Beyond spectral tuning: human cone visual pigments adopt different transient conformations for chromophore regeneration

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Abstract

Human red and green visual pigments are seven transmembrane receptors of cone photoreceptor cells of the retina that mediate color vision. These pigments share a very high degree of homology and have been assumed to feature analogous structural and functional properties. We report on a different regeneration mechanism among red and green cone opsins with retinal analogs using UV–Vis/fluorescence spectroscopic analyses, molecular modeling and site-directed mutagenesis. We find that photoactivated green cone opsin adopts a transient conformation which regenerates via an unprotonated Schiff base linkage with its natural chromophore, whereas red cone opsin forms a typical protonated Schiff base. The chromophore regeneration kinetics is consistent with a secondary retinal uptake by the cone pigments. Overall, our findings reveal, for the
first time, structural differences in the photoactivated conformation between red and green cone pigments that may be linked to their molecular evolution, and support the proposal of secondary retinal binding to visual pigments, in addition to binding to the canonical primary site, which may serve as a regulatory mechanism of dark adaptation in the phototransduction process.

**Keywords**

Color vision  
Ligand binding  
Visual phototransduction  
Retinal  
G-protein coupled receptors

**Abbreviations**

9CR  9-*cis*-Retinal  
11CR  11-*cis*-Retinal  
ATR  All-*trans*-retinal  
DM  *n*-Dodecyl-β-D-maltoside  
GPCR  G-protein coupled receptor  
ECL  Extracellular loop  
PBS  Phosphate buffered saline  
PSB  Protonated Schiff base  
SB  Schiff base  
TM  Transmembrane

**Electronic supplementary material**

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**Introduction**

Human color perception is mediated by cone photoreceptor cells which are mainly located in the fovea of the retina [1, 2]. Bright light activates the photosensitive opsin pigments which are embedded in the outer segment membrane discs of cone retinal cells thereby initiating the complex process of
photopic vision with a fast response [3]. The cone visual pigments correspond to blue opsin (SWL, short wavelength), green opsin (MWL, medium wavelength) and red opsin (LWL, long wavelength). In the genome, the red cone opsin gene (OPN1LW) is arranged proximal to a green cone opsin gene (OPN1MW) in an array at Xq28 with 98 % identical nucleotide sequences [4, 5] presenting a 96 % protein sequence identity and differing merely in 15 amino acids. This very high sequence identity has prompted the assumption that both green and red pigments share the same structural and functional features. The blue pigment, however, shows a lower degree of 43 % homology to the red and green opsins [5]. Thus, aside from their specific role in spectral tuning into different absorption wavelengths, the differences in structure–function relationships caused by this minor sequence variance have not been previously elucidated. Furthermore, most of the mechanistic insights into cone phototransduction are extrapolated from the rod photoreceptor cell system, responsible for dim-light scotopic vision, and its visual pigment rhodopsin which has been much more extensively studied [6].

The cone opsin pigments belong to the superfamily of G-protein-coupled receptors (GPCRs) consisting of a heptahelical transmembrane apoprotein and a photosensitive chromophore, the vitamin A derivative 11-cis-retinal (11CR). In its inactive dark-adapted state, the retinal chromophore is covalently bound to these visual pigments through its aldehyde group via a protonated Schiff base (PSB) linkage to the ε-amino group of Lys (numbering corresponding to red/green human opsins, with the Ballesteros–Weinstein general scheme [7] as superscript) in the seventh transmembrane (TM7) helix. The PSB is stabilized by a salt bridge with the carboxylic group of the counterion, Glu129 at TM3 [8]. A PSB is a structural feature of visual pigments and its corresponding visible spectrum would show an absorption maximum at 440 nm. However, the characteristic differences in visual sensitivities are due to specific amino acid variations in the retinal binding pocket which result in different absorption maxima—as a result of the so-called opsin shift effect—corresponding to 560 nm for the red cone pigment 530 nm for the green cone pigment, and 420 nm for the blue cone pigment [9–11].

The first step in the phototransduction process involves photon absorption by the visual pigments that causes 11CR isomerization to all-trans-retinal (ATR) [12] leading to the formation of active metarhodopsinII (MetaII) conformation which binds and activates the G-protein transducin [13]. Photoactivation of visual pigments alters the pKa of the PSB and induces deprotonation which is
the primary molecular event of the visual transduction cascade after retinal isomerization [14, 15]. Subsequent breakage of the PSB linkage results in retinal release from the retinal binding pocket. The MetaII conformation of cone opsins is short lived compared to that of rhodopsin [16–19].

In a recent report on the detailed comparison of photoactivated properties of rhodopsin and red cone opsin, we could conclude that there is a difference in retinal analog accessibility in the regeneration process of red cone opsin and rhodopsin [20] which was proposed to be relevant to the fine-tuning of ligand–receptor interactions in GPCRs [21]. We have now extended our approach by comparing the structural and functional features of the closely related red and green cone opsins upon photoactivation. To this aim we have focused on the detailed analysis of the regeneration ability, and subsequent dark adaptation, of red and green cone opsins with retinal analogs immediately after illumination, including the kinetics of retinal binding and release in the presence of exogenously added retinal.

Materials and methods

Materials

The red and green cone opsin genes, cloned into pMT4 plasmid vector, were kindly provided by Prof. Kevin D. Ridge. Dulbecco's modified Eagle medium (PAA Laboratories), supplemented with fetal bovine serum (Sigma), l-glutamine (Sigma), and penicillin–streptomycin (Sigma), was used to culture COS-1 cells (American Type Culture Collection no. CRL-1650). 11CR was provided by the National Eye Institute, National Institutes of Health. Purified mAb rho-1D4 was obtained from Cell Essentials and was coupled to CNBr-activated Sepharose beads (Sigma). n-Dodecyl-β-D-maltoside (DM) was purchased from Affymetrix. The nonamer-peptide H-TETSQVAPA-OH was obtained from Unitat de Tècniques Separatives i Síntesi de Pèptids, Universitat de Barcelona. 9CR, hydroxylamine, protease inhibitor cocktail, and phenylmethanesulfonyl fluoride (PMSF) were purchased from Sigma, and polyethyleneimine (PEI) was purchased from Polysciences. Quartz absorption cuvettes with light path 10 mm (Hellma Analytics, Germany). Oligonucleotides were purchased from Sigma (Spain).

Expression and purification of red and green cone pigments

Red, or green, cone opsin was expressed in transiently transfected COS-1
cells by chemical transfection using PEI reagent. Cells were harvested 48–60 h after transfection and regenerated with 10 µM 11CR in PBS buffer (137 mM NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, and 1.8 mM KH₂PO₄, pH 7.4) by overnight incubation at 4 °C. Regenerated cells were subsequently solubilized using 1 % DM with PMSF and protease inhibitors, and the pigments were purified by immunoaffinity chromatography using Sepharose coupled to rho-1D4 antibody. The bound cone opsin pigments were eluted in PBS containing the nonamer peptide and 0.05 % DM.

**UV–Vis spectrophotometry of cone pigments**

Purified cone pigments were spectroscopically characterized using a Varian Cary 100 Bio spectrophotometer (Varian), equipped with a water-jacketed cuvette holder connected to a circulating water bath. Temperature was controlled by a peltier accessory connected to the spectrophotometer and the temperature was maintained at 20 °C during all experiments. All spectra were recorded in the 250–650 nm range for green and 250–700 nm for red cone opsin, with a bandwidth of 2 nm, a response time of 0.5 s, and a scan speed of 400 nm/min.

**Fluorescence spectroscopy of cone pigments**

QuantaMaster 4 spectrofluorimeter (Proton Technology International) was employed to measure the Trp fluorescence emission corresponding to retinal release and uptake processes. The excitation wavelength was 295 nm, and the emission wavelength was 330 nm, measuring 1 point per second for 2 s, followed by a 28 s pause (with a beam shutter to prevent sample photobleaching by the fluorimeter lamp). The excitation slit settings were 0.5 nm, and emission was 10 nm. Trp fluorescence was monitored over time in the dark until a steady baseline was obtained. Then the sample was photobleached (>495 nm) for 30 s using a Dolan-Jenner MI-150 fiber-optic illuminator, and the change in fluorescence was recorded. For regeneration experiments, a 2.0- to 2.5-fold molar concentration of retinal over pigment was used which accounts for a maximum of 1 % of the protein sample volume.

**Site-directed mutagenesis**

Phe309Tyr was introduced into human green cone opsin gene using site-directed mutagenesis kit (Quickchange, Stratagene) using the mutagenic primers: forward primer -5′GCCCTGCGGCTACTTTGCCCCAAAAGTGGGC3′, reverse primer -5′GGCATTTTTGGCAGTAGGGCCGCGCCAGGC-3′.
Molecular models of red and green opsin

Homology models for the dark state and active human red and green opsin were constructed, respectively, based on the crystal structure of bovine rhodopsin (PDB id: 1GZM) [22] and the opsin soaked with ATR (PDB id: 2X72) [23]. Modeller 9.8 was used for this purpose [24]. All models were energy minimized using AMBER program [25], employing the Amberff99SB-ILDN forcefield [26].

We computed interaction potential maps [27] using them as implementation in MOE (Molecular Operating Environment (MOE), 2013.08; Chemical Computing Group Inc., 1010 Sherbooke St. West, Suite #910, Montreal, QC, Canada, H3A 2R7, 2015.) in order to identify possible secondary binding pockets for retinal. Flexible docking of the second retinal molecule was done using AutoDock Vina [28].

Results

Regeneration mechanism of green and red cone opsin with retinal analogs

The dark-state UV–Vis spectrum of purified green cone pigment showed two characteristic bands with absorption maxima at 530 and 280 nm (Fig. 1a, solid line) corresponding to the amount of regenerated green cone pigment present and the total opsin pigment, respectively. The regeneration experiment was carried out by adding an aliquot of a concentrated stock sample of 11CR yielding increased absorbance at 380 nm corresponding to the free 11CR added (Fig. 1a, dotted line). No absorbance increase at 530 nm (Fig. 1a, dashed line) was observed after illumination which suggested that the photoactivated conformation of green cone opsin was not able to regenerate with 11CR. In contrast, a parallel experiment performed with 9CR, showed a regenerated band at ~495 nm with 85 % maximal regeneration (Fig. 1b).

Fig. 1

Regeneration of photoactivated green and red cone pigments with 11CR and 9CR. a, b Immunopurified human recombinant green cone pigment was analyzed by UV–visible spectrophotometry in the dark (1, solid line) and after addition of exogenous ligand, i.e., 2.5-fold of either 11CR (a) or 9CR (b) over opsin (2, dotted line). The pigment was then illuminated for 30 s (>495 nm) and a spectrum was recorded after 30 min (3, dashed line). c, d Red cone opsin was spectroscopically measured in an analogous way to green cone opsin and spectra were measured in the dark (1, solid line), after 11CR (c) or 9CR (d) addition (2,
Photoactivated red cone opsin regenerates via a PSB with both 9CR and 11CR

An analogous regeneration experiment was carried out with purified red cone opsin. In both experiments, 9CR and 11CR can bind opsin after illumination and regenerate the visible chromophore with PSB linkage indicated by the presence of absorbance increase in the visible region above 440 nm. In particular, complete regeneration with 11CR was achieved as indicated by the re-appearance of the typical absorption maximum at 560 nm (Fig. 1c), whereas ~70% regeneration with 9CR was recovered with an absorption maximum at ~525 nm (Fig. 1d). The shifted absorption maximum of the protein is due to the specific absorbance properties of 9CR.

Photoactivated green cone opsin regenerates with 11CR by means of an unprotonated SB linkage

The lack of absorbance increase in the visible region suggested no 11CR binding and prompted the question whether green cone opsin could be regenerating with 11CR by means of an unprotonated SB linkage in which the absorption maximum of the regenerated pigment would be shifted to the UV region [29]. To test this hypothesis, two parallel UV–Vis spectroscopic experiments were carried out with the same green opsin sample that was...
illuminated and split into two aliquots. One of the aliquots was acidified immediately after photobleaching, whereas the other was acidified 30 min later (Fig. 2a). The difference spectrum between the two aliquots showed a band at ~440 nm (Fig. 2b) which clearly indicated that the photoactivated green cone opsin regenerates via an unprotonated SB linkage with 11CR. The same experiment was carried out for red cone opsin, in which 11CR regeneration after illumination can be detected in the visible region, and sample acidification shifted the absorption band to 440 nm as well (Fig. S1).

In order to rule out that the differences in regeneration mechanisms may be due to the different stability of red and green cone opsins in the detergent solution, the reactivity of dark-state red and green cone opsin pigments with hydroxylamine (which can access the cone opsin retinal binding site and sequester the bound chromophore to form retinaloxime) was tested (Fig. S2). Both pigments showed similar reaction kinetics indicating that differences in stability were not the cause of the observed different regeneration mechanisms.

**Fig. 2**

Photoactivated green cone opsin regenerates with 11CR by means of an unprotonated Schiff base linkage. **a** The UV–vis spectrum of purified green cone pigment was measured in the dark (1, *solid line*). After addition of 2.5-fold 11CR, it was illuminated for 30 s (λ > 495 nm) (2, *dotted line*). Then, the sample was split into two aliquots in which one was acidified with 2 N H₂SO₄ immediately (3, *dashed line*) and the other 30 min (4, *dot-dashed line*) later. **b** A difference spectrum was obtained by subtracting the spectrum of the sample acidified after 30 min and the sample acidified immediately after photoactivation. **c** Green cone pigment, in a fluorimetric micro-cuvette was illuminated for 30 s (λ > 495 nm) after a stable baseline was obtained. Then, 2.5-fold of 11CR to the concentration of the pigment was added immediately after photoactivation and mixed thoroughly. The *dotted line* represents the fast regeneration process immediately after 11CR addition.
We also measured the photobleaching process by means of fluorescence spectroscopy by monitoring the Trp fluorescence increase upon green cone opsin illumination (see “Methods: Materials and methods”). In the dark state, the fluorescence from Trp residues at the retinal binding pocket is quenched by the β-ionone group of the retinal. Upon photoactivation, the exit of retinal from opsin results in a fluorescence increase that (as the cone opsins exhibit a fast MetaII decay kinetics) reaches a maximum within a second [20, 30]. Purified green cone pigment sample was illuminated after obtaining a stable baseline and the sudden increase in fluorescence indicated fast retinal release and MetaII decay processes. Subsequently, immediate addition of exogenous 11CR to the illuminated sample resulted in a decrease in the fluorescence signal confirming binding of 11CR to photoactivated green cone opsin (Fig. 2c) as previously reported [20].

Molecular modeling was carried out to understand the structural features behind such remarkable functionally relevant differences between red and green cone opsins. Figure 3a shows a sequence alignment between the human red and green cone opsins (including bovine rhodopsin as a reference), excluding the N- and C-terminus (Fig. 3a). The two photoreceptors have same exact number of residues and differ in only 15 residues, with most changes located in the central portion of the transmembrane (TM) helices 5 and 6. The similarity is remarkable in the retinal binding pocket, where only position...
309 in TM7 is different (Tyr and Phe in red and green opsins, respectively). Assuming a similar movement of this residue with receptor activation, as observed in the crystal structures of rhodopsin, Tyr309 would form, in red cone opsin, a hydrogen bond with Thr201 in ECL2 (Fig. 3b). This interaction, which is not possible in green cone opsin, could underlie the observed differences in spectral and retinal regeneration properties. The involvement of the 309 residue in spectral tuning of human red and green cone pigments has been previously reported [31].

**Fig. 3**

Phe309 regulates Schiff base protonation during 11CR regeneration of photoactivated green cone opsin. a Sequence alignment between the human red and green opsins and bovine rhodopsin (in gray), excluding the N- and C-terminus. Residues highlighted in blue indicate differences between red and green opsins (numbered according to the general scheme [7]). b Molecular model of the inactive red opsin showing the proposed interaction between Tyr309 in TM7 and Thr201 in ECL2. The cartoon of TM7 and the conformation of Tyr309 in the active state are shown with a transparent representation, and the arrow indicates the movement. c 11CR regeneration experiment was carried out with purified F309Y green cone opsin mutant. After measuring a dark-state spectra (1), 11CR was added prior to illumination (2) and the regenerated spectrum was recorded after 30 min (3) showing a band at 530 nm and the sample was acidified (4). c inset. A closer look at the visible region of dark (1) and (2) regenerated spectra. d The difference spectrum, obtained by subtracting the acidified spectrum and the regenerated spectrum, shows a clear difference band at 530 nm
To validate the modeling observation, Phe309Tyr mutation was introduced into green cone opsin and the 11CR regeneration experiment was carried out with the purified mutant pigment which showed a regenerated peak at 530 nm (Fig. 3c). The difference between the acidified and the regenerated spectra showed a clear band at 530 nm representing the PSB in 309Tyr mutant of green cone opsin, and the difference band at 440 nm corresponds to the amount of net SB from the regenerated pigment (Fig. 3d). The fact that the amplitude of the 530 nm band is lower than that of WT pigment regenerated with 11CR, suggests that other residues, like Ala285 (Thr in red cone opsin), close to the β-ionone ring and Phe277 (Tyr in red cone opsin), close to Trp288 are responsible for the spectral differences in green and red opsins. However, the role of Tyr309 in modulating the process of protonation of photoactivated green cone pigment during 11CR regeneration has been substantiated.

**Red and green cone opsins chromophore regeneration kinetics**

The conformation of opsin and the photochemistry of the chromophore determine the spectral properties of cone visual pigments and, in turn, the observed spectral behavior could provide hints about the structural details of the receptor proteins [32]. Illumination causes a decrease in absorbance of the
Regenerated pigment band in the visible region which would be apparently parallel to an absorbance decrease at 380 nm. The 380-nm absorbance decrease, upon chromophore regeneration, could reflect either 11CR binding to, or ATR release from, the opsin receptor. Each data point corresponding to the absorption maximum in the visible region and the retinal region of each regenerated spectra was plotted versus time. The data could be fit to the curves reflecting a fast regeneration process that rapidly reached a plateau.

Regeneration kinetics of green and red cone opsin with 9CR
In red and green cone opsins, the kinetics of the 9CR regeneration process gave $t_{1/2}$ of 1.31 ± 0.39 min and 0.66 ± 0.23 by red and green cone opsin, respectively (Fig. 4a). The decrease in absorbance at 380 nm parallels the visible increase and represents entry of retinal into photoactivated opsin during the regeneration process (Fig. 4b) as expected. Parallel experiments were carried out with purified rhodopsin, in which the regeneration with 11CR and 9CR followed single saturation patterns which are significantly slower than those observed for cone opsins, and can be correlated with absorbance changes at 380 nm (Fig. S3).

Fig. 4
Regeneration and retinal release kinetics of green and red cone opsin with 9CR. 

a The absorbance increase at the $\lambda_{\text{max}}$ in the visible region of each regenerated red/green pigment was plotted and fit with a saturation curve. Chromophore regeneration with 9CR was measured for red and green cone opsins for 30 min. The dark-state spectral values of red and green cone opsin were normalized to have equivalent dark absorbance and the regenerated absorbances were also normalized by the same factor. b Absorbance changes at 380 nm for the regenerated samples were plotted and fit to a single exponential decay curve

Secondary retinal uptake kinetics by cone visual pigments after regeneration
Analogous regeneration experiments were carried out with 11CR but only for red cone opsin as green cone opsin showed no absorbance increase in the visible region upon regeneration. A similar pattern to that observed for 9CR was obtained in the case of red cone opsin regenerating with 11CR with a $t_{1/2}$ of $0.70 \pm 0.3$ min, and the regeneration with 11CR was found to be even faster than with 9CR (Fig. 5a). This behavior is in agreement with the complete fast regeneration observed for red cone opsin with 11CR (Fig. 1c).

**Fig. 5**

Regeneration and secondary retinal uptake kinetics of red and green cone opsin with 11CR. **a** 2.5-fold of 11CR to the concentration of red cone opsin was added in the dark and illuminated for 30 s ($\lambda > 495$ nm). UV–visible spectra of the sample were obtained every min for 30 min. The $\lambda_{\text{max}}$ at the visible region of each regenerated spectrum was plotted and fit with a saturation curve. **b** In the regeneration experiments with 11CR, the regeneration was followed at 380 nm and the absorbance change at this wavelength was monitored for 30 min and the data fit to a sigmoidal curve. **c** Molecular model of the active red opsin showing the potential coexistence of two retinal molecules one in the primary site (as observed in the rhodopsin structure with PDB id 2X72 [49] in pale yellow) and another in the proposed secondary sites (showing representative poses obtained from flexible docking; in *multiple colors*). TM helices are represented as cylinders and side-chains of neighboring residues are shown with sticks.
Cone opsin regeneration is usually quantified by measuring the absorbance increase at the characteristic visible band, but in our case this was not possible for green cone opsin chromophore regeneration with 11CR. In order to estimate the regeneration kinetics, we measured the decrease in absorbance at 380 nm with time. This would reflect retinal entry into the binding site of cone opsins and was expected to parallel the regeneration process. In this regard, the change in absorbance at 380 nm after illumination was plotted and the data could be unexpectedly fit to a sigmoidal curve (Fig. 5b) which suggested entry of 11CR into visual pigments at a slower rate, with $t_{1/2}$ of 11.38 ± 2.34 min for red, and 13.12 ± 0.78 min for green cone opsin, respectively. This behavior is in contrast to the faster binding kinetics of cone opsins regenerated with 11CR (Fig. 5a). This fast regeneration which is complete within 1.0 min suggests that the slower process observed (Fig. 5b) cannot be due to retinal binding to the 11CR canonical pocket, but suggested additional uptake of 11CR into a secondary binding site facilitated by the presence of 11CR excess in the protein milieu.

In the search for a possible secondary binding pocket for 11CR, we computed interaction potential maps for a hydrophobic probe in both inactive and active models of red and green opsins (see “Materials and methods”). The maps
revealed a cavity between TMs 1 and 7 in the active models of red and green opsins as well as in rhodopsin with sufficient size to accommodate a second retinal molecule. Flexible docking calculations revealed possible binding modes of retinal penetrating through TMs 1 and 7 (Fig. 5c) placing the aldehyde group of retinal close to His197, Thr201, Ser202 in ECL2 and the \( \beta \)-ionone ring in the mouth of the hydrophobic pocket between TM1 and TM7, flanked by residues Thr56 and Leu302. Because TM7 moves with activation, increasing the size of the cavity, only the active structures seem to allow binding of a second retinal in this region. This cavity is compatible with the proposed entry/exit channels for retinal in rhodopsin [33] and with the presence of hydrophobic molecules observed in the crystal structures of rhodopsin, as we previously noted [34].

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We interpret this phenomenon as a slower binding of a second 11CR molecule to the regenerated pigment to a site with lower affinity than that at the canonical ligand binding pocket (Fig. 6). This secondary 11CR uptake might be potentially coupled with a delay in ATR release from the protein. We propose that this slower secondary retinal uptake is a distinct phenomenon to that of the fast regeneration of red and green cone opsins observed in a ~1 min time frame with 11CR. Such a behavior was not observed when photoactivated red and green cone opsins were regenerated with 9CR (Fig. 4b) suggesting that it is specific in the case of the native 11CR chromophore.

Fig. 6
Photoactivated intermediate conformations and 11CR regeneration of red and green cone pigments. a Cone pigments heterologously expressed in mammalian cells were regenerated with 11CR and immunopurified. b Exogenous 11CR (2.5 molar ratio over opsin) was added in the dark, to the immunopurified regenerated sample, and the sample was subsequently illuminated (\( \lambda \) > 495 nm). This causes the isomerization of opsin-bound 11CR to ATR resulting in a protein conformational change that facilitates fast pre-binding of free 11CR to a secondary, lower affinity, retinal binding site followed by chromophore regeneration. c Fast chromophore regeneration would be achieved by translocation and covalent binding of this pre-bound 11CR to the chromophoric binding pocket by displacing ATR which in turn would move out from the protein. Further, the secondary uptake of 11CR by the visual pigment, with a slower kinetics, may be due to the conformational change of the regenerated protein coupled with the reduced availability of bulk 11CR that would synergistically decelerate 11CR binding to the secondary site. d Under
Discussion

Specific amino acids at the retinal binding site, as well as their interactions with the retinal chromophore and surrounding water molecules, modulate spectral tuning of visual opsins [35–37]. Cone opsin pigments have been much less characterized than the rod photoreceptor rhodopsin due to their lower availability and their higher conformational instability. With only 4% variance between red and green cone opsin sequences, these were believed to have analogous structure–function relationships, other than the well-established wavelength absorption fine-tuning in the visible region of the spectrum.

Green cone opsin regenerates through an unprotonated SB intermediate conformation that eventually has to become protonated in vivo to regenerate the 530-nm absorbing chromophore. This novel intermediate conformation may have a decreased pKa of the SB nitrogen, in the case of green cone opsin, that could imply electrostatic changes at the SB linkage vicinity including changes in the SB-counterion distance. Though the mechanism that would convert this unprotonated SB intermediate conformation to the typical native green cone opsin pigment is unclear, the native membrane environment could potentiate the reaction by shortening the life-time of the transient intermediate we observe in our in vitro system. Green cone cells are less abundant in the human retina (the ratio of red cone cells to green cone cells in the human retina mosaic is estimated to be 1.5–4.5:1) but their response is believed to be analogous to that of red cone cells [38]. This different cone cells population, and the structural differences herein reported, could be linked to specific subtle differential roles for red and green cone visual pigments in color vision.

A clear difference is found in the intermediate conformation after
photoactivation of red and green cone opsin pigments that may be related to differences arisen in the molecular evolution of these two genetically closely related visual pigments. Analogous transient intermediate conformations of the ligand-free opsin, during the regeneration process, have also identified for rhodopsin [39]. The unprotonated SB conformation detected could be involved in the dark adaptation process of the photoactivated receptor, in an excess of retinal environment, and consequently play a role in the molecular basis of color vision by differentially modulating the chromophore regeneration and functionality of green and red cone photoreceptor cells.

We believe that our spectral changes are not reflecting potential retinal binding to other regions on the surface, or external side of the protein, for several reasons. First, the secondary binding observed is not covalent, so binding to external Lys residues is excluded. It should also be pointed out that retinal binding to the protein surface is unlikely to cause the spectral change at 380 nm observed. The absorbance decrease at 380 nm could be the result of an altered extinction coefficient resulting from the second retinal bound to hydrophobic helices and the extracellular domain of the receptor. This secondary retinal uptake, after retinal binding to its primary site, could be coupled with ATR exit from the regenerated cone opsins. The exit of the isomerized ATR could share the same channel as the entry of the secondary retinal molecular and this would justify the slower kinetics of the sigmoidal curves obtained (Fig. 5).

Furthermore, such non-specific retinal-protein surface interactions would not be isomer specific but, in our case, only 11CR can induce the spectral behavior observed. Thus, the secondary binding site proposed shows specificity towards 11CR, but not 9CR, and such ligand specific accessibility would be difficult to explain if the secondary retinal binding would happen at the surface of the protein. The geometry of 11CR would favor accessibility into the opsin secondary binding site in contrast to the 9CR geometry. This would reflect that 11CR has been evolutionarily selected to act as the natural chromophore in vision. All these arguments reinforce our proposal of secondary retinal binding at the proposed site by our molecular modeling analysis. Notably, the two phase kinetics observed from the fluorescence regeneration experiment (Fig. 2c) in the case of green cone opsin provides further support to this interpretation. The fast component would represent regeneration with 11CR at the primary site, and the second slower component would be related to the secondary retinal uptake.
Our findings are in agreement and support the results of previous studies proposing that more than one retinoid can interact with visual opsins [40] which strongly suggests the presence of functionally relevant secondary binding sites of retinoids in opsin [41]. Furthermore, previous experimental evidence showed an important role for non-covalent 11CR binding to opsin prior to regeneration [42] and in its ability to bind to a selective conformation of the protein among various conformational modes of visual pigments prior and after photoactivation [39].

In addition to the secondary 11CR binding event, changes at 380 nm could also be reflecting changes in ATR release that could alter the rate of retinoid recycling thus regulating dark-state adaptation after photoactivation for cone opsins. Retinoid binding protein, ABCA4, is an ATP dependent transporter of ATR from the luminal side to the cytoplasmic side of photoreceptor cells. At the cytoplasmic side, ATR is converted to all-trans-retinol by RDH8 which is then released to Müller cells or retinal pigment epithelium (for the retinoid cycle) and further converted to 11CR which in turn diffuses into photoreceptor cell and results in dark adaptation of visual pigments [43–45]. Stargardt disease is an autosomal recessive disorder characterized by mutations at ABCA4, in which vision loss is associated with accumulation of ATR and its derivatives that can be toxic by causing oxidative stress that can result in damage of the retinal cells [46, 47]. It is also known that transportation of ATR by ABCA4 is a multi-step process and that the substrate for ABCR4 is N-retinylidene-phosphotidyl ethanolamine (NRPE) formed from the interaction of ATR with phosphotidyl ethanolamine. NRPE is a reversible product, and the conversion process of RDH8 from ATR to all-trans-retinol is slow [43, 48]. Hence, the controlled release of ATR after photoactivation, coupled to 11CR binding to a secondary binding site other than the canonical retinal binding pocket, would help regulating the effective transport of ATR, from cone opsins. This may suggest a temporary entrapment of ATR to the same or other secondary binding site, and this may be connected to a slower movement of the hydrophobic retinal molecule along the ligand channel proposed between openings A and B which would be perpendicular to the helices axis. A secondary retinal binding site in cone visual pigments can accommodate a retinal molecule in addition to the one bound to the classical retinal binding pocket [40].

We propose that this second binding site (specific for 11CR and not for 9CR) may function as a retinal buffer, in cone opsins, and may possibly have an allosteric role although this may need further investigation. The different behavior for 11CR and 9CR is also evident
from the faster initial velocity for the chromophore regeneration observed with 11CR than 9CR (Figs. 4, 5). This secondary binding site may allow retinal pre-binding and subsequent fast transfer of this retinal molecule to the primary retinal binding pocket for covalent binding and dark-adapted pigment regeneration ready for another light stimulus.

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**Electronic supplementary material**

Below is the link to the electronic supplementary material.

Supplementary material 1 (DOCX 61253 kb)

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