked Rho* hydrolysis directly by monitoring the opsin-SB-bound and -released atRAL over time. The rate of hydrolysis was faster with Rho* in ROS membranes than in DDM detergent micelles, demonstrating how the nature of the lipid environment can alter molecular dynamics. Our measured E_A of 17.7 kcal/mol for Rho* hydrolysis in ROS membranes at pH 7.4 is in agreement with early measurements by photocalorimetry, which produced values of 22.1 kcal/mol at pH 8.0 and 11.7 kcal/mol at pH 5.4 for Rho* in ROS membranes (31). The lower E_A of hydrolysis at pH 5.4 also aligned with the results of our Rho* hydrolysis experiments with ROS, which showed hydrolysis occurring to a greater extent at lower pH, thereby supporting the longstanding notion that hydrolysis proceeds via the MII signaling state of Rho. Our computational results further support the favorability

of hydrolysis proceeding via the deprotonated SB of MII over the protonated SB of MI or MIII. Our computational studies confirmed the key influence of the protonation states of the Glu¹¹³ and Glu¹⁸¹ residues on the energetic landscape of the reaction, with the hydrolysis process being exothermic only at lower pH.

The formation of *N*-ret-PE also displayed pH dependency, with faster rates and higher steady states as pH increased, likely because of a greater proportion of free PE available for nucleophilic attack on the C¹⁵ aldehyde carbon of atRAL to initiate the SB condensation reaction. Therefore, pH changes had opposite effects on the rate of Rho* hydrolysis versus the rate of *N*-ret-PE formation; at lower pH, there was more Rho* hydrolysis but less *N*-ret-PE formation. Furthermore, the overall rate of *N*-ret-PE formation from released atRAL was slower than the rate of Rho* hydrolysis at low pH ($k^{\text{hyd}} = 0.001537 \text{ s}^{-1}$ vs. $k^{\text{NretPE}} = 0.001062 \text{ s}^{-1}$). Prior to our examination of the kinetics of *N*-ret-PE formation, we hypothesized that PE could potentially play a role in facilitating the hydrolysis of the atRSB of Rho* by serving as a proximal acceptor of the released atRAL. However, based on our pH studies of the Rho* hydrolysis and the *N*-ret-PE condensation reaction, the rate of hydrolysis did not appear to be influenced by the subsequent formation of *N*-ret-PE with the atRAL from Rho* hydrolysis. Instead, the hydrolysis of the atRSB relied on the protein dynamics of Rho, reflecting the dependency of the hydrolysis kinetics on pH-driven changes in the metarhodopsin equilibrium.

At physiological pH, the rate of *N*-ret-PE formation was an order of magnitude faster than the rate of Rho* hydrolysis ($k^{\text{hyd}} = 0.001479 \text{ s}^{-1}$ vs. $k^{\text{NretPE}} = 0.01819 \text{ s}^{-1}$). The observations of *N*-ret-PE formation from exogenous or Rho*-derived atRAL and their faster rate of formation as compared with Rho* hydrolysis are significant to our understanding of the formation of toxic bis-retinoids that are abundant with Stargardt disease and age-related macular degeneration (19, 32, 33). This notion is consistent with hydrolysis being rate limiting such that soon after atRAL is released, it can be captured by PE.

Furthermore, our results show that exogenous or Rho* hydrolysis-derived *a*RAL react reversibly with PE to reach a steady-state mole fraction of *N*-ret-PE of 0.4. Therefore, after intense light exposure to the retina, close to half of all the resultant *a*RAL released from Rho* has the propensity to form *N*-ret-PEs, which serve as the starting point to drive the formation of bisretinoids. Although bisretinoids were not observed under the conditions and timescale of our studies of Rho* hydrolysis, their formation could be facilitated by any number of factors, including delayed reduction (e.g., by intense illumination and depletion of NADPH), attenuated phagocytosis, and genetic alterations of pertinent visual cycle proteins and enzymes (34, 35). Longer durations of light exposure and incubation periods with released *a*RAL or accelerated supply of 11*c*RAL that is constitutively supplied by the retinal pigment epithelium could also lead to bisretinoid formation (36). However, *N*-ret-PE formation was recently shown to not always lead to a pathologic outcome but could very well serve as a nonclassical means of chromophore regeneration via production of 9-*cis* or 11-*cis* retinal from *N*-ret-PE photoisomerization (18).

Another physiological role that we hypothesized for *N*-ret-PEs was as a shuttle for *a*RAL delivery to RDHs at the membrane periphery. However, RDHs were found to exhibit activity selectively on free *a*RAL, with little effect on the proportion of *a*RAL sequestered as *N*-ret-PE adducts. Following Rho* hydrolysis, RDH activity led to conversion of nearly all of the released *a*RAL to *a*ROL and thereby near complete attenuation of *N*-ret-PE formation. Such an observation highlights the significance of RDHs in the prevention of the formation and accumulation of *N*-ret-PEs after light exposure. Our method using NaBD₄ therefore allowed us to capture outer-segment membrane-bound RDH activity and can be broadly used to study other redox biochemical reactions involving functional groups, such as aldehydes or ketones, that are susceptible to borohydride/deuteride chemistry. In short, transformation of *a*RAL via the visual cycle is essential for sustainability of our vision. These reactions are carried out in the membrane environment and involve very active aldehydes (*a*RAL and 11*c*RAL) that undergo several enzymatic and nonenzymatic transformations. Using rapid quantitative chemical and analytical methods, we are able now to follow each step quantitatively with preservation of stereospecificity.

Recently, in a collaboration with Chen et al. (37), we described a partially overlapping method that used state-of-the-art native MS to investigate the first steps of the visual cycle and phototransduction in ROS membranes. Native MS enables real-time monitoring of events that cause changes in molecular mass, such as SB formation and retinal release upon hydrolysis of Rho; formation of retinyl-phospholipid adducts; protein-protein, protein-ligand, and protein-lipid interactions; and so on. The method described in the current study requires simpler instrumentation, which is available in many more laboratories and institutions. In addition, the method in this report allows for nonvolatile buffers and salts, including NADPH, which is required for reduction of *a*RAL by RDHs. Furthermore, the isomeric composition of the retinylidene adduct of opsin could be determined, in addition to identifying the specific Lys residue where the SB adduct occurs. Thus, the current method nicely complements the capabilities of the native MS method for investigating the visual cycle.

In this study, we captured and quantified the kinetics and energetics of the critical steps of our vision. We demonstrate that the rate-limiting step is hydrolytic release of *a*RAL from Rho*, and this reaction is dependent on the presence of native

lipids. This methodology should be applicable to investigate the photochemistry occurring with other retinylidene-opsins, including cone opsins, melanopsin, channel Rho, and nonvisual pigments, like RGR or peropsin. Furthermore, our method of capturing bound retinal chromophore could also translate to detecting whether chromophore analogs successfully form an SB linkage in the chromophore binding pocket of various opsin proteins.

Materials and Methods

Proteinase K Digestion of Rho from ROS Membranes. ROS membranes from InVision Bioresources were prepared from fresh bovine retinas using sucrose gradient isolation as described previously and washed with hypotonic buffer (20 mM HEPES; pH 7.4) (38). In the dark room under dim red light, one part ROS membranes (2 mg/mL or 50 μ M Rho) was treated with three parts saturated solution of NaBH₄ granules (~100 μ M) in cold *i*PrOH for immediate reduction of the SB and isolation of Rho by precipitation. Cold ethanol (EtOH) can also be used in place of *i*PrOH. After centrifugation at 20,000 \times *g* to pellet the protein precipitate, the pellet was washed with cold methanol (MeOH) followed by cold water. The protein pellet was then resuspended in proteinase K buffer (4 M urea, 100 mM bis-Tris propane [BTP; pH 7.8], 100 mM CaCl₂). Subsequently, proteinase K (20 mg/mL) was added at ~10 \times the weight of the Rho substrate. Then, the digestion mixture was incubated at room temperature for 24 h, with occasional mixing by brief gentle spinning on a vortex mixer. The digest was passed through a BioPureSPN C18 spin column for desalting. The column was washed with 20% acetonitrile (ACN) in water with 0.1% formic acid (FA), and peptides were eluted using 60% ACN. The eluted N^ε-retinyl-peptide products were separated using a Dionex UHPLC with an XBridge C18 column and a 40-min gradient of 20 to 60% ACN in water with 0.1% FA at a flow rate of 0.3 mL/min. N^ε-retinyl-peptide products were detected by UHPLC absorbance at 330 nm and identified by MS/MS with CID fragmentation using an LTQ XL spectrometer.

Pronase Digestion of Rho from ROS Membranes. In the dark room under dim red light, one part ROS membranes (2 mg/mL or 50 μ M Rho) was treated with three parts saturated solution of NaBH₄ granules in cold *i*PrOH (~100 μ M) for immediate reduction of the SB and isolation of Rho by precipitation. After centrifugation at 20,000 \times *g* to pellet the protein precipitate, the pellet was washed with cold MeOH followed by cold water. The protein pellet was then resuspended in pronase buffer (100 mM BTP [pH 7.8], 100 mM CaCl₂). Subsequently, pronase (20 mg/mL in water) was added at ~10 \times the weight of Rho substrate. Then, the digestion mixture was incubated at 8 to 10 $^{\circ}$ C for 24 h with gentle agitation using a shaker. The digest was passed through a BioPureSPN C18 spin column for desalting. The column was washed with 20% can in water, and digest products were eluted using 60% ACN. The eluted N^ε-retinyl-Lys products were separated using a Dionex UHPLC with an XBridge C18 column and a 30-min gradient of 20 to 50% ACN in water with 0.1% FA at a flow rate of 0.3 mL/min. N^ε-retinyl-Lys products were detected by UHPLC absorbance at 330 nm and identified by MS/MS with CID fragmentation using an LTQ XL spectrometer.

Separation and Identification of *N*-ret-PE Species from ROS. A suspension of ROS membranes (2 mg/mL or 50 μ M Rho) in 20 mM HEPES-buffered solution (at pH 5.6, 7.4, or 8.3) was either supplied with 2.5 molar equivalents of *a*RAL relative to the amount of Rho or illuminated to supply *a*RAL via Rho* hydrolysis. For illumination, a 565-nm fiber light set to an intensity of 625 μ W was applied to ROS for 10 s; three parts of a saturated solution of NaBH₄ (~100 μ M) in cold *i*PrOH was added to one part ROS membranes to solubilize lipids while immediately halting hydrolytic release of *a*RAL from Rho* and the progression of SB formation between *a*RAL and PE. After centrifugation at 20,000 \times *g* to pellet the denatured membrane proteins, the solubilized lipids within the supernatant were analyzed by LC-MS/MS. *N*-retinyl-PEs were separated using an XBridge C18 column and a mobile phase consisting of water with 0.1% FA (solvent A) and MeOH with 0.1% FA (solvent B). The conditions were as follows: a 10-min gradient of 95 to 100% solvent B, followed by 20 min of isocratic 100% solvent B at a flow rate of 0.3 mL/min. The separated *N*-retinyl-PE

products were detected by UHPLC absorbance at 330 nm and identified by MS/MS with CID fragmentation using an LTQ XL spectrometer.

Quantum Chemical and Kinetic Analyses. Details of the quantum chemical and kinetic analyses of the Rho* hydrolysis can be found in the *SI Appendix*. Details of the analyses of N-ret-PE formation and RDH activity can also be found in the *SI Appendix*.

Data, Materials, and Software Availability. All study data are included in the article and/or *SI Appendix*.

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