

Acetylcholine Receptor Gating at Extracellular Transmembrane Domain Interface: the Cys-Loop and M2–M3 Linker

Archana Jha, David J. Cadugan, Prasad Purohit, and Anthony Auerbach

Department of Physiology and Biophysics, State University of New York at Buffalo, Buffalo, NY 14214

Acetylcholine receptor channel gating is a propagated conformational cascade that links changes in structure and function at the transmitter binding sites in the extracellular domain (ECD) with those at a “gate” in the transmembrane domain (TMD). We used Φ -value analysis to probe the relative timing of the gating motions of α -subunit residues located near the ECD–TMD interface. Mutation of four of the seven amino acids in the M2–M3 linker (which connects the pore-lining M2 helix with the M3 helix), including three of the four residues in the core of the linker, changed the diliganded gating equilibrium constant (K_{eq}) by up to 10,000-fold ($P272 > I274 > A270 > G275$). The average Φ -value for the whole linker was ~ 0.64 . One interpretation of this result is that the gating motions of the M2–M3 linker are approximately synchronous with those of much of M2 (~ 0.64), but occur after those of the transmitter binding site region (~ 0.93) and loops 2 and 7 (~ 0.77). We also examined mutants of six cys-loop residues (V132, T133, H134, F135, P136, and F137). Mutation of V132, H134, and F135 changed K_{eq} by 2800-, 10-, and 18-fold, respectively, and with an average Φ -value of 0.74, similar to those of other cys-loop residues. Even though V132 and I274 are close, the energetic coupling between I and V mutants of these positions was small (≤ 0.51 kcal mol⁻¹). The M2–M3 linker appears to be the key moving part that couples gating motions at the base of the ECD with those in TMD. These interactions are distributed along an ~ 16 -Å border and involve about a dozen residues.

INTRODUCTION

The acetylcholine receptor (AChR) is a large, five-subunit membrane protein that isomerizes between C(losed)- and O(pen)-channel conformations (Edelstein and Changeux, 1998; Unwin, 2000; Karlin, 2002; Lester et al., 2004; Sine and Engel, 2006). This “gating” reaction, which couples a change in structure (and affinity) at two transmitter binding sites with a change in structure (and conductance) at a narrow region of pore, involves the motions of hundreds of residues spread throughout this ~ 300 -kD protein. The diliganded C \leftrightarrow O transition can occur in ~ 1 μ s (Maconochie et al., 1995; Chakrapani and Auerbach, 2005), a time scale that is shorter than the resolution of the patch clamp but long compared with many different kinds of protein motion (Jardetzky, 1996; Karplus and Kuriyan, 2005; Boehr et al., 2006). To understand this complex and relatively slow chemical reaction it is useful to illuminate the dynamics of the short-lived intermediate states that link stable C with stable O. We will refer to this ensemble of brief intermediates as the transition region (TR) of the reaction.

Although the structure or function of the TR cannot be probed directly, it is possible to infer its characteristics from the rate constants of the diliganded gating reaction. A rate-equilibrium free energy relationship (REFER) is a log-log plot of the opening rate constant vs. the equilibrium constant (opening rate constant/closing rate constant) for a mutational series. This relationship is often approximately linear with a slope (Φ) that is a

fraction between 0 and 1. Φ values for diliganded gating have been measured for >100 different AChR residues, and the emergent map, although incomplete, shows a distinct pattern. There is, approximately, a coarse-grained and decreasing gradient in Φ between the transmitter binding site and gate (Grosman et al., 2000b; Purohit et al., 2007).

Although there are likely to be many possible conformational trajectories through the TR ensemble, a simple model that assumes a single pathway through a one-dimensional, sequential series of intermediate-state transitions accounts surprisingly well for the map of Φ (Zhou et al., 2005). With this model, Φ values provide the relative timing of the gating motion of individual residues, with the limits 1 and 0 marking “early” and “late” (Auerbach, 2007). Alternative interpretations have been proposed, namely that fractional Φ values reflect fractional structures (C-like vs. O-like) of individual side chains at the TR, or heterogeneous structures in the TR arising from multiple pathways. With the interpretation that Φ provides temporal information, the clustering of Φ values in the AChR implies that there are about a dozen nanometer-sized domains comprising residues that exhibit temporally correlated gating motions (“ Φ blocks”), and that these domains move back and forth (on ~ 50 -ns time scales) to link the structural

Correspondence to Anthony Auerbach: auerbach@buffalo.edu

Abbreviations used in this paper: AChR, acetylcholine receptor; ECD, extracellular domain; REFER, rate-equilibrium free energy relationship; TMD, transmembrane domain; TR, transition region; wt, wild type.

TABLE I
M2–M3 Linker Sequence Alignment of Mouse AChR Subunits

Subunit							
$\alpha 1$	A	V	P	L	I	G	K
$\alpha 2$	V	I	P	L	I	G	E
$\alpha 3$	V	I	P	L	I	G	E
$\alpha 4$	V	I	P	L	I	G	E
$\alpha 6$	V	I	P	L	V	G	E
$\alpha 7$	S	V	P	L	I	A	Q
$\alpha 9$	–	V	P	L	I	G	K
$\alpha 10$	–	V	P	L	I	G	K
$\beta 1$	A	V	P	L	I	I	I
$\beta 2$	D	V	P	L	V	G	K
$\beta 3$	V	I	P	L	I	G	E
$\beta 4$	D	I	P	L	I	G	K
δ	A	I	P	L	V	G	K
γ	A	V	P	L	I	S	K
ϵ	S	V	P	L	L	G	R

In the $\alpha 1$ subunit, the M2–M3 linker is A270–K276.

changes at the transmitter binding sites that regulate affinity for agonists with those at a distant “gate” that regulate ion flux. We use this conceptual framework, namely that of brownian movement along a linear sequence of intermediate steps (a “brownian conformational wave”; Auerbach, 2005) for understanding the detailed mechanism of AChR gating.

TR structural transitions occurring near the interface of the ECD and the TMD (mainly in the α subunits) are components of this “wave.” Unwin (2005) proposed that there is an isomerization of backbone bonds in the M2–M3 linker at residue G275, and Lummis et al. (2005) suggested that in the related 5HT_{3A} receptor an isomerization of a proline in this linker is essential for efficient gating. Lee and Sine (2005) provided evidence that the perturbation of a salt bridge at this interface, between R209 in the “preM1” segment and E45 in loop 2, is an important dynamic gating event. Based on cryo-EM analyses of unliganded-closed *Torpedo* AChRs, Unwin et al. (2002) hypothesized that in diliganded C \leftrightarrow O gating the α subunits undergo a rotation of the inner β sheets of the ECD, which then perturbs loop 2, the top of the M2 (pore-lining) helix, and the gate. Here, we describe the results of Φ -value analyses of residues in the M2–M3 linker and the cys-loop (loop 7), both of which are located at the ECD–TMD interface region. In two companion papers, we describe related observations regarding the effects of mutations of residues in the preM1 ($\beta 10$ –M1) linker, the site of the putative salt bridge (Purohit and Auerbach, 2007a), and residues in the β -core of the ECD (Purohit and Auerbach, 2007b).

The 4-Å structure of *Torpedo* AChRs (Unwin, 2005) shows that in the α subunits the linker between the M2 and M3 helices consists of four amino acids: P272–L273–I274–G275. Mutation of the homologous proline in 5HT_{3A} receptors (to six different side chains) yields nonfunc-

tional channels that are trafficked normally to the membrane and have wild-type antagonist-binding properties (Lummis et al., 2005), but the substitutions of unnatural amino acids that can undergo a backbone cis–trans isomerization are functional. This suggests that a change in the angle of the M2–M3 backbone is necessary for efficient gating. In AChRs, a full cis–trans isomerization of P272 is not essential for gating because a serine substitution only modestly alters the gating equilibrium constant (K_{eq} ; Lee and Sine, 2005). Nonetheless, these same studies show that AChR residue P272 moves in the TR (because a glycine substitution reduces K_{eq} by ~ 150 -fold) and interacts energetically with V46 in loop 2 (by ~ 1.6 kcal/mol).

The flanking regions of the M2–M3 linker, which include residues S269 and A270 in M2 and residues K276 and Y277 in M3, also participate in the gating reaction. Mutation of α S269 increases K_{eq} mainly by increasing the channel opening rate constant (Grosman et al., 2000a; Mitra et al., 2005), and mutation of α Y277 increases K_{eq} mainly by decreasing the channel closing rate constant (Cadugan and Auerbach, 2007). This suggests that the gating motions in the α subunit at the top of M2 occur before those at the top of M3. Below, we describe the consequences of α A270 and α K276 mutations.

The M2–M3 linker has also been shown to participate in diliganded gating of various neuronal AChRs (Campos-Caro et al., 1996; Rovira et al., 1998), glycine receptor $\alpha 1$ homomers (Lynch et al., 1997), and GABA_C receptor $\rho 1$ homomers (Kusama et al., 1994). In chimeric $\alpha 7$ -5HT_{3A} receptors, whole-cell current amplitudes are increased when an Asp side chain is inserted in the flanking M2 region, probably because of an increase in K_{eq} (Castillo et al., 2006). In GABA_A receptors, a charge reversal at K279 (α S266 in AChRs, in M2) decreases K_{eq} , and energetic coupling is apparent between this residue and D57 in loop 2 (Kash et al., 2003). Further evidence of the importance of the M2–M3 linker (plus flanking regions) in gating is that mutation causes human disease (Shiang et al., 1993, 1995; Elmslie et al., 1996; Croxen et al., 1997).

The “cys-loop” ($\beta 6$ – $\beta 7$ linker, or loop 7) is composed of 13 residues bracketed by a disulfide bond (α C128– α C142). In the mouse α -subunit, the cys-loop sequence is: EIIVTHFPFDEQN. Previously, it was shown that mutations of several cys-loop residues (H134, F135, F137, D138, and Q140) increase or decrease K_{eq} , mainly by changing the channel-opening rate (Chakrapani et al., 2004). This Φ -value places the gating motion of the cys-loop before that of M2, and approximately synchronous with that of loop 2. The cys-loop myasthenic mutation V132L reduces the K_d for ACh (by ~ 30 -fold) at one transmitter binding site but has only a small effect on K_{eq} (Shen et al., 2003). Mutation of residue D138 (to K and A) yields nonfunctional AChRs, whereas an E substitution generates only a small (less than fivefold) change in K_{eq} (Chakrapani et al., 2004; Xiu et al., 2005). In other receptors, it has been shown that the fast activation of whole cell currents

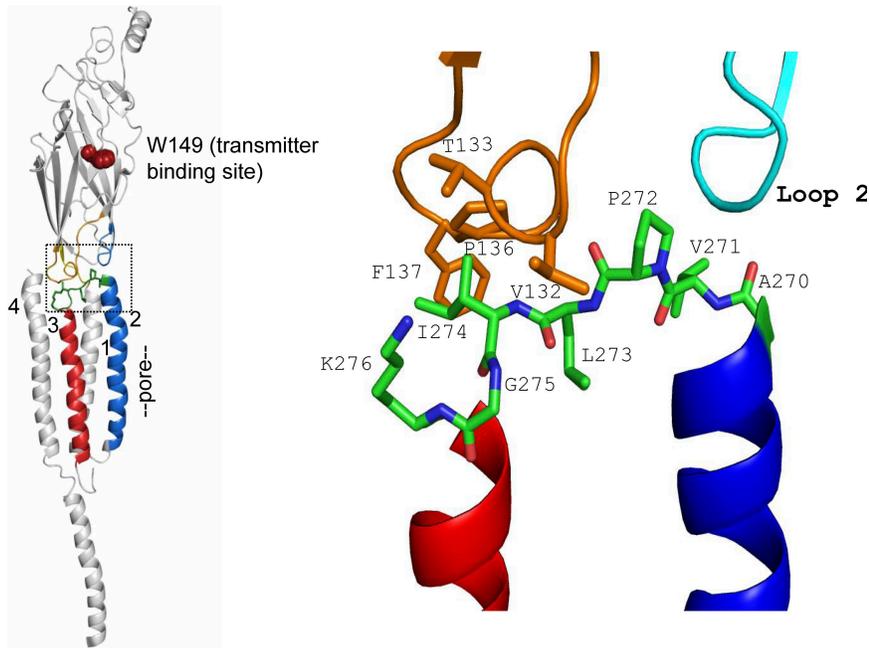


Figure 1. The α_e subunit M2–M3 linker (from 2bg9.pdb; Unwin, 2005). Left, in each AChR subunit the extracellular domain (ECD) is mostly β -sheet and the transmembrane domain (TMD) has four α helices. Residue W149 (brown) marks the transmitter binding site, M2 (blue) lines the pore and is linked to M3 (red) at the ECD/TMD interface (boxed). Right, expanded view of the boxed area. M2–M3 linker residues A270–K276 are colored by element: green, carbon; red, oxygen and blue, nitrogen. The colored domains are: orange, cys-loop; cyan, loop 2. M1, M4, loop9, and the preM1 linker have been removed for clarity. The core sequence of the M2–M3 linker is PLIG (272–275).

from chimeras of $\alpha 7$ (ECD) and glycine (TMD) receptor increases when the $\alpha 7$ cys-loop residues are changed to their glycine receptor counterparts, or when the glycine receptor M2–M3 linker residues are changed to their $\alpha 7$ counterparts (Grutter et al., 2005). An interaction between F135 in the cys-loop and L270/I271 in the M2–M3 linker has been predicted from MD simulations (Cheng et al., 2007). Mutation of D149 in GABA_A receptors decreases the amplitudes of whole cell currents, and the mutation of D148 in glycine receptors increases the response EC_{50} , in both cases by charge-dependent mechanisms (Kash et al., 2003; Schofield et al., 2003).

We have quantified the effects of mutations on the diliganded AChR opening and closing rate constants for every residue in the α -subunit M2–M3 linker (positions 270–276) plus six cys-loop residues. The results add to the map of AChR Φ values and shed light on some of the structural transitions of the TR. We conclude that at the ECD–TMD interface the transfer of energy between adjacent Φ blocks occurs along a broad, ~ 16 -Å boundary and requires the motion of multiple residues.

MATERIALS AND METHODS

Mutagenesis and Expression

Mutants of mouse AChR subunit cDNAs were made by using the QuickChange Site-directed Mutagenesis Kit (Stratagene) and were verified by nucleotide sequencing. Human embryonic kidney fibroblast cells (HEK 293) were transiently transfected using calcium phosphate precipitation. HEK cells were treated with 0.875 mg DNA per 35-mm culture dish in the ratio of 2:1:1:1 (α : β : δ : ϵ) for ~ 16 h. Most electrophysiological recordings were made ~ 24 h later. We quantified the gating rate constants for a total of 37 different mutants, at 11 different amino acid positions in the M2–M3 linker region and cys-loop.

Electrophysiology

Recordings were performed in cell-attached patch configuration at 22°C. The bath and pipette solutions were Dulbecco's phosphate-buffered saline containing (in mM): 137 NaCl, 0.9 CaCl₂, 2.7 KCl, 1.5 KH₂PO₄, 0.5 MgCl₂, and 8.1 Na₂HPO₄ (pH 7.3). Pipettes made from borosilicate capillaries were coated with Sylgard (Dow Corning Corp.). The average pipette resistance was 10 M Ω . The pipette potential was held at +70 mV, which corresponds to a membrane potential of approximately -100 mV. Single-channel currents were recorded using a PC-505 amplifier (Warner Instrument Corp.) with low-pass filtering at 20 kHz. The currents were digitized at a sampling frequency of 50 kHz using a SCB-68 acquisition board (National Instruments Corp.) and QUB software (www.qub.buffalo.edu).

Agonist (acetylcholine or choline) was added to the pipette solution at a concentration that is approximately five times the closed-conformation equilibrium dissociation constant (K_d ; 500 μ M or 20 mM, respectively). The choice of agonist was determined by the effect of the mutation. To estimate the gating rate constants it is necessary to identify clusters of openings that arise from a single AChR (see below) and then to measure the lifetimes of open and shut intervals within those clusters. At high agonist concentration, the ability to satisfy both of these criteria depends on the gating equilibrium constant K_{eq} . More specifically, because the closing rate constant (k_c) is approximately the same for all agonists (Grosman et al., 2000b), the magnitude of K_{eq} depends mainly on the magnitude of the opening rate constant (k_o). In wild-type (wt) AChRs under our experimental conditions, k_o with choline (~ 120 s⁻¹; see below) or ACh ($\sim 48,000$ s⁻¹) is near the lower or upper limit of our estimation capability, respectively. Thus, choline was used to activate constructs in which K_{eq} was larger than in the wt (gain-of-function mutants) and ACh was used to activate constructs in which K_{eq} was smaller than in the wt (loss-of-function mutants). The opening rate constants for un- and monoliganded AChRs are much smaller than that for diliganded AChRs, so even though the concentration of agonist was only five times the K_d , essentially all of the openings were from fully liganded receptors.

Rate Constant Determination

The analyses were done by using QUB software. At 500 μ M ACh or 20 mM choline openings occur in clusters with long gaps between

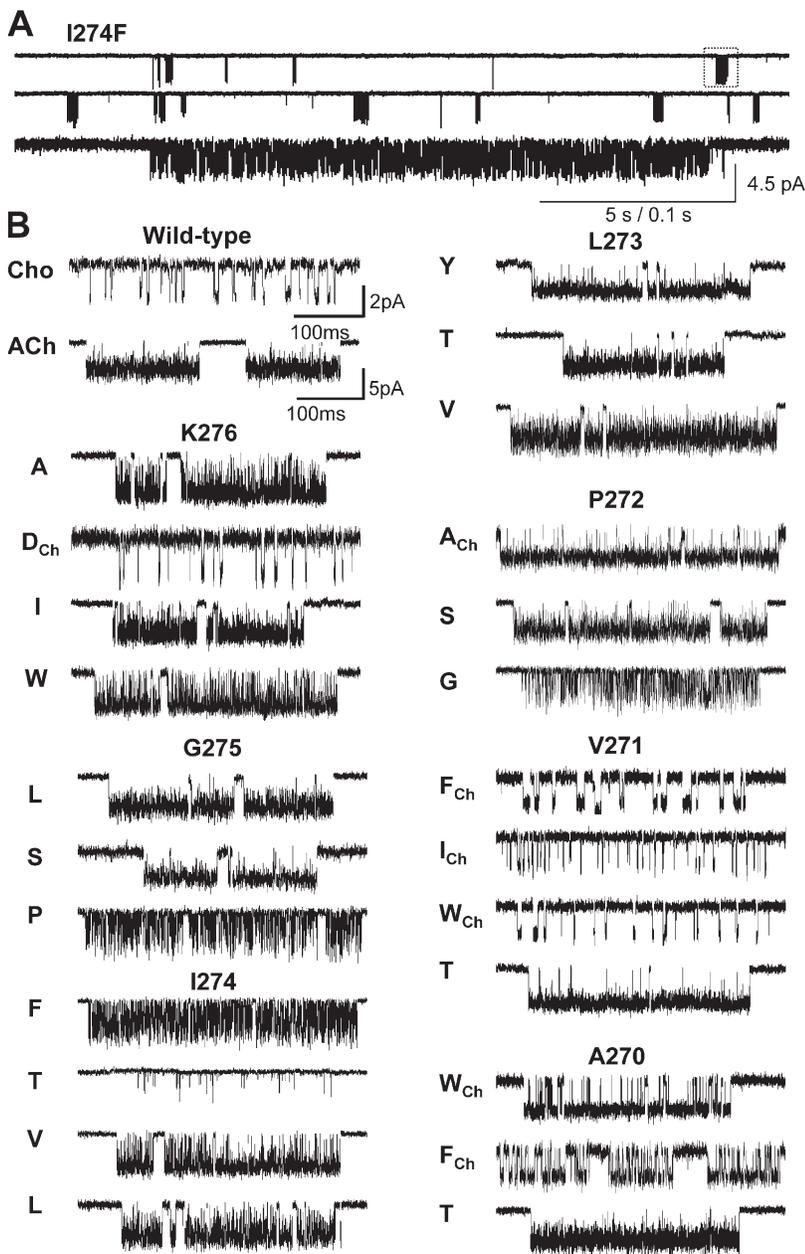


Figure 2. Example single-channel current clusters for different M2-M3 linker constructs. (A) Continuous recording at low time resolution (I274F activated by 500 μ M acetylcholine). Openings are down. The long gaps between clusters of opening reflect times when all channels in the patch are desensitized. Each cluster mainly reflects C \leftrightarrow O gating of a single AChR. Below, one cluster shown on an expanded time scale. (B) Example clusters of different constructs. Currents were activated either by 20 mM choline (indicated by a subscript) or by 500 μ M ACh.

clusters reflecting epochs when all of the AChRs in the patch are “desensitized.” Clusters of individual-channel, diliganded C \leftrightarrow O activity were usually selected by eye. We sometimes tested the eye-selection procedure by using a critical time (t_{crit}) of 50 ms to define a cluster, with equivalent results. Clusters were idealized into noise-free intervals after filtering digitally (12 kHz) by using the segmental k-means algorithm (Qin, 2004) with a C \leftrightarrow O model (starting rate = 100 s^{-1}). The opening and closing rate constants were estimated from the interval durations by using a maximum-interval likelihood algorithm (Qin et al., 1997) after imposing a dead time correction of 25 μ s. Usually, the rate constants were estimated by using a two-state, C \leftrightarrow O model because the log likelihood of the fit did not increase after adding additional C or O states. In \sim 25% of patches a second closed state was connected to the O state to accommodate a relatively rare (<5% of closed intervals within clusters) and short-lived (\sim 2 ms) nonconducting state (Elenes and Auerbach, 2002).

Historical reports from our lab of k_o for wt AChRs activated by choline range from $53 \pm 8 s^{-1}$ (mean \pm SEM; Zhou et al., 1999) to $257 s^{-1}$ (in 140 mM KCl; Grosman and Auerbach, 2000). This degree of variance is larger than typical for other constructs (Table III). More recently, Mitra et al. (2004) report $120 \pm 6 s^{-1}$ (mean \pm SEM) and Corradi et al. (2007) report $95 s^{-1}$ (no error given) for k_o of wt AChRs activated by 20 mM choline. We suspect that some of the variance in the k_o estimate with choline arises from difficulties in identifying single-channel, low open probability clusters, which may be contaminated by sojourns in desensitized states or by openings from overlapping clusters (see above). In support of this hypothesis, we note that mutant constructs showing higher cluster open probabilities have similar variances for k_o regardless of whether ACh or choline was used as the agonist. For example, the k_o estimates in three patches for α A270F activated by choline were 1601, 1846, and 1706 s^{-1} ($1717 \pm 70 s^{-1}$; mean \pm SEM), whereas those for G275P activated by ACh were

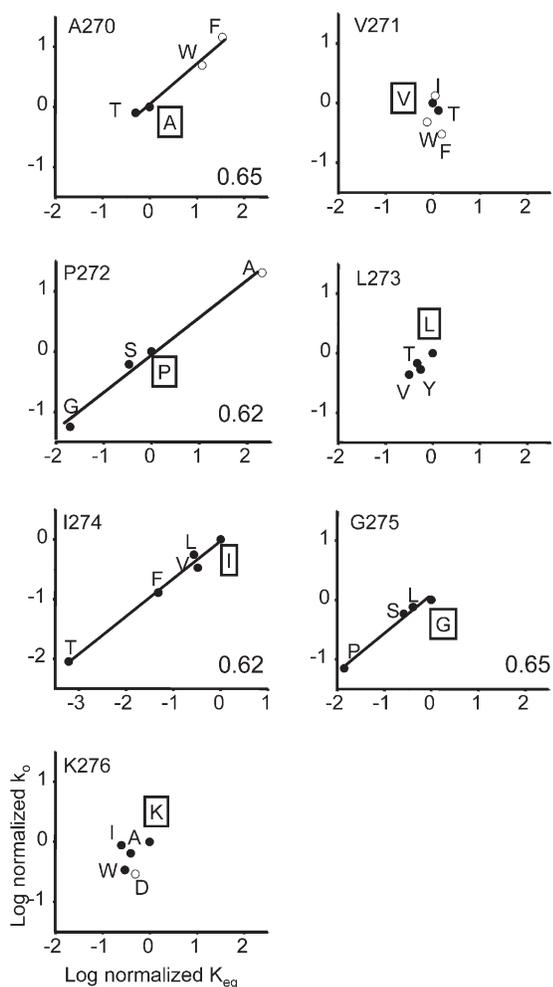


Figure 3. Rate-equilibrium free energy relationships (REFERs) for all seven M2–M3 linker residues. Φ (indicated in the lower right corner of each plot; see Table II) is estimated as slope of a linear fit. The wild-type side chain is boxed. The agonist was either ACh (500 μ M, closed circles) or choline (20 mM, open circles). Both the opening rate and equilibrium constants have been normalized (mutant/wt). Φ values could not be determined for positions V271, L273, and K276 because the change in K_{eq} was too small. The average Φ -value for the M2–M3 linker is ~ 0.64 .

3217, 3756, and 3195 s^{-1} (3380 ± 183). Regardless of the source, the effect of the uncertainty in k_o for the wt/choline construct on the Φ estimate is not major. For instance, normalizing k_o for mutants of α A270, a residue that has a “typical” REFER (150-fold range in K_{eq}), using 53, 120, and 250 s^{-1} as the wt/choline value yields Φ estimates of 0.69 ± 0.05 , 0.65 ± 0.07 , and 0.56 ± 0.08 , respectively.

The K_d for acetylcholine was estimated only for the α I274F construct (Fig. 4). Open and closed interval durations were obtained at three different ACh concentrations (30, 50, and 100 μ M). The two agonist binding sites were assumed to be equivalent and independent (Salamone et al., 1999), and the interval durations at all three concentrations were fitted together by using a $C \leftrightarrow AC \leftrightarrow A_2C \leftrightarrow A_2O$ kinetic model (A = agonist; dead time = 35, 75, and 35 μ s, respectively) that had four rate constants as free parameters: single-site association (k_+ , scaled by [A]), single-site dissociation (k_-), k_o , and k_c .

TABLE II
 Φ Values

Position	Φ (\pm SD)	Fold-change in K_{eq}
A270	0.65 (0.07)	150
V271	–	2
P272	0.62 (0.05)	11850
L273	–	4
I274	0.62 (0.04)	2014
G275	0.65 (0.06)	88
K276	–	5
V132	0.75 (0.08)	2820
T133	–	6
H134	0.71 (0.03)	10
F135	0.75 (0.12)	18
F137	–	4

The fold-change in K_{eq} is for the most extreme constructs at each position. The REFERs are shown in Figs. 3 and 5. Φ values could not be determined accurately for positions where the fold-change in K_{eq} was < 10 -fold.

REFER Analysis

Φ was estimated as the slope of the rate-equilibrium free energy relationship (REFER), which is a plot of $\log k_o$ vs. $\log K_{eq}$ (see Figs. 3 and 5).

The apparent values of k_o and k_c both underestimate the true opening and closing rate constants. k_o is underestimated because the binding sites are not fully saturated. The factor by which k_o is underestimated (F) is a function of the agonist concentration ([A]) and the equilibrium dissociation constant (K_d ; Colquhoun and Hawkes, 1981): $F = ([A]/2K_d)/(1 + ([A]/2K_d))$. In our experiments $[A]/K_d = 5$ for both ACh and choline, hence $F \approx 0.71$ for both agonists. Assuming that the mutations did not alter K_d , the normalized (wt/mut) apparent k_o values for both agonists could therefore combined without correction in the REFER without adding bias to the Φ estimate.

The observed closing rate constant (k_c^{obs}) underestimates the true value because of channel block by the agonist. The equilibrium dissociation constant for this block (K_B) is ~ 12 mM for choline (Purohit and Grosman, 2006) and ~ 2 mM for ACh (Auerbach and Akk, 1998), thus the value $[A]/K_B$ was different for 20 mM choline (~ 1.67) vs. 0.5 mM ACh (~ 0.25). Using the approximation that unblock from the closed channel is negligible (Purohit and Grosman, 2006), we calculate the underestimation factor as $(1 + [A]/K_B)$. Thus, k_c^{obs} underestimates k_c by a factor of ~ 2.67 in the choline experiments but only by 1.25 in the ACh experiments. To eliminate this agonist bias in the K_{eq} estimate and the REFER slope, k_c^{obs} was multiplied by the appropriate factor in order obtain a corrected estimate, k_c^{cor} . Note that because the lifetime of the agonist-blocked channel is shorter than our time resolution, the apparent amplitudes of the choline- vs. ACh-induced currents were reduced approximately by the same factor as the underestimation of k_c .

For residues that showed both an increase and a decrease in K_{eq} upon mutation, the rate constants for AChRs activated by choline or by ACh were combined into the same REFER after normalizing both k_o and K_{eq} by the corresponding wt values. In our REFERs, the wt values used for k_o and K_{eq} were 120 s^{-1} and 0.046 for AChRs activated by choline and 48,000 s^{-1} and 28.2 for AChRs activated by ACh. The normalized rate and equilibrium constants with either agonist fall approximately on the same line. Similarly, Φ values for the δ S268 mutant family were the same regardless of whether the AChRs were activated by choline or acetylthiocholine (Grosman et al., 2000a), and the Φ -value for M4 residue T422 was

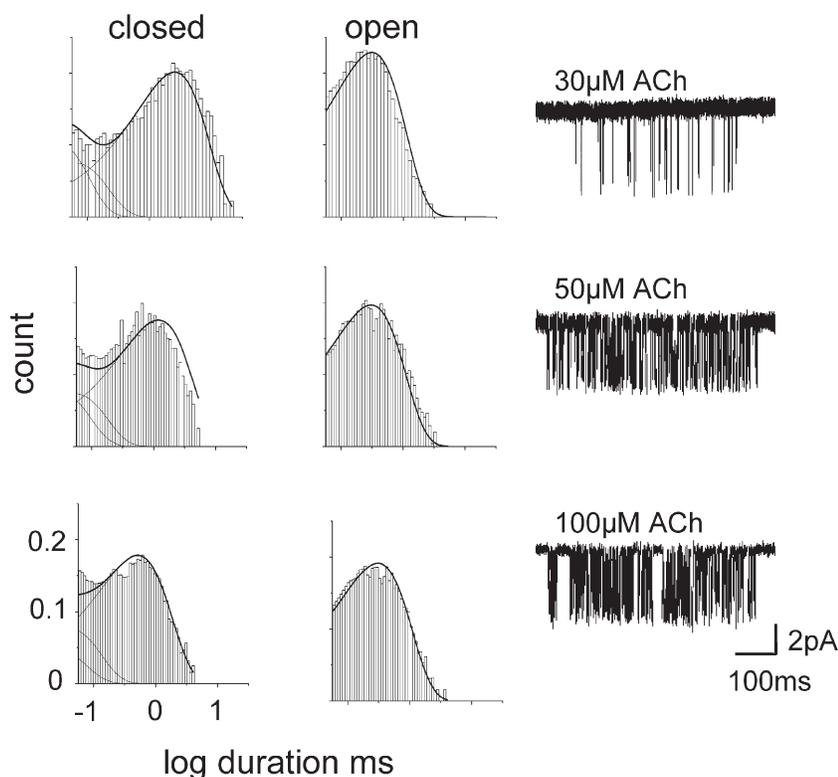


Figure 4. The mutation I274 shows wild-type binding properties. Interval durations were obtained at three different ACh concentrations (30, 50, and 100 μM) and the association and dissociation rate constants were estimated by fitting with a kinetic scheme that assumed two equal and independent transmitter binding steps followed by a single gating step (dead time = 35, 75, and 35 μs , respectively). Left, interval duration histograms and square-root probability density functions (solid lines) calculated from the globally optimized rate constants. Right, an example cluster at each concentration. Total number of events analyzed were 48,776. The optimal rate constants were: k_+ (single-site association) = 141 $\mu\text{M}^{-1}\text{s}^{-1}$, k_- (single-site dissociation) = 14,781 s^{-1} . We calculate K_d (k_-/k_+) = 105 μM . For comparison, one estimate for wild-type AChRs is k_+ = 169 $\mu\text{M}^{-1}\text{s}^{-1}$ and k_- = 16,904 s^{-1} , K_d = 100 μM (Akk et al., 1996). There is no significant effect of this mutation on ACh binding to closed AChRs.

the same with choline (on an δS268N background, 0.54 ± 0.02 ; Mitra et al. 2004) and ACh (Bouzat et al., 2000). These results strongly support the notion that a change in agonist (or an independent background mutation) is like a primary mutation insofar as it can be treated as a simple perturbation in the energy landscape of the gating reaction (rather than as something that creates an entirely new reaction pathway). We document this observation further in Purohit and Auerbach (2007b).

The slope of the REFER was estimated by an unweighted, linear fit in OriginPro 7.0. We did not weight the residuals by using the errors on either axis because it was previously shown that for REFERs of M2 residues, this procedure changed only the standard deviation of the Φ estimate and not the mean (Purohit et al., 2007). For a linear, sequential, bounded reaction mechanism, Φ provides relative temporal information regarding the movement of the perturbed side chain in the gating reaction (1 is “early,” 0 is “late,” and the same is “synchronous”) (Auerbach, 2007).

We used a weighted, k-means cluster analysis algorithm (in QUB) to determine the number (n) of distinct populations of Φ -value and to assign each residue to one of these populations (Fig. 6). Φ values were segregated into populations and a minimum sum squared deviation (SSQ) was computed. We tested values of n up to and including 6, and, in order to find the global minimum SSQ, for each trial we used 50 different random starting assignments. The weight for each Φ -value was the inverse of the SEM obtained from the unweighted linear fit of the REFER.

All structures were displayed by using PYMOL (DeLano Scientific).

Online Supplemental Material

The supplementary animation shows the timing of domain motions in C–O gating. The frame sequence was generated by simulation (QUB) using the model $C \leftrightarrow X_1 \leftrightarrow X_2 \leftrightarrow X_3 \leftrightarrow X_4 \leftrightarrow O$, where C is stable-closed, O is stable-open, and X is a short-lived, nonconducting intermediate state. The rate constants (ms^{-1}) for the individual steps were: $C \rightarrow X_1 = 5.3$, $X_1 \rightarrow X_2 = X_2 \rightarrow X_3 = X_3 \rightarrow X_4 =$

$X_4 \rightarrow O = 6000$, $X_1 \rightarrow C = 2370$, $X_2 \rightarrow X_1 = 5598$, $X_3 \rightarrow X_2 = 4716$, $X_4 \rightarrow X_3 = 12000$, and $O \rightarrow X_4 = 9.09$. Each of the bitmaps were made by using PYMOL (the domain motions are arbitrary) and assembled into a movie by using Videomach.

RESULTS

M2–M3 Linker

In all vertebrate α_1 subunits the residues between the C terminus of M2 and the N terminus of M3 (positions 270–276) are completely conserved: AVPLIGK. Table I gives the sequence alignment for other mouse AChR subunits, and Fig. 1 shows the structure of this M2–M3 linker region in the *Torpedo* α subunit (2bg9.pdb; Unwin, 2005). The core of the M2–M3 linker is the sequence PLIG (residues 272–275). In mouse, in 16 AChR subunit sequences the Pro and Leu are completely conserved and the Ile and Gly residues are present in 12 and 13 sequences, respectively.

Previously, S269 (in M2) and Y277 (in M3) were found to have significantly different Φ values, 0.65 ± 0.06 (Mitra et al., 2005) and 0.34 ± 0.11 (Cadugan and Auerbach, 2007). We attempted to measure Φ for all seven of the intervening residues (Figs. 2 and 3; Table II). At three positions, 11 different mutants exhibited approximately wild-type gating kinetics (V271 \rightarrow F, I, T, and W; L273 \rightarrow T, V, and Y; K276 \rightarrow A, D, I, and W) and for these residues a Φ -value could not be estimated. At the remaining four positions at least one mutation caused a >10 -fold change in K_{eq} and a Φ -value could be estimated. The order of sensitivity, based on the magnitude of the

TABLE III
Rate and Equilibrium Constants

Construct	k_o (s^{-1})	k_c^{obs} (s^{-1})	k_c^{cor} (s^{-1})	K_{eq}	Agonist	Fold-change in K_{eq} (mut/wt)	n
wt ^a	120	834	2583	0.046	Cho	1	
wt ^b	48000	1700		28.2	ACh	1	
A270T	25332 (1645)	2692 (611)	3365 (764)	7.53 (1.8)	ACh	0.27	3
A270W	591 (38)	319 (79)	852 (212)	0.69 (0.18)	Cho	15.0	4
A270F	1717 (70)	346 (36)	924 (96)	1.86 (0.15)	Cho	40.4	3
V271W	58 (5.2)	532 (36)	1420 (97)	0.04 (0.002)	Cho	0.87	3
V271I	160 (13)	988 (72)	2638 (193)	0.06 (0.007)	Cho	1.3	3
V271T	36283 (3899)	972 (71)	1215 (89)	29.9 (5.4)	ACh	1.06	2
V271F	36 (2)	161 (9)	430 (25)	0.08 (0.006)	Cho	1.73	3
P272G	2737 (274)	4821 (141)	6026 (364)	0.45 (0.03)	ACh	0.02	4
P272S	29600 (1751)	3252 (141)	4065 (995)	7.28 (2.2)	ACh	0.26	4
P272A	2431 (42)	83 (131)	222 (46)	10.9 (1.3)	Cho	237	4
L273V	20967 (931)	2649 (146)	3311 (182)	6.3 (1.0)	ACh	0.22	3
L273T	32496 (1345)	2486 (313)	3108 (391)	10.45 (2.0)	ACh	0.37	3
L273Y	25825 (3001)	1791 (586)	2239 (732)	11.53 (4.0)	ACh	0.40	3
I274V	16057 (455)	1770 (278)	2213 (347)	7.25 (1.2)	ACh	0.25	3
I274L	26707 (1291)	3494 (257)	4368 (321)	6.11(0.68)	ACh	0.21	3
I274F	6210 (499)	4608 (448)	5760 (561)	1.07 (0.18)	ACh	0.04	3
I274T	439 (59)	25085 (1093)	31354 (1366)	0.014 (0.002)	ACh	0.0005	5
G275L	36716 (1028)	3189 (293)	3986 (366)	9.21 (0.60)	ACh	0.32	3
G275S	28365 (3174)	3915 (710)	4894 (888)	5.79 (0.52)	ACh	0.21	2
G275P	3380 (183)	8510 (439)	10638 (549)	0.32 (0.03)	ACh	0.01	3
K276D	35 (3)	492 (43)	1314 (116)	0.03 (0.004)	Cho	0.65	2
K276A	31689 (4998)	2778 (219)	3473 (273)	9.12 (0.83)	ACh	0.33	3
K276W	16628 (2353)	1942 (35)	2428 (43)	6.84 (0.88)	ACh	0.24	3
K276I	42316 (4227)	5985 (418)	7481 (522)	5.65 (0.69)	ACh	0.20	3
V132A	11949 (1755)	2658 (192)	3322 (240)	3.59 (0.79)	ACh	0.13	2
V132I	3093 (263)	1579 (270)	1973 (337)	1.57 (0.14)	ACh	0.06	3
V132H	1436 (232)	1860 (201)	2325 (252)	0.62 (0.05)	ACh	0.20	3
V132F	135 (38.5)	10250 (795)	12812 (994)	0.01 (0.004)	ACh	0.0004	2
T133A	16211 (1320)	1940 (143)	2425 (178)	6.68 (0.59)	ACh	0.24	3
T133F	7321 (776)	1364 (117)	1705 (146)	4.3 (0.50)	ACh	0.15	3
T133V	15793 (1971)	2259 (1190)	2824 (1488)	5.59 (3.11)	ACh	0.20	2
T133H	16751	1915	2394	6.99	ACh	0.25	1
T133I	17257 (1666)	2127 (312)	2659 (390)	6.5 (1.61)	ACh	0.23	2
T133S	21966	1658	2072	10.6	ACh	0.37	1
H134S ^c	18474		2506	7.37	ACh	0.26	
H134R	14730 (385)	2496 (172)	3120 (215)	4.79 (0.4)	ACh	0.17	4
H134Q	8884 (659)	2599 (248)	3249 (310)	2.79 (0.3)	ACh	0.1	3
F135T	8978 (1227)	2833 (784)	3541 (980)	2.53 (0.32)	ACh	0.09	3
F135Y	99	669	1786	0.05	Cho	1.08	1
F135L	7080 (794)	3552 (331)	4441 (413)	1.59 (0.33)	ACh	0.06	2
F137L	248 (44)	875 (165)	2336 (441)	0.11 (0.04)	Cho	2.4	3
F137Y	69 (5)	783 (57)	2091 (152)	0.03 (0.004)	Cho	0.66	3

k_o , apparent opening rate constant; k_c^{obs} , apparent closing rate constant; k_c^{cor} , closing rate constant after correction for channel block; n , number of patches. Values are mean (\pm SEM). No rate constants are reported for P136G, A, S, and Y because no currents were detected for these constructs (5 patches each, 10 min/patch).

^aFrom Mitra et al. (2005).

^bFrom Chakrapani and Auerbach (2005).

^cFrom Chakrapani et al. (2004).

largest excursion in K_{eq} for a point mutation (in both α subunits), was P272 (A \rightarrow G) > I274 (I \rightarrow T) > A270 (F \rightarrow T) > G275 (G \rightarrow P) (Table III). At A270 and P272 the mutations could either increase or decrease K_{eq} , whereas

at the other positions the mutations only decreased K_{eq} or had no effect (Fig. 3). In summary, the P I and G of the core, and the A at the terminus of M2, were sensitive to mutation.

Fig. 3 plots the experimentally determined kinetic parameters in the form of rate-equilibrium free energy relationships (REFERs), the slope of which (around $K_{eq} = 1$) is called Φ . The Φ values for all four mutation-sensitive residues ranged from 0.62 ± 0.05 (P272) to 0.65 ± 0.07 (L270) (Table II). Overall, the mean value was 0.64, which is the same as those for many residues in the M2 helix (S269, V259, L258, L257, and F256) (Grosman et al., 2000a; Mitra et al., 2005; Purohit et al., 2007). The Φ -value was approximately independent of the agonist used to activate the protein (see the REFERs for A270 and P272, Fig. 3).

Fig. 4 shows currents elicited by different concentrations of ACh for one M2–M3 linker mutation, I274F. The mutation had a large effect on K_{eq} but essentially no effect on K_d , the equilibrium dissociation constant for ACh binding to closed AChRs.

Cys-loop

In the *Torpedo* AChR structure model, atoms from three cys-loop residues (Fig. 1) contact ($<4 \text{ \AA}$) residue I274 in the M2–M3 linker: V132, T133, and P136. We examined the effects of mutations of these three residues, plus H134, F135, and F137 (Table III).

Fig. 5 A (right) shows a REFER analysis for four mutations of position V132 (I, H, A, and F). All of the mutations decreased K_{eq} (by up to 2,800-fold, for V→F) and the Φ -value was 0.75 ± 0.08 (Table II). Fig. 5 B (right) shows that six mutations of position T133 were examined (V, A, S, H, I, and F) but the net effect on K_{eq} was only approximately sixfold, so no estimate of Φ was obtained. Fig. 5 C (right) and D (right) show the results for H134 (Q, R, and S) and F135 (Y, L, and T). Here, at least one mutation decreased $K_{eq} >10$ -fold, and corresponding Φ values for these two positions were 0.71 ± 0.03 and 0.75 ± 0.12 , respectively. No functional AChRs could be recorded whatsoever for G, A, S, and Y mutations of position P136 (five patches each, ~ 10 min per patch). We do not know if these AChRs fail to express or express and fail to gate. Two mutations of F137 (L and Y) exhibited a less than threefold change in K_{eq} (Table III).

Φ Populations and Coupling

Fig. 6 shows the results of using a statistical test (k-means) to ascertain the number of populations of Φ for residues in loop 2, the cys-loop, the M2–M3 linker, and the M2 helix of the α -subunit, and to assign each residue to one of these populations. The most likely number of populations is two. Residues in loop 2 and the cys-loop belong with the group having a mean Φ of 0.77 ± 0.02 , and those in the M2–M3 linker and M2 group belong to the group having a mean Φ of 0.63 ± 0.02 .

We next measured the coupling energy between I274V in the M2–M3 linker and V132I in the cys-loop.

We chose these mutations because each causes a substantial decrease in K_{eq} (3.9- and 17.9-fold, respectively), and because in 2bg9.pdb the distance between these two residues is $<4 \text{ \AA}$. Combined, these two mutations caused a 161-fold decrease in K_{eq} , whereas independent action predicts that would be only a 69-fold decrease in K_{eq} . This result indicates that there is a modest degree of coupling between the I274 and V132 side chains ($+0.51 \text{ kcal/mol}$).

DISCUSSION

In the α -subunit, about half of the residues in the M2–M3 linker and the cys-loop that have been examined so far substantially change K_{eq} . This indicates that these amino acids experience a change in energy between C and O, which implies a change in structure, which implies a gating movement. Clearly, the ECD–TMD interface is a dynamic zone in the diliganded gating TR. The biggest observed energy changes occur in I274 and P272 in the M2–M3 linker and in V132 in the cys-loop. Note that the cys-loop analysis is not yet complete because four residues have not yet been examined in detail (E129, I130, I131, and E139).

The M2–M3 linker and many M2 residues have approximately the same Φ -value, ~ 0.64 (Grosman et al., 2000a; Mitra et al., 2005; Purohit et al., 2007). This group constitutes the third Φ -block to move in the α -subunit in the channel-opening process. $\Phi \sim 0.93$ for the first block (the region surrounding the transmitter binding sites) and $\Phi \sim 0.77$ for the second block (cys-loop and loop 2). If Φ values provide the relative timing of gating motions, then this pattern suggests that in the TR the motions of the M2–M3 linker and many positions in M2, including residues near equatorial “gate” and the intracellular domain, are approximately synchronous and occur after the motions of the binding sites and loop 2 and the cys-loop. Note that Φ values for M2 residues 260–268 have not yet been reported so we cannot be certain that the M2–M3 linker and the gate motions occur by the movement of a contiguous structural element.

The average Φ -value for V132, H134, and F135 was 0.73, which makes these residues members of the second α -subunit gating block ($\Phi \sim 0.77$; Fig. 6). Additional members of this Φ -block include other cys-loop residues (D138, Q140), residues in loop 2 (E45, V46, N47, Q48), and Y127 on β -strand 6. In the AChR α -subunit, the 0.77 Φ -block is a slab ($\sim 6 \text{ nm}^3$) that is interposed between the transmitter binding site and the top of the TMD (Fig. 7). Because V132 is close to P272 and I274 in the M2–M3 linker, these amino acids may participate in the propagation of the conformational wave between the 0.77 and 0.63 Φ blocks.

There is an abrupt, step decrease in Φ (~ 0.3 units) between G275 in the M2–M3 linker and Y277 in the M3 helix.

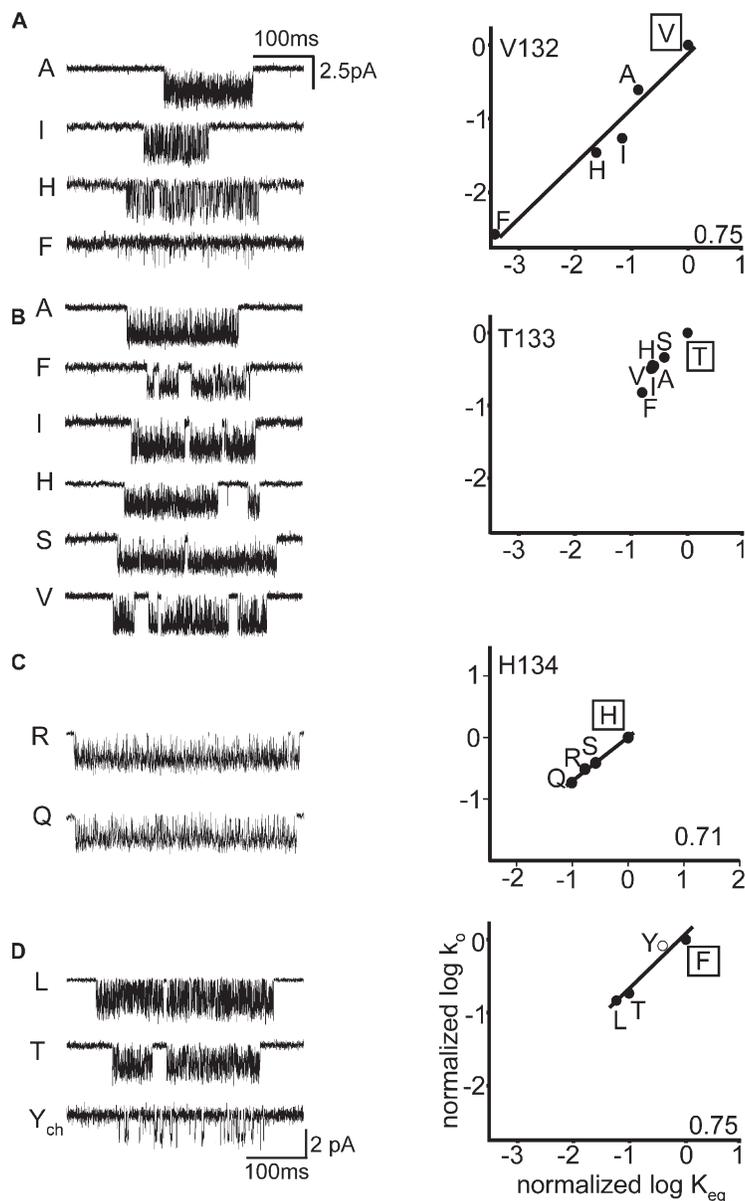


Figure 5. Kinetic analysis of cys-loop residues. (A) Example clusters and REFER for α V132. The Φ -value was 0.75. (B) Example clusters and REFER for α T133. The change in K_{eq} was too small to allow an estimation of Φ . (C) Example clusters and REFER for α H134. The Φ -value was 0.71. (D) Example clusters and REFER for α F135. The Φ -value was 0.75. The wild-type side chain is boxed. The agonist was either ACh (500 μ M, closed circles) or choline (20 mM, open circles).

The intervening residue, K276, is not sensitive to mutation (to A, D, I, or W) and therefore does not experience a change in energy between C and O. In 2bg9.pdb the atoms of residues G275 and Y277 are separated by $<4 \text{ \AA}$, yet G275 and its neighbor I274 are part of the $\Phi = 0.63$ block, whereas Y277 is part of the $\Phi = 0.31$ block. Hence, by definition, G275 and Y277 are at a Φ -block boundary. The breakpoint in Φ at K276 suggests that this residue is a point of flexure about which the M2 and M3 helices move, asynchronously, during gating. A similar pattern of a nonmoving residue interposed between amino acid members of two different Φ blocks (also have a different in Φ of ~ 0.3 units) was previously observed in the M2 helix of the δ and α subunits (Cymes et al., 2002; Mitra et al., 2005). A nonmoving residue at a Φ -block boundary may be a common, but not absolute, feature of the Φ -map.

Insofar as a change in K_{eq} indicates movement and Φ gives relative temporal information, we can place the cys-loop and M2–M3 linker movements into the overall framework for the α -subunit channel-opening process (see Video 1, available at <http://www.jgp.org/cgi/content/full/jgp.200709856/DC1>). First, residues near the transmitter binding sites (loops A, B, and C) move ($\Phi \approx 0.93$; purple; Grosman et al., 2000b; Chakrapani et al., 2003) followed by residues in loop 2 and the cys-loop ($\Phi \approx 0.77$; orange; Chakrapani et al., 2004). The next group to move is the $\Phi \approx 0.63$ block (green), which includes the M2–M3 linker and much of M2 (Grosman et al., 2000a; Mitra et al., 2005; Purohit et al., 2007). This map of Φ values is consistent with the notion that during channel opening the presence of an agonist triggers the gating motions of loops A, B, and C,

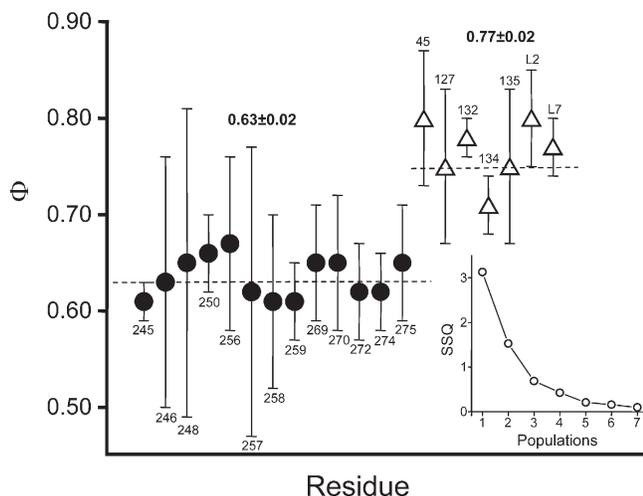


Figure 6. Cluster analysis of Φ values. The Φ values for the indicated α -subunit residues (in the M2 helix, M2–M3 linker, cys-loop, and loop 2) were grouped into populations by using a segmental k-mean algorithm (see Materials and methods). Inset, the sum squared deviation (SSQ) decreases sharply between 1 and 2 populations and gradually thereafter, indicating that the most likely number of Φ populations is two. The mean \pm SEM values for these populations (dashed lines) are shown. Each residue was assigned either to the $\Phi = 0.77$ population (open triangles) or the $\Phi = 0.63$ population (filled circles). The error bars on Φ are \pm SD. Y127 and residues in loop 2 and the cys-loop belong to the 0.77 population, and many residues in M2 and the M2–M3 linker belong to the 0.63 population. L2 and L7 are the mean Φ values for loop 2 and cys-loop residues measured by Chakrapani et al. (2004). Other sources: M2 (Mitra et al., 2005; Purohit et al., 2007), E45 (Purohit and Auerbach, 2007a), and Y127 (Purohit and Auerbach, 2007b).

which trigger those in the cys-loop and loop 2, which trigger those in the M2–M3 linker and in many M2 residues. Subsequent α -subunit motions are three residues near the middle of M2 (Purohit et al., 2007) and in M4 ($\Phi \approx 0.54$; blue; Mitra et al., 2004; Purohit et al., 2007), followed by two M2 equatorial residues plus M3 and preM1 ($\Phi \approx 0.31$; red; Mitra et al., 2005; Cadugan and Auerbach, 2007; Purohit et al., 2007; Purohit and Auerbach, 2007a). We imagine that the Φ -block gating motions are sequential and stochastic, and that the structural transitions of the TR occur as a random walk, with the conformational dynamics of AChR gating characterized by brownian motion of approximately nanometer-sized domains. We estimate that the time scale of the individual Φ -block motions is in the ~ 50 -ns range (Auerbach, 2005).

The results shed some light on the mechanisms that have been proposed for the propagation of the AChR channel opening conformational cascade between ECD and the TMD (the cys-loop/loop 2 and M2–M3 linker/M2 Φ blocks). The structure-motivated suggestion of a steric interaction (“pin-into-socket”) between loop 2 and M2 (V46 \leftrightarrow S269; Miyazawa et al., 2003) is supported by three functional observations: (1) Φ values decrease abruptly between these two residues (0.78 and 0.65,

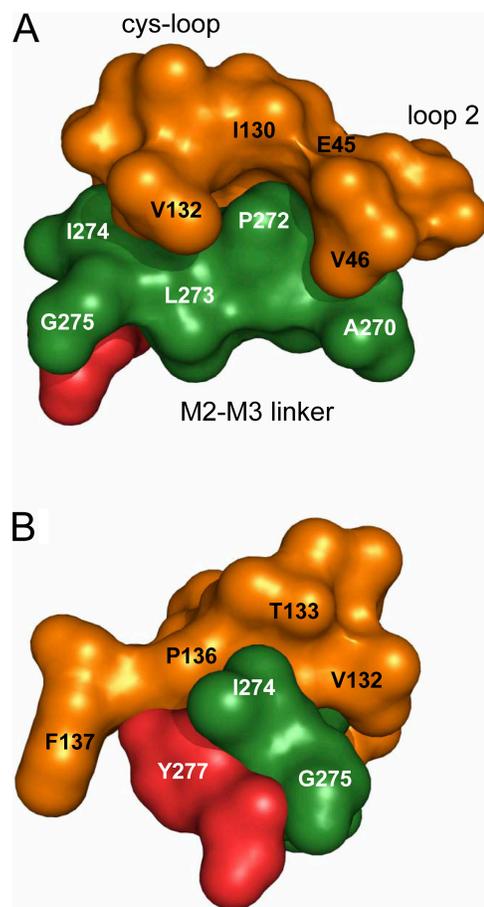


Figure 7. The ECD–TMD interface. The M2–M3 linker residues that were sensitive to mutation are labeled (P272 > I274 > G275 > A270). (A) Side view, from the middle of the ε -subunit. The lumen of the pore is to the right and the membrane bilayer is to the left. Orange, cys-loop (left) and loop 2 (right). Green, M2–M3 linker. Red, residue Y277. The N terminus of the M3 helix is Y277 and the C-terminus of the M2 helix is A270 (see Fig 1). (B) Radial view, from the bilayer.

respectively), (b) the hydrophobicity of the side chains at both positions is correlated with the magnitude of the change in K_{eq} (Chakrapani et al., 2004), and (3) the effects of mutations at these two sites are coupled energetically (Lee and Sine, 2005). In addition, we find that A270 (in M2) is also sensitive to mutation, and, because A270 and V46 are close in the *Torpedo* structure, we propose that this residue is an additional candidate for coupling ECD and TMD gating motions. However, the mutation-induced changes in K_{eq} at P272 or I274 were larger than those at S269 and A270, by more than a factor of 10 (Table III). These sensitive residues are far from the site of the “pin-into-socket” interaction, which suggests that the V46 \leftrightarrow S269/A270 coupling energies are only part of the story and that other forces at other sites also contribute to the total ECD \leftrightarrow TMD interaction energy.

With regard to the hypothesis that the isomerization of an M2–M3 linker proline or glycine is an important

event in gating, the observation that P, S, and L substitutions at G275, and G, S, and A substitutions at P272 yield functional AChRs indicates that a substantial change in the angle of backbone atoms at either of these two positions is not an absolute requirement for the propagation of the opening conformational change between the ECD and TMD. It is therefore unlikely that a full cis-trans isomerization of the M2-M3 linker backbone is the “trigger” for gating in muscle AChRs, although a partial isomerization of these bonds may occur in the C↔O process.

The M2-M3 linker and cys-loop are key structural elements governing the propagation of the channel-opening and -closing conformational changes between the ECD and TMD. Although we still do not know the precise molecular forces that underpin the ECD↔TMD connection, the results indicate that they are spread along a broad, ~16-Å boundary that projects radially from the lumen of the pore (residues S269 and A270) to the lipid bilayer. This interface contains atoms from loop 2 and the cys-loop in the ECD, and (at least) M2, M3, and M4 in the TMD, and includes more than a dozen amino acids (Fig. 7). Our results do not support the notion that a single interaction is the “key” event in the C↔O transition. Rather, we think that at the ECD-TMD hydrophobic interactions, charge/dipolar interactions, and, perhaps, small changes in the angles of backbone bonds all contribute to the energy barrier that separates C from O. Finally, it is important to note that residues in the M2-M3 linker and cys-loop in other subunits, and in nearby segments in all subunits (loop 9, the β10-M1 linker, and M1-M4) may also serve to link ECD-TMD gating motions in the AChR.

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