Functional equivalence of the nicotinic acetylcholine receptor transmitter binding sites in the open state

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Functional equivalence of the nicotinic acetylcholine receptor transmitter binding sites in the open state

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The subunits of the muscle-type nicotinic acetylcholine receptor (AChR) are not uniformly oriented in the resting closed conformation: the two α subunits are rotated relative to their non-α subunits. In contrast, all the subunits overlay well with one another when agonist is bound to the AChR, suggesting that they are uniformly oriented in the open receptor. This gating-dependent increase in orientational uniformity due to rotation of the α subunits might affect the relative affinities of the two transmitter binding sites, making the two affinities dissimilar (functionally non-equivalent) in the initial ligand-bound closed state but similar (functionally equivalent) in the open state. To test this hypothesis, we measured single-channel activity of the G153S gain-of-function mutant receptor evoked by choline, and estimated the resting closed-state and open-state affinities of the two transmitter binding sites. Both model-independent analyses and maximum-likelihood estimation of microscopic rate constants indicate that channel opening makes the binding sites’ affinities more similar to each other. These results support the hypothesis that open-state affinities to the transmitter binding sites are primarily determined by the α subunits.

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1. Introduction

One of the central aims in nicotinic acetylcholine receptor (AChR) physiology and pharmacology is to understand the dynamic interaction between ligands and the transmitter binding sites (TBSs) in the major AChR conformations: the resting closed state (also referred to as the resting state), the open ion-conducting state, and the desensitized closed state (also referred to as the desensitized state) [1]. To develop small-molecule agonists and antagonists with therapeutic potential, it would be useful to understand the state-dependent structural contributions to ligand binding at the TBSs. Biophysical, biochemical, and electrophysiological methods have been used to investigate agonist and antagonist binding to the AChR [1–6], including cryo-electron microscopy of the intact AChR [7], X-ray crystallography of acetylcholine-binding proteins (AChBPs) that are homologous to the intact receptor’s TBSs [8–10], X-ray crystallography of the mouse AChR α subunit extracellular domain [11], and X-ray crystallography of a pentameric prokaryotic ligand gated channel structurally similar to AChR models [12]. These structural studies have provided high-resolution models of the agonist-free and agonist-bound receptor; however, there are currently no high-resolution structures of the intact AChR in the functionally relevant agonist-bound resting and agonist-bound open states [2]. The desensitized closed conformations that predominate when agonist is continuously present for long durations (~100 ms or more) are more accessible (and interesting in their own right), but are likely to differ significantly in structure from the resting closed state [13]. Capturing the molecular details of ligand interactions in the transient open state therefore remains a challenging task.

One major conformational change that has been proposed to occur when the AChR opens is the rotation of the α subunits. The muscle-type AChR is a pentameric complex made up of two α subunits, one β subunit, one δ subunit, and either one ε or one γ subunit, arranged in clockwise order αβδεγ(γ) when viewed from the synaptic cleft. Cryo-electron microscopy experiments have shown that in the resting closed state, the α subunits overlay well with one another, and the non-α subunits overlay well with one another; however, the α subunits do not overlap well with the non-α subunits [7]. This non-equivalence of subunit orientation in the AChR tertiary structure occurs because the α subunits’ interior β-sheets are rotated approximately 10°. In the open state, a 9 Å-resolution model suggests the α subunits rotate such that all subunits overlay well, and the open AChR is highly uniform with respect to subunit orientations (Fig. 1) [14]. Molecular dynamics has also furnished useful insights into mechanisms of channel gating [15]. In the case of the AChR, molecular dynamics studies support the proposal that twisting or tilting reorientations of the M1 and M2 helices occur in the gating conformational change for muscle type [16] and α7 [17] receptors.

If tertiary subunit orientations directly affect the structure of the transmitter binding sites, the state-dependent change in the uniformity of subunit orientations would predict a state-dependent change in the uniformity of binding site affinities, because the α subunits make up part of the TBSs. The adult muscle-type AChR’s two
transmitter binding sites (TBSSs) are situated at the α-δ and α-ε subunit interfaces in the ligand-binding domain. Each TBS has a “principal face” made up of Loops A–C and a “complementary face” made up of Loops D–G [1]. The δ and ε subunits contribute the complementary components at their respective TBSs, and the α subunits contribute the principal components at both TBSs. State-dependent conformational changes in the α subunits may therefore affect the TBS’s resting-state and open-state binding affinities.

There are three plausible hypotheses for how the gating conformational change might affect the relative affinities of the two TBSs. First, the difference in the TBS’s open-state affinities (measured as $j_o$, the dissociation constant for binding to the open conformation) might be larger than the difference in their resting-state affinities (measured as $K_d$, the more familiar dissociation constant for binding to the resting closed conformation). Because the two TBSSs differ structurally in their complementary faces, this observation would suggest that the complementary faces increase their contributions to agonist binding in the open state. Second, the difference in affinities might remain the same in the resting and open states. This observation would suggest that increased open-state affinity is due to improved binding to both faces. Third, the difference in open-state affinities might be smaller than the difference in their resting-state affinities. Because the two TBSSs have identical principal faces, this observation would suggest that the principal faces increase their contributions to binding in the open state. We hypothesize that the third possibility is correct: the resting AChR TBSS are structurally and functionally non-equivalent, while the open AChR, in which subunit orientations are more uniform, has TBSSs that are equivalent with respect to open-state affinities. Formally, the subunit orientations need not be correlated with the affinities of the TBSSs, but the importance of the subunit rotations for gating suggests that a structure–function relationship may exist in which the rotation of the two α subunits affects the two different agonist binding site environments.

Previous work suggests that the difference in resting closed state affinities between the sites can be significant, but that differences in desensitized state affinities tend to be small. Functional studies on intact receptors from a variety of species and subtypes have shown that the resting-state binding sites are functionally non-equivalent with respect to their affinities and association/dissociation kinetics [18–23]. Although the two TBSSs of the adult-type AChR have similar resting-state affinities for acetylcholine [19], they have distinctly dissimilar affinities for other agonists such as epibatidine [24]. In the desensitized state, ligand-binding and stopped-flow fluorescence experiments on intact receptors indicate that the differences between the two sites are much smaller [25,26]. However, information about agonist binding in the transient open state has not been easily accessible in functional studies, and it is not clear what differences there may be between the open and desensitized states.

To test the hypothesis that the two TBSSs have equivalent binding affinities in the open state, we carried out an analysis of single-channel conformational changes for the αG153S gain-of-function mutant and αG153S/wild-type hybrid channels. The αG153S mutant [20,27] permits a larger range of activity to be studied than the wild-type, and it preserves the topography of subunit–subunit interactions in the wild-type receptor by virtue of having the mutated residue in the α subunit. The kinetics of activation of this mutant using acetylcholine [19] and choline have previously been reported [19,20,27], but the thermodynamics of binding to the open state were not investigated. The low-efficacy agonist choline was used in combination with the αG153S mutant because it stimulates gating that is not faster than conventional patch-clamp recording bandwidth [28–30], while retaining the gating mechanism observed for stronger agonists [31]. For choline-evoked αG153S AChR currents, a model-free analysis of open probabilities and dwell times supports the hypothesis that the TBSSs are functionally non-equivalent in the resting, diliganded (but non-desensitized) closed state but equivalent in the open state. Maximum-likelihood model fitting of single-channel activity was then used to determine resting-state and open-state affinities. The results of the model fitting support the hypothesis that the thermodynamics of binding reactions differ substantially for the two TBSSs in the ligand-bound resting state but become equivalent in the open state. One possible explanation for the functional equivalence of binding sites in the open state is that open-state affinities are primarily determined by the α subunits, a conclusion that is consistent with existing structural data.

2. Materials and methods

2.1. AChR expression

Cell culture reagents were from Invitrogen (Carlsbad, CA). Plasmids for expression of $\alpha$, $\beta$, $\delta$, and $\varepsilon$ subunits were generously provided by Professor Anthony Auerbach (SUNY Buffalo, Buffalo, NY) [19,32]. The αG153S mutation was engineered by site-directed mutagenesis using a Qiagen QuickChange Kit (Valencia, CA) [19,20,27]. Plasmid sequences were confirmed by dideoxy sequencing at the MIT Biopolymers Laboratory (Cambridge, MA).

HEK-293 human embryonic kidney cells (ATCC CRL-1573) were maintained in Dulbecco’s Minimum Essential Media (DMEM) supplemented with 10% Fetal Bovine Serum (FBS) at 37 °C in a 5% CO2 humidified atmosphere. Cells were transfected at 40–60% confluency using the method of calcium phosphate precipitation according to previously published protocols [19]. For one 35 mm dish, a total of 3.5 μg of plasmid DNA was used in a mass ratio of 2:1:1:1 of $\alpha$:$\beta$:$\delta$:$\varepsilon$ subunits. Media was changed 8–30 h after addition of DNA, and patch-clamp experiments were conducted 24–72 h after media change.

2.2. Patch-clamp recordings

Choline chloride was obtained from Sigma (St. Louis, MO). Single-channel recording was performed in the cell-attached mode according to previously published protocols [33,34]. The bath solution was Dulbecco’s Phosphate Buffered Saline (DPBS) containing (in mM): 137 NaCl, 0.9 CaCl2, 0.5 mM MgCl2, 2.7 KCl, 1.5 KH2PO4, 8.1 Na2PO4, pH 7.3.
Pipette solutions were DPBS supplemented with choline. Membrane potentials were typically — 30 to — 40 mV, and pipettes were held at a command voltage of — 70 mV during recording. Single-channel currents were amplified with an Axopatch 200B (Axon Instruments, Foster City, CA) and recorded through a low-pass Bessel filter at 10 kHz. Data were digitized at a sampling rate of 20—100 kHz using a NI 6040 E Data Acquisition Board (National Instruments, Austin, TX). Data were recorded using QuB software (www.qub.buffalo.edu) [35—39] and resampled to 20 kHz during idealization for consistency when necessary. The baselines of single-channel records were adjusted manually using QuB. A 5 kHz Gaussian digital filter was applied, and records were idealized using either the segmental k-means or half-amplitude algorithms in QuB [35]. All records were examined visually in their entirety, and misidealizations were corrected manually.

2.3. Kinetic analysis

Analysis of single-channel clusters was performed as previously described [19,34,40]. At high agonist concentrations, single-channel activity occurs as clusters of openings and closings that represent activity from one AChR. Each cluster is a series of openings flanked by long closed durations in which all channels are desensitized. These flanking desensitized durations are longer than a critical time (τ_{crit}). The value of τ_{crit} is assigned between the major closed component (non-desensitized sojourns) and its successor in closed-time distribution. The major closed component scales with agonist concentration and reflects transitions between AChR resting and open conformations, including binding and gating steps. The value of τ_{crit} chosen to minimize the percentage of misclassified events, and the fraction of misclassified events was typically less than 5%. Performing the kinetic analysis on clusters of events allows transitions between open states and resting (non-desensitized) closed states, rather than desensitized closed states, to be selected for measurement of rate constants. Clusters with multiple-conductance levels (more than one channel) or fewer than five events were excluded. Homogeneous clusters were used if their activity fell within 1—2 standard deviations (SDs) of the mean current amplitude, mean closed time, and mean open time [19]. Single-channel clusters recorded at 0.25, 0.5, 1, 2, and 5 mM choline were globally fitted to AChR gating models (Schemes 1 and 2), using the maximum interval likelihood algorithm with missed event correction. A dead time of 2.5 times the sampling interval was imposed. Fitting was repeated multiple times (10—50 repetitions) for each model with rate constant perturbations to provide higher confidence that the fitted model did not represent erroneous convergence. An additional gap state has been previously described and used in AChR gating models [19]; however, a gap state did not significantly improve fitting here. Other models incorporating choline-dependent block or additional dilligand open states also did not improve fitting.

2.4. Justification for assuming the AChR obeys detailed balance

Detailed balance requires that the AChR operate at thermodynamic equilibrium in the absence of a source of additional free energy. For the AChR, which has no enzymatic component, this free energy can only come from the ion gradient. Binding of a permeant ion to the channel pore might affect gating and cause violations of detailed balance. However, AChR permeant ion block is not detected in single-channel recordings, suggesting the ion gradient is not coupled to AChR gating. Consistent with this hypothesis, previous studies have not detected the temporal asymmetries in bursts of openings that would be expected for violations of detailed balance [41].

3. Results

To determine if the TBSs are functionally non-equivalent in the resting closed state and functionally equivalent in the open state, we performed kinetic analysis on hybrid channels formed from mixing wild-type and G153S mutant α subunits and analyzed open lifetimes of fully mutant channels without assuming a specific kinetic model. Two types of hybrid channel can be functionally expressed, since the binding sites are functionally non-equivalent, the hybrids are expected to exhibit distinct kinetic properties, while if the sites are functionally equivalent, the hybrids are expected to exhibit similar or indistinguishable kinetic properties. As a further test of the functional equivalence between TBSs, we estimated the affinities of the resting closed state and open states using maximum likelihood model fitting and rate estimation.

3.1. Analysis of hybrid channels

To determine the state-dependent changes in functional equivalence with respect to the TBSs, we first analyzed the activity of hybrid channels containing a wild-type and a mutant α subunit. Functional non-equivalence of the TBSs in the resting closed state has been established previously through biochemical and electrophysiological studies [18—24,42], but functional equivalence of the TBSs in the open

![Scheme 2. AChR kinetic model permitting non-equivalence of the binding sites. Low- and high-affinity sites and associated rate constants are denoted by superscript "L" and "H", respectively.](image)
state has not been investigated. One method for examining functional equivalence of the TBSs is the analysis of hybrid channels formed from mixtures of wild-type and mutant α subunits. For example, it has previously been reported that the TBSs are non-equivalent in the resting closed state and contribute unequally to gating [42]. Our results corroborate previous reports indicating the resting TBSs are functionally non-equivalent and provide new evidence that the open-state TBSs are functionally equivalent.

Currents were recorded from cells expressing both the wild-type α and mutant αG153S subunits, along with wild-type β, δ, and ε subunits. Because muscle-type AChRs contain two α subunits, channels can assemble in which both binding sites are wild-type, both sites are mutated, the α-δ site is mutated (hybrids), or the α-ε site is mutated (hybrids) [42]. Currents were recorded at 20 mM choline (4 patches) to saturate the receptors [28]. Four distinct modes of channel activity were observed in the distribution of open probabilities (Fig. 2). For equivalent sites that contribute equally to gating, single-channel activity from the two hybrid populations would be indistinguishable, as has been observed for transmembrane-region gating, single-channel activity from the two hybrid populations would be indistinguishable, as has been observed for transmembrane-region mutants [43–45]. In this case, three modes of activity would be observed: wild-type, hybrid, and mutant. In contrast, for non-equivalent sites that contribute unequally to gating, two distinguishable subsets of hybrid channel activity are predicted [42,45], leading to a prediction of four distinct modes of activity. Four modes of activity were observed, and a segmental k-means algorithm was used to fit the open probabilities to four Gaussian curves: wild-type, Pα = 0.12 ± 0.02; hybrid, Pδ = 0.33 ± 0.04; hybrid, Pε = 0.54 ± 0.04; and mutant, PG153S = 0.80 ± 0.03 (mean ± standard error) (Fig. 2). These results show that the TBSs contribute unequally to gating with the αG153S mutant in the presence of choline, consistent with previous results using cd200N and acetycholine [42].

To determine the functional equivalence of the TBSs in the open state, gating rate and equilibrium constants of the four populations were determined (Table 1). Although two hybrid activities were observed, the hybrid closing rate constants are equal, consistent with functionally identical open-state TBSs. Only one open lifetime was necessary to fit the experimental distribution of hybrid openings. However, two distinct open lifetimes might be too similar in magnitude to distinguish. The ratio of the two hybrid lifetimes is 1.1 ± 0.1 (mean ± standard error), indicating that if there are two distinct lifetimes, they are unlikely to differ by more than 20–30%.

Because the currents analyzed above were recorded at 20 mM choline to saturate receptors, unresolved fast blockade prolongs the apparent open events, resulting in underestimation of the diliganded closing rate α2 by 2-fold [28]. However, this phenomenon occurs for all four receptor populations. It therefore does not affect the number of kinetic components observed.

### Table 1 Maximum likelihood rate estimation of hybrid activity

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<th>θ2 (s−1)</th>
<th>α2 (s−1)</th>
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<td>Wild-type</td>
<td>0.15 ± 0.01</td>
<td>130 ± 40</td>
<td>900 ± 100</td>
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<tr>
<td>Hybrid</td>
<td>0.46 ± 0.04</td>
<td>250 ± 60</td>
<td>500 ± 100</td>
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<tr>
<td>Hybrid</td>
<td>1.1 ± 0.1</td>
<td>500 ± 100</td>
<td>500 ± 100</td>
</tr>
<tr>
<td>Mutant</td>
<td>2.8 ± 0.2</td>
<td>1300 ± 400</td>
<td>500 ± 200</td>
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3.2. Analysis of open lifetimes

To further test the functional equivalence of the open state in a model-independent analysis, we analyzed unliganded, monoligated, and diliganded open lifetimes of the αG153S mutant receptor. The hypothesis of functional non-equivalence in the open-state binding sites predicts that there are four AChR open states: an unliganded state, two distinct hybrid monoliganded states, and the diliganded state. For functionally equivalent open-state binding sites, only three open states are predicted, since the monoliganded states will be identical.

To determine the number of open states and their associated lifetimes, single-channel activity was measured at low choline concentrations (numbers of patches: 1 at 1 nM, 7 at 10 nM, 7 at 100 nM, 5 at 1 μM, 6 at 10 μM; 26 patches total). At low choline concentrations, single-channel activity is not clustered, and resting closed and desensitized sojourns cannot be distinguished. Thus, only open dwell-time distributions and their exponential open time components were analyzed (Fig. 3).

Open events could be classified into short (0.14 ± 0.02 ms, mean ± SD), intermediate (0.6 ± 0.1 ms), and long (1.2 ± 0.2 ms) lifetimes, consistent with prediction of three distinct states for equivalent open-state binding sites. The relative amplitudes of these open lifetimes were concentration dependent (Fig. 3). The short-lifetime open state steadily decreased in relative amplitude with increasing concentration. For the intermediate-lifetime open state, the relative amplitude exhibited a bell-shaped concentration dependence with a maximum at ~10−5 M. The long-lifetime open state was absent at very low concentrations and steadily increased in relative amplitude with increasing concentration. The magnitude of the three open lifetimes and the concentration dependencies of their relative amplitudes indicate that they correspond to openings of the unliganded, monoligated, and diliganded states.
diliganded αG153S AChR. The shortest lifetime is consistent with the reported unliganded lifetime [27] and can be assigned to the unliganded open state, in accord with the observation that increasing choline concentrations decrease the relative amplitude of this kinetic component. The longest lifetime is similar to the lifetime observed at saturating choline concentrations (Table 1) and can be assigned to the diliganded open state, consistent with the observation that increasing choline concentrations favor the longest open lifetime. The moniliganded open state is unlikely to have a lifetime shorter than the unliganded open state or longer than the diliganded open state. The intermediate open lifetime therefore can be assigned to a moniliganded open state, consistent with the observed bell-shaped concentration dependence.

A histogram of the open lifetimes observed in all low choline concentrations clearly shows three peaks (Fig. 3). A Gaussian was fitted to each peak, and the channel closing rate constants were determined: the unliganded closing rate constant \( \alpha_0 \) (7000 ± 1000 s\(^{-1}\)), the moniliganded closing rate constant \( \alpha_1 \) (1900 ± 400 s\(^{-1}\)), and the diliganded closing rate constant \( \alpha_2 \) (1100 ± 200 s\(^{-1}\)). It is not surprising that the unliganded and moniliganded open states are easily observable, as other gain-of-function mutants have shown a propensity for opening with fewer than two agonist molecules bound [46].

Notably, only one moniliganded open lifetime was observed, consistent with the hypothesis that the open-state TBSs are functionally identical. Additional moniliganded open states were not necessary given the number of events fitted. These data do not rule out the hypothesis that there are two distinct moniliganded open states, but these states would have lifetimes that differ by 10% or less (as judged by least-squares fitting of the histogram in Fig. 3).

### 3.3. Maximum-likelihood model fitting and rate estimation

To investigate the functional equivalence of the TBSs with respect to resting closed-state and open-state affinities, maximum likelihood model fitting and rate estimation was carried out. We examined two commonly used types of AChR kinetic models: a model which requires functionally equivalent binding sites (Scheme 1) [47] and a model which does not require functionally equivalent binding sites (Scheme 2) [19,20,48,49]. Both models are physically reasonable, and it is important to note that Scheme 2 allows for but does not require the possibility of non-equivalent affinities. Scheme 1 is often a good model for adult muscle-type AChR activity, because the two TBSs have similar resting closed-state agonist affinities for acetylcholine. However, several observations suggest Scheme 2 may be appropriate for analyzing our system. First, the αG153S mutant receptor appears to have distinct resting closed-state affinities for the two TBSs [19]. Second, mutant cycle analysis and the rate-equilibrium free energy relationship suggest that the αG153S mutation does not cause binding site coupling or change the allosteric gating mechanism (Supplementary information). Third, time constant analysis of resting closed times measured from clusters evoked at 0.25 mM choline were fitted best with four closed components, indicating two distinct binding steps may be kinetically distinguishable (Supplementary information). We performed maximum likelihood fitting of both models to establish that Scheme 2 is appropriate for estimating resting closed-state and open-state affinities. While recent results support the hypothesis that a second short-lived closed state (or “flip” state) is an intermediate on the gating pathway [50], the simpler model of Scheme 2 is sufficient to describe the data recorded under our experimental conditions, so we have chosen not to use a more complex model. Since use of the “flip” model yields gating rate constants that are relatively insensitive to structural perturbations in agonists, the open-state affinities we estimate here might be considered apparent affinities for the flip state if interpreted in the context of the “flip” model. Regardless of the model used, however, we have measured affinities in two distinct and pharmacologically important conformations.

To enable the fitting of the multi-state models employed [19,47], model-independent measurements of \( \beta_2 \), \( \alpha_0 \), and \( \alpha_2 \) were used to constrain the fits. To determine diliganded gating rate constants (\( \beta_2 \) and \( \alpha_2 \)), clustered single-channel activity was recorded at high concentrations of choline (numbers of patches: 4 at 0.25 mM, 4 at 0.5 mM, 4 at 1 mM, 5 at 2 mM, 4 at 5 mM; 21 total) and the dose-response curves were analyzed (Fig. 4A). The apparent opening rate,
\( \beta_2' \), is the reciprocal of the major intracluster resting closed-time component that scales with agonist concentration. The dose response of \( \beta_2' \) was fitted to a Hill equation to estimate the true microscopic opening rate constant \( (\beta_2 = 2200 \pm 700 \text{ s}^{-1}) \) (Fig. 4B) [19,34,51]. The diliganded closing rate constant, \( \alpha_2 \), was measured from the dose response of the mean open time \( \tau_O \). Choline is useful as a low-efficacy agonist that induces resolvable gating events, but fast open-channel blockade also occurs at high concentrations of this agonist.

When fast blockade is present, the observed mean intracluster open time is \( \tau_O = \alpha_2^{-1} + (\alpha_2 \cdot K_{\text{block}})^{-1} \cdot [\text{choline}] \) (Fig. 4C) [34,52,53]. Using this relationship, the diliganded closing rate constant \( (\alpha_2 \approx 1300 \pm 100 \text{ s}^{-1}) \) and choline blocking equilibrium constant \( (K_{\text{block}} \sim 10^{-2} \text{ M}) \) were determined. Fast choline block of the mutant is not different from the wild-type channel \( (K_{\text{block}} \sim 10^{-20} \text{ mM}) \), as expected from the location of residue G153 in the binding site, far from the ion conduction pore. The concentration dependence of \( \tau_O \) was observed to be shallow and linear, indicating that fast choline block is not severe up to 5 mM and is unlikely to skew rate constant estimation. Monoliganded openings still occur frequently at these concentrations, and they are included in the models for fitting. Only one monoliganded lifetime was observed above, and the monoliganded closing rate was therefore constrained to \( \alpha_1 = 1900 \pm 400 \text{ s}^{-1} \) for both monoliganded open states in Scheme 2 during fitting. Unliganded openings are not

### Table 2

Maximum interval likelihood rate estimation

<table>
<thead>
<tr>
<th>Scheme 1</th>
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<tr>
<td>( k_+ )</td>
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*Association rate constants are in M\(^{-1}\)·s\(^{-1}\), and all other rate constants are s\(^{-1}\). Dissociation constants are in \( \mu \text{M} \), and equilibrium gating constants are unitless. Rate constants were fixed at measured values during fitting as explained in the text.*
To understand the differences in binding and dissociation kinetics, we investigated the association and dissociation rates of choline binding to the α1 subunit of the AChR. Our data show that the rates of choline association and dissociation differ substantially and are state-dependent.

The kinetics of choline association and dissociation differ substantially. The association rates differ by orders of magnitude, with higher rates observed for the open state compared to the resting state. This is consistent with previous studies that have shown differential agonist binding to the resting and open states of the AChR.

We used the maximum interval likelihood algorithm in the QuB software suite to fit our data to Scheme 1 or Scheme 2. Scheme 2 improved the log-likelihood score by 18936 compared to fitting Scheme 1 with only three additional free parameters. This difference is significant by the Schwartz criterion and likelihood-ratio tests. Both models were reason-ably well fitted to the observed dwell time distributions.

4. Discussion

4.1. State-dependent changes in binding site organization

Single-channel kinetic analysis of the αC153S AChR activated by choline has demonstrated that the two TBSs have non-equivalent resting-state affinities but equal open-state affinities. Structural data suggest that the channel subunits are non-uniformly oriented in the resting AChR: the two α subunits are rotated relative to the non-α subunits. In contrast, all subunits have equivalent conformations in the open AChR, and the ligand-binding domains are equivalent with respect to the subunit orientation. Our results suggest that there is a correlation between the subunit orientation, the structural organization of the two TBSs, and the function of the TBSs.

Regardless of the state of the receptor, the differences in binding affinity and selectivity between the two TBSs may be attributed largely to residues on the complementary face in Loops D–G. Because the principal face is formed by α subunits at both binding sites, binding interactions between loops A–C (which come from the α subunit) and the agonist are expected to be similar for the two sites. The complementary face is formed by the δ subunit at one TBS and the ε subunit at the other TBS. The non-equivalence of resting-state agonist affinities suggests that the complementary faces contribute appreciably to ligand–channel interactions. Several residues on the complementary face have been implicated in determining differential agonist and competitive antagonist affinities and specificities between the two sites in the closed AChR.

We observe that the TBSs have equal open-state affinities, suggesting that their principal faces increase their fractional contributions to agonist binding in the open state compared to the resting closed state. This is consistent with previous studies that have shown differential agonist binding to the resting and open states of the intact receptor. The agonist-bound AChBP structures are thought to mimic the desensitized state of the intact receptor’s TBSs, but it is not clear what differences exist between the open-state and desensitized state TBSs. However, our results are consistent with the hypothesis that structural models of the agonist-free resting closed state and the agonist-bound desensitized state are similar to the agonist-bound resting and open states, respectively. In X-ray crystal structures of carbamylcholine- or nicotine-bound AChBP, the agonist-buried surface area is approximately 2-fold greater for the principal face than the complementary face, suggesting that the agonist makes a larger number of contacts with the principal face. The bound agonist may also be stabilized by closure of Loop C and compaction of the aromatic “cage”, both α subunit components. Our results are therefore also consistent with structural similarity between the open and desensitized states of the TBSs.

Both of these states are high affinity conformations with slow rates of agonist dissociation (0.1–10 s⁻¹) and the binding affinities for the two TBSs in the...
Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.bbamem.2009.01.009.

References

2. C.A. Cay, J.L. Vake, Gating of nicotinic ACh receptors; new insights into structural transitions triggered by agonist binding that induce channel opening, J. Physiol. 584 (2007) 727–733.
22. Y. Purohit, C. Grosman, Block of muscle nicotinic receptors by choline suggests that the activation and desensitization gates act as distinct molecular entities, J. Gen. Physiol. 127 (2006) 703–717.
33. A. Auerbach, C.J. Lingle, Heterogeneous kinetic properties of acetylcholine receptor channels in Xenopus myocytes, J. Physiol. 378 (1986) 119–140.
35. G. Akk, S. Sine, A. Auerbach, Binding site contributions are unequal to the gating of mouse nicotinic alpha D200N acetylcholine receptors, J. Physiol. 496 (1996) 185–196.