Cellular/Molecular

An Arginine Involved in GABA Binding and Unbinding But Not Gating of the GABA<sub>A</sub> Receptor

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GABA<sub>A</sub> receptor function can be conceptually divided into interactions between ligand and receptor (binding) and the opening and closing of the ligand-bound channel (gating). The relationship between binding, gating, and receptor structure remains unclear. Studies of mutations have identified many amino acid residues that contribute to the GABA-binding site. Most of these studies assayed changes in GABA dose–response curves, which are macroscopic measures that depend on the interplay of many processes and cannot resolve individual microscopic transitions. Understanding the microscopic basis of binding and gating is critical, because kinetic transition rates predict how receptors will behave at synapses. Furthermore, microscopic rates are directly related to the molecular interactions underlying receptor function.

Here, we focused on a residue (β<sub>2</sub>-R207) previously identified as lining the GABA-binding site that, when mutated to cysteine, greatly reduces apparent GABA affinity and was predicted to affect both binding and gating. To better understand the role of β<sub>2</sub>-R207, we expressed α<sub>1</sub>β<sub>2</sub> and α<sub>1</sub>β<sub>2</sub>-R207C receptors in human embryonic kidney 293 cells and studied receptor kinetics using fast solution applications. The mutation accelerated deactivation by 10-fold, without altering desensitization in the presence of saturating GABA. Maximum open probability and single-channel open times were also unaltered by the mutation, but the GABA-binding rate was reduced eightfold. Therefore, the effects of this mutation in a predicted binding site residue are solely attributable to changes in GABA-binding and unbinding kinetics, with no changes in channel gating. Because β<sub>2</sub>-R207 stabilizes GABA in the binding pocket, it may directly contact the GABA molecule.

Key words: kinetics; HEK293 cells; mutation; single channel; agonist; antagonist

Introduction

During fast synaptic transmission, neurotransmitter activates receptors in the postsynaptic membrane. The amplitude and time course of the subsequent postsynaptic conductance depend on the two interrelated processes of binding (the interaction of receptor and ligand) and gating (transitions of the bound receptor between open, closed, and desensitized states). Binding is mediated by molecular interactions between the ligand and amino acid residues that line the ligand-binding pocket, that then trigger gating.

A description of binding-pocket structure is rapidly emerging for the superfAMILY of ligand-gated ion channels, which includes the nicotinic acetylcholine, glycine, serotonin type 3, and GABA<sub>A</sub> receptors. Site-directed mutagenesis, cysteine-scanning mutagenesis, and photoaffinity labeling have shown that the ligand-binding pocket is at the interface between two subunits, with each subunit contributing multiple discontinuous “loops” (for review, see Galzi, 1994). For example, in the GABA<sub>A</sub> receptor, the α and β subunits each contribute three regions to the binding pocket (Sigel et al., 1992; Amin and Weiss, 1993; Smith and Olsen, 1994; Westh-Hansen et al., 1997, 1999; Boileau et al., 1999, 2002; Wagner and Czajkowski, 2001; Holden and Czajkowski, 2002; Newell and Czajkowski, 2003).

To evaluate the role of individual residues in binding or gating, most previous studies have examined the effects of mutations on ligand dose–response curves. This approach measures macroscopic parameters such as the concentration of ligand that gives the half-maximal response (EC<sub>50</sub>), which is itself determined by the more fundamental microscopic processes of ligand binding–unbinding, channel opening–closing, and desensitization–resensitization. Dose–response curves cannot directly reveal these microscopic processes (Colquhoun, 1998). For wild-type GABA<sub>A</sub> receptors, significant progress toward understanding microscopic kinetics has been made using single-channel recording and rapid ligand application techniques to develop kinetic models of receptor function and to estimate individual rate constants (Twyman et al., 1990; Celentano and Wong, 1994; Jones and Westbrook, 1995; Lavoie and Twyman, 1996; Jones et al., 1998; Haas and Macdonald, 1999; Burkat et al., 2001). Although the combination of kinetic approaches with mutagenesis would obviously be more powerful than either method alone, only a few studies have examined the effects of mutations on microscopic kinetics (Chang and Weiss, 1999; Bianchi and Macdonald, 2001; Scheller and Forman, 2002).

In this study, we applied the techniques of rapid ligand application, single-channel recording, and kinetic modeling to clarify...
the biophysical role that the arginine at position 207 of the β2 subunit of the rat GABA<sub>A</sub> receptor plays in channel function. This residue is likely to face into the GABA-binding pocket, and its mutation to cysteine causes a 70-fold increase in the EC<sub>50</sub> for GABA (Wagner and Czajkowski, 2001). The present experiments demonstrate that the mutation, β2-R207C, causes a 20-fold increase in the ligand-unbinding rate, an eightfold decrease in the ligand-binding rate, and has no effect on any of the rates associated with gating. Therefore, β2-R207 is important for stabilizing the agonist–receptor complex, and may enjoy a direct interaction with the GABA molecule.

Materials and Methods

Cell culture and transfection. Human embryonic kidney (HEK-293) cells were cultured in Minimum Essential Medium with Earle's salts (Invitrogen, Gaithersburg, MD) containing 10% fetal bovine serum (Sigma-Aldrich, St. Louis, MO) in a 37°C incubator under a 5% CO<sub>2</sub> atmosphere. When the cells reached 60–90% confluence in 60 mm culture dishes, they were transfected with 1 μg each of α<sub>1</sub>-FLAG and β<sub>2</sub> (wild type or R207C) rat cDNAs subcloned into the pCEP4 vector (Invitrogen) using the LipofectAMINE2000 reagent (Invitrogen) according to the supplied protocol. At 20–28 hr after transfection, the cells were replated on 12 mm sterile glass coverslips, transferred to a 30°C incubator, and recorded at 24–96 hr after replating. Transfected cells were visualized by incubation with anti-FLAG antibody-coated beads (Sigma-Aldrich), which recognize the flag epitope sequence (DYKDDDDK) inserted between the sixth and seventh amino acid of the mature α<sub>1</sub> subunit. The flag epitope insertion has been previously not to alter function (Connolly et al., 1996).

Patch-clamp electrophysiology. Recordings from outside-out patches excised from HEK cells were made using borosilicate glass pipettes filled with (in ms): 140 KCl, 10 EGTA, 2 MgATP, 20 phosphocreatine, and 10 HEPES, pH 7.3. For macroscopic current recording, patches were voltage clamped at −60 mV and placed in the stream of a multibarreled flowpipe array (Vitrodynamics, Rockaway, NJ) mounted on a piezoelectric bimorph (Vernontron, Bedford, OH). A computer-controlled current source drove the bimorph to move solution interfaces over the patch with 10–90% exchange times of ~200 μsec, as measured by the liquid junction current at the open pipette tip after each experiment. GABA and SR-95531 were dissolved in the perfusion solution, which contained (in ms): 145 NaCl, 2.5 KCl, 1 MgCl<sub>2</sub>, 10 HEPES, 4 mM Glucose, pH 7.4, 315 mOsm. For extracellular solutions that contained >10 mM K<sup>+</sup>-gluconate was added to maintain a constant Cl<sup>−</sup> driving force. GABA and SR-95531 were obtained from Sigma-Aldrich Chemicals. Currents were low-pass filtered at 2–5 kHz with a four-pole Bessel filter and digitized at a rate no less than twice the filter frequency. Data were collected using an Axopatch 200B amplifier and Digidata 1320A digitizer, controlled by AxoGraph software (Axon Instruments, Foster City, CA) running on a Macintosh G4 (Apple Computers, Cupertino, CA). Curve fitting was also performed using AxoGraph software.

Single-channel currents were recorded from patches held at −80 mV, sampled at 20 kHz, and filtered at 2 kHz. Before analysis, the current records were digitally refiltered using a 1 kHz Gaussian filter. Events were detected using amplitude windows on the basis of a 15 pS conductance level, with homegrown routines in Matlab 5.2 (Mathworks, Natick, MA). Events that included multiple openings were discarded. Open time distributions were fit by the maximum likelihood method, with corrections for missed events, and the number of open times was determined using the log-likelihood ratio test (Colquhoun and Sigworth, 1995). Only events lasting ≥200 μsec were accepted for analysis.

Peak open probability analysis. Nonstationary variance analysis (Sigworth, 1980) was performed on currents recorded from a series of saturating GABA pulses (2 msec, 10 sec intervals) from which ensemble mean current (I<sub>me</sub>) and variance (σ<sup>2</sup>) were calculated for each data point. The mean current was divided into 100 equally sized bins, and the corresponding variances were averaged. Plots of binned variance versus current were fit with the equation: σ<sup>2</sup> = I<sub>me</sub> − I<sup>2</sup>N<sup>−1</sup>, where I is the single-
channel current, and $N$ is the number of channels. Variance resulting from slow drift (i.e., rundown or run-up) was corrected by local linear fitting of the drift, calculating the variance attributable to this trend at each point, and subtracting the drift variance (scaled by the square of normalized current amplitude) from the calculated variance before fitting. This method yields accurate estimates of $r$ and $N$ when tested on simulated data with drift.

**Kinetic modeling.** Kinetic modeling was performed with homewritten software using the Q-matrix method (Colquhoun and Hawkes, 1995a,b). During optimization, the rate constants $k_{i\rightarrow i}$, $\alpha$, $k_{i\rightarrow i\text{on-ant}}$, and $k_{i\rightarrow i\text{off-ant}}$ were constrained to values obtained from experiments presented in this study. The unconstrained rate constants were optimized by a simultaneous fit to current responses from wild-type and mutant receptors to 2 and 500 mM saturating (10 mM) GABA pulses and a paired pulse protocol (2 msec, 10 mM pulses separated by 0.1, 0.3, and 1 sec). After simultaneous optimization, all of the rate constants for $k_{i\rightarrow i}$, $d_{r}$, and $p$ were fixed, and the model was optimized to fit individual responses to 2 msec (six patches) and 500 msec (five patches) pulses of 10 mM GABA. The final values for $k_{i\rightarrow i}$, $d_{r}$, and $p$ are the means of the values collected from these 11 individual optimizations (Fig. 6). Optimization used a simplex algorithm to minimize the amplitude-weighted sum of squared errors between actual and simulated currents. In all cases, significant differences were tested using a two-tailed unpaired Student’s $t$ test at a significance level of $p = 0.05$.

**Results**

**Mutation of $\beta_{2}$-R207 alters GABA$_{A}$ receptor kinetics**

We initially characterized the effect of the $\beta_{2}$-R207C mutation on receptor kinetics by observing macroscopic deactivation (relaxation after removal of agonist) and desensitization (relaxation in the presence of saturating agonist). After a brief (2 msec) pulse of saturating GABA, currents from $\alpha_{2}\beta_{2}$-R207C receptors deactivated 10 times faster than currents from wild-type receptors (Fig. 1A, Table 1). Because the time course of deactivation depends on the fraction of time spent in bound states and thus predict a rightward shift in the macroscopic dose–response curve. Alternatively, faster deactivation could be attributable to increased desensitization, which would increase the fraction of time in bound states and thus predict a leftward shift. Figure 1C illustrates that the $\beta_{2}$-R207C mutation caused a 40-fold rightward shift in the dose–response curve for peak current and a 340-fold rightward shift for current measured at 500 msec. Therefore, the mutation reduces the fraction of time spent in bound states, consistent with an accelerated GABA-unbinding rate.

Another phenomenon that depends on the interplay between gating and binding is paired-pulse depression. When pairs of brief (2–3 msec) saturating GABA pulses were given to $\alpha_{2}\beta_{2}$ receptors at variable intervals, the second pulse evoked a smaller peak response that recovered as the inter-pulse interval grew longer (Fig. 2A). In patches containing $\alpha_{2}\beta_{2}$-R207C receptors, the depression of the second pulse was significantly reduced, and the receptors recovered more quickly than wild type (Fig. 2B,C). This result could imply that the desensitized state in the mutant receptor has a slower entry rate or a shorter lifetime. However, it is difficult to reconcile this interpretation with the fact that the mutation does not affect desensitization during long pulses (Fig. 1). Furthermore, it has been shown that paired-pulse depression of neuronal GABA$_{A}$ receptors is similarly attenuated when using $\beta$-alanine, an agonist that has a faster unbinding rate than GABA but does not affect gating (Jones and Westbrook, 1995; Jones et al., 1998). Therefore, reduced paired-pulse depression in $\alpha_{2}\beta_{2}$-R207C receptors is also likely to be caused by an increase in the GABA-unbinding rate.

**The mutation does not alter peak open probability or single-channel open times**

The hypothesis that the $\beta_{2}$-R207C mutation does not affect gating is supported by the fact that desensitization during long GABA pulses is indistinguishable between mutant and wild-type receptors. Alternatively, it is possible that the $\beta_{2}$-R207C mutation causes precisely balanced shifts in multiple gating rates so that there is no obvious effect on macroscopic desensitization. We tested whether this might be the case by measuring the peak open probability in response to a saturating GABA pulse ($P_{\text{o}}$). Like macroscopic desensitization, $P_{\text{o}}$ depends on the interplay of several gating processes. We measured $P_{\text{o}}$ by nonstationary variance analysis (Sigworth, 1980) and found no significant difference between mutant and wild-type receptors (Fig. 3), further supporting the idea that the $\beta_{2}$-R207C mutation does not affect gating.

To further test whether the $\beta_{2}$-R207C mutation affects gating,
0.8 msec, 0.06 (six patches; mean ± SD). Wild-type and α₂β₂-R207C receptors both displayed two conductance states. Wild-type conductances were 14.6 ± 0.6 pS (20%) and 10.5 ± 0.8 pS (six patches). R207C conductances were 12.9 ± 0.5 pS (23%) and 9.3 ± 0.4 pS (five patches; mean ± SD). We do not believe that these minor, but significant (p < 0.01), differences are caused by the mutation. Rather, they most likely occur because of the lower chloride concentration in the mutant recording solutions ([Cl]wt, 90 mM; [Cl]mut, 40 mM), which was reduced to accommodate the addition of 100 mM GABA (see Materials and Methods).

**Mutation of β₂-R207 reduces the binding rate of GABA**

When comparing α₁β₂ and α₂β₂-R207C receptors, we could not detect any change in gating by examining desensitization, Pmax, or mean open times. Because the mutation does not affect gating, it appears that the observed increase in deactivation rate and reduction in paired-pulse inhibition must be attributable to an increase in the GABA-unbinding rate (koff-GABA). One might expect a binding site mutation that effects the unbinding rate to simultaneously alter the binding rate (kon-GABA) (Jones et al., 1998). However, none of the data presented thus far can provide a measure of kon-GABA.

To measure kon-GABA, we performed experiments in which the competitive antagonist SR-95531 and GABA were simultaneously applied and allowed to compete with each other to occupy the receptor. In this “race” experiment, the percentage of channels occupied by each ligand at the peak of current depends on their relative binding rates. Therefore, if the SR-95531-binding rate (kon-SR) is known, the ratio of the peak current in GABA alone to the peak current in GABA plus SR-95531 (Ipeak-off-GABA) can be used to calculate kon-GABA (Clements et al., 1992; Jones et al., 1998). We determined kon-SR using the equation kon-SR = Kd-SRkon-GABA, where Kd-SR and kon-GABA (the dissociation constant and microscopic unbinding rate for SR-95531) were directly measured using the deconvolution method developed by Jones et al. (2001). This method exploits the fact that, after equilibration in competitive antagonist, the current evoked by a jump into GABA is shaped by the convolution of the time course of antagonist unbinding and the waveform of the control current (evoked with no pre-equilibration in antagonist). Therefore, the time course of antagonist unbinding can be obtained by deconvolving the control current from the current recorded after antagonist equilibration (Fig. 5A–C). The time course of antagonist unbinding is fit with an exponential function giving kon-GABA and the percentage of receptors occupied by antagonist at equilibrium (see Fig. 5 legend). Kd-SR is obtained by repeating this experiment with different concentrations of SR-95531 and plotting the antagonist occupancy versus concentration. We found that the mutation had no effect on the binding rate, unbinding rate, or dissociation constant for SR-95531 (kon-SR = 11 ± 3 sec⁻¹, Kd-SR = 210 nM, kon-GABA = (5.5 ± 1.7) × 10⁻⁷ M⁻¹s⁻¹, n = 11; α₂β₂-R207C: Kd-SR = 16 ± 6 sec⁻¹, Kd-SR = 180 nM, kon-SR = (8.9 ± 3.4) × 10⁻⁷ M⁻¹s⁻¹, n = 11; mean ± SD).

However, during race experiments that measure the ratio of current activated by coapplication of GABA and SR-95531 to that activated by GABA alone, α₂β₂-R207C receptors required ~10-fold higher concentrations of GABA to achieve the same current ratio (Ipeak-off-GABA) observed in α₁β₂ receptors (Fig. 5D, E). This difference indicates that the binding rate of GABA, relative to that of SR-95531, is much slower in mutant than in wild-type receptors. From these Ipeak-off-GABA values and the SR-95531-binding rates measured above, we computed kon-GABA using Equation 12 from Jones et al. (1998) and found that the mutation caused an eightfold decrease
in the GABA-binding rate \(k_{\alpha_2} = (6.85 \pm 2.8) \times 10^6 \text{ M}^{-1} \text{s}^{-1}, n = 14\); \(\alpha_2\beta_2\)-R207C = \(0.8 \pm 0.3\) \(\times 10^6 \text{ M}^{-1} \text{s}^{-1}, n = 13\); mean \(\pm\) SD; \(p < 0.001\).

An interesting result observed during these experiments is that when GABA and SR-95531 were coapplied, there was an apparent increase in the rate of macroscopic desensitization that was more pronounced in the mutant receptor. It is possible that SR-95531 causes this effect by altering receptor gating. However, a simpler explanation is that as the system goes to equilibrium, some receptors unbind GABA, allowing SR-95531 to occupy a larger percentage of receptors than it did during the peak current. To test this idea, we simulated race experiments for both wild-type and mutant receptors using a model in which SR-95531 acts as a pure competitive antagonist and has no effect on GABA-mediated gating (see model and rates in Fig. 6). As seen in actual currents, the presence of antagonist increased apparent desensitization of simulated currents (Fig. 5 D, E, insets), demonstrating that this phenomenon occurs even if SR-95531 has no effect on channel gating.

The effects of the mutation can be explained solely by altering binding and unbinding rates

If, as the data suggest, mutation of \(\beta_2\)-R207 affects only agonist binding and unbinding, then a kinetic model that describes \(\alpha_2\beta_2\) receptors should be convertible into one that describes \(\alpha_2\beta_2\)-R207C receptors simply by adjusting \(k_{\text{on,GABA}}\) and \(k_{\text{off,GABA}}\) while leaving all other rates unchanged. A kinetic model that accurately describes the behavior of \(\alpha_2\beta_2\) receptors was developed using the model shown in Figure 6A (Jones et al., 1998). Before optimization, \(k_{\text{on}}\) was fixed to its measured rate, \(\alpha_2\) was set to the fastest measured closing rate, and \(\alpha_2\) was set to the intermediate closing rate. The longest open state, which accounts for \(\sim\)5% of all openings, was ignored. The remaining rate constants were allowed to vary while the model was optimized to simultaneously fit responses to short, long, and paired pulses of saturating GABA that were scaled to the measured value of \(P_{\text{o-max}}\) (see Materials and Methods). The optimized rate constants are shown in Figure 6B, and simulated currents generated with this model are overlaid on actual patch responses in Figure 6C. The model was then fit to currents recorded from \(\alpha_2\beta_2\)-R207C with \(k_{\text{on,GABA}}\) constrained to the value measured for mutant receptors. After optimization, the only unconstrained rate constant that was significantly different between the mutant and wild-type models was \(k_{\text{off,GABA}}\), which shifts from 19 sec\(^{-1}\) in wild-type receptors to 400 sec\(^{-1}\) in mutant receptors. As an independent check on the quality of the models, we simulated dose-response curves, which had not been used during optimization. The resulting EC\(_{50}\) values were: \(\alpha_2\beta_2\), 12 \(\mu\)M at peak and 0.8 \(\mu\)M at 500 msec; \(\alpha_2\beta_2\)-R207C, 620 \(\mu\)M at peak and 180 \(\mu\)M at 500 msec. In all cases, these numbers are within 25% of the actual values (10% on average), verifying that these models are good predictors of receptor behavior. In summary, all kinetic effects of the \(\beta_2\)-R207C mutation can be explained solely by an eightfold decrease in \(k_{\text{on,GABA}}\) coupled with a 20-fold increase in \(k_{\text{off,GABA}}\).

**Discussion**

Determining whether an amino acid residue is involved in ligand binding or channel gating through mutational analysis has proven to be a nontrivial problem. Mutational effects are commonly quantified by shifts in EC\(_{50}\) or \(K_I\). However, because these measures depend on both binding and gating, it is often impossible to distinguish between alterations in binding or gating using them (Colquhoun, 1998). Changes in maximum open probability are potentially useful for diagnosing gating changes. Unfortunately, the notoriously variable expression in heterologous systems makes it difficult to control for the number of receptors, so that only very large (\(>10\)-fold) shifts in peak current are reliably detectable. Despite these difficulties, techniques have been developed that can potentially identify binding or gating changes using EC\(_{50}\) and peak current measurements. For example, the relative efficacy of a partial agonist has been used as an internal control for receptor expression, and changes in relative efficacy have thus been interpreted as being caused by gating changes (Maksay et al., 2000; O’Shea and Harrison, 2000; Wagner and Czajkowski, 2001). In contrast, evidence for binding changes has included: (1) a change in the potency of GABA that is not accompanied by a change in the potency of a modulator that opens the channel but does not bind at the GABA-binding site, and (2) large changes in GABA potency in the absence of large changes in maximal cur-
rent (Amin and Weiss, 1993). Although these methods can suggest that a change in binding or gating has occurred, they cannot rule out whether both have changed or reveal which microscopic transitions were affected.

The limitations inherent in using doseresponse curves can be overcome by adopting methods that explicitly relate a structural change to changes in elementary functional properties, namely the conformational states of the receptor and the transition rates between them. To this end, we applied single-channel recording, rapid ligand application, and kinetic modeling to quantify the effects of mutating a predicted binding site residue on individual binding and gating processes. Our results lead to three conclusions: the \( \beta_2\)-R207C mutation (1) has no effect on gating, (2) greatly increases the likelihood that the GABA molecule will leave the binding site, and (3) greatly reduces the likelihood that GABA will bind in the first place.

**Mutation of \( \beta_2\)-R207C does not affect gating**

Multiple lines of evidence indicate that the mutation does not effect gating. Macroscopic desensitization and \( P_{o,\text{max}} \) both of which depend on the interplay of multiple gating processes, are not affected by the \( \beta_2\)-R207C mutation. It is possible that a mutation could induce a balanced shift in multiple microscopic gating rates so that neither of these macroscopic parameters appears to be affected. However, because individual transition rates influence desensitization and \( P_{o,\text{max}} \) to different degrees, it seems unlikely that alterations in multiple gating rates would exactly balance out so that both of these processes appear unchanged. In addition, mutant and wild-type receptors have indistinguishable single-channel open time distributions, demonstrating that the channel-closing rate is unaffected. This conclusion, of course, assumes that the measured events are actual openings and not bursts that appear to be openings because their closures are too brief to measure. Because the small conductance of these channels necessitates filtering at 1–2 kHz, the distributions in Figure 4 probably do contain at least some bursts. Ironically, this potential source of error further strengthens our conclusion that gating is not affected by the mutation. If the distributions are contaminated by bursts, yet are unaffected by the mutation, then the burst durations must be unaffected. Because burst duration depends on multiple gating steps (e.g., opening, closing, and desensitization) rather than merely channel closure, unchanged burst distributions imply that none of the intraburst-gating steps are affected.

We previously reported that this mutation affects gating as well as binding (Wagner and Czajkowski, 2001). This conclusion was reached by observation in oocytes, that the mutation altered the relative efficacy of the partial agonist piperidine-4-sulphonic acid (P4S; \( \alpha_1\beta_2\): \( I_{\text{peak-P4S}}/I_{\text{peak-GABA}} = 0.5; \alpha_1\beta_1\beta_2\)-R207C: \( I_{\text{peak-P4S}}/I_{\text{peak-GABA}} = 0.21 \)). This result conflicts with the previous finding that the mutation has no effect on gating. It could be that the mutation only alters P4S-mediated gating but not GABA-mediated gating. However, a simpler explanation is that the apparent efficacy change in oocytes is actually an artifact of slow solution exchange. When agonist is rapidly applied, the resulting current peaks within the first few milliseconds. In a typical oocyte experiment, the GABA concentration probably requires more than 1 sec to reach peak [Wagner and Czajkowski (2001) used a flow rate of 70 \( \mu l/sec \) and a bath volume of 100 \( \mu l \), which predicts an exchange time constant of 1.4 sec, assuming good mixing]. The simulations in Figure 7, A and B, show that slow application results in severe underestimation of peak current and falsely indicates a reduction in peak open probability for \( \alpha_1\beta_2\)-R207C receptors, effects not observed in rapid application experiments. In addition, simulations of slow application using the increased unbinding rate, but no gating changes as observed in the present study, showed a false change in the relative efficacy of P4S (Fig. 7C) (\( \alpha_1\beta_2\): \( I_{\text{peak-P4S}}/I_{\text{peak-GABA}} = 0.5; \alpha_1\beta_1\beta_2\)-R207C: \( I_{\text{peak-P4S}}/I_{\text{peak-GABA}} = 0.33 \)). In other words, a change in the relative efficacy of a partial agonist can lead to erroneous conclusions about gating if solution exchange is not sufficiently rapid (i.e., milliseconds or less for fast ligand-gated channels). Therefore, our previous prediction that \( \beta_2\)-R207C affected gating was an error, probably because of the fact that slow ligand application precludes accurate measurement of efficacy or even relative efficacy.

![Figure 5](image)

Figure 5. \( \beta_2\)-R207C reduces the GABA-binding rate. A, B, Competitive antagonist unbinding slows the rise of agonist activated current. Currents were evoked by 10 \( \mu M \) (wild type; A) or 100 \( \mu M \) (mutant; B) GABA alone in outside-out patches with and without pre-equilibration in 1 \( \mu M \) SR-95531. The unbinding rate of the competitive antagonist can be extracted from the data shown in A and B by deconvolving the control and experimental responses. C, Plots showing the deconvolution of currents with antagonist pre-equilibration from those without pre-equilibration. The data shown is the mean ± SEM of multiple deconvolutions (wild type, \( n = 5 \), mutant, \( n = 4 \)). The solid lines are curves fit to the data resulting from the deconvolutions with the equation \( A(t) = [P_o - (P_o - P_t)e^{-t/\tau_{\text{off}}}]N \), where \( A(t) \) is the fraction of receptors not occupied by SR-95531, \( P_o \) and \( P_t \) are the probabilities of being available initially (at \( t = 0 \)) and at steady state (at \( t = \infty \)), \( \tau_{\text{off}} \) is the time constant of antagonist unbinding at each site, and \( N \) is the number of binding sites. Best fits were always obtained with \( N = 1 \). A, A race experiment (see Results) on a patch containing wild-type receptors in which the current evoked by simultaneous application of 1 \( \mu M \) GABA and 100 \( \mu M \) SR-95531 is compared with the current evoked by 1 \( \mu M \) GABA alone. E, A race experiment on mutant receptors using 10 \( \mu M \) GABA and 100 \( \mu M \) SR-95531. The current ratios from D and E were used to compute \( K_{\text{m, GABA}} \), using Equation 12 from Jones et al. (1998). Inset traces in D and E are the result of simulations of the race experiments shown using the model and rate constants from Figure 6.
**β2-R207C alters both GABA binding and unbinding**

Because deactivation is accelerated in α4β2-R207C receptors while gating is unaltered, the mutation must increase the microscopic unbinding rate. An increased unbinding rate is also supported by the result that mutant receptors are less susceptible to, and recover more quickly from, paired-pulse depression. An increase in unbinding rate would have this effect by reducing the current peaks. This holds in wild-type receptors for which GABA and SR-95531 both have relatively slow unbinding rates. However, GABA unbinds from α4β2-R207C receptors much more quickly, which may give SR-95531 extra opportunities to compete with the agonist before the current peaks. In this case, the race equation yields a lower limit on k_{off,GABA}. Fortunately, only ~5% of singly bound GABA receptors will unbind rather than binding a second GABA or entering de-sensitized state after the channel closes. Finally, simulations demonstrate that an increase in k_{off,GABA} paired with a decrease in k_{on,GABA} are the only changes necessary to account for the kinetic effects of the mutation.

The case for a mutation-induced decrease in the GABA-binding rate is even more straightforward, because this rate was measured in race experiments. However, there are some assumptions that must be true in order for the I_{race} method of determining k_{on,GABA} to give accurate values. The original equation used to calculate k_{on,GABA} from I_{race} and k_{on,SR} (Jones et al., 1998) was derived under the assumption that there is only one binding site for SR-95531, and that SR-95531 binding precludes GABA binding at that site only. Under that scheme, if there are two GABA-binding sites, one molecule of GABA and one of SR-95531 could be bound at the same time, and the channel would be prevented from opening. The assumption of a single SR-95531-binding site was supported by Jones et al. (1998, 2001). In contrast, it remains unknown whether or not SR-95531 and GABA can co-occupy the receptor. In fact, the kinetic model used here does not allow co-occupancy (Fig. 6A). When the race equation is rederived under this assumption, the resulting k_{on,GABA} is half of the value obtained from the original equation, because GABA has two chances to bind for every chance had by SR-95531. The resulting simulations produced EC_{50} values that agreed with the experimentally determined values to within 10% on average (see Results). In contrast, the alternate assumption, that SR-95531 and GABA can co-occupy the receptor, resulted in a five-fold larger average discrepancy. These findings imply that binding of one SR-95531 molecule precludes GABA binding at both sites, not only to the site occupied by the antagonist. Because the two GABA-binding sites probably do not overlap physically, this mechanism requires some sort of allosteric exclusion between the two sites. However, such allosteric exclusion in no way contradicts the idea that SR-95531 is competitive, because competition is formally defined as mutually exclusive binding of agonist and antagonist, regardless of the physical arrangement of their binding sites (Segel, 1976).

Another assumption that must hold in order for the race equation to accurately estimate k_{on,GABA} is that any given receptor only binds agonist or antagonist once before the current peaks. This holds in wild-type receptors for which GABA and SR-95531 both have relatively slow unbinding rates. However, GABA unbinds from α4β2-R207C receptors much more quickly, which may give SR-95531 extra opportunities to compete with the agonist before the current peaks. In this case, the race equation yields a lower limit on k_{off,GABA}. Fortunately, only ~5% of singly bound GABA receptors will unbind rather than binding a second GABA or entering gating steps [i.e., k_{off}/(k_{on} + [GABA] + k_{on} + β2 + d_{2})], and of those, only ~30% will unbind before the peak of current (t_{off}, 0.24 msec; time of peak, ~1 msec), approximately one-third of which will then bind SR-95531. Therefore, we expect rapid unbinding in the mutant receptor to cause an insignificant (<1%) underestimate of k_{on,GABA}.

In neuronal GABA_{A} receptors, decreasing the length of full agonists decreases affinity by simultaneously speeding unbinding and slowing binding (Jones et al., 1998). This correlation between length and affinity was explained by a simple physical model, in which the binding site must undergo an energy-requiring motion during binding. The affinity reduction of β2-R207C receptors is also attributable to a simultaneous speeding of unbinding and...
arginine side chain and the negatively charged carboxylate group on the GABA molecule. However, if $\beta_2$-R207 were solely responsible for stabilizing the carboxylate group, we would then expect the cysteine mutation to completely abolish GABA binding. Therefore, other residues, and possibly other arginines, must also be involved. Indeed, such an interaction has already been postulated to occur between the arginine at position 66 of the $\alpha_1$ subunit and the GABA carboxylate group (Holden and Czajkowski, 2002). Like $\beta_2$-R207, $\alpha_1$-R66 faces into the binding pocket, and its mutation to cysteine causes a 300-fold rightward shift in GABA-EC_{50}. There is also a third arginine, at position 131 of the $\alpha_1$ subunit, that faces into the binding pocket and the mutation to cysteine of which causes a 70-fold shift in GABA-EC_{50} (Holden and Czajkowski, 2004). Because mutation of each of these arginines similarly reduces GABA affinity, although no single mutant completely abolishes binding, all three of these residues may collaborate to stabilize the GABA carboxylate group. It might seem surprising that side chains from different subunits would share an interaction with the same end of the GABA molecule. Nonetheless, this hypothesis appears to be supported by a molecular model of the extracellular domain of the GABA$_A$ receptor derived from the crystal structure of the homologous acetylcholine-binding protein (Brejc et al., 2001; Cromer et al., 2002; Holden and Czajkowski, 2002). In this model, $\beta_2$-R207, $\alpha_1$-R66, and $\alpha_1$-R131 can come together to form a triangle of positive charge that spans a narrow part of the agonist-binding pocket.

We propose that this “crown of arginines” forms a region of positive charge that accommodates the negatively charged carboxylate group of the GABA molecule. Such an interaction would explain why single mutations among these three arginines reduce affinity but do not abolish binding altogether. This arrangement would imply that the GABA molecule is oriented vertically within the binding site (i.e., perpendicular to the plane of the plasma membrane), with the amino group facing away from the arginine crown (Holden and Czajkowski, 2002). In addition, antagonists such as SR-95531 have bulky aromatic groups attached to the amino end of the GABA-like moiety, which would also face away from the arginine crown, allowing these bulky groups to interact with structures outside the core of the binding site. This could explain why mutations in the arginine crown have little effect on SR-95531 interactions, whereas some mutations above (e.g., $\alpha_1$-R119C) or below (e.g., $\alpha_1$-D62C) the core of the binding site can be protected from interacting with sulphydryl-specific reagents by SR-95531 but not by GABA (Holden and Czajkowski, 2001, 2004). In contrast, kinetic studies suggest that competitive antagonist affinity is dominated by hydrophobic interactions (Jones et al., 2001), consistent with the large increases in SR-95531 EC_{50} caused by mutation of aromatic residues $\alpha_1$-F64, $\beta_2$-Y97, or $\beta_2$-Y205 (Wagner and Czajkowski, 2001; Boilleau et al., 2002; Holden and Czajkowski, 2002). These aromatic residues may help to stabilize the crown against electrostatic repulsion between arginines, possibly through cation-pi interactions. Similar bonds with the amino group of GABA and antagonists, and additional hydrophobic bonds with antagonists, might contribute to binding of these ligands. Additional studies on the microscopic kinetics of binding site mutations will be a valuable complement to existing structural methods in piecing together the interactions between ligands and the GABA$_A$ receptor.

**References**


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