Initial Coupling of Binding to Gating Mediated by Conserved Residues in the Muscle Nicotinic Receptor

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We examined functional consequences of intrasubunit contacts in the nicotinic receptor α subunit using single channel kinetic analysis, site-directed mutagenesis, and structural modeling. At the periphery of the ACh binding site, our structural model shows that side chains of the conserved residues αK145, αD200, and αY190 converge to form putative electrostatic interactions. Structurally conservative mutations of each residue profoundly impair gating of the receptor channel, primarily by slowing the rate of channel opening. The combined mutations αD200N and αK145Q impair channel gating to the same extent as either single mutation, while αK145E counteracts the impaired gating due to αD200K, further suggesting electrostatic interaction between these residues. Interpreted in light of the crystal structure of acetylcholine binding protein (AChBP) with bound carbamylcholine (CCh), the results suggest in the absence of ACh, αK145 and αD200 form a salt bridge associated with the closed state of the channel. When ACh binds, αY190 moves toward the center of the binding cleft to stabilize the agonist, and its aromatic hydroxyl group approaches αK145, which in turn loosens its contact with αD200. The positional changes of αK145 and αD200 are proposed to initiate the cascade of perturbations that opens the receptor channel: the first perturbation is of β-strand 7, which harbors αK145 and is part of the signature Cys-loop, and the second is of β-strand 10, which harbors αD200 and connects to the M1 domain. Thus, interplay between these three conserved residues relays the initial conformational change from the ACh binding site toward the ion channel.

INTRODUCTION

Activation of post-synaptic receptors is accomplished by allosteric communication between the neurotransmitter binding site and the distal ion channel. The local conformational change due to neurotransmitter binding is amplified in a cascade that culminates in the global conformational change that opens the channel. For the Cys-loop superfamily of post-synaptic receptors, atomic-scale insight into the neurotransmitter binding site emerged from the x-ray structure of the homologous acetylcholine binding protein (AChBP; Brejc et al., 2001) and from homology models derived from it (Le Novère et al., 2002; Molles et al., 2002; Schapira et al., 2002; Sine et al., 2002a). Additionally a 4Å resolution structural model of the binding, channel, and cytoplasmic domains was generated from electron microscopy of two-dimensional arrays of Torpedo receptors (Miyazawa et al., 2003; Unwin, 2005). At the interface between binding and pore domains, a network of loops was shown to couple neurotransmitter binding to channel gating (Kash et al., 2003; Bouzat et al., 2004; Chakrapani et al., 2004), but coupling structures near the neurotransmitter binding site remain unknown.

In the adult muscle nicotinic receptor, the two ACh binding sites are formed at interfaces between an α subunit and a neighboring ε or δ subunit where multiple recognition domains from each subunit converge (for reviews see Corringer et al., 2000; Karlin, 2002; Sine, 2002). The α subunit contributes recognition domains A–C, while the non-α subunit contributes domains D–G. Conserved aromatic residues from domains A–D form an aromatic cage that coordinates the positively charged agonist (Celie et al., 2004), whereas nonconserved residues in domains D–G endow the two binding sites with selectivity for agonists and competitive antagonists (Sine, 2002). Domains C and F are the most peripheral of the recognition domains, and of these, domain C exhibits the greatest displacement on binding the agonist (Celie et al., 2004; Gao et al., 2005).

To identify structures that propagate the local conformational change elicited by ACh away from the binding site, we examined our homology model of the receptor ligand binding domain (Sine et al., 2002a). We looked for conserved residues at the periphery of the binding site near the mobile domain C and found a cluster of three conserved residues, αK145, αD200, and αY190, whose side chains potentially interact through electrostatic forces. Our patch clamp recordings show that structurally conservative mutations of each residue profoundly impair gating of the channel. The common contributions to channel gating and direct linkages to the binding site and binding-pore interface suggest

Abbreviations used in this paper: AChBP, acetylcholine binding protein; Btx, bungarotoxin; CCh, carbamylcholine.
interplay among these three residues initiates the allosteric cascade from the binding site to the channel gate.

**MATERIALS AND METHODS**

**Construction of Wild-type and Mutant AChRs**

Human α, β, ε, and δ subunit cDNAs were obtained as previously described (Ohno et al., 1996) and subcloned into the mammalian expression vector pRBG4 (Lee et al., 1991). Site-directed mutations were introduced using the Quick-Change site-directed mutagenesis kit (Stratagene). The presence of each mutation and the absence of unwanted mutation were determined by sequencing the entire cDNA insert.

**Mammalian Cell Expression**

BOSC 23 cells (Pear et al., 1993; Wang et al., 2000), a variant of HEK 293 clonal fibroblasts, were transfected with mutant or wild-type cDNAs using calcium phosphate precipitation as previously described (Lee and Sine, 2004). Patch clamp and [125I]α-bungarotoxin (Btx) binding measurements were made 2 and 3 d after transfection, respectively.

**Patch-clamp Recordings**

Recordings were obtained in the cell-attached configuration (Hamill et al., 1981) at a membrane potential of ~70 mV and a temperature of 21°C. Bath and pipette solutions contained 142 mM KCl, 5.4 mM NaCl, 1.8 mM CaCl2, 1.7 mM MgCl2, 10 mM HEPES (pH 7.4). Single channel currents were recorded using an Axopatch 200B (Axon Instruments Inc.). Data were obtained from two to four different patches for each ACh concentration. Recordings were accepted for analysis only when the frequency of occurrence of clusters of events was low enough to be sure they originated from a single receptor channel. Currents were low pass filtered at 100 kHz and recorded to hard disk at 200 kHz using the program Acquire (Bruxton Co.), using the option that allows high bandwidth sampling of periods of channel activity while keeping track of the intervening quiescent periods.

**Single-channel Kinetic Analysis**

Digitized current signals were analyzed as recently described (Lee and Sine, 2004) using a 10-kHz digital Gaussian filter, cubic spline interpolation of the signal, a dead time of 10 μs, the half-amplitude threshold for detection and correction of the measured dwell times at threshold for the effects of the Gaussian filter (Colquhoun and Sigworth, 1983). Dwell time histograms were constructed using a logarithmic abscissa and square root ordinate and fitted to the sum of exponential components by maximum likelihood using the program TACFit (Bruxton Co.). Clusters of events corresponding to a single channel were identified as a series of closely spaced openings preceded and followed by closed intervals greater than a specified duration. This duration was taken as the point of intersection of the closed time component that depended on ACh concentration with the succeeding closed time component due to desensitization. For analyses that included desensitized states, the closed duration for defining clusters was taken as the point of intersection of the closed time due to intermediate onset desensitization and the succeeding component determined by slow desensitization and the number of channels in the patch. Clusters of events with five or more openings and cluster mean open probabilities and cluster mean closed times within two standard deviations of the respective means were accepted for further analysis. Kinetic analysis was conducted simultaneously on data obtained across a range of ACh concentrations, termed global analysis, using MLL software (Qub suite, State University of New York, Buffalo), which corrects for missed events and gives error estimates of the fitted parameters (Qin et al., 1996). A dead time of 22 μs was imposed on all datasets. For each type of receptor studied, global analysis included data from two to four patches for each ACh concentration. ACh concentrations spaced at half-log unit intervals of the concentration, and a 10- to 333-fold range of the ACh concentration.
ACh Binding Measurements

The total number of $^{125}\text{I}$/H9251-Btx binding sites on the cell surface of transfected BOSC 23 cells and ACh competition against the initial rate of $^{125}\text{I}$/H9251-Btx were determined as previously described (Sine, 1993). ACh competition measurements were analyzed according to the following form of the Hill equation:

$$1 - Y = \frac{1}{1 + ([\text{ACh}] / K_{\text{app}})^{n_H}}.$$

where $Y$ is fractional occupancy by ACh, $n_H$ is the Hill coefficient, and $K_{\text{app}}$ is the apparent dissociation constant.

RESULTS

Triad of Conserved Residues at the Periphery of the ACh Binding Site

Structures of AChBP and our homology model of the nicotinic receptor ligand binding domain show that recognition domain C is strategically positioned to transmit local conformational change due to ACh binding to structures linked to the ion channel (Fig. 1 B). Domain C comprises the loop spanning $\beta$-strands 9 and
10 of the α subunit and links directly to the M1 transmembrane domain via β-strand 10. It also overlays the Cys-loop, which articulates with the linker joining the M2 and M3 transmembrane domains (Kash et al., 2003; Bouzat et al., 2004; Unwin, 2005). Inspection of the AChBP structure, with and without carbamylcholine (CCh) bound, reveals two salt bridges that have counterparts in our structural model of the receptor ligand binding domain. At binding sites without CCh bound, a salt bridge forms between K139 and D194 of AChBP, whereas at binding sites with CCh bound, a salt bridge forms between K139 and Y185 (Celie et al., 2004; Fig. 1 C). In our structural model of the receptor ligand binding domain, residues equivalent to these, αK145, αD200, αY190, are close together in three-dimensional space and are conserved across all α1 subunits (Figs. 1, A and D), suggesting key structural or functional contributions. Thus the x-ray structure of AChBP, our structural model of the receptor ligand binding domain and residue conservation together suggest that αK145, αD200, and αY190 are candidates for transmitting local conformational change at the ACh binding site to structures linked to the ion channel.

Single Channel Kinetics of the Adult Human Receptor

To provide a frame of reference for assessing the effects of receptor mutations, we recorded an independent set of ACh-evoked single channel currents for the adult human receptor under experimental conditions identical to those used to study the mutant receptors. We co-expressed α, β, ε, and δ subunits in BOSC 23 cells and recorded single channel currents using the cell-attached configuration of the patch clamp with defined concentrations of ACh in the patch pipette (see MATERIALS AND METHODS). Currents recorded at desensitizing concentrations of ACh consist of a series of clusters of closely spaced current pulses separated by prolonged quiescent periods (Sakmann et al., 1980; Sine and Steinbach, 1987; Colquhoun and Ogden, 1988). Channel events within each cluster correspond to activation of a single receptor channel, and exhibit an ACh-dependent shortening of closed periods between successive openings, with little ACh dependence of the open periods (Fig. 2). The corresponding closed dwell time histograms show a shift from long to short time of a major component of closings as the ACh concentration is increased, whereas the major component of openings is concentration independent.

To estimate rate constants for elementary steps underlying activation of a single receptor channel, we analyzed the global set of dwell times according to kinetic schemes (Fig. 3). Scheme I depicts receptor activation as sequential binding of ACh to the resting, closed receptor, followed by opening and then block by ACh of the doubly occupied open receptor. Scheme II contains all of Scheme I but includes two desensitized states connected to the open state (Elene and Auerbach, 2002; Lee and Sine, 2004). Scheme II is used to quantitatively account for the small fraction of closings due to fast and intermediate onset desensitization that are short enough to be indistinguishable from closings due to receptor activation. A third scheme depicting independent binding of ACh to each site, although able to describe data from the wild-type receptor (Hatton et al., 2003; Lee and Sine, 2004), is not applied here because it did not permit adequate definition of all the rate constants for the mutant receptors examined in this study.

Maximum likelihood fitting based on Scheme I yields probability density functions that describe the global set of closed and open dwell times for the adult human receptor (MATERIALS AND METHODS; Fig. 2). Rate constants underlying ACh binding, channel opening, and ACh blocking steps are all well defined by the analysis, and show rapid and efficient opening of the doubly occupied receptor, distinguishable association and dissociation rate constants for each binding site, and low affinity block of the open channel by ACh (Table I). The fitted rate constants are within the range of values previously reported for the adult human receptor (Ohno

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**Figure 3.** Kinetic schemes for receptor activation. Receptors in the resting closed state are symbolized by R, the open state by R*, and blocked state by R<sub>b</sub>. ACh (A) associates with rate constants k<sub>a1</sub> and k<sub>a2</sub>, and dissociates with rate constants k<sub>d1</sub> and k<sub>d2</sub>. Scheme II incorporates fast and intermediate onset desensitized states with onset rate constants k<sub>a11</sub> and k<sub>a12</sub> and recovery rate constants k<sub>a11</sub> and k<sub>a12</sub>.

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**Scheme I.**

\[
\begin{align*}
A + R & \overset{k_{a1}}{\underset{k_{d1}}{\rightleftharpoons}} A + AR \\
\end{align*}
\]

**Scheme II.**

\[
\begin{align*}
A + R & \overset{k_{a1}}{\underset{k_{d1}}{\rightleftharpoons}} A + AR \\
\end{align*}
\]
et al., 1996; Wang et al., 1999; Hatton et al., 2003; Lee and Sine, 2004).

Maximum likelihood fitting based on Scheme II, which incorporates brief and intermediate duration desensitized states, also describes the distributions of closed and open dwell times, and yields essentially the same rate constants for binding and gating steps obtained using Scheme I (Table I). The similar results obtained with Schemes I and II show that the presence of closings due to desensitization does not affect the estimates of rate constants for receptor activation when Scheme I is applied to the wild-type human receptor (Lee and Sine, 2004).

**Functional Contribution of αK145**

αK145 is a potential link between the binding site and the ion channel because it is proximal to the binding site, it is physically linked to the Cys-loop, it is proximal to electron-rich side chains that link to M1 via β-strand 10, and it is conserved across α1 subunits. In the agonist-free structure of AChBP, the residue equivalent to αK145, K139, forms a salt bridge with D194, whereas in the CCh-bound structure, K139 forms a salt bridge with the key binding site tyrosine, Y185 (Celic et al., 2004). To investigate the functional contribution of αK145, we generated a series of mutations of αK145, coexpressed each with complementary β, ε, and δ subunits, and recorded ACh-evoked single channel currents.

Substitution of Ala for αK145 profoundly impairs receptor activation at all concentrations of ACh, with activation episodes due to a single receptor consisting of predominantly single openings separated by prolonged closings (compare Figs. 2 and 4). The corresponding closed time histograms span a wide range of times at all ACh concentrations examined, and the major peak of closings shows an ACh-dependent shift toward shorter times. Maximum likelihood fitting of Scheme I to the global dataset for αK145A well describes the closed and open dwell time histograms (Fig. 4). The fitted rate constants reveal nearly a 200-fold slowing of the channel opening rate constant and no change of the channel closing rate constant (Table I). For αK145A, adequate definition of the rate constants underlying ACh binding steps required the approximation that the set

| Table I | Kinetics of Activation of AChRs Expressed in BOSC 23 Cells |
|---------|------------------------------------------------------------|
| Receptor | $k_{-1}$ | $k_{1}$ | $K_i$/$\mu$M | $k_{-2}$ | $k_{2}$ | $K_i$/$\mu$M | $\beta_1$ | $\alpha_1$ | $\Theta_1$ | $\beta_2$ | $\alpha_2$ | $\Theta_2$ | $k_{1s}$ | $k_{1b}$ | $K_o$/mM |
| Wild-type | 346 | 3480 | 10 | 92 | 19000 | 207 | 65 | 3500 | 0.02 | 43000 | 2140 | 20.1 | 33 | 3 ± 3 | 162000 | 4.9 |
| Wild-type | ±39 | ±429 | ±2 | ±355 | ±6 | ±302 | ±1046 | ±41 | ±3790 |
| αK145A | 154 | 4900 | 32 | 80 | 9870 | 123 | ND | ND | – | 240 | 1900 | 0.13 | 18 ± 1 | 128000 | 7.1 |
| αK145Q | ±37 | ±390 | ±2 | ±297 | ±6 | ±436 | ±1133 | ±43 | ±3700 |
| αD200N | 105 | 7150 | 68 | 53 | 14300 | 269 | ND | ND | – | 734 | 1670 | 0.44 | 30 ± 2 | 106000 | 3.5 |
| αD200N+ | ±7 | ±475 | ±3 | ±950 | ±69 | ±15 | ±9600 |
| αD200Q | 275 | 4200 | 15 | 137 | 8450 | 61 | 80 | 14100 | 0.018 | 635 | 2600 | 0.24 | 28 ± 2 | 105000 | 3.8 |
| αD200Q+ | ±33 | ±520 | ±16 | ±1050 | ±9 | ±1420 | ±112 | ±40 | ±1700 |
| αK145E | ±16 | ±410 | ±8 | ±850 | ±5 | ±900 | ±90 | ±19 | ±11900 |
| αK145E | ±3 | ±300 | ±1 | ±600 | ±50 | ±15 | ±1100 |
| αY190F | – | – | – | – | – | – | – | – | – | 105 | 7600 | ±0.14 | 17 ± 2 | 130000 | 7.6 |
| αY190F | ±1 | ±300 | ±1 | ±600 | ±1 | ±90 | ±5900 |
| αY190Q | ±31 | ±600 | ±122 | ±40 | ±8100 | ±90 | ±122 | ±27 | ±3 | 114000 | 4.2 |
| αY190Q | ±8 | ±1000 | ±5 | ±1000 | ±86 | ±31 | ±20100 |
| αT292V | ±9 | ±102 | ±34 | ±3050 | ±183 | ±1730 | ±20100 |
| αT292V | ±3 | ±200 | ±5 | ±440 | ±37 | ±30 | ±20500 |
| αK145A | ±3 | ±300 | ±3 | ±640 | ±860 | ±32 | ±2500 |
| αK145A | ±4 | ±23 | ±1 | ±52 | ±1 | ±52 | ±2600 |

Values are from global fits of Scheme I to data obtained over a range of ACh concentrations, with error limits given (see MATERIALS AND METHODS). Rate constants are in units of $\mu$m$^{-1}$s$^{-1}$ for association rate constants, and s$^{-1}$ for all others. The channel gating equilibrium constants ($\Theta_i$) are $\beta_i$/$\alpha_i$. $K$s are dissociation constants, $k/k_{-1}$.

*Fits according to Scheme II, which contains two desensitized states linked to the open state (Fig. 3 and text). Fitted desensitization rate constants, ($k_{s1}/k_{-s1}$) and ($k_{s2}/k_{-s2}$) in units of s$^{-1}$, are $D_1$ (19/39), $D_2$ (24/660).

*Fits according to the subset of doubly occupied states in Scheme I. ND indicates parameters were not defined.

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of elementary association and dissociation rate constants are equal at each binding site. Presumably this approximation was required because information on the first binding step was limited: clusters of events corresponding to a single receptor could only be identified at ACh concentrations >30 μM, which may exceed the dissociation constant for ACh binding to the first binding site, and brief, mono-liganded openings were not present. Nevertheless, our analysis suggests that the rate constants underlying ACh binding are only modestly affected by K145A (Table I). Thus substitution of Ala for K145 at the periphery of the ACh binding site predominantly impairs elementary steps underlying channel gating.

**Functional Consequences of K145Q**

Because the Lys to Ala substitution represents a substantial structural change, we examined functional consequences of the more structurally conservative mutation K145Q. The Gln substitution neutralizes the positive charge but maintains polarity and moderate size of the side chain. Receptors containing K145Q show impaired channel gating similar to K145A, but individual activation episodes show three kinetic modes distinguished by low, intermediate, and high open probability (P_open; Fig. 5). These kinetic modes were observed only at high concentrations of ACh (300 μM to 1 mM), and although they were present at all high concentrations, the contributions of each mode varied from patch to patch. We therefore describe detection of the three kinetic modes for one patch to exemplify our treatment of the overall data for K145Q.

We identified clusters of events corresponding to a single channel using our standard cluster discrimination method (MATERIALS AND METHODS), and generated histograms of mean cluster P_open and mean cluster duration.

**Figure 4.** ACh-evoked single channel currents and kinetic analysis of receptors containing the mutation αK145A. Currents recorded at the indicated ACh concentrations are shown at a bandwidth of 10 kHz. Dwell time histograms are shown with pdfs generated from the Scheme 1 rate constants (Table I) overlaid.
closed time ($\tau_C$) for αK145Q (Fig. 5; Milone et al., 1998). The distribution of $P_{\text{open}}$ shows a major and a minor peak, while the distribution of $\tau_C$ shows a major peak and a secondary tail of long $\tau_C$. Analysis of cluster properties reveals that distinct portions of the two distributions are correlated: the peak of high $P_{\text{open}}$ corresponds to clusters with short $\tau_C$ (filled bars), and the tail with long $\tau_C$ corresponds to clusters with low $P_{\text{open}}$ (open bars). The remainder of the distributions consists of clusters with intermediate $P_{\text{open}}$ and $\tau_C$ (gray bars). Clusters from wild-type receptors subjected to the same analysis show a predominant mode of high $P_{\text{open}}$ and short $\tau_C$.

We then separated clusters of events for each of the kinetic modes for αK145Q receptors, and combined one mode at a time with the kinetically homogeneous data obtained at lower ACh concentrations. Maximum likelihood fitting showed that only clusters with intermediate $P_{\text{open}}$ and $\tau_C$ were compatible with data obtained at lower ACh concentrations. Fitting Scheme I to the intermediate mode dataset (30 μM to 1 mM ACh) shows that substitution of Gln for αK145 profoundly impairs the efficiency of channel gating; the corresponding probability density functions adequately describe the distributions of closed and open dwell times (Fig. 6). The effects of αK145Q on channel gating are similar to those of αK145A, with the channel opening rate constant slowed 60-fold and the channel closing rate constant relatively unchanged (Table I). Adequate definition of rate constants underlying ACh binding steps again required the approximation that the set of elementary association and dissociation rate constants are equal at each binding site. Given this approximation, the analysis suggests αK145Q has relatively little effect on agonist binding (Table I). Thus, located at the periphery of the ACh binding site, αK145 contributes primarily to gating of the receptor channel, fulfilling an important criterion as a transduction element.

Functional Contribution of αD200

αD200 is a potential electrostatic partner of αK145, and in the mouse receptor was found to contribute predominantly to channel gating (Akk et al., 1996). To determine whether αD200 contributes to channel gating in the human muscle receptor, and further, whether its functional contribution mirrors that of αK145, we recorded currents from adult human receptors harboring the mutation αD200N. Like mutations of αK145, αD200N profoundly impairs receptor activation at all concentrations of ACh; individual clusters consist of predominantly single openings interspersed by long closings (Fig. 7). Maximum likelihood fitting according to Scheme I reveals a predominant effect on channel gating, with the channel opening rate constant slowed nearly 70-fold and the channel closing rate constant slightly increased (Table I). As observed for mutations of αK145, adequate definition of rate constants underlying ACh binding steps required the approximation that the set of elementary association and dissociation rate constants are equal at each binding site. The rate constants underlying ACh binding are relatively unaffected by αD200N, as observed for the same mutation in the mouse receptor (Table I). Thus the consequences of αD200N for channel gating are highly similar between mouse and human receptors, and, moreover, mirror the consequences of mutations of αK145. The similar effects of mutations of αK145 and αD200.
Initial Binding to Gating Link Mediated by Triad of Conserved Residues

on channel gating and proximity of the two side chains in our structural model collectively suggest these residues are interdependent and constitute part of an intermediate link between binding and gating domains.

Interdependence of αK145 and αD200

To gain additional evidence for interdependence of αK145 and αD200, we combined the mutations αK145Q and αD200N into a single α subunit and recorded ACh-evoked single channel currents from the resulting double mutant receptors. The current traces again show impaired receptor activation at all ACh concentrations (Fig. 8), with predominantly single openings separated by long closings, similar to either single mutation (compare Figs. 6, 7, and 8). Unlike receptors containing αK145Q alone, only a single kinetic mode was distinguished for the double mutant receptor, simplifying the analysis. Maximum likelihood fitting of Scheme I to the global set of dwell times reveals a 60-fold slowing of the channel opening rate constant and little change of the channel closing rate constant when compared with the wild-type receptor (Table I). These results are similar to those for either single mutation, showing qualitatively that αK145Q and αD200N are interdependent in contributing to channel gating. Double mutant cycles analysis (Horovitz and Fersht, 1990) applied to the set of gating equilibrium constants for wild-type, αK145Q, αD200N, and double mutant receptors yields a free energy of interaction of −2.6 kcal/mol. Rate constants underlying ACh binding steps are altered twofold or less by the double mutation, similar to αD200N alone, but this conclusion remains provisional because the assumption of equivalent binding sites was required to obtain well-defined rate constants.
Thus, the largely selective perturbation of channel gating by the double mutation quantitatively mirrors that for either single mutation, further suggesting \( \alpha K145 \) and \( \alpha D200 \) are interdependent in coupling agonist binding to channel gating.

**Charge Reversal of \( \alpha K145 \) and \( \alpha D200 \)**

To test for electrostatic interaction between \( \alpha K145 \) and \( \alpha D200 \), we generated the single mutants \( \alpha K145E \) and \( \alpha D200K \) and the corresponding double mutant and recorded ACh-evoked single channel currents from the resulting mutant receptors (Fig. 9). The single mutation \( \alpha D200K \) slows the channel opening rate constant 400-fold and increases the channel closing rate constant nearly fourfold, yielding a 1,400-fold decrease of the channel gating equilibrium constant (Table I). For \( \alpha D200K \), well-defined single receptor clusters could not be discerned at ACh concentrations <1 mM, preventing estimation of rate constants underlying ACh binding. Consequences of the single mutation \( \alpha K145E \) mirror those of other mutations of \( \alpha K145 \); compared with the wild-type receptor, the channel opening rate constant slows 100-fold, the channel closing rate constant slows almost twofold, and the ACh dissociation rate constants are only modestly affected. The rate constants for ACh association, on the other hand, slow three- to fivefold compared with other mutations of \( \alpha K145 \) (Table I).

Compared with \( \alpha D200K \) alone, restoring the local negative charge by combining \( \alpha K145E \) markedly increases the frequency of channel openings within single receptor clusters (Fig. 9), and counteracts the slowing of the channel opening rate constant and the increase of the channel closing rate constant (Table I). The extent to which the gating rate constants are restored is only partial, perhaps because residues flanking each member of the putative salt bridge are not compatible with the reversed charge. The net result of
charge reversal is a 14-fold increase of the channel gating equilibrium constant compared with αD200K alone. Double mutant cycles analysis applied to the set of four gating equilibrium constants yields a free energy of interaction of $-4 \text{kcal/mol}$. Further, the slow association of ACh caused by αK145E alone is restored to the range of values observed for other mutations of αK145 (Table I).

Finally, we examined the single mutation αK145D and found that it prevented expression of functional channels and cell-surface α-Btx binding sites. However, αK145D is rescued by the charge reversal mutation αD200K. Full kinetic analysis of the double mutation reveals an eightfold increase of the channel gating equilibrium constant compared with αD200K alone, due to a twofold increase of the channel opening rate constant and a fourfold slowing of the channel closing rate constant (Table I). Changes in rate constants underlying ACh binding remain unknown because these could not be measured for either single mutation. However, the ACh association rate constants for the double mutation increase twofold compared with αK145E alone, while the ACh dissociation rate constants slow twofold compared with αK145E. The overall results from charge-neutralization and charge-reversal experiments provide strong evidence that αK145 and αD200 interact through electrostatic forces to couple agonist binding to channel gating.

**Functional Contributions of αY190 and αT202**

Our structural model predicts that two more residues are proximal to αK145 and αD200: αY190 and αT202.
We therefore generated structurally conservative mutations of each residue, αY190F and αT202V, and recorded single channel currents from the resulting mutant receptors. For receptors containing αY190F, the ACh concentration had to be increased to 3 mM to obtain well-defined clusters of single receptor currents; the corresponding intracluster events consist of single brief openings separated by prolonged closings (Fig. 10). The availability of data at only a single high concentration of ACh limited kinetic fitting to the subset of doubly occupied gating steps in Scheme I, but nevertheless reveals a 1,000-fold slowing of the channel opening rate constant with little change in the channel closing rate constant (Table I). These effects of αY190F on channel gating are analogous to those observed for the same mutation in the fetal mouse muscle receptor (Chen et al., 1995), but the 800-fold decrease of the gating equilibrium constant for the adult human receptor exceeds the 200-fold decrease observed for the fetal mouse receptor.

The kinetics of receptor activation are only modestly affected by αT202V. Rate constants for elementary

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**Figure 9.** Charge reversal of αK145 and αD200. ACh-evoked single channel currents and kinetic analysis are shown for the single mutant and corresponding double mutant receptors. Currents recorded at the indicated ACh concentrations are shown at a bandwidth 10 kHz. Dwell time histograms are shown with pdfs generated from the Scheme I rate constants (Table I) overlaid. Note only the indicated saturating concentration of ACh produced clear-cut clusters of events for αD200K, allowing assignment of only channel gating and channel blocking rate constants. However the full set of Scheme I rate constants was obtained for αK145E and the charge reversal mutant (Table I). The charge reversal mutation increases the frequency of openings compared with αD200K alone.
binding or gating steps change twofold or less compared with the wild-type receptor (Table I). Thus despite its conservation across α1 subunits and proximity to αD200 and αK145, αT202 does not contribute substantially to either ACh binding or channel gating.

**Interdependence of αK145 and αY190**

The agonist-bound conformation of AChBP shows that K139 and Y185 form a salt bridge (Celie et al., 2004). To determine whether residues in the receptor equivalent to these, αK145 and αY190, are interdependent, we combined αK145Q and αY190F into a single α subunit and recorded ACh-evoked single channel currents from the resulting double mutant receptors. The current traces again show impaired receptor activation at 1 mM ACh, with brief openings separated by long closings, similar to either single mutation (Fig. 10). Unlike receptors containing αK145Q alone, only a single kinetic mode could be distinguished for the double mutant receptor. Further, well-defined clusters of single receptor openings could only be discerned at ACh concentrations from 100 μM to 1 mM, restricting the
Maximum likelihood fitting shows that compared with the wild-type receptor, the double mutation slows the channel opening rate constant 200-fold without affecting the channel closing rate constant (Table I). Double mutant cycles analysis applied to the set of gating equilibrium constants for wild-type, K145Q, Y190F, and double mutant receptors yields a free energy of interaction of 2.9 kcal/mol. Thus K145Q and Y190F are interdependent in coupling agonist binding to channel gating.

### Pairwise Interdependence Assessed by Steady-state Binding of ACh

Most mutations that affect receptor function alter steady-state binding of ACh (examples include Chen et al., 1995; Ohno et al., 1996; Sine et al., 2002b). Although measurements of steady-state ACh binding contain contributions from closed, open, and desensitized states of the receptor, each of which differs in affinity for the agonist (Katz and Thesleff, 1957; Sine and Taylor, 1979; Jackson, 1989), binding measurements can be used as an overall index of receptor function. We therefore sought further evidence for pairwise interdependence by measuring steady-state binding of ACh to receptors containing single and double mutations. We first studied the charge neutral mutations αK145Q and αD200N and the corresponding double mutation. Relative to the wild-type receptor, the single mutations αK145Q and αD200N increase the apparent dissociation constant for ACh (Kapp) 10- and 26-fold, respectively, qualitatively consistent with expectations from impaired channel gating observed by single channel recording (Fig. 11; Table II). The double mutation shows a virtually identical binding profile to that of αD200N alone, and double mutant cycles analysis applied to Kapp yields an interaction free energy of 1 kcal/mol for this charge neutralization experiment. Like Kapp interaction free energy based on Kapp arises from multiple processes: ACh binding, desensitization, and channel gating. One of these processes is channel gating, which contributes 2.6 kcal/mol, but the other processes likely contribute. We next measured steady-state ACh binding for the charge reversal mutations αK145E and αD200K and the corresponding double mutation. The single mutations αK145E and αD200K increase Kapp 100- and 860-fold, respectively, again qualitatively consistent with profoundly impaired channel gating (Fig. 11; Table II). However, charge reversal by the double mutation strongly counteracts the effects of either single mutation, yielding an overall 20-fold increase of Kapp compared with the wild-type receptor. Double mutant cycles analysis applied to Kapp for this charge reversal experiment reveals an interaction free energy of 4.8 kcal/mol, which is similar to the contribution of 4 kcal/mol observed for channel gating alone. The mutation αK145D does not produce functional receptors, but when combined with αD200K, yields a Kapp coincident with that obtained for αK145E plus αD200K (Fig. 11; Table II).

The double mutation containing αK145Q and αY190F failed to bind α-Btx, preventing assessment of interdependence by steady-state ACh binding. Nevertheless the overall results provide strong evidence for electrostatic interaction between αK145 and αD200 and interdependence of αK145 and αY190. A three-way electrostatic interplay of charged residues thus emerges as a compelling candidate for an initial link between ligand binding and channel gating domains owing to the proximity of the triad to the ACh binding site, interdependence of residues in the triad, connections to the Cys-loop and M1 domain, and the predominant contributions of all three residues to channel gating.

### Subunit Specificity of the Mutations

The three residues studied here are unique to and conserved across α1 subunits. However, 5 subunits also contain conserved Asp and Lys at positions equivalent to αD200 and αK145, respectively. To determine whether
this residue pair in the δ subunit contributes to receptor function, we generated the mutations δK147Q and δD214N, coexpressed each with complementary α, β, and ε subunits, and measured steady-state ACh binding. Neither mutation affects K_app for ACh binding (Table II), showing that if these two residues form a salt bridge, it is unlikely to contribute to receptor function.

**DISCUSSION**

We show that three conserved residues at the periphery of the ACh binding site, αK145, αD200, and αY190, contribute profoundly to gating of the receptor channel. One of these residues, αY190, contributes to the aromatic cage that coordinates the quaternary ammonium moiety of the agonist (Celie et al., 2004), and its hydroxyl group projects away from the binding site toward the other two members of the triad (Fig. 1). Of the other two residues, αK145 is the most likely contact for αY190, but this contact corresponds to the agonist-bound conformation. Evidence for this contact comes from the AChBP crystal structure with CCh bound, which shows a salt bridge between the equivalent residues K139 and Y185 (Celie et al., 2004), and from the interdependent contributions of αK145 and αY190 to gating shown here. Support for electrostatic contact between αK145 and αD200 comes from the AChBP crystal structure without bound agonist, which shows a salt bridge between the equivalent Lys and Asp residues (Brejc et al., 2001), and from our data demonstrating interdependence of αK145 and αD200 in contributing to channel gating.

Our findings combine with inter-atomic distances in AChBP to suggest how interplay among these three residues couples local conformational change due to agonist binding to gating of the channel. Our inferences draw from two sets of atomic coordinates of AChBP with bound CCh deposited in the protein data bank (PDB code 1UV6): a pentameric unit with one molecule of CCh bound and another unit with two molecules of CCh bound. Inspection of the three subunit interfaces to which CCh is bound shows that recognition domain C is drawn inward toward the center of the binding site, whereas inspection of one of the unoccupied interfaces shows that domain C is extended outward from the site. Molecular dynamics simulation of AChBP and measurements of intrinsic tryptophan fluorescence reveal similar changes in the conformations of domain C with and without bound agonist (Gao et al., 2005). In the following mechanistic interpretation, we refer to these conformations as agonist-free and agonist-bound.

To illustrate our structural interpretation, the agonist-free and -bound conformations of recognition domain C of AChBP are compared (Fig. 12). The agonist-bound conformation is that of the crystal structure of AChBP with bound CCh (Celie et al., 2004). The agonist-free conformation is that generated by prolonged molecular dynamics simulation of AChBP (Gao et al., 2005), which is similar to the extended conformation of a subunit of AChBP at an unoccupied interface (see PDB entry 1UV6). ACh is shown docked in the orientation obtained by computational docking to the HEPES-bound crystal structure of AChBP (Brejc et al., 2001; Gao et al., 2005). In the agonist-free conformation, the Tyr at the position equivalent to αY190 is 8 Å away from the Lys equivalent to αK145, while residues equivalent to αK145 and αD200 form a salt bridge. In the agonist-bound conformation, the Tyr equivalent to αY190 moves to within 2–3 Å of the Lys equivalent to αK145, drawing recognition domain C inward, and allowing the residue equivalent to αD200 to move away from the Tyr/Lys pair. Thus by analogy to AChBP, in the resting state of the receptor, αK145 and αD200 are proposed to pair through electrostatic forces and αY190 is out of register (Fig. 11). However when the agonist binds, αY190 is drawn toward the αK145/αD200 pair and, through electrostatic forces, pulls αK145 away from αD200, allowing both residues to relax to new positions. These local inter-residue displacements could propagate to the channel via β-strands 7 and 10 of the ligand binding domain. β-Strand 7 harbors αK145 and forms part of the signature Cys-loop, which interacts with the linker spanning the M2 and M3 transmembrane domains (Kash et al., 2003; Bouzat et al., 2004), while β-strand 10 contains αD200 and links directly to the M1 transmembrane domain.

Although all three residues of the triad contribute profoundly to channel gating, each has a distinct role in linking the binding site to the channel. αY190 is the clear link to the ACh binding site, as shown by site-
domain C through aromatic-cation interactions, allowing Y185 to establish a salt bridge with K139 and weaken the K139/D194 salt bridge. The resulting displacements of β-strands 7 and 10 are proposed to propagate to channel and trigger opening. Putative salt bridges are indicated by dashes.

directed labeling (Abramson et al., 1989) and its location in the aromatic cage that coordinates the agonist (Celie et al., 2004). Its function is to help draw recognition domain C toward the center of the binding site when the agonist is bound and, in the process, form a salt bridge with K145. Accordingly, removal of the aromatic hydroxyl group by mutation to Phe would prevent formation of the salt bridge and disrupt coupling of binding to gating. Likewise, substitution of Gln, Ala, or Glu for K145 would remove the electrostatic partner for Y190, also disrupting coupling. By analogy to AChBP in the agonist-free conformation, D200 is proposed to form a salt bridge with K145, positioning K145 for contact with the incoming Y190 and maintaining β-strands 7 and 10 in their closed state conformations. When the agonist binds, Y190 draws K145, positioning K145 for the incoming D200, allowing β-strands 7 and 10 to move to their active conformations. Substitution of Asn for D200 would prevent stabilization of K145 for the incoming Y190, and break the link between β-strand 10 and the binding site, disrupting coupling.

One of the main lines of evidence that the triad identified here is an initial link in the binding to gating cascade is that mutation of each residue profoundly impairs gating of the channel. However two additional observations require explanation. First, the allosteric link between binding and gating domains is not completely disrupted by any of the mutations, as ACh can still trigger opening of the channel, although with greatly reduced efficiency. The most likely explanation is that there are additional allosteric links that remain intact in each mutant receptor and account for the residual ACh-evoked currents. Second, the extent to which gating is disrupted differs by as much as 10-fold among the three mutations. If each member of the triad is essential for coupling, the extent to which gating is disrupted should be the same following mutation of each member. However in addition to coupling binding to gating, the triad may also contribute to the allosteric equilibrium between resting and open channel states in the absence of agonist (Jackson, 1989). This contribution to the allosteric set point likely involves dynamic interplay among the three residues, such that changes in the set point differ when one of the members is structurally altered. Thus for each mutation, the gating equilibrium for di-liganded receptors could be altered by differing amounts, owing to different contributions to the allosteric set point.

Other residues that contribute to the allosteric set point have been identified in the ligand binding domains of members of the Cys-loop receptor superfamily. In the nicotinic receptor α1 subunit, mutation of Asp 97 markedly increases the frequency of channel opening events in the absence of ACh (Chakrapani et al., 2003). In the GABA<sub>A</sub> receptor ρ1 subunit, mutation of Tyr 107 increases the resting membrane conductance in a manner that was sensitive to the specific GABA receptor inhibitor picrotoxin (Torres and Weiss, 2002). Similarly in the GABA<sub>B</sub> receptor β<sub>3</sub> subunit, mutation of Glu 155 increases the resting conductance in a picrotoxin-sensitive manner (Newell et al., 2004). In all three cases, the allosteric set point was increased in favor of the open channel state, whereas mutation of any residue of the triad in the present study impedes opening of the channel. Thus among functionally key residues in the ligand binding domain of the receptor, a balance is likely established between residues that promote and those that impair channel gating.

The triad of residues identified here is found in all species of nicotinic receptor α<sub>1</sub> subunits, suggesting
they contribute to agonist binding transduction in all muscle type nicotinic receptors (for subunit sequences see Le Novere and Changeux, 2001). Similarly, the triad is also present in all neuronal nicotinic α subunits (α2–α10), except α6, which contains Asp at the position equivalent to α190, and α10 which contains Thr at the position equivalent to α145. However, neither of these exceptions likely forms the principal face of the binding site; α5 appears to be a structural subunit, requiring both neuronal α and β subunits to form functional receptors (Ramirez-Latorre et al., 1996), and α10 requires coexpression of the α9 subunit (Sgard et al., 2002). On the other hand, the triad is not present in GABAA, glycine, and 5-HT receptors. Thus among the Cys-loop superfamily of receptors, the initial coupling mechanism is likely to be highly similar for muscle and neuronal nicotinic receptors, but the more evolutionary distant receptors evolved different structures to relay the initial consequences of agonist binding. This structural divergence among receptors activated by different neurotransmitters is not surprising because the different agonist structures inherently require different chemical counterparts in the binding site.

In summary, our observations lead to identification of a triad of conserved residues as an initial link that couples agonist binding to channel gating. All three residues of the triad localize to the same three-dimensional region of our structural model of the receptor ligand binding domain, and all three residues are conserved across α subunits. Also, equivalent residues in AChBP change conformation when the agonist is bound. Finally, structurally conservative mutations of each residue predominantly impair channel gating, and two pairs within the triad, α145/α200 and α145/α190, are interdependent in contributing to gating. Our findings suggest future studies aimed at identifying additional links that couple agonist binding to channel gating.

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