Molecular Dynamics Study of the 13-cis form (bR$_{548}$) of Bacteriorhodopsin and Its Photocycle

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ABSTRACT  The structure and the photocycle of bacteriorhodopsin (bR) containing 13-cis,15-syn retinal, so-called bR$_{548}$, has been studied by means of molecular dynamics simulations performed on the complete protein. The simulated structure of bR$_{548}$ was obtained through isomerization of in situ retinal around both its C$_{13}$-C$_{14}$ and its C$_{15}$-N bond starting from the simulated structure of bR$_{568}$ described previously, containing all-trans,15-anti retinal. After a 50-ps equilibration, the resulting structure of bR$_{548}$ was examined by replacing retinal by analogues with modified ß-ionone rings and comparing with respective observations. The photocycle of bR$_{548}$ was simulated by inducing a rapid 13-cis,15-anti → all-trans,15-syn isomerization through a 1-ps application of a potential that destabilizes the 13-cis isomer. The simulation resulted in structures consistent with the J, K, and L intermediates observed in the photocycle of bR$_{568}$. The results offer an explanation of why an unprotonated retinal Schiff base intermediate, i.e., an M state, is not formed in the bR$_{548}$ photocycle. The Schiff base nitrogen after photoisomerization of bR$_{548}$ points to the intracellular rather than to the extracellular site. The simulations suggest also that leakage from the bR$_{548}$ to the bR$_{568}$ cycle arises due to an initial 13-cis,15-anti → all-trans,15-anti photoisomerization.

INTRODUCTION

Bacteriorhodopsin (bR) is a protein composed of seven ß-helices that spans the purple membrane of Halobacterium halobium and that functions as a light-driven proton pump. It is a member of the retinal protein family, which encompasses proteins with a retinal chromophore bound within the protein interior via a protonated Schiff base linkage to a lysine side chain. The retinal isomer composition in bR is 0.66 13-cis and 0.34 all-trans in the dark-adapted form of the pigment (DA), a ratio that is altered in mutants of bR$_{548}$ and in bacterial rhodopsins of other species. The two isomers denoted bR$_{548}$ (13-cis) and bR$_{568}$ (all-trans) absorb at 548 and 568 nm, respectively. Retinal in the bR$_{568}$ pigment exists in the all-trans,15-anti configuration, and in the bR$_{548}$ pigment in the 13-cis,15-syn configuration as suggested first in Orlandi and Schulten (1979) and observed in Harbison et al. (1984), Smith et al. (1984), and Livnah and Sheves (1993). Both the bR$_{568}$ and the bR$_{548}$ isomers undergo a characteristic photocycle (see Fig. 1) initialized in both cases by a photoisomerization involving rotation around the C$_{13}$-C$_{14}$ double bond. However, proton pumping is restricted to the photocycle of bR$_{568}$. The photocycle of bR$_{548}$ does not involve a vectorial proton translocation. The photocycle of native bR$_{548}$ leaks into the bR$_{568}$ form whereas the bR$_{568}$ photocycle replenishes only the bR$_{548}$ form. As a result, under the influence of light, any mixture of bR$_{568}$/bR$_{548}$ adopts to the bR$_{568}$ pigment, which is accordingly called the light-adapted form (LA). For recent reviews, see Khorana (1988), Birge (1990), Mathies et al. (1991), Lanyi (1992), Oesterhelt et al. (1992), and Ebrey (1993).

Three main questions arise concerning bR$_{548}$ and its photocycle. What factors determine the ratio of the bR$_{568}$ and the bR$_{548}$ forms? Why does the photocycle of bR$_{548}$ not pump protons and what does this fact tell us about the mechanism of proton pumping in bacteriorhodopsin? By what pathway is 15-syn retinal of bR$_{548}$ transformed to 15-anti retinal of bR$_{568}$ during light adaptation? The availability of the structure of bacteriorhodopsin (bR$_{568}$) reported by Henderson et al. (1990) provides an opportunity to answer these questions. A promising approach is provided by molecular dynamics simulations based on this structure.

Unfortunately, the structure in Henderson et al. (1990) is of relatively low resolution (e.g., water bound in the protein is not resolved) and also does not entail the interhelical loops. As a result, the loops needed to be added, the structure refined, and water added before molecular dynamics simulations could be carried out. In this paper, we extend previous molecular dynamics simulations (Nonella et al., 1991; Zhou et al., 1993; Humphrey et al., 1994) that focused on bR$_{568}$ to study the bR$_{548}$ pigment and its photocycle. As a test of the proposed bR$_{548}$ structure obtained, we examine the replacement of retinal by various analogues as well as investigate the properties of the D85A mutant of bR$_{548}$. The results of our simulations are compared with the relevant experimental observations (Rath et al., 1993; Ottolenghi and Sheves, 1989, Steinberg et al., 1991).

MATERIALS AND METHODS

The original structure of bR, as determined by Henderson et al. (1990) with electron microscopy, resolved the membrane-spanning helical residues; the interhelix loops had been added to this structure and equilibrated by molecular dynamics simulations in Nonella et al. (1991). The backbone atoms

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of the resulting simulated model had a RMSD of 2.8 Å from the electron microscopy model structure. A first attempt to place water into bacterio- rhodopsin had been made by Zhou et al. (1993). Significant improvements of the simulated bR358 structure, reported in Humphrey et al. (1994), resulted from a refinement procedure that included an all-atom force field and hydrogen bonding forces as well as from a renewed attempt to place water molecules inside bR. This refined bR358 structure shows close agreement with the Henderson structure (Henderson et al., 1990), the root mean square deviation (RMSD) for the Cα backbone atoms of the membrane-spanning part of bR measuring only 1.8 Å. An explicit representation of all of the hydrogen atoms has also been incorporated into the simulated structure. The most important feature of the refined model is that it reproduces well the interactions in the retinal binding site as observed through reconstitution of bR with retinal analogues. This supports the assumption that structural details at a resolution in the 1-Å range are properly represented, but, of course, there exists a large probability that the structure also contains major faults. The refined model of bR358 has been used as a starting point of our simulations of bR358.

The program X-PLOR (Brünger, 1988), with the CHARMM energy function (Brooks et al., 1983) was used for all molecular dynamics simulations reported here. All simulations used a cutoff distance of 8 Å and a dielectric constant of ε = 1. All atoms, including hydrogens, are described explicitly. We also use an explicit hydrogen-bonding term in the energy function provided by X-PLOR/CHARMM. The parameters and charges used in the simulations are, respectively, the parma11h3x.pro parameters and the topal1h6x.pro charges (Brooks et al., 1983; Brünger, 1988), except for retinal.

Equilibrium bond lengths, angles, and torsional angles assumed for the retinal chromophore are those determined by recent x-ray crystallography studies (Santantiero and James, 1990). The charges and force constants used for the protonated Schiff base of retinal are those also used in Humphrey et al. (1994).

The protonation states of titratable groups in bR are as given in the standard X-PLOR amino acid topology files, except for Asp-96 and Asp-115, which are assumed protonated according to the observations in Gerwert et al. (1989) and Engelhard et al. (1990). As suggested previously (Bashford & Gerwert, 1992; Zhou et al., 1993), Arg-82 has been moved up to a position facing toward the cytoplasmic side of the protein. In this position, Arg-82 is part of the Schiff base counterion complex. All simulations, except where otherwise stated, were performed at 300 K.

Inducing the light → dark adaptation

The 13-cis,15-syn isomer of bacteriorhodopsin, i.e., of bR358, was simulated through isomerization of in situ retinal around both its C15-C14 and its C17-N bond starting from the simulated model of bR358 reported in Humphrey et al. (1994) containing all-trans retinal. Isomerization around these bonds is accomplished through modification of the potential function governing the respective dihedral angles. The potential used in X-PLOR is

$$E_{dih} = k_\phi [1 + \cos(n\phi + \delta)]$$

(1)

where ϕ is the actual torsion angle, kϕ is an energy constant, n is the periodicity, and δ is a phase shift. In the electronic ground state, n = 2 and δ ~ 180° are suitable parameters for the conjugated carbon chain of retinal. This implies that each bond can assume either a trans or a cis configuration. The isomerization all-trans,15-anti → 13-cis,15-syn was induced by changing the torsional potential in Eq. 1 for the C15-C14 and the C17-N bond such that the all-trans,15-anti configuration became unstable and the 13-cis,15-syn stable. During this procedure the potentials adopted for the C15-C14 and the C17-N bond were

$$E_{dih}^{13-14} = k_\phi^{13-14} [1 - \cos(\phi)]$$

(2)

$$E_{dih}^{15-N} = k_\phi^{15-N} [1 - \cos(\phi)]$$

respectively. These potentials have a maximum at the trans position ϕ = 180° and a minimum at the cis position ϕ = 0°. kϕ13-14 and kϕ15-N were taken to be 2.5 kcal/mol to ensure that the all-trans,15-anti → 13-cis,15-anti transition occurs in a few picoseconds, i.e., rapidly, without, however, increasing the temperature of retinal significantly. The maximum of the new torsional potential is Emax (LA → DA) = 5 kcal/mol. The procedure was followed by equilibration at 300 K for 50 ps. The equilibrated retinal-protein system was identified with bR358 and served as the starting structure for the study of the photocycle of bR358.

Inducing the photoisomerization

The initial photoisomerization step of the photocycle of bR358 was induced by instantaneous change of the torsional potential (Eq. 1) for the C17-C14 bond such that the 13-cis configuration became unstable and the respective trans configuration stable. The potential used for this purpose was

$$E_{dih}^{13-14} = k_\phi^{13-14} [1 + \cos(\phi)]$$

(3)

This potential has a maximum at the 13-cis position ϕ = 0° and a minimum at the trans position ϕ = 180°. For kϕ13-14, a value of 22.5 kcal/mol had been assumed such that the torsional potential at its maximum measures $E_{max}^{13-14}$ (photo) = 45 kcal/mol, i.e., a value close to the energy of a 548-nm photon. The reader should note that this value is much larger than the value of $E_{max} = 5$ kcal/mol(LA → DA) assumed for the transformation from light-adapted to dark-adapted bR.

The primary reaction of the 13-cis photocycle is very fast. The time needed to reach the intermediate J is only 1.2 ps (Zinth et al., 1988). This implies that the crossing between potential energy surfaces of the S0 and S1 states of the retinal chromophore occurs close to the maximum of the ground state potential surface (Schulten and Tavan, 1978). It has been suggested that retinal relaxes in ~200 fs (Mathies et al., 1991) to the excited state minimum near the ground state isomerization barrier, with a torsion around the C15-C14 bond of ~90°, then crosses to the ground state potential surface and completes the photoisomerization. In our simulations, this process was described by the single potential surface (Eq. 3) that, hence, represents in its higher energy range the excited state potential and, in its lower energy range, the ground state potential. Photoisomerization of the C15-C14 bond is achieved by placing the initial 13-cis state at the maximum of the potential (Eq. 3). Subsequent simulations then monitor the relaxation of retinal down the potential surface (Eq. 3). The simulations account for inertial effects as well as for the steric interactions that resist the isomerization enforced by the potential (Eq. 3).

Corotation of the C14-C15 bond

We started the simulated photoisomerization from the 13-cis,15-syn retinal configuration. Previous studies (Nonella et al., 1991; Zhou et al., 1993) showed that concomitant rotations around the C14-C15 bond can occur and determine the geometry of retinal after the photoisomerization. Because
little is known regarding the excited state torsional barrier for this bond, we
carried out simulations with different respective barriers. The potential
describing the torsional angle of the C_{14}-C_{15} bond is

$$E_{\text{bar}}^{\text{14-15}} = k^{14-15} [1 - \cos(2\phi)],$$  \hspace{1cm} (4)

where the parameter $k^{14-15}$ is chosen according to the torsional energy barrier
desired. In one simulation, we chose a torsional barrier (of the C_{14}-C_{15} single
bond) of 2 kcal/mol to enable essentially free rotation around this bond in
the excited state such that both the 13-cis $\rightarrow$ all-trans and the 13-cis $\rightarrow$ 14-cis
isomerizations are feasible. In another simulation, we enforced a 13-cis,
15-syn $\rightarrow$ 15-syn isomerization by assuming a barrier of 10 kcal/mol,
preventing rotation around the C_{14}-C_{15} bond.

**Sampling photoisomerization processes**

In the present work we have addressed also the question of connectivity
between the cycles of brS_{548} and brS_{556}. We have explored at which state of
the brS_{56} cycle leakage to the brS_{56} cycle might occur. To account for the
fact that the probability of leakage between the cycles is smaller than unity
we simulated photoisomerization processes for different (random) initial
velocities (starting from the same structure of brS_{56}) to account for the
possibility that the photoisomerization of brS_{56} might not always involve the
same geometrical transformation.

**Simulated annealing for the L intermediate**

An attempt has been made to simulate the L_{556} state of the brS_{56} photocycle,
which is generated within 10 ns. Because such a long time scale is not easily
accessible for molecular dynamics simulations we resorted to the so-called
annealing technique to model the L_{556} intermediate. Simulated
annealing constitutes a systematic technique that has both a solid theoretical
foundation (van Laarhoven and Aarts, 1987) and a broad range of successful
applications, in particular, in the area of biomolecular structure refinement
(Br"unger, 1991). Our annealing procedure has encompassed the slow cooling
protocol consisting of two steps. First, starting at a temperature of 500
K the protein was cooled down to 300 K through coupling the protein motion
to a heat bath by means of the t-coupling method (friction coefficient = 0.1
fs$^{-1}$) (Berendsen et al., 1984). Every 100 fs the temperature of the heat bath
was lowered by 10 K until a final temperature of 300 K was reached. The
same procedure was then repeated starting from 400 K. The resulting structure
was finally equilibrated for 10 ps at 300 K.

**Simulating retinal analogues**

Retinal modifications were performed on the equilibrated structure of brS_{56}.
The modifications were accomplished with the PATCH facility of X-PLOR;
initial coordinates were assigned using the positions of replaced atoms and
using the HBUILD command to generate coordinates for explicitly added
hydrogens. For all simulations, including retinal analogues, we carried out
an energy minimization after the modifications, followed by 5 ps of equi-
libration and 5 ps of dynamics.

**RESULTS**

**Construction of brS_{548}**

The refined simulated model of brS_{56} reported in Humphrey et al. (1994), containing all-trans retinal, was used as a starting
point. In simulation A1, the isomerization from all-
trans,15-anti to 13-cis,15-syn retinal was enforced as de-
scribed in Materials and Methods. Fig. 2 presents the dihedral angles of the C_{13}-C_{14} and C_{15}-N bonds during the thermal
isomerization. It can be seen that the isomerization is com-
pleted after $\sim$2 ps. The results also show that the rotations

![FIGURE 2 Variation of the dihedral angles of the C_{13}-C_{14} and C_{15}-N bonds of retinal during simulation A1.](image)

around the C_{13}-C_{14} and C_{15}-N bonds are strongly correlated
with each other; i.e., the sum of the dihedral angles of the C_{13}-C_{14} and
C_{15}-N bonds tends to be zero, corresponding to a bicycle pedal motion (Warshel, 1976). Such motion had been suggested for the transition from brS_{56} to brS_{56} in Or-
landi and Schulten (1979) on account of a quantum chemical prediction of a low (22 kcal/mole) barrier.

The thermal isomerization described was followed by a
50-ps equilibration (simulation A2). The resulting structure,
corresponding to brS_{549}, is depicted in Figs. 3-5. Fig. 3 provides
a view of the protein backbone, water, and retinal. The
Schiff base region of the proposed structure is shown in Fig. 4,
along with a stereo view in Fig. 5. These figures show the
relative positions of active groups in the binding site, as well
as the suggested positions and orientations of water mole-
cules in this region. Hydrogen bonds are shown as dashed
lines together with the associated distances from hydrogen to
hydrogen bond acceptor, to judge the strength of the respec-
tive bonds. The hydrogen bonds, near the Schiff base nitrogen,
form a network involving five proximate and two distant
water molecules and protein residues Asp-85, Asp-212, Arg-
82, Tyr-185, and Tyr-57. The oxygens of Asp-85 are located at
an average distance of 5.6 Å and 5.0 Å from the Schiff base
hydrogen, whereas both oxygens of Asp-212 are at a distance of
$\sim$4.2 Å (Fig. 6 a). Two water molecules connect the Schiff
base NH to the carboxylate of Asp-85. Asp-212 is strongly
hydrogen bonded to Arg-82, Tyr-57 (through a water mole-
cule), Tyr-185, and Trp-86 (not shown) and to a water mole-
cule. The picture is similar to the binding site of the simu-
lated model of brS_{56} (Humphrey et al., 1994) in which Asp-
212 is hydrogen bonded to various protein residues in con-
trast to Asp-85, which is connected only to Arg-82. Tyr-57,
besides being connected to Asp-212 through a water mole-
cule, is connected as well to Asp-85 and Arg-82 through
other water molecules.

Fig. 6 presents some dynamical properties of the suggested
brS_{548} model. Fig. 6 a shows the distances of the carboxylate
oxygens of residues Asp-85 and Asp-212 to the Schiff base
proton during the last 5 ps of equilibration. Asp-212 is seen
to remain rather immobile, with both oxygens maintaining a distance of \( \sim 4.2 \, \text{Å} \). Asp-85 exhibits larger fluctuations, but its motion is still relatively small, with distance values of 4.6 and 5.6 Å for the two oxygens. The relative immobility of residues Asp-85 and Asp-212 provides a good indication of the stability of retinal’s counterion complex in the simulated model.

Similar to the bR<sub>588</sub> model (Humphrey et al., 1994), the bR<sub>548</sub> model suggested here exhibits a distinctive twist of retinal’s backbone. The overall twist of 55° is a result of torsions around retinal’s single bonds. The largest twist of 30° is observed around the C<sub>12</sub>-C<sub>13</sub> bond. Fig. 6b presents the fluctuations of the dihedral angles of the C<sub>6</sub>-C<sub>7</sub>, C<sub>10</sub>-C<sub>11</sub>, and C<sub>14</sub>-C<sub>15</sub> bonds during the last 5 ps of simulation A2. It should be noticed that fluctuations increase going toward the β-ionone ring. Fluctuations of the C<sub>14</sub>-C<sub>15</sub> torsion are very small; this bond stays essentially planar. The C<sub>5</sub>-C<sub>7</sub> torsion angle, which determines the rotation of the β-ionone ring relative to the retinal plane, is seen to fluctuate in a wide range of angles, deviating up to 60° from the planar conformation, around an average value of 20°.

An interesting difference between bR<sub>588</sub> and bR<sub>548</sub> emerged in the vibrational spectra observed by Fourier transform infrared (Roepe et al., 1988), Fourier transform Raman (Sawatzki et al., 1990), and resonance Raman (Smith et al., 1987; Noguchi et al., 1990) spectroscopy. bR<sub>548</sub> exhibits stronger intensities of the C<sub>14</sub>-H hydrogen-out-of-plane modes than bR<sub>588</sub>. This difference had been attributed to a twist of the C<sub>14</sub>-C<sub>15</sub> bond in bR<sub>548</sub>. Our results do not support this explanation. The C<sub>14</sub>–C<sub>15</sub> bond is found to be planar. However, we find a strong steric interaction between the hydrogen on C<sub>14</sub> and the two hydrogens on C<sub>16</sub> of the Lys-216 side chain (see Fig. 11).

Additional differences between bR<sub>588</sub> and bR<sub>548</sub> have been observed in solid-state nuclear magnetic resonance (NMR) spectroscopy (Smith et al., 1989). First, the chemical shifts of <sup>13</sup>C<sub>12</sub> and <sup>13</sup>C<sub>15</sub> in the case of bR<sub>588</sub> is further upfield than in the case of bR<sub>548</sub>. Second, the opposite holds for <sup>13</sup>C<sub>14</sub>, which exhibits an upfield chemical shift for bR<sub>548</sub> relative to
bR$_{568}$. The first difference has been explained by a stronger interaction between the Schiff base and the associated counterions. This explanation is corroborated by our simulations presented here and in Humphrey et al., (1994), which suggest that in bR$_{548}$ the carboxylate moieties of Asp-85 and Asp-212 are closer by $\sim 1$ Å to the Schiff base as compared with bR$_{568}$. Studies with model compounds indicated as well that stronger electrostatic interactions between the positively charged Schiff base linkage and its counterion induces downfield shifts in the $^{13}$C$_{13}$ NMR (Albeck et al., 1992). It had been suggested that the second difference results from steric interactions between C$_{14}$H and the C$_{14}$H of Lys-216. This explanation is also in harmony with our results reported here and in Humphrey et al. (1994) in that we find for bR$_{548}$ a close proximity (2.8 Å) between C$_{14}$ and C$_{15}$. The upfield shift that was measured by $^{13}$C$_{14}$ NMR is much stronger than that observed in model compounds (Livnah and Sheves, 1993). This difference can originate from a twist around the C$_{14}$-C$_{15}$ bond that the 13-cis,15-syn retinal protonated Schiff base experiences in solution but that is prevented in the protein. A twist around the C$_{14}$-C$_{15}$ bond should weaken the interaction between C$_{14}$H and the C$_{14}$H of Lys-216, thereby reducing the upfield shift effect observed in the $^{13}$C$_{14}$ NMR spectrum.

**Examination of the structure of bR$_{548}$**

Various experimental data concerning the spectral properties of bR reconstituted with different 13-cis-retinal analogues are available (Sheves et al., 1984). In addition, in several artificial pigments (Steinberg et al., 1991) as well as in some mutants of bR, the ratio of all-trans,15-anti and 13-cis,15-syn retinal in light-adapted and dark-adapted bR has been determined. These observations can be used to check the validity of the suggested bR$_{568}$ and bR$_{548}$ structures.

**Simulation of the D85A mutant**

It was observed by resonance raman studies (Rath et al., 1993) that mutation of Asp-85 to alanine significantly lowers the C = N stretching frequency of the protonated Schiff base in bR$_{548}$ from 1642 cm$^{-1}$ to 1616 cm$^{-1}$. This effect is accompanied by an unusually small deuterium effect (6 cm$^{-1}$) on the C = N stretching frequency. Both effects can be attributed to weak hydrogen bonding to the Schiff base NH group prevailing in this mutant of bR$_{548}$. It is currently believed that the C = N stretching frequency is significantly controlled by its coupling with N/H bending, and a weak hydrogen bonding to NH will induce a lower C = N stretching frequency (Aton et al., 1980; Kakitani et al., 1983; Baasov et al., 1987). In native bR, the C = N stretching frequency of both trans and 13-cis isomers is similar, pointing to a similar environment of the N-H bond. However, the resonance raman experiment suggests quite a different environment in D85A mutants of these isomers.

Simulation B1 modeled the effect of an Asp-85 $\rightarrow$ Ala mutation. Residue modification was followed by a 10-ps equilibration of the protein. A detailed picture of the mutant active site is given in Fig. 7. A comparison of Figs. 4 and 7 reveals considerable changes in the active site of the D85A mutant as compared with native bR$_{548}$. Structural changes not only involve the nearest neighbors of Ala-85 but also extend over the whole binding site. The hydrogen-bonding network within the active site is completely modified. It is suggested that the hydrophilic groups surrounding the binding site are no longer bridged together to form an integrated counterion complex of the Schiff base; instead, the hydrogen-bonding network of the mutant consists of a few water molecules that, together with Asp-212, form an isolated hydrogen-bonding cluster. The mutation affects the protein backbone as well. The backbone of helix G is found to be distorted from a standard $\alpha$-helical configuration, with the carbonyl of Asp-212 lacking the usual hydrogen-bonding pattern within the helix. This carbonyl is hydrogen bonded to water molecule W3, which breaks the standard hydrogen bond between the CO group of Asp-212 and the NH group of Lys-216. The most distinctive feature of the simulated D85A mutant model is that there is a very weak hydrogen bonding of the Schiff base NH group to a water molecule. This hydrogen bonding
is weaker than that observed in the model of bR₅₄₈ and provides an explanation for the low C=N stretching frequency and the small deuterium effect observed in the resonance raman studies of bR₅₄₈ mutant D85A. Simulations of the D85A mutant of bR₅₄₈ (not shown) indicate only minor changes in the hydrogen bonding strength of water molecule to the NH moiety for the mutant, in keeping with the resonance raman results that show a moderate shift in the C=N stretching frequency from 1642 cm⁻¹ to 1630 cm⁻¹ (Rath et al., 1993).

**Retinal analogues simulations**

Experiments with modified retinal chromophores have proven to be a powerful tool for obtaining information on the steric interactions of retinal with the surrounding protein environment (for review see Ottolenghi and Sheves, 1989). In the present study we focus on the retinal β-ionone ring modifications, which provide clues for orientation and location of the retinal ring within the binding site. Fig. 8 presents the four retinal analogues considered. Studies of artificial pigments derived from the retinal analogues reveal that introduction of bulky substituents at the ring C₄ position blue-shifted the absorption maximum of both trans and 13-cis isomers of bacteriorhodopsin considerably (Sheves et al., 1984; Steinberg et al., 1991). In contrast, a methyl substitution at the C₂ position do not cause any significant change in the absorption maximum (Sheves and Friedman, unpublished results). Simulations of the artificial pigments derived from the analogues depicted in Fig. 8 and comparison with the spectral shifts observed for the respective pigments should provide a measure of the consistency and accuracy of the simulated bR₅₄₈ model suggested here. Results of these simulations are given in Fig. 9.

**4-Dimethylamino modification**

Simulations C1A and C1B were carried out to determine the effect of substitution of the retinal C₄ position with a dimethylamino group (Fig. 8 a) and resulted in two structures, bRₐ₃ and bRₐ₄. The two structures arise as the C₄ position has two hydrogen atoms and, thus, there are two possibilities for attachment that are inequivalent as a result of the protein chiral discrimination. We define the plane of retinal facing toward the extracellular side as α and that facing to the cytoplasmic side as β. bRₐ₃ corresponds to the retinal analogue substituted at the α-position and bRₐ₄ to substitution at the β-position.

Fig. 9 a summarizes the results of the simulations of bRₐ₃ and bRₐ₄. Shown in this figure are the torsional angles of bonds along the retinal backbone. For the native pigment, retinal is in a 13-cis,15-syn configuration, with the C₆-C₇, C₈-C₉, C₁₀-C₁₁, and C₁₂-C₁₃ bonds slightly twisted, giving rise to the corkscrew configuration of retinal in the simulated bR₅₄₈ model discussed above. The changes of retinal’s conformation in going from bR₅₄₈ to bRₐ₃ are quite noticeable. Substitution of the C₄ α-position by a dimethylamino group leads to a considerable twist of 20° around the C₁₃-C₁₄ double bond (rotation barrier, 20 kcal/mol) and around the C₁₄-C₁₅
The C₄ positions of retinal are achiral; however, after incorporation into the protein binding site, the two enantiomers are different, and chiral discrimination by the protein is possible. Thus, in general, substitution at the C₄ position should result in two different enantiomers that one, indeed, often observed through two spectral maxima of the reconstituted bR (Ottolenghi and Sheves, 1989). When a single spectral maximum is observed, one expects only one enantiomer to reconstitute. When bR is reconstituted with 13-cis,15-syn retinal with a dimethylamino group at the C₄ position, a single pigment absorbing at 460 nm is observed (Sheves et al., 1984). We suggest that this absorption corresponds to the simulated bRₐ₄ Pigment. The large degree of the retinal twist caused by the β-position substitution could explain the blue-shifted 460-nm peak. The considerable twist of the double bonds and overall shift of the retinal molecule observed for bRₐ₄ indicates that binding of the dimethylamino group to the α-position requires a conformational change, too large to allow this retinal analogue to reconstitute with bacteriophorhopsin. The proximity of Trp-138 prevents attachment of the bulky dimethylamino group to the α-position, but this group can be accommodated at the β-position at the price of some twists in retinal.

4-Methyl modification

Simulations C2A and C2B modeled the effect of methyl group substitutions at the C₄ position of retinal (Fig. 8 b) and resulted in the simulated structures bRₐ₄ and bR₂₄. Attachment at the α-position corresponds to bRₐ₄, attachment at the β-position corresponds to bR₂₄. The results of the simulations are summarized in Fig. 9 b. In the case of bR₂₄, the system stays very close to the bR₄₈ conformation. Slight rotations around different bonds essentially compensate each other, leaving the overall geometry and orientation of retinal virtually unchanged. The retinal geometry and orientation in the case of bR₂₄ are quite different from that in bR₄₈. The C₈-C₉ bond rotates by 45° from the original conformation, developing the twist with absolute value of 30°. The C₁₂-C₁₃ bond develops an even larger twist of 50°. It is suggested that these major rotations around two single bonds lead to a retinal conformation that is noticeably more twisted as compared with the native chromophore.

Experimental studies with artificial pigments revealed that substitution of a methyl group at the C₄ position resulted in two pigments, which absorbed at 550 and 460 nm (Ottolenghi and Sheves, 1989). A possible explanation for the two observed pigments is that each pigment consists of a different enantiomer. The fact that bR₂₄ contains very large rotations about retinal’s single bonds, particularly about the C₈-C₉ and C₁₂-C₁₃ bond, suggests that the respective enantiomer corresponds to the strongly blue-shifted 460-nm peak. The simulated bR₂₄ model shows very little conformational change from the native pigment model and suggests that the corresponding enantiomer produces the 550-nm peak. The close proximity of Trp-138 to the α-position of the C₄ (3.4 Å from the C₈₁ atom of Trp-138 to the α-hydrogen

![Comparison of dihedral angles along retinal’s backbone for the retinal analogues shown in Fig. 8: (a) 4-dimethylamino analogue; (b) 4-methyl analogue; (c) 2-methyl analogue; and (d) cleaved-ring analogue.](image-url)
of \( C_4 \) could account for this effect, namely, that substitution at the \( \alpha \)-position results in a steric hindrance that causes a twist in retinal, whereas substitution at the \( \beta \)-position does not lead to strong steric interactions.

**2-Methyl modification**

Methyl group substitutions at the \( C_2 \) position of retinal resulted in the simulated models \( bR_{C3A} \) and \( bR_{C3B} \) (Fig. 8 c). As for the previous analogues, \( bR_{C3A} \) corresponds to an attachment at the \( \alpha \)-position and \( bR_{C3B} \) to an attachment at the \( \beta \)-position. Fig. 9 c summarizes the results of the respective simulations. No significant changes in the dihedral angles of retinal’s backbone are found in this case, which is consistent with the observation that the spectra are not affected in these analogues. \( bR_{C3B} \), in particular, has a retinal conformation essentially identical with that of the simulated \( bR_{548} \) model.

In the case of \( bR_{C3A} \), the only significant change is an additional rotation of the \( C_{12} - C_{13} \) single bond by 27° relative to its twist in the native pigment. However, it is proposed that this additional twist is offset by smaller rotations around the \( C_5 - C_7, C_8 - C_9, \) and \( C_{10} - C_{11} \) single bonds in the opposite direction. Thus, the overall retinal conformation in the case of \( bR_{C3A} \) also stays close to that of \( bR_{548} \).

**Ring cleavage modification**

Simulations were carried out to model the effect of cleaving the ring between the 3- and 4-carbons and between the 1- and 6-carbons, resulting in structure \( bR_{C4} \) shown in Fig. 8 d. The resulting dihedral angles of retinal’s backbone are summarized in Fig. 9 d. The retinal chromophore, in the case of \( bR_{C4} \), is found to be essentially unaltered as compared with the simulated native pigment. It is suggested that slight rotations around different single bonds induced by the ring cleavage counterbalance each other. The overall retinal shape stays practically unchanged. Results of our simulations are in keeping with a slightly blue-shifted absorption maximum of 535 nm observed for the pigment.

**Simulation of the \( bR_{548} \) photocycle**

**Primary photoisomerization**

We simulated the initial photoisomerization step of the \( bR_{548} \) photocycle by instantaneously changing the torsional potential of the \( C_{13} - C_{14} \) bond as described in Materials and Methods. We carried out three simulations, simulations D1 and D3 with a torsional barrier of 10 kcal/mol for the \( C_{13} - C_{14} \) single bond and simulation D2 with a barrier of 2 kcal/mol for this bond. All simulations were started from the 13-cis,15-syn configuration of retinal and lasted 1 ps. The resulting simulated structures were both equilibrated for 10 ps using the torsional potentials for retinal in the ground state. Simulations D1 and D2 were carried out at 300 K, whereas simulation D3 was carried out at 77 K.

Fig. 10 a presents the time dependence of the dihedral angles of the \( C_{13} - C_{14}, C_{14} - C_{15}, \) and \( C_{15} - N \) bonds during simulation D1. One observes that the whole isomerization process can be divided into two stages. During the first stage, which lasts \( \sim 0.6 \) ps, retinal isomerizes around the \( C_{13} - C_{14} \) bond by \( \sim 150^\circ \). The simulated \( bR \) structure produced after 0.6 ps of the photoisomerization is ascribed to a J intermediate. The strain in the binding site of retinal, caused by the photoisomerization, is further relaxed during the next picosecond of the molecular dynamics simulations when retinal rotates...
by another 30° around the C\textsubscript{13}-C\textsubscript{14} bond. The simulated structure obtained after this relaxation is assigned to a K_{590} intermediate. In this model the retinal protonated Schiff base proton points up towards the cytoplasmic side of a membrane, with the Schiff base hydrogen forming a hydrogen bond with a water molecule on the cytoplasmic site of the Schiff base linkage. The occurrence of two different kinetics for the dihedral dynamics in our simulations appears in agreement with the observation of an early J intermediate and a later K intermediate in a 13-cis cycle (Zinth et al., 1988).

Fig. 10 \textit{b} presents the time dependence of the dihedral angles for simulation D2. No essential difference is observed between the simulations D1 and D2 for the behavior of the dihedral angles of the C\textsubscript{13}-C\textsubscript{14}, C\textsubscript{14}-C\textsubscript{15}, and C\textsubscript{15}-N bonds. Even though the rotation around the C\textsubscript{14}-C\textsubscript{15} bond in the case of simulation D2 has only a 2 kcal/mol barrier, no corotation around this bond is observed. Apparently, the corotation is prevented through steric hindrances due to the syn configuration of the C\textsubscript{15}-N bond. This behavior is distinctly different from that found in the simulated bR_{586} photocycle in which a corotation around the C\textsubscript{14}-C\textsubscript{15} bond is possible and arises, in fact, in the case of low respective barriers (Nonnella et al., 1991; Zhou et al., 1993; unpublished results).

\textit{J} intermediate

The simulated structure obtained after 0.6 ps of the photoisomerization process is defined here as that of the \textit{J} intermediate. In our simulations this intermediate is unstable and converts to a new intermediate (see below) as a result of torsions of retinal’s double bonds and a lack of hydrogen bonding stabilizing the momentary position of the protonated Schiff base, structural features that can be discerned from Fig. 11. The dihedral angles of retinal’s backbone in the simulated model are shown in Fig. 12. This figure suggests that the C\textsubscript{13}-C\textsubscript{14} bond is twisted by 25° and the C\textsubscript{15}-N bond is twisted by 45°; these strong torsions explain the lack of stability as well as the resonance raman spectra that suggest that the retinal chromophore is twisted (Noguchi et al., 1990).

\textit{K}_{590} intermediate

Our simulations suggest (Fig. 10) that, in ~1.2 ps after photoisomerization, retinal’s dihedral angles assume essentially stationary values corresponding to an intermediate with a retinal all-trans,15-syn geometry that is stable on the time scale (20 ps) of the simulation. We identify this state with the K_{590} intermediate that is observed experimentally within 3 ps after light absorption (Zinth et al., 1988). The proposed model of the intermediate is presented in Fig. 11. The Schiff base NH group points to the cytoplasmic side and engages in hydrogen bonding with a nearby water. Figs. 11 and 12 show that the backbone of retinal in the K_{590} state is torsionally more relaxed than the J intermediate. The suggested geometry is consistent with the blue shift of the spectrum of K_{590} with respect to the spectrum of J. The absorption maximum, however, is more red-shifted than the spectrum of the in initial bR_{548}, a property that can be attributed to the larger separation between the protonated Schiff base and the negatively charged Asp-85 and Asp-212 residues (see Fig. 11). The torsional angles given in Fig. 12 indicate that K_{590} in the vicinity of the bonds C\textsubscript{6}-C\textsubscript{9} and C\textsubscript{12}-C\textsubscript{13} is considerably less planar than bR_{548}; this result is consistent with the observation of strong hydrogen-out-of-plane bands for K_{590} (in a 13-cis photocycle of an artificial pigment derived from a 13-demethylretinal) as reported in Noguchi et al. (1990).
Application of the annealing scheme described in Materials and Methods leads to convergence to a third structure that is presented in Figs. 11 and 12. Assuming that the annealing scheme effectively bridges a longer time scale, we interpret this structure as the L_{610} intermediate that is observed to form within 5 ns after light absorption. The dihedral angles proposed by our simulations for L_{610} in Fig. 12 show that the retinal chromophore at the L_{610} stage is essentially planar, a feature that contributes to a red-shifted spectrum relative to the spectrum of K_{550}. We note that the spectrum of this late nanosecond intermediate of the bR_{548} cycle is significantly red-shifted relative to the spectrum of the late microsecond intermediate of the bR_{568} photocycle, i.e., of L_{550}. In addition to the larger torsions observed in the molecular dynamics simulations of the L_{550} structure (unpublished results), the main difference between the two intermediates is the direction in which the N-H bond points. In L_{610} of the bR_{548} photocycle, the N-H bond points to the cytoplasmic side, which results in a larger distance between the protonated Schiff base and Asp-85 and Asp-212. Our results suggest that this large distance is the major cause for the red shift in L_{610}.

Fig. 11 shows clearly that the simulated L_{548} intermediate features a protonated Schiff base with its proton pointing towards the cytoplasmic side, participating in a hydrogen bond network on this side, and not engaging in any hydrogen bonding to Asp-85 or Asp-212. This result provides an explanation for the fact that an unprotonated intermediate, i.e., an M intermediate, is not formed in the bR_{548} photocycle; instead of passing through an unprotonated intermediate, the bR_{548} cycle decays back from L_{610} to bR_{548} with a halftime of 40 ms (Kalisky et al., 1977; Hofrichter et al., 1989). We note that an M intermediate was observed in the photocycle of bR_{548} measured at pH values higher than 9 (Drachev et al., 1993) and in an artificial pigment derived from 13-demethyl14-F retinal (Steinberg et al., submitted for publication). It is possible that both cases induce a different isomerization or have different proton transfer pathways.

**Low temperature isomerization**

Many experimental studies of the bR_{548} photocycle were conducted at low temperatures (Iwasa et al., 1981; Roepe et al., 1988; Balashov et al., 1991). We have carried out, therefore, a simulation of the bR_{548} photoisomerization at T = 77K. The resulting motion of the dihedral angles during the first 2 ps of the photocycle are presented in Fig. 10 c. Similar to the cycle at T = 300 K, the dihedral angle of the C_{13}-C_{14} bond rotates by 140° within ~0.5 ps. However, the remaining rotation of this bond by 40° requires ~1.5 ps, which is noticeably slower than the relaxation of this angle in the T = 300 K case. As kinetics of the J to K transition is temperature dependent, this transition appears to be thermally activated. Formation of the J intermediate is observed to be temperature independent and can be considered as a purely photochemical process.

**Leakage from 13-cis to the trans cycle**

Bacteriorhodopsin in the dark is found to form a mixture of bR_{548} and bR_{568} in a 1:2 ratio. After exposure to light, bR is found only in the bR_{568} state. This conversion has been attributed to a leakage from the bR_{548} cycle to the bR_{568} cycle, every absorption of a photon by bR_{548} leads with approximately 10% probability to bR_{568} (Korenstein and Hess, 1977). A number of pathways had been proposed for the light adaptation, with various steps of the 13-cis photocycle as possible branching points (Sperling et al., 1977; Kalisky et al., 1977; Sperling et al., 1979; Iwasa et al., 1981; Varo and Bryl, 1988; Bryl et al., 1992). It is conventionally assumed that the leakage occurs during the post-K stages of the bR_{548} cycle through a 15-syn → 15-anti isomerization leading directly to bR_{568}. Here we investigate another leakage pathway that involves a 13-cis,15-syn → all-trans,15-anti photoisomerization. For this purpose we lowered, by a small margin, namely 2.5 kcal/mol, the isomerization barrier of the C_{13}-N bond as described in Materials and Methods. Sampling 10 photoisomerization processes (see Materials and Methods), we found that in 7 cases an isomerization to the J and K states occurs as described above, but in 3 cases a 13-cis,15-syn → all-trans,15-anti isomerization took place. The time evolution of the dihedral angles for 3 sample cases is presented in Fig. 13, in which case c corresponds to a leakage reaction. These findings suggest the possibility that light adaptation of bR involves a 13-cis,15-syn → all-trans,15-anti photoisomerization. It is interesting to observe in Fig. 13 c that the rotation around the C_{13}-N bond occurs essentially after the rotation around the C_{13}-C_{14} bond is nearly completed. We note that similar simulations that were carried out for the all-trans photocycle did not lead to coisomerization of the C≡N bond.

**DISCUSSION AND CONCLUSIONS**

Before the present investigation was undertaken, the retinal isomeric state of bR_{548} was already well known, namely, 13-cis,15-syn (Harbison et al., 1984; Smith et al., 1984; Livnah and Sheves, 1993). One expected that this retinal isomer would fit well into the binding site. However, the detailed conformation of retinal in bR_{548} and its effect on the counterion complex were uncertain. The photocycle of bR_{548} had been studied as well, but it was not understood why bR_{548} does not pump protons. Our simulations allowed us to address these issues. Below we discuss first the properties of the structure of bR_{548} and then its photocycle.

**Structure of bR_{548}**

Bacteriorhodopsin in the bR_{548} state in the dark converts within ~30 min to the bR_{548} state, assuming a 1:2 equilibrium between the bR_{568} and the bR_{548} forms (Ohno et al., 1977). This contrasts with the behavior of retinal in solution in which a trans configuration is more stable (Sheves and
The spectral (NMR, vibrational, electronic) differences between br<sub>568</sub> and br<sub>548</sub> have been mainly ascribed to the twist of the C<sub>14</sub>-C<sub>15</sub> bond (Smith et al., 1987, 1989; Roepe et al., 1988; Sawatzki et al., 1990; Noguchi et al., 1990). However, our simulations suggest another explanation. A torsion around the C<sub>14</sub>-C<sub>15</sub> bond does not arise; rather, the simulations suggest that the differences between br<sub>568</sub> and br<sub>548</sub> are the result of the distance between the protonated Schiff base linkage and Asp-85 and Asp-212 as well as the result of different steric interactions between C<sub>14</sub>-H and C<sub>c</sub>-H<sub>2</sub> of Lys-216. The simulations also relate successfully to the outcomes of experiments in which bacteriorhodopsin is reconstituted with various analogues of 13-cis retinal (Ottolenghi and Sheves, 1989). We conclude from this that the binding site of the ring moiety, possibly also a larger part of the site, is reproduced in our simulations with an accuracy of ∼1 Å.

A particularly promising source of information on the properties of dark-adapted bacteriorhodopsin are artificial pigments, mutants, and different wildlife forms (Mukohata et al., 1991) that show vastly different ratios of all-trans and 13-cis retinal after dark adaptation. As a first step to study such effects, we have investigated the D85A mutant. Resonance Raman measurements of the C=--N stretching frequency of this mutant indicate a weakened, relative to br<sub>548</sub>, hydrogen bonding of the Schiff base linkage. This is corroborated by the results reported here; as shown in Fig. 7, the mutation changes the hydrogen bonding pattern of the counterion complex such that bonding to the Schiff proton of water molecules becomes very weak. In the case of br<sub>568</sub> the mutant does not show such a large effect either in the observations or in the simulations. These results further support the suggestion that the Schiff base hydrogen in br<sub>568</sub> and br<sub>548</sub> is not hydrogen bonded directly to Asp-85 or Asp-212 but, rather, through water molecules.

**Photocycle of br<sub>548</sub>**

The key finding of our simulations is that the photocycle of br<sub>548</sub> leads to an L<sub>610</sub> intermediate for which the Schiff base proton points towards the cytoplasmic side and loses contact with Asp-85, the normal proton acceptor. An intact hydrogen bonding network between the Schiff base proton and Asp-85 has been found for the simulated L<sub>550</sub> intermediate of the br<sub>548</sub> photocycle, which allows proton transfer from the Schiff base to Asp-85 (unpublished results). However, our simulations suggest that such a network does not exist for the L<sub>610</sub> intermediate after photosomerization of br<sub>548</sub>. This could explain why the photocycle of br<sub>548</sub> lacks an M intermediate.

The simulations of the isomerization process also shed light on the question of how the photocycle of br<sub>548</sub> leaks into the photocycle of br<sub>568</sub> which requires a rotation around the C<sub>15</sub>-N bond. It is observed that, after light absorption, ∼10% of the isomerized br<sub>548</sub> experiences an isomerization around two bonds producing the all-trans,15-syn isomer. The question arises as to how this conversion is realized. Some ex-
Experimental data indicate that conversion from the br548 to br508 photocycle occurs early in the excited state (Korenstein and Hess, 1977; Varo and Bryl, 1988; Balashov et al., 1988; Gergely et al., 1994). Our simulations support the notion that the branching pathway is a 13-cis,15-syn → all-trans,15-anti isomerization during the initial photoexcitation of br548. A possibility of a similar photoisomerization around two double bonds induced by a single photon absorption had been suggested earlier (Warshel, 1976) and has recently been demonstrated experimentally in visual pigment isomerization (Zhu and Liu, 1993).

We like to stress that the present theoretical treatment of br is mainly lacking the quantitative quantum chemical description of the ground and excited state potential surfaces of retinal. We consider simulations that combine the required quantum chemical calculations with a molecular dynamics description as the most important issue for future study. These types of calculations would dramatically improve all facets of the modeling presented in this paper. It is believed (Balashov et al., 1993) that the dark adaptation process in br occurs via corotation around the C13-C14 and the C14-N bonds and is catalyzed via protonation of the Asp-85 residue. Only rigorous quantum chemical calculations in which charges of various protein groups are explicitly included in the electronic Hamiltonian of retinal can be expected to describe reasonably well such phenomena. The same holds for modeling the photoisomerization process, in which case the charged side groups near retinal are expected to govern excited state processes (Kobayashi et al., 1990; Song et al., 1993). Obviously, one wishes to predict quantitatively the spectra of in situ retinal for br508, br548, and photocycle intermediates. By combining molecular dynamics and semi-empirical quantum techniques, some attempts have already been made to describe spectra and photodynamics of in situ retinal (Warshel et al., 1991). The ultimate goal, however, is a combination of ab initio quantum chemistry and molecular dynamics to reproduce consistently and quantitatively observed spectra, spectral widths, and dielectric relaxation times. Quantitative agreement between calculated and observed electronic spectra could provide a convenient means to prove the validity of models for the intermediates in br photocycles.

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