

# Complex Photochemistry within the Green-Absorbing Channelrhodopsin ReaChR

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**ABSTRACT** Channelrhodopsins (ChRs) are light-activated ion channels widely employed for photostimulation of excitable cells. This study focuses on ReaChR, a chimeric ChR variant with optimal properties for optogenetic applications. We combined electrophysiological recordings with infrared and UV-visible spectroscopic measurements to investigate photocurrents and photochemical properties of ReaChR. Our data imply that ReaChR is green-light activated ( $\lambda_{\text{max}} = 532$  nm) with a non-rhodopsin-like action spectrum peaking at 610 nm for stationary photocurrents. This unusual spectral feature is associated with photoconversion of a previously unknown light-sensitive, blue-shifted photocycle intermediate L ( $\lambda_{\text{max}} = 495$  nm), which is accumulated under continuous illumination. To explain the complex photochemical reactions, we propose a symmetrical two-cycle-model based on the two  $C_{15}=N$  isomers of the retinal cofactor with either *syn*- or *anti*-configuration, each comprising six consecutive states D, K, L, M, N, and O. Ion conduction involves two states per cycle, the late M- ( $M_2$ ) with a deprotonated retinal Schiff base and the consecutive green-absorbing N-state that both equilibrate via reversible reprotonation. In our model, a fraction of the deprotonated M-intermediate of the *anti*-cycle may be photoconverted—as the L-state—back to its inherent dark state, or to its M-state pendant ( $M'$ ) of the *syn*-cycle. The latter reaction pathway requires a  $C_{13}=C_{14}$ ,  $C_{15}=N$  double-isomerization of the retinal chromophore, whereas the intracircular photoconversion of M back to D involves only one  $C_{13}=C_{14}$  double-bond isomerization.

## INTRODUCTION

Channelrhodopsins (ChRs) are photosensitive ion channels, naturally occurring in the eyes of motile green algae (1–4). ChRs unify the ability of light perception with a regulated ion conductance in one small protein, which can be coopted to serve as an ideal actuator for optogenetic applications. Hence, ChRs have become wide-spread tools in the field of neuroscience, where they are used to manipulate the membrane voltage of neurons and to noninvasively evoke or inhibit action potentials with high spatial and temporal precision (4–9). Today, the most widely applied ChR variants are still channelrhodopsin-2 from *Chlamydomonas reinhardtii* (CrChR2 or simply ChR2) and its gain-of-func-

tion mutant H134R (10–12), which are excited most effectively with blue light of 460 nm. However, due to deeper penetration and reduced scattering in tissues, the use of yellow or even red light would be advantageous. This holds particularly true for in vivo application in rodents, where noninvasive delivery of weakly scattered light is a demanding challenge. Over the last few years, a couple of red-shifted ChRs were discovered, modified, and established as optogenetic tools. In 2008, the native green-absorbing channelrhodopsin-1 from *Volvox carteri* (VcChR1) with  $\lambda_{\text{max}} = 535$  nm was used to trigger action potentials in neurons as a response to orange light (13). Subsequent mutational and helix shuffling approaches produced variants with improved expression, larger photocurrents in mammalian cells, and action spectra that were red-shifted compared with ChR2 (14,15). One of the chimeric constructs was the CrChR1/VcChR1 hybrid C1V1<sup>ET</sup> (E122T) enabling fast AP firing even upon stimulation with 630 nm light (14).

In 2013, the red-activatable ChR (ReaChR) was introduced as an improved VcChR1 variant with the N-terminus of CrChR1 (amino acid 1–94), helix 6 from VcChR2, and

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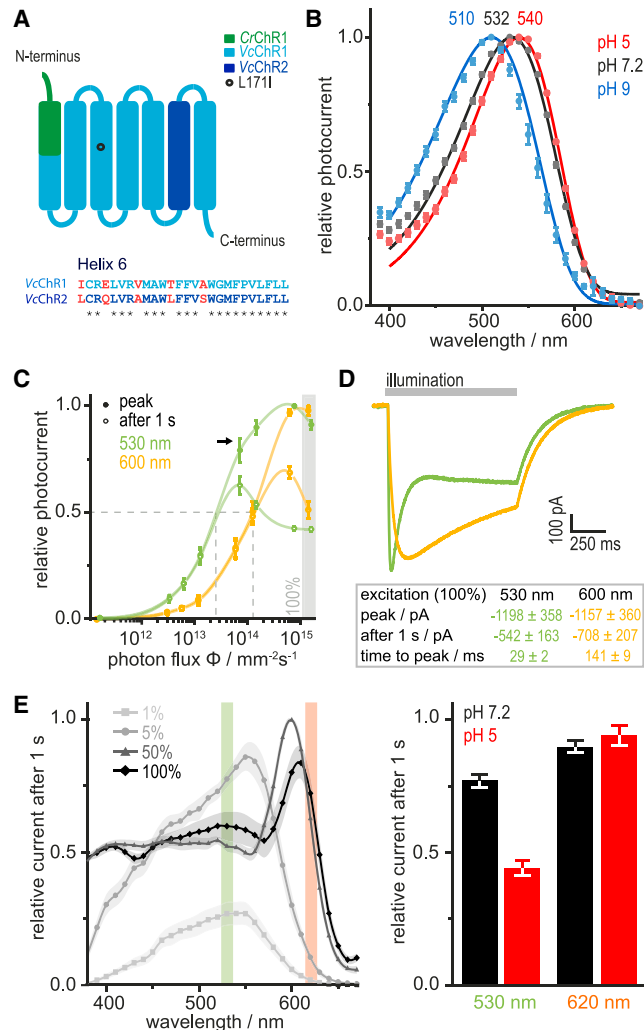
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the L171I mutation in helix 3 (*CrChR2* I131) to reduce desensitization (16) (see Fig. 1 A). In comparison with C1V1<sup>ET</sup>, ReaChR shows improved expression, faster channel closing rates, and larger stationary currents with an action spectrum that substantially deviates from rhodopsin absorption respective shape and side maximum at 630 nm (16). Additionally, the authors demonstrated deep brain tissue activation through the intact skull of awake mice



**FIGURE 1** Photocurrent properties of ReaChR in HEK293 cells. (A) Helix architecture of the chimeric ReaChR consisting of VcChR1, helix 6 (H6) of VcChR2, and the N-terminal sequence of *CrChR1*. H6 of VcChR1 and VcChR2 only differ in five amino acids. (B) Action spectra (10 ms activation) of ReaChR at different external pH (pH<sub>e</sub>) values (mean ± SE, n = 7–12). (C) Light titration curve after excitation with green (530 nm; intensities from  $6.03 \times 10^{-5}$  mW mm<sup>-2</sup> (0.01%) to  $0.61$  mW mm<sup>-2</sup> (100%)) or orange light (600 nm;  $6.65 \times 10^{-5}$  to  $0.47$  mW mm<sup>-2</sup>); photocurrents are normalized to the highest peak response of each cell (mean ± SE, n = 6), lines were added for visual guidance. (D) Comparison of photocurrents evoked with 530 and 600 nm at the same cell (equal photon flux, n = 6). (E) Stationary action spectra at 1, 5, 50 and 100% light intensity (left, mean ± SE, n = 5–6). At pH 5.0 (red bars, right), reduction of the stationary current is more pronounced at 530 nm compared to 620 nm, implying that the photoreactive intermediate is accumulated more at pH 5.0 (mean ± SE, n = 6–8).

without removal or surgical thinning of the cranium. In a subsequent study, ReaChR was employed to excite central nervous system neurons of *Drosophila melanogaster* to study behavior of freely moving flies (17). Remarkably, ReaChR possesses a significant propensity for two-photon absorption most likely due to a high 2P cross section, which was exploited for in-depth excitation of acute brain slices (18) and manipulation of transgenic *D. melanogaster* with near-infrared light (19). Moreover, Sengupta et al. (20) highlighted the enormous potential of ReaChR to restore vision in blind *retinitis pigmentosa* mice. Its superior properties mentioned before render ReaChR a versatile and frequently used red-light-activated optogenetic tool for in vitro and in vivo applications.

Although ReaChR is successfully used in optogenetic experiments, many aspects of its basic photochemistry were not reported in the initial publication and are still unknown, representing the starting point for this investigation. Within the last decade, photochemical reactions as they occur in ChRs have been studied in great detail with different spectroscopic and electrophysiological methods leading to the development of complex photocycle models. Spectroscopic data of ChR2 under single-turnover conditions could be properly described by a single reaction cascade comprising one dark state and four spectrally distinct intermediates (P500, P390, P520, P480) (21–23). However, explanation of photocurrents under extended illumination demands a more complex two-cycle model with at least two closed and two open states with different conductances and ion selectivities (24–27). More recently, spectroscopic data of various recombinant ChRs supported the two-cycle-model (28–33). Bruun et al. (30) reported that both all-*trans* and 13-*cis* retinal isomers of the two dark states may directly photoconvert into each other at low temperature without passing through the photocycle. Here, we present a comprehensive electrical and spectroscopic analysis of ReaChR allowing us to explain the unusual action spectrum by secondary photochemical reaction(s).

## MATERIALS AND METHODS

### Molecular biology

For spectroscopic experiments, DNA encoding ReaChR (GenBank: KF448069.1, 1–345 amino acids) with a C-terminal 1D4 epitope (T-E-T-S-Q-V-A-P-A) was inserted into the pMT4 expression vector (34) between *EcoRI/NotI* (Thermo Fisher Scientific, Waltham, MA), while for patch-clamp recordings ReaChR was C-terminally fused to the mCerulean3 fluorophore (35) and cloned into the pEGFP-N1 vector between *HindIII/XbaI* (Thermo Fisher Scientific). The mutations C168A and C168S were created via site-directed mutagenesis (QuikChange; Agilent Technologies, Santa Clara, CA).

### Patch-clamp recordings

HEK293 cells were seeded onto coverslips ( $1.5 \times 10^5$  cells per 40 mm dish; TPP Techno Plastic Products, Trasadingen, Switzerland) and supplemented

with 1  $\mu\text{M}$  all-*trans* retinal (Sigma-Aldrich, Munich, Germany). 48–56 h before recordings cells were transiently transfected with the respective plasmid DNA using FuGENE HD transfection reagent (Promega, Madison, WI). Measurements were performed in whole-cell mode at a holding potential of  $-60$  mV. Signals were amplified with an AxoPatch 200B and digitized using a DigiData 1400 (Molecular Devices, Sunnyvale, CA). Light was delivered by a Polychrome V (TILL Photonics, Planegg, Germany) or a Xenon lamp and regulated with a programmable shutter, while photon densities were adjusted using a neutral density filter wheel and/or neutral density filters. Data was acquired with the Clampex software and evaluated using Clampfit 10.4 (Molecular Devices). If not stated otherwise, pH was intra- and extracellularly adjusted to 7.2 (internal: 110 mM NaCl, 2 mM  $\text{MgCl}_2$ , 2 mM  $\text{CaCl}_2$ , 1 mM KCl, 1 mM CsCl, 10 mM HEPES, 10 mM EGTA; external: 140 mM NaCl, 2 mM  $\text{MgCl}_2$ , 2 mM  $\text{CaCl}_2$ , 1 mM KCl, 1 mM CsCl, 10 mM HEPES).

## Expression and purification

HEK293T cells were maintained in Dulbecco's Modified Eagle Medium (DMEM; Biochrom, Berlin, Germany) supplemented with 10% fetal bovine serum (DMEM-S; Biochrom) at 37°C and 5%  $\text{CO}_2$ . For transfection, cells were grown on 100 mm culture dishes (Corning; Sigma-Aldrich) to 70–80% confluency and treated with TurboFect (Thermo Fisher Scientific) according to the manufacturer's protocol. Cells were harvested 36–48 h posttransfection in the presence of protease inhibitor (complete EDTA-free; Roche, Mannheim, Germany). Protein purification was achieved under safelight ( $>600$  nm) at 4°C. During whole-cell treatment with 1% (w/v) *n*-dodecyl  $\beta$ -D-maltopyranoside (DDM; Glycon, Luckenwalde, Germany) in Dulbecco's phosphate buffered saline (DPBS), pH 7.4 (Biochrom), and 10  $\mu\text{M}$  all-*trans* retinal (Sigma-Aldrich) for 4–6 h, the photoreceptors were solubilized and subsequently purified by 1D4 immunoaffinity chromatography. For this purpose, 1D4-antibody-resin was prepared according to Oprian et al. (36). Immobilized protein was eluted with DPBS buffer containing 400  $\mu\text{M}$  1D4-peptide (T-E-T-S-Q-V-A-P-A, GenScript, NY) and 0.03% DDM, whereas for pH-dependent measurements concentrated ReaChR samples were diluted in 10 mM citric acid pH 5.0, 100 mM NaCl, 0.03% DDM, and 10 mM TRIS pH 9.0, 100 mM NaCl, 0.03% DDM, respectively. For retinal extraction experiments, protein was expressed in *Pichia pastoris* and purified in buffer containing 10 mM TRIS (pH 7.4), 100 mM NaCl, and 0.03% DDM as described in the literature (23,37,38).

## Ultraviolet-visible absorption spectroscopy

Absorption spectra and slow kinetic traces were recorded by a Cary 50 Bio spectrophotometer (Varian, Palo Alto, CA) at 22°C. Initial dark state (IDA) spectra were recorded with protein purified under safelight. ReaChR C168S was illuminated with a 530 nm light of  $1 \times 10^{20}$  photons  $\text{m}^{-2} \text{s}^{-1}$  (Luxeon LED; Phillips, Amsterdam, the Netherlands) while photon flux was decreased to  $5.6 \times 10^{19}$  photons  $\text{m}^{-2} \text{s}^{-1}$  for dual-color experiments. If necessary, brighter blue-light pulses ( $1 \times 10^{20}$  photons  $\text{m}^{-2} \text{s}^{-1}$ ) were applied using an LED (400 nm; Thorlabs, Newton, NJ). Flash-photolysis experiments and data processing were performed as described in Wietek et al. (39). For this purpose, samples with an optical density of  $A_{280} = 1.0$  were excited with green flashes (530 nm,  $\sim 10$  ns, 5 mJ/flash). Global analysis is based on a sequential model with four or five spectral components. Absolute spectra of ReaChR wild-type at pH 5.0 were calculated based on the IDA spectrum (see Fig. 2 A) and the corresponding non-normalized evolutionary associated difference spectra (EADS) (data not shown; compare to Fig. S1). We assumed 10% photoreceptor activation by the laser flash. Cryostatic ultraviolet-visible (UV-vis) measurements were performed using a sample chamber placed inside a cryostat (DN; Oxford Instruments, Abingdon, UK) according to the following protocol: After 2 min of illumination ( $\sim 530$  nm) and succeeding 30 min of dark adap-

tation at minimum 20°C, sample was cooled down to the desired temperature. Before the next measuring cycle, the sample was heated up again to a minimum of 20°C. Recorded spectra were scaled by normalizing to the dark state chromophore absorption band.

## Fourier-transform infrared spectroscopy

Rehydrated samples of ReaChR at pH 7.4 in DPBS were placed between two  $\text{BaF}_2$  windows and analyzed by an ifs66v/s Fourier-transform infrared (FTIR) spectrometer (Bruker Optics, Karlsruhe, Germany) with a liquid nitrogen-cooled mercury cadmium telluride detector (Kolmar Technologies, Newburyport, MA) and an  $1850 \text{ cm}^{-1}$  optical cutoff filter. Cryostatic FTIR measurements were conducted similar to cryo-UV-vis measurements. Corresponding spectra were recorded with a 200 kHz sampling rate and with a spectral resolution of  $2 \text{ cm}^{-1}$ . Difference spectra were corrected for baseline drifts using a spline algorithm and the baseline-correction mode implemented in the OPUS 6.5 software package (Bruker Optics). The photostationary state was defined as the steady state of the kinetics of the strong absorption band at  $1660 \text{ cm}^{-1}$ . Spectra were scaled by normalizing to the amide II band in the absolute spectra at  $\sim 1550 \text{ cm}^{-1}$ .

## Retinal extraction and separation by high-pressure liquid chromatography

Purified protein was illuminated for 10 min with green light ( $\sim 530$  nm LED) and kept in the dark for 10 min. Chromophore extraction was achieved by addition of ice-cold ethanol (delay time  $\leq 2$  s). After 2 min, the extracted retinal was dissolved in heptane containing 7% diethyl ether (w/w). After 2 min, phase separation was achieved by centrifugation (1 min, 2000 rpm). All working steps, except sample illumination, were conducted under dim red light and at room temperature. Separation of the retinal isomers in the heptane/diethyl ether mixture was performed using a LC-20AD high-pressure liquid chromatography (HPLC) device (Shimadzu, Kyoto, Japan) equipped with a ReproSil 70 Si, 5  $\mu\text{m}$  column (Dr. Maisch, Ammerbuch-Entringen, Germany). Isomer ratio was determined by peak integration. The chromatogram was corrected for baseline-drifts.

## RESULTS

### Photocurrent properties in HEK293 cells

In 2013, Lin et al. (16) reported unusual, non-rhodopsin-like action spectra for ReaChR with maximal activity at 590 nm excitation for the peak and even 630 nm for the stationary photocurrent. However, subsequent studies on ReaChR revealed that this ChR is most sensitive to green light (40,41). Based on this discrepancy, we reinvestigated ReaChR by electrophysiological recordings in HEK293 cells. First, flash-induced photocurrents (10 ms excitation) were recorded at different wavelengths and three extracellular pH-values ( $\text{pH}_e = 5.0, 7.2, \text{ and } 9.0$ ) revealing maximal sensitivity to 532 nm light ( $\text{pH}_e 7.2$ ), and small shifts after  $\text{pH}_e$  change. These action spectra were rhodopsin-shaped without any particular unusual characteristics (Fig. 1 B). Upon 1 s illumination, the shape of the photocurrent depends on the intensity of the actinic light as known from CrChR1 (25), CrChR2 (4), and relatives. Titration of the 530 nm activation light (Fig. 1 C, green) revealed that peak photocurrents were half-maximal at a photon irradiance of  $2.68 \times 10^{13} \text{ mm}^{-2} \text{ s}^{-1}$  ( $0.010 \text{ mW mm}^{-2}$ ), while

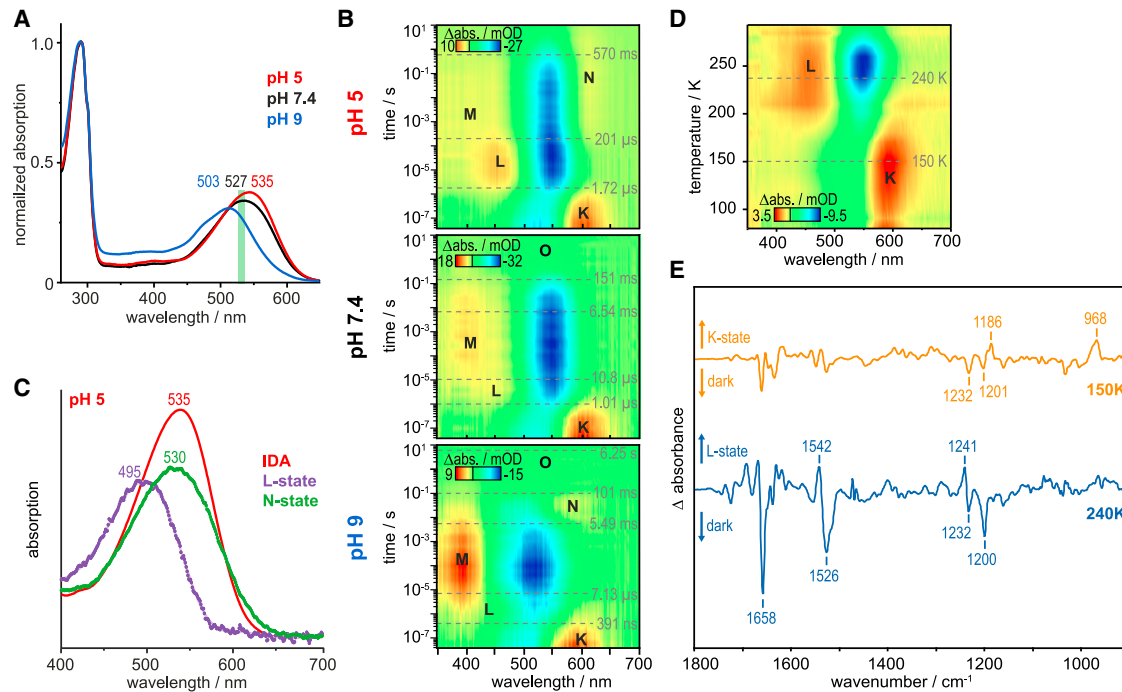


FIGURE 2 Spectral properties of recombinant ReaChR. (A) IDA spectra of purified ReaChR at pH 5.0, 7.4, and 9.0 (normalized to 280 nm). (B) Transient absorption changes induced by green laser flashes (10 ns, 530 nm, 5 mJ) were measured in saline detergent (0.03% DDM) solution at pH 5.0 (*top*, 100 mM NaCl, 10 mM citric acid), 7.4 (*middle*, DPBS), and 9.0 (*bottom*, 100 mM NaCl, 10 mM TRIS). Half-life times ( $t_{1/2}$ ) are highlighted in the contour plots. Five photocycle-intermediates K, L, M, N (not seen at pH 7.4), and O were identified. L and M are enriched at pH 5 and 9, respectively. Acidic pH decelerates photocycle kinetics. (C) Calculated absolute spectra of the L- and N-states of ReaChR at pH 5.0. Dots resemble raw data points and solid lines were added for visual guidance. (D) Cryostatic UV-vis measurements (*light-dark*, pH 7.4) of ReaChR under continuous illumination reveal maximal accumulation of the K- and L-states at 150 and 240 K, respectively. (E) FTIR difference (*light-dark*) of the photostationary states under continuous illumination at 150 (*orange*) and 240 K (*blue*) at pH 7.4. Imposing HOOP vibration at 968(+)  $\text{cm}^{-1}$  points toward a distorted 13-*cis* chromophore in the K-state, while cofactor strain is almost completely relieved in the upcoming L-state.

stationary currents saturated at much lower light intensities and were half-maximal already at  $0.005 \text{ mW mm}^{-2}$ . Interestingly, the curve for the peak current decreased (i.e., kinked) at light intensities already a decade away from saturation (*black arrow*). Second, the stationary currents decreased again at higher intensities, which was much more pronounced in ReaChR compared with CrChR2 (10). In comparison, the peak current sensitivity at 600 nm (Fig. 1 C, *orange*) is shifted to four-times higher intensities ( $I_{50} = 0.042 \text{ mW mm}^{-2}$ ;  $1.27 \times 10^{14} \text{ photons mm}^{-2} \text{ s}^{-1}$ ), in agreement with a reduced absorption coefficient at 600 nm as seen from the flash-induced action spectrum for  $\text{pH}_e$  7.2 (Fig. 1 B). However, the reduced slope of the stimulus response curve at intensities 10-fold below saturation is not visible for 600 nm activation (Fig. 1 C). Fig. 1 D shows a direct comparison of photocurrents recorded from the same cell upon activation with 530 and 600 nm light at equal photon irradiance (100%).

In accordance with the unusual shape of the light titration curves at high photon densities, action spectra of the steady-state currents were strongly intensity dependent. At low light intensities of 1 and 5% the spectra peaked at 550 nm, whereas maxima were shifted to 600 nm for 50%

and 610 nm for 100% activation light, revealing a complex secondary photochemistry at high photon irradiance (Fig. 1 E).

### Spectral properties of recombinant ReaChR

To determine the absorption properties of the dark state, recombinant ReaChR was purified under safelight ( $>600 \text{ nm}$ ) from HEK293T and absorption spectra were recorded (Fig. 2 A). The IDA showed maximal absorption at 535 nm at pH 5.0, 527 nm at pH 7.4, and 503 nm at pH 9.0, the latter with a shoulder at  $\sim 450 \text{ nm}$  not seen in the action spectra. The extinction coefficient of chromophore absorption increased gradually upon acidification. Next, recombinant ReaChR was exposed to brief green (530 nm) laser pulses of  $\sim 10 \text{ ns}$  and absorption changes between 40 ns and 10 s after flash were monitored (Fig. 2 B). Normalized EADS (Fig. S1) and their half-life times ( $t_{1/2}$ ) were extracted using a global fit routine. Dissected intermediates were designated according to the bacteriorhodopsin (BR) nomenclature. The dark state (D) is converted into a distinctive red-shifted K-state (seen at  $\sim 600 \text{ nm}$ ) that rises faster than our time-resolution and decays with  $t_{1/2} = 1.01$  or



1.72  $\mu\text{s}$  at pH 7.4 and 5.0, respectively. At pH 9.0, K-state decay is 2.5 times faster ( $t_{1/2} = 391$  ns) compared to pH 7.4. At pH 5.0 this early photoproduct evolves into a blue-shifted L-state with maximal absorption at 460 nm and shows roughly 20 times prolonged lifetime compared to neutral pH. Such an L-state was not reported for ChR2 (22). The subsequent well-known M-intermediate with deprotonated chromophore prevails over several orders of time and exhibits a broad unstructured absorption profile with small amplitude at pH 5.0 and 7.4 but high population at pH 9.0. From previous studies on ChR2, it is known that, as for BR, M is actually composed of an early  $M_1$  and a late  $M_2$  state (42–44). The decay of M is biphasic, and accelerated at pH 9.0 (Fig. 2 B, bottom). At pH 5.0 and 9.0, but not at pH 7.4, a late photointermediate appears with positive absorption at  $\sim 600$  nm on a millisecond timescale, referred to as N-state. At pH 9.0, the N-intermediate decays with  $t_{1/2} = 101$  ms to a state with a spectral signature hardly discriminable from the dark state spectrum. This intermediate, designated as O-state, finally recovers to D with  $t_{1/2} = 6.25$  s at pH 9.0 (Fig. 2 B, bottom). In neutral and acidic conditions these last transitions are either poorly resolved or/and highly decelerated, thus not completed during the observation window of 10 s (Fig. 2 B, middle and top). To disentangle photocycle intermediates, absolute spectra of the L- and N-state at pH 5.0 were determined by adding the appropriate nonnormalized EADS (data not shown; and compare to Fig. S1, top) to the IDA spectrum (Fig. 2 A). The calculated L-state spectrum shows maximal absorption at 495 nm while N-state peaks at 530 nm similar to the dark state but with a broader absorption profile toward longer wavelengths (Fig. 2 C).

To investigate structural features associated with the newly identified L-state, ReaChR was analyzed at cryogenic conditions by UV-vis and FTIR spectroscopy. The temperature-versus-wavelength plot shows that up to 200 K, the K-state is enriched, whereas between 200 K and room temperature, L is the dominant photoproduct (Fig. 2 D). FTIR spectra were recorded at 150 and 240 K (Fig. 2 E). The K-minus-D difference spectrum exhibits relatively small changes with the exception of a typical band pattern within the retinal fingerprint region with two negative bands at 1232 and 1201  $\text{cm}^{-1}$  and a positive band at 1186  $\text{cm}^{-1}$ , indicative of a net isomerization of the retinal from all-*trans* to 13-*cis* (45). A pronounced hydrogen out-of-plane (HOOP) vibration at 968(+)  $\text{cm}^{-1}$  indicates a distorted 13-*cis* chromophore structure (46,47) in the early state, which relaxes in the upcoming L state. During transition from K- to L-, the amide I region gains intensity with a strong negative band at 1658  $\text{cm}^{-1}$ , suggesting major backbone rearrangements probably associated to channel pre-gating. The L spectrum possesses a unique band at 1241(+)  $\text{cm}^{-1}$ , which is caused either by stretching vibrations of  $C_{12}$ – $C_{13}$  of the retinal (48) or N–H in-plane bending of the retinal Schiff base (RSB) (49). The blue-shifted

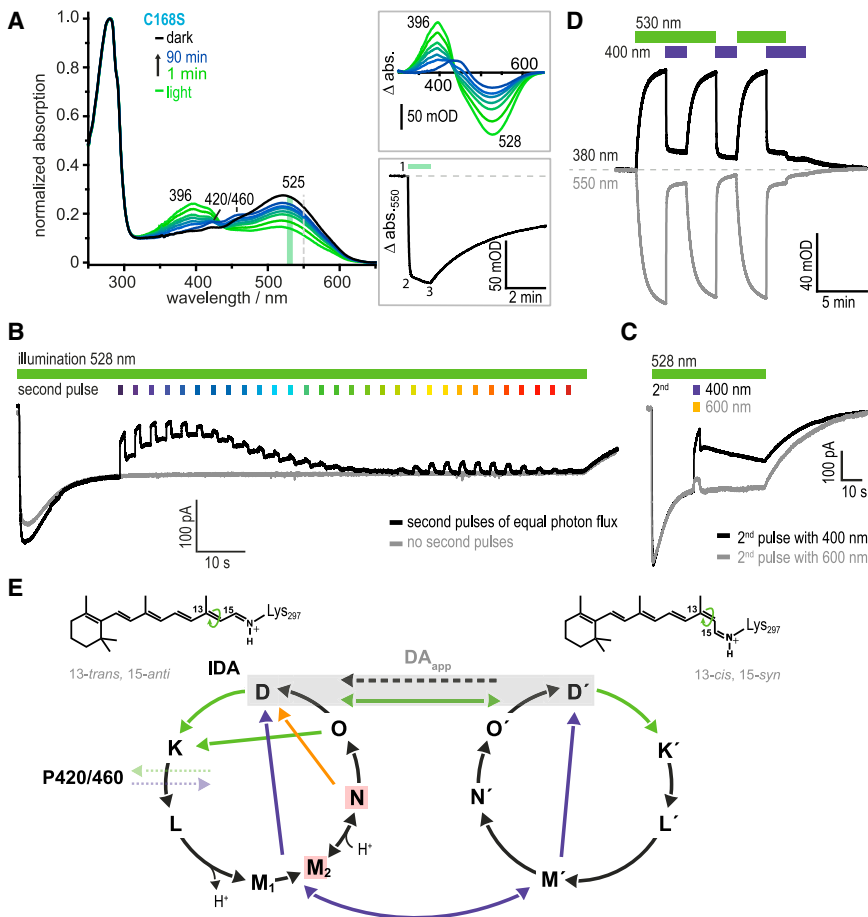
L-state absorption maximum (Fig. 2 C) is reflected in the upshift of the C=C stretching mode from 1526(–) to 1542(+)  $\text{cm}^{-1}$  in the FTIR spectrum of the D to L transition (50,51). The presence of a strong positive ethylenic band hints at a protonated RSB ( $\text{RSBH}^+$ ) in L, as only small positive ethylenic bands were observed in states with a deprotonated RSB in channelrhodopsin-1 from *Chlamydomonas augustae* (CaChR1) (52,53) and CrChR2 (54).

### Photoconversion of intermediate states

To investigate the conducting state(s) and subsequent late photointermediates, we generated slow-cycling mutants of ReaChR by modification of the DC-pair residues C168 and D196 (CrChR2 C128 and D156), which play a crucial role for the channel gating kinetics in CrChR2 (55,56). We introduced C168A and C168S and confirmed the slowed photocurrent decline by electrical analysis, as illustrated in Fig. S2. Wild-type photocurrent decayed with  $\tau_{\text{off}} = 181 \pm 22$  ms (monoexponential fit), whereas the closing kinetics for C168A and C168S were two orders of magnitude slower with  $\tau_{\text{off}} = 19.7$  and 28.1 s, respectively (Fig. S2). On average, the C168S photocurrent amplitudes were  $<20\%$  of the wild-type and C168A currents at similar conditions, thus we used the C168A mutant for further electrical analysis.

Next, we recorded absorption spectra of purified ReaChR-C168S in the dark and after illumination with 530 nm light (60 s). The IDA spectrum shows maximal absorption at 525 nm, while upon illumination it converts to the UV-absorbing M-state (396 nm) (Fig. 3 A and top inset), which decays biexponentially with time constants in the early to late minute range. Additionally, slowly appearing photoproducts are seen at 420 and 460 nm (P420/460) (Fig. 3 A and top inset), which do not thermally reconvert to the IDA during the entire observation window of 90 min but are fully photoconvertible by UV-light (data not shown). These P420/460 side products are accumulated solely under prolonged illumination (Fig. 3 A, bottom inset, 2  $\rightarrow$  3) and were assigned to inactive side products of the photocycle in analogy to our conclusions drawn for ChR2-C128T (54). Strikingly, even extended green illumination (530 nm light for 60 s) bleaches the absorption at  $\sim 520$  nm to maximally  $\sim 55\%$ , suggesting that either a fraction of the protein is not photoactive, or more likely, a green-absorbing photocycle intermediate as the N-state is still prevailing in continuous light.

With the slow-cycling mutants at hand we electrically analyzed the photoconversion of the conducting state(s) as previously reported for slow-cycling ChR2- and C1V1-variants (15). Continuous illumination of ReaChR-C168A with 528 nm light accumulated the open state(s) after some inactivation and established a steady-state current level. Subsequently, more intense light pulses of different wavelengths but equal photon flux were applied and showed that the steady-state photocurrent can be reduced by UV and orange



**FIGURE 3** Photoconversion of intermediate states and photocycle model. (A) Absorption spectra of ReaChR-C168S recorded for the IDA and after excitation (60 s, 530 nm). Difference spectra were calculated (*light-dark*, *top inset*). Upon prolonged illumination, M-state ( $\lambda_{\text{max}} = 396$  nm) is accumulated predominantly, which completely decays within 90 min, but substantial absorption at  $\sim 520$  nm ( $\sim 55\%$ ) indicates the presence of another green-absorbing intermediate (N-state). Chromophore bleaching monitored at 550 nm involves two processes (*bottom inset*), whereby the faster component is attributed to M-state formation (1 $\rightarrow$ 2) and the slower one to formation of minor blue-shifted species seen at 420 and 460 nm (P420/460) (2 $\rightarrow$ 3). (B) Photocurrents of HEK cells expressing ReaChR-C168A under continuous illumination with green light and additional stronger light pulses (380–670 nm) inducing current inactivation. The 400 nm light effectively inactivates currents with slow recovery, whereas inactivation at 600 nm is smaller and instantaneously reversible with green light. (C) These current traces show more clearly that application of a single blue-light pulse on top of green background light causes current inactivation and slow recovery, whereas a 600 nm light pulse causes smaller inactivation but fast recovery. (D) Recombinant ReaChR-C168S illuminated with green (530 nm, weak) and/or blue (400 nm, strong) light, and rise and decay of the M- (380 nm) and dark states (550 nm) were monitored by single kinetic traces. Green-light illumination induces formation of M and dark state (D) bleaching in parallel, while blue light reverses this process, indicative for an M $\rightarrow$ D shortcut. (E) Proposed symmetrical two-cycle model of ReaChR. One cycle is based on the 13-*trans*, 15-*anti* retinal configuration (*anti*-cycle) while the other starts from 13-*cis*, 15-*syn* (*syn*-cycle); the equilibrium of both forms the apparent dark state (DA<sub>app</sub>, *gray background*), while the IDA is supposed to consist of 100% all-*trans* retinal. Upon illumination with green light, D converts to a red-shifted K-like intermediate, succeeded by the blue-shifted states L and M. The deprotonated M-state decays biexponentially (M<sub>1</sub> and M<sub>2</sub>) and is reprotonated during N-state formation. N decays to an intermediate that is spectrally not distinguishable from the dark state (O-state). (*Colored arrows*) Proposed photoinduced pathways between the intermediates and both cycles; (*black arrows*) thermal conversions. The conducting states are assigned to the M<sub>2</sub>- and N-equilibrium (*red box*). The *syn*-cycle consists of the same spectral states marked by a hyphen and undergoes the same protonations; corresponding shortcuts and the M-substates are omitted for clarity. Upon prolonged illumination of slow-cycling mutants, side-products (P420/460) are accumulated, which could be photoconverted by UV-light (data not shown).

light. While the inactivation with orange light was smaller and immediately reversed by the green basal illumination, UV or blue-light pulses inactivated the protein to  $<50\%$  of the steady-state current level (Fig. 3 B) and only a small fraction recovered under green background light within 1 s. The full stationary current level was restored only after a longer time in the minute range under green-light illumination (Fig. 3 C), suggesting that ReaChR was converted into a low- or nonconducting photocycle product by UV and blue light. To elucidate underlying photochemistry under continuous illumination, recombinant ReaChR-C168S was exposed to a series of parallel and sequential light pulses of green (530 nm) and UV (400 nm) light (Fig. 3 D), while M-state rise and decay was monitored by absorbance changes at 380 nm and dark state absorption was monitored at 550 nm. Under permanent green illumination, M was accumulated but was almost completely con-

verted to D when UV-light was interspersed, although the full recovery after cessation of light was slow and occurred on timescales of minutes.

## DISCUSSION

Lin et al. (16) described the VcChR1 variant ReaChR that exhibits large photocurrents and shows steady-state current action spectra that substantially differ from rhodopsin absorption spectra. Such spectra are either caused by multiple overlaying photoreceptors—which can be excluded here—or an extended secondary photochemistry. This is the reason why *in vivo* action spectra that led to the original identification of rhodopsin as the phototaxis photoreceptor in *Chlamydomonas* were recorded at low light intensities (threshold spectra) (57) or as flash spectra for the correlation between photocurrents and photoreceptor species (58).

We confirmed the non-Gaussian-shaped action spectra for ReaChR, but this only holds true for extended illumination periods of a few seconds and high intensities. Low intensity or flash action spectra resemble the absorption spectra of recombinant protein (compare to Figs. 1 B and 2 A) and are similar to the parent VcChR1 (13) and the related C1V1-hybrid (15,41). The reduced stationary photocurrents under continuous green light can be explained in three ways, namely by photochemical conversion of one or more photocycle intermediates back to the dark state(s), photochemical conversion between intermediates of the *syn*- and *anti*-cycle, and conversion of one or more photocycle intermediates into an inactive or less active side product. This interpretation raises the question of the nature of the activated photointermediate and of the secondary photoproducts that are formed more efficiently compared to other ChRs.

Photochemical back reactions between photocycle intermediates and dark states were previously proposed for wild-type ChR2 (22,23) and its step-function opsin (SFO) variants (54–56), although the practical relevance of this secondary photoreaction is low for wild-type ChR2. For ChR2-C128T, side reactions into inactive, slowly recovering species that are not part of the bona fide photocycle were also previously reported (54,59). Within our study, we identified six states for ReaChR designated as the following: D- (dark), K-, L-, M- ( $M_1$  and  $M_2$ ), N-, and O-states that are expected to be consecutive in analogy to BR, with D- and O- having similar spectral properties (Fig. 2 B). Remarkably, a state corresponding to the L-state has not been identified for CrChR2 and CaChR1 (22,23,60), but was just recently reported for channelrhodopsin-2 from *Platymonas subcordiformis* (PsChR2) (32,33) and the anion-conducting channelrhodopsin-1 from *Proteomonas sulcata* (PsACR1) ( $P_{460}$ ) (39) and *Guillardia theta* (GtACR1) (61). The hypsochromic shift from the K-intermediate absorbing at 600 nm (difference spectrum, Fig. 2 B) to the L-state at 495 nm observed at pH 5.0 (absolute spectrum, Fig. 2 C) is not a result of the protonation state of the RSB but rather from an altered distance between RSBH<sup>+</sup> and counterion-complex upon chromophore relaxation. While the 13-*cis* retinal molecule is twisted and distorted in the early K-state, the cofactor strain is almost relieved in the upcoming L-state, as deduced from the lack of the HOOP band in the L-state IR-spectrum (Fig. 2 E). Most likely, chromophore relaxation is achieved by rotation of the C<sub>13</sub>=C<sub>14</sub> and C<sub>15</sub>=N bonds, as it occurs in BR (62,63). As a consequence, the RSBH<sup>+</sup> is shifted closer to the water-counterion complex compared to the configurations both in K and D, thus stabilizing the electronic ground state of L and leading to a blue-shifted absorption spectrum (62).

The proposition of the L-state as the main photoreactive intermediate under continuous illumination is further supported by two findings: First, the L-state is strongly accumulated under continuous illumination compared to

single-turnover conditions (compare to Fig. 2, B, middle, and D). Second, photochemical inactivation of the photocurrents under continuous green light at pH 5.0 is even more efficient compared to pH 7.2 (Fig. 1 E, bar diagram), in line with an enrichment of L at low pH as seen for the recombinant protein in single flash experiments (Fig. 2 B, top). Because calculated absolute spectra of N- and dark states are almost identical (Fig. 2 C), L-state conversion by blue/green light should be efficient whereas N to D conversion appears less efficient because green light activates N and D simultaneously in the opposite direction.

It might be worthwhile to recapitulate that ReaChR-C168S accumulates mostly M (probably  $M_2$ ) during extended illumination, similar to what was reported for the SFOs ChR2-C128S or D156A (14,55), but substantial absorption at 520 nm remains (Fig. 3 A); this supports the idea of an equilibrium between M and N, as proposed for ChR2 (P390 and P520) (22,23). The observed small and fully reversible inactivation with orange light (600 nm) (Fig. 3 C) is explained by the red tail of the N-state (Fig. 2 C). The last question remaining is why 400 nm bleaching is not immediately reversed by green background light. Similar observations were previously reported for the SFO C128T (59) and the stable step-function opsin ChR2-C128S/D156A (14). The slow recovery of the steady state is explained by the two-cycle model based on the two C<sub>15</sub>=N retinal isomers. Assuming a symmetrical model with similar spectral intermediates in each cycle, photoactivation of the dark state D with 13-*trans*, 15-*anti* conformation produces K, L, M, and N intermediates with 13-*cis*, 15-*anti*. Thus, we consider this cycle as *anti*-cycle because the C<sub>15</sub>=N bond should be *anti* throughout the whole cycle. Accordingly, the cycle starting from a dark state with 13-*cis*, 15-*syn* (D') is designated *syn*-cycle. Hence, either a photochemical transition from an M-state of the *anti*-cycle to an M-state of the *syn*-cycle occurs or M is converted into an inactive UV-absorbing state, possibly with another *cis*-isomer than 13-*cis*, e.g., 9- and 11-*cis*, which are formed in the ChR2-C128T mutant upon prolonged illumination (28).

Recently, we have reported for the bimodal switchable Histidine Kinase Rhodopsin-1 from *C. reinhardtii* that UVA-light may not only cause a sole isomerization of the RSB around C<sub>13</sub>=C<sub>14</sub> bond, it can also promote a simultaneous isomerization of both the C<sub>13</sub>=C<sub>14</sub> and the C<sub>15</sub>=N double bond, preferentially if the RSB chromophore is deprotonated (64). Such a double isomerization would convert the 13-*cis*, 15-*anti* M-state into a 13-*trans*, 15-*syn* M-state that thermally converts to 13-*cis*, 15-*syn*, the expected dark state of the *syn*-cycle. Because the open state(s) of the *syn*-cycle is assumed to be less conductive with altered ion selectivity in CrChR2 (27), and both the absorption cross section and the quantum efficiency for a protonated 13-*cis*, 15-*syn* chromophore is anticipated to be reduced compared to a 13-*trans*, 15-*anti* RSBH<sup>+</sup>, the slow recovery

of the steady-state photocurrent after a UV-light pulse could be explained by the transition from the *anti*- to the *syn*-cycle. However, recombinant ReaChR-C168S shows a long-lasting chromophore bleaching and conversion of the dark state(s) into P420/460, which are not representatives of the *syn*- or *anti*-cycle. P420/460 recover to the dark state only upon UV-illumination (data not shown). Thus, we consider them as photochemical side products as previously proposed for ChR2-C128T (54).

### ReaChR photocycle model

Based on our study, we conclude that the non-rhodopsin action spectrum for steady-state currents of ReaChR is caused by a complex orchestration of secondary photochemical reactions starting from the newly identified L-state and its subsequent M-states. We propose a symmetrical two-cycle model as recently discussed for CrChR2 (30) with two dark states (D and D') establishing the apparent dark state (DA<sub>app</sub>), and two open states (N and N') with different conductivities and selectivities, which are in a fast equilibrium with the preceding M<sub>2</sub>-state (Fig. 3 E), to describe photochemical events as they occur under both single-turnover conditions and continuous illumination. Under continuous green illumination, the initial dark state (IDA) consisting of 100% all-*trans* retinal is converted to an equilibrium between the two cycles (i.e., conducting states), which is shifted when a second blue- or green-light pulse is applied. Remarkably, blue light drives both intra- and intercircular shortcuts. While the intracircular shortcut is based on C<sub>13</sub>=C<sub>14</sub> isomerization, the latter one is assumed to be based on a C<sub>13</sub>=C<sub>14</sub>, C<sub>15</sub>=N double-isomerization of the retinal chromophore and a conversion between M to M' or vice versa. However, strictly speaking, a single C<sub>15</sub>=N *anti/syn* or *syn/anti* isomerization cannot be excluded. In such hypothetical cases, isomerization would produce 13-*cis*, 15-*syn* or 13-*trans*, 15-*anti* retinal Schiff base species that both are expected to be protonated under formation of the O-states. Based on our experiments, it is unclear whether the cycle crossover is purely light-induced or occurs partially thermally. Thermal branching was reported to happen highly efficiently in other microbial rhodopsins such as BR and *Anabaena* sensory rhodopsin (65), thus could be also possible for ChRs and should be part of further investigations. More evidence for a two-cycle model was given by Szundi et al. (32,33), who reported that the cation-conducting PsChR2 runs through two cycles in parallel, while 30% of excited protein follow cycle 1 and 70% cycle 2. Retinal extraction and HPLC experiments of ReaChR wild-type in the DA<sub>app</sub>-state revealed a proportion of 78:22 of all-*trans*:13-*cis* retinal (Fig. S3). This ratio is in line with the composition in CrChR2 (28,45,54) and would allow the assignment of a 15-*anti* cycle with the C<sub>15</sub>=N bond being always in *anti*-configuration (previously named “*trans*-cycle”, according to the 13-*trans* conformation of

the dark state) and a *syn*-cycle with the C<sub>15</sub>=N conformation always in *syn*-conformation. However, an exact assignment would require Raman or NMR experiments, respectively. A better understanding of side reactions into an inactive and only slowly recovering state also requires more extensive studies, which are not trivial due to the heterogeneity of the photoproduct mixture. In general, a deeper knowledge of the photochemistry of ReaChR—a frequently used optogenetic tool—allows for a purposeful implementation within neurophysiological experiments, especially in cases where complex illumination protocols are requested.

### SUPPORTING MATERIAL

Three figures are available at [http://www.biophysj.org/biophysj/supplemental/S0006-3495\(17\)30155-8](http://www.biophysj.org/biophysj/supplemental/S0006-3495(17)30155-8).

### AUTHOR CONTRIBUTIONS

B.S.K. expressed and purified protein, and performed and analyzed UV-vis spectroscopic measurements; C.G. conducted and evaluated electrophysiological experiments; J.C.D.K. performed FTIR and cryo-UV-vis measurements as well as retinal extraction and HPLC experiments; F.S. and T.P.S. provided material and experimental expertise; P.H. and F.J.B. designed and interpreted experiments; and B.S.K. and P.H. wrote the article with further contributions from all authors. All authors reviewed the results and approved the final version of the article.

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**Supplemental Information**

**Complex Photochemistry within the Green-Absorbing Channelrhodopsin ReaChR**

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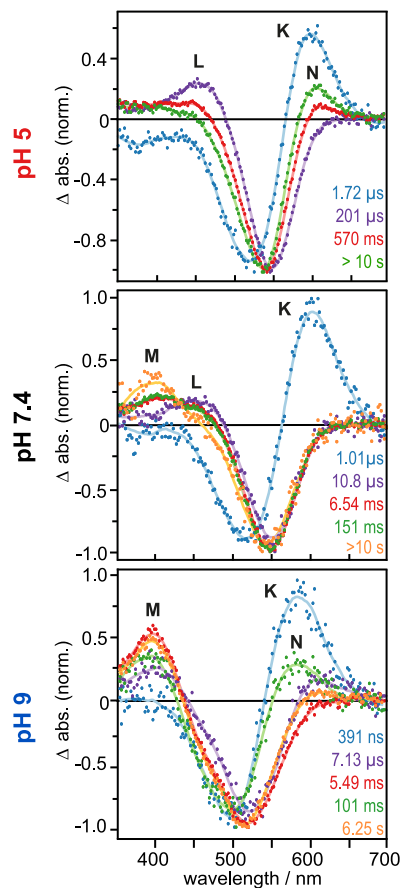


FIGURE S1 Normalized Evolutionary Associated Difference Spectra (EADS) of ReaChR wild-type at pH 5 (*top*), 7.4 (*middle*) and 9 (*bottom*). EADS and their half-life times ( $t_{1/2}$ ) are derived from a global fit routine. For illustration purposes, solid lines were added to the EADS. Photocycle intermediates are indicated by capital letters.

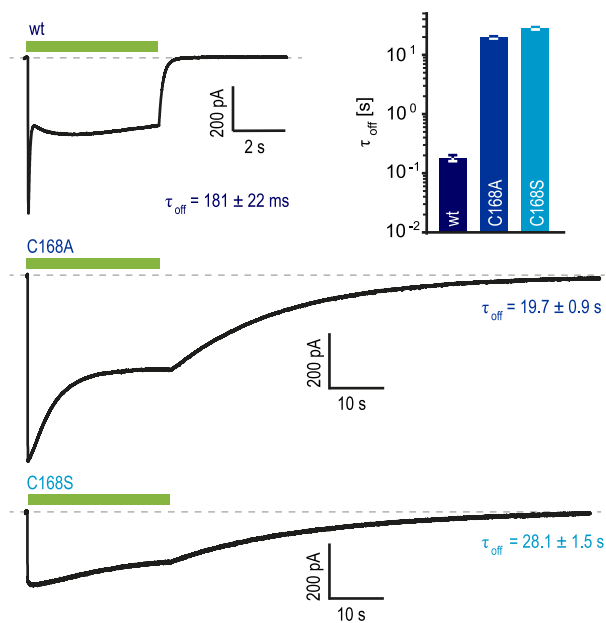


FIGURE S2 Slow-cycling mutants. Representative photocurrents of the C168A and C168S mutant after illumination with 530 nm light (25 s) and wild-type (wt) ReaChR (5 s, 530 nm). Bar graph summarizes monoexponentially fitted current decays (top right, mean  $\pm$  SE.;  $n = 6-7$ ).



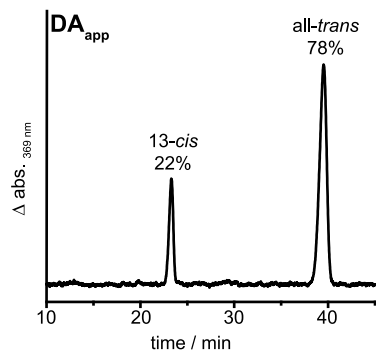


FIGURE S3 High-pressure liquid chromatography (HPLC) of retinal isomers isolated from ReaChR wild-type at pH 7.4 (room temperature). Before retinal extraction the sample was illuminated for 10 min ( $\sim 530$  nm LED) followed by 10 min dark adaptation (apparent dark state, DA<sub>app</sub>). The retinal isomers eluted after 23.3 min (13-cis) and 39.5 min (all-trans). The ratio of 78:22 all-trans:13-cis retinal was determined via peak integration.