Block of AMPA Receptor Desensitization by a Point Mutation outside the Ligand-Binding Domain

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Desensitization of ionotropic glutamate receptors (GluRs), specifically the AMPA receptor subtype, shapes the postsynaptic response at certain synapses in the brain. All known mechanisms that alter desensitization, either pharmacological or mutational, are associated with the ligand-binding domain. Here we report that substitution of a conserved positively charged arginine (R) with a negatively charged glutamate in the linker between the pore-forming M3 segment and the S2 lobe, a region outside the ligand-binding domain, blocks desensitization in homomeric AMPA receptors composed of GluR-B subunits. A charge-reversing substitution of a glutamate adjacent to this conserved R enhanced desensitization, consistent with these effects attributable to electrostatics. Homologous substitutions of the conserved R in GluR-B, Glu-A, and the kainate receptor GluR-6 subunits produced comparable but less visible effects on desensitization. Subunit specificity was also apparent for accessibility of substituted cysteines in the M3–S2 linker, suggesting that this part of the channel is not structurally identical in different GluRs. Additionally, reactivity with a sulfhydryl-specific reagent was state dependent, suggesting that the conformations of the nonconducting closed and desensitized states are different at the level of the M3–S2 linker. Our results therefore represent the first identification of elements outside the ligand-binding domain affecting desensitization in non-NMDAR subunits and suggest that electrostatic interactions involving charged residues in the M3–S2 linker influence channel gating in a subunit- and subtype-specific manner.

**Key words:** channel gating; desensitization; ligand-binding domain; M3 segment; M3–S2 linker; substituted cysteine accessibility method

**Introduction**

In the continued presence of glutamate, ionotropic glutamate receptors (GluRs), specifically AMPA receptor (AMPA) and kainate receptor (KAR) subtypes, undergo fast and strong desensitization, a process in which the associated ion channel enters into a nonconducting (desensitized) state. At present, all known mechanisms that alter desensitization in non-NMDA receptors (non-NMDARs), either pharmacological (Partin et al., 1995) or mutational (Stern-Bach et al., 1998; Banke et al., 2001; Partin, 2001), are associated directly with the ligand-binding domain. Structurally and functionally, the ligand-binding domain of GluRs is organized as a pair of dimers (Armstrong and Guoaux, 2000; Mansour et al., 2001; Robert et al., 2001) with desensitization possibly reflecting dimerization of a dimer (Sun et al., 2002). Hence, binding of cyclothiazide (CTZ) (Partin et al., 1995) as well as a leucine-to-tyrosine substitution (Stern-Bach et al., 1998) apparently block desensitization by stabilizing a dimer interface.

In contrast to the ligand-binding domain, little is known about the contribution of the ion channel to the process of desensitization. In GluR subunits, the ligand-binding domain is formed by the proximal part of the N terminus (S1 lobe) and the large loop between transmembrane segments M3 and M4 (S2 lobe) (Fig. 1A). All four hydrophobic segments (M1–M4) apparently contribute to the ion channel (Kuner et al., 1996, 2001; Beck et al., 1999), with M3 representing the major structural element lining the pore and involved in channel opening and closure (Kohda et al., 2000; Jones et al., 2002; Sobolevsky et al., 2002, 2003). Accordingly, the linker between M3 and the ligand-binding domain, the M3–S2 linker, transfers the conformational changes in the ligand-binding domain to M3, but its structural and functional contribution to gating in non-NMDARs is unknown.

Figure 1B shows a sequence alignment of the M3–S2 linker and adjacent regions in non-NMDAR subunits. This linker is positioned downstream from SYTANLAAF, the most highly conserved motif in GluRs (Kuner et al., 2003), and is also conserved to some extent. Notable in this regard is the presence in all subunits of a highly conserved positively charged arginine (R). This arginine and an adjacent negatively charged glutamate (E) in AMPAR subunits influence Ca$^{2+}$ permeation (Jatzke et al., 2003) and presumably are positioned in the ion conduction pathway. Recent evidence has also suggested that charged residues in linker regions in GABA<sub>A</sub> (Kash et al., 2003) and voltage-gated channels (Larson and Elinder, 2000) affect channel gating. We therefore investigated the functional significance of the conserved arginine...
as well as other charged residues in the M3–S2 linker using site-directed mutagenesis, fast agonist application, and chemical modification. We find that charged residues in the M3–S2 linker are involved in electrostatic interactions that affect channel desensitization in a subunit- and subtype-specific manner.

Materials and Methods

Molecular biology and heterologous expression

AMPAR subunits are named using the nomenclature of Seeburg (1993) with the flip/flop cassette indicated by the i/o subscript. Numbering of amino acids is for the mature protein. The KAR subunit, GluR-6, was fully edited within the M1 segment (V and C) (Kohler et al., 1993). Point mutations in GluR-A, or GluR-B, either with glutamine (Q) or asparagine (N) at the Q/R site, and GluR-6 subunits were generated by a QuickChange site-directed mutagenesis kit (Stratagene, La Jolla, CA) or by other PCR-based methods using Platinum Pfx DNA polymerase (Invitrogen, Carlsbad, CA) or Pfu poly DNA polymerase (Stratagene). Positive clones were subcloned back into the respective wild-type clone present either in the eukaryotic expression vector pRK or a pSP64T-derivated vector optimized for expression in Xenopus laevis oocytes. All constructs were sequenced over the entire length of the replaced fragment. cRNA was transcribed and capped for each expression construct using SP6 RNA mMessage kit (Ambion Inc., Austin, TX) and examined electrophoretically on a denaturating agarose gel. RNA concentrations were determined by ethidium bromide stain of the gel relative to an RNA molecular weight marker. Dilutions of RNA (0.01–0.1 μg/μL) were prepared to achieve optimal expression.

Channels were expressed transiently either in human embryonic kidney 293 (HEK 293) cells or in Xenopus laevis oocytes. HEK 293 cells were transfected with GluR subunits using FuGene 6 (Roche Molecular Biochemicals, Indianapolis, IN). A vector for enhanced green fluorescent protein (pEGFP-C1; Clontech, Palo Alto, CA) was cotransfected at a ratio of 1:9 (pEGFP-C1:GluR subunit). Oocytes were prepared, injected, and maintained as described (Wollmuth et al., 1996; Sobolevsky et al., 2002). Recordings were made 1–6 d after transfections or injections.

Current recording

Currents from HEK 293 cells were recorded at room temperature (20–23°C) using an EPC-9 amplifier with PULSE software (HEKA Elektronik, Lambrecht, Germany), low-pass filtered at 2.9 kHz, and digitized at 10 kHz. Pipettes had resistances of 2–4 MΩ when filled with the pipette solution and measured in the Na+ reference solution. We did not use series resistance compensation. External solutions were applied using a piezo-driven double-barrel application system. One barrel contained the external solution, and the other barrel contained the same solution with added glutamate (1–3 ms, unless otherwise noted). The open tip response (10–90% rise time) of the application system was ~350 μs.

Our standard intracellular (pipette) solution consisted of (in mM): 140 KCl, 10 HEPES, and 1 BAPTA, pH 7.2 (KOH). The standard external solution consisted of (in mM): 140 NaCl, 10 HEPES, 1.8 CaCl2, and 1 MgCl2, pH 7.2 (NaOH). In some instances, we added CTZ (30–200 μM; stock solution was in 100 mM NaOH) to the external solutions for AMPARs to minimize desensitization. Unless otherwise noted, all chemicals were obtained from Sigma (St. Louis, MO) or J. T. Baker Company (Phillipsburg, NJ).

We recorded whole-cell currents in Xenopus oocytes at room temperature (20–23°C) using a two-electrode voltage clamp (TEV-200A; Dagan Corp., Minneapolis, MN) with Cell Works software (npi Electronic, Tamm, Germany). Microelectrodes were filled with 3 M KCl and had resistances of 1–4 MΩ. To minimize solution exchange rates, we used a narrow flow-through recording chamber with a small volume of ~70 μL. The external solution consisted of (in mM): 115 NaCl, 2.5 KCl, 0.18 CaCl2, and 10 HEPES, pH 7.2 (NaOH). Glutamate (1 mM), CNQX, CTZ, and 2-(trimethylammonium)ethyl methanethiosulfonate (MTSET) were applied with the bath solution. MTSET was purchased from Toronto Research Chemicals (Toronto, Ontario, Canada).

Experimental protocols

Kinetics. To measure kinetics, we recorded currents in outside-out patches excised from HEK 293 cells, although sometimes (where noted) in the whole-cell mode. In some instances, specifically when current amplitudes were small, we averaged 3–10 current records. To determine the rate and extent of desensitization, we rapidly applied glutamate for 100–1000 msec typically at ~60 mV. Time constants of desensitization (τdes) were determined by fitting the current decay with a single exponential function. We contrasted the extent of desensitization in several different ways. On the basis of steady-state (Iss) and peak (Ip) current amplitudes recorded in the same solution, we calculated the percentage of desensitization (%des = 100 × (1 – Ip/Iss)). We also used these amplitudes to calculate the plateau/peak ratio (plateau/peak = Ip/Ip). Alternatively, we calculated the percentage desensitization using CTZ as a reference (%desCTZ), comparing steady-state current amplitudes in the control solution (Icss) with peak amplitudes recorded in the same solution.
but with added CTZ ($I_{norm}$ [(%des]CTZ) = 100 × (1 − $I_{p}/I_{p(CTZ)}$)]. Peak current amplitudes measured in a solution without ($I_{p}$) or with ([$I_{p(CTZ)}$) added CTZ were used to calculate changes in current amplitudes (%des) of 100 × $I_{p(CTZ)}$/$I_{p}$).

Recovery from desensitization ($\tau_{rec}$) was estimated using a two-pulse protocol. After a base application of glutamate (100 or 200 msec in length), a 5 msec test application of glutamate was applied at increasing times thereafter. The peak current during the test application ($I_{p}$) was normalized either to the peak current of the base application ($I_{norm} = I_{p}/I_{p}$) or to only the desensitizing portion of the current during the base application ($I_{norm} = (I_{p} − I_{ss})/(I_{p} − I_{ss})$). Normalized currents (expressed as a percentage, %des = 100 × $I_{norm}$) were plotted as a function of time after the end of the base application and fitted with a single exponential to derive $\tau_{rec}$.

Glutamate concentration dependence. Concentration–response curves were measured in the whole-cell mode in the presence of 30 μM CTZ. Solutions containing various concentrations of glutamate (0.001–10 mM) were applied to cells held at −60 mV. Current amplitudes normalized to the maximal response were plotted as a function of concentration and fitted with the Hill equation, $1/(1 + ([Glu]/EC50)^n)$, where $EC_{50}$ is the concentration to achieve half-maximal response, and $n$ is the Hill coefficient.

**Substituted cysteine accessibility method.** Cysteine-substituted GluR channels were probed from the extracellular side of the membrane with the positively charged MTSET. MTSET-containing solutions were prepared, stored, and applied as described previously (Sobolevsky et al., 2002). Accessibility of substituted cysteines was determined using steady-state reaction protocols (Sobolevsky et al., 2002).

**Data analysis**

All curve fitting was done using Igor Pro (WaveMetrics, Lake Oswego, OR). Results are reported and shown graphically as mean ± SEM. An ANOVA was used to test for statistical differences, with the Tukey test used for multiple comparisons. Significance was assumed if $p < 0.05$. In figures, and unless otherwise noted, filled bars indicate values statistically different from those for wild type.

**Results**

**Opposite charge substitutions of the conserved arginine in the M3–S2 linker**

To study gating in AMPAR and KAR subtypes, we recorded glutamate-activated currents from wild-type and mutant channels in outside-out patches isolated from HEK 293 cells using rapid solution exchange. Figure 2A–C (left traces) shows currents recorded from wild-type GluR-A1, GluR-B1, and GluR-6 channels. During the glutamate application (filled bars, 100 msec), the current rapidly reached peak amplitude and then strongly decayed, reflecting the process of desensitization. For wild-type channels, the rate of desensitization ($\tau_{des}$) was fast with its extent, expressed as a percentage (%des; see Materials and Methods), always >96%. Mean values for $\tau_{des}$, %des, plateau/peak ratio, and the time constant for recovery from desensitization ($\tau_{rec}$) for wild-type channels including GluR-B1 are summarized in Table 1. These values are comparable with those reported previously (Mosbacher et al., 1994; Dingledine et al., 1999; Sun et al., 2002).

Figure 2A–C (right traces) shows currents for mutant channels, where the conserved positively charged R in the M3–S2 linker is replaced with the negatively charged E (R $\rightarrow$ E substitution). Surprisingly, the extent of desensitization, especially for the AMPAR subtypes, was greatly attenuated. The strongest effect was observed for GluR-B1 (R624E), in which only ∼15% of the peak current desensitized compared with ∼97% in wild-type GluR-B1 (Table 1). The R $\rightarrow$ E substitution in GluR-A1 (%des = 58%), GluR-B1 (%des = 58%), and GluR-6 (%des = 91%) produced less dramatic reductions in the extent of desensitization. However, the plateau/peak ratio was increased to approximately the same extent (20–30 times) for all subunits (Table 1), indicat-
desensitization, GluR-Bi (R628E) channels show an increased apparent glutamate affinity. Consistent with the idea that R → E substitution disrupts desensitization, GluR-Bi (R628E) channels show an increased apparent glutamate affinity (EC_{50} = 24 μM) compared with that for wild type (EC_{50} = 134 μM) (Fig. 3B).

Figure 3. The R → E substitution disrupts desensitization in GluR-Bi channels. A, Superimpositions of glutamate-activated currents for wild-type (left traces) and mutant (R → E; right traces) GluR-Bi channels recorded either in the absence or in the presence of 30 μM CTZ. Currents were elicited by a 100 msec application of glutamate (3 mM; filled bar) at a holding potential of −60 mV. B, Glutamate concentration response curves for wild-type (wt; open symbols) and mutant (R → E; filled symbols) GluR-Bi channels. The solid lines are a fitted Hill equation with an EC_{50} of 134 ± 3 μM (n = 5) for wild type and 24 ± 2 μM (n = 4) for GluR-Bi (R628E). C, Average extent of desensitization, %des, for wild-type and GluR-Bi (R628E) channels measured in the absence or presence of 100 μM EDTA. Currents were recorded either in our standard external solution containing 1.8 mM Ca^{2+} and 1 mM Mg^{2+} or in the same solution but with 100 μM EDTA and no added divalent cations.

Table 1. Desensitization properties of wild-type and mutant AMPAR and KAR channels

<table>
<thead>
<tr>
<th>Subunit</th>
<th>τ_{des} (msec)</th>
<th>%des</th>
<th>Plateau/peak</th>
<th>τ_{rec} (msec)</th>
</tr>
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<tbody>
<tr>
<td>A_wt</td>
<td>3.2 ± 0.2</td>
<td>97.9 ± 0.6</td>
<td>0.021 ± 0.006</td>
<td>155 ± 11</td>
</tr>
<tr>
<td>A_R628E</td>
<td>5.8 ± 0.3</td>
<td>57.6 ± 4.0</td>
<td>0.424 ± 0.040</td>
<td>127 ± 6</td>
</tr>
<tr>
<td>B_wt</td>
<td>5.6 ± 0.3</td>
<td>96.5 ± 0.8</td>
<td>0.035 ± 0.008</td>
<td>21.8 ± 2.0</td>
</tr>
<tr>
<td>B_R628E</td>
<td>10.8 ± 1.1</td>
<td>143 ± 1.5</td>
<td>0.853 ± 0.015</td>
<td>ND</td>
</tr>
<tr>
<td>B_R628Q</td>
<td>1.8 ± 0.2</td>
<td>98.4 ± 0.5</td>
<td>0.016 ± 0.005</td>
<td>24.3 ± 1.9</td>
</tr>
<tr>
<td>B_R624E</td>
<td>3.3 ± 0.2</td>
<td>58.0 ± 3.4</td>
<td>0.420 ± 0.034</td>
<td>23.9 ± 2.2</td>
</tr>
<tr>
<td>6_wt</td>
<td>3.4 ± 0.1</td>
<td>99.6 ± 0.2</td>
<td>0.004 ± 0.002</td>
<td>1920 ± 110</td>
</tr>
<tr>
<td>6_R623E</td>
<td>6.8 ± 0.7</td>
<td>90.5 ± 3.1</td>
<td>0.095 ± 0.021</td>
<td>1070 ± 120</td>
</tr>
<tr>
<td>6_R631R</td>
<td>1.9 ± 0.1</td>
<td>99.9 ± 0.1</td>
<td>0.001 ± 0.001</td>
<td>1050 ± 40</td>
</tr>
<tr>
<td>6_R634R</td>
<td>3.5 ± 0.3</td>
<td>99.8 ± 0.1</td>
<td>0.002 ± 0.001</td>
<td>1620 ± 110</td>
</tr>
</tbody>
</table>

The time constants of desensitization (τ_{des}) and recovery from it (τ_{rec}) were obtained from single exponential fits (for details, see Materials and Methods). %des and plateau/peak ratio are based on peak and steady-state current amplitudes. Values shown are the mean ± SEM (n = 4–10). We could not determine (ND) the recovery from desensitization for GluR-Bi (R628E) because of the small desensitizing current component. wt, Wild type.

The R → E substitution introduces an additional negative side chain in the M3–S2 linker and underlies surface charge-like effects on Ca^{2+} permeability (Jatzke et al., 2003). The nondesensitizing phenotype therefore might arise because of divalent ions such as Ca^{2+} or Zn^{2+} more strongly interacting with this highly negative element. However, the extent of desensitization in GluR-Bi (R628E) as well as wild-type channels was not significantly changed when recorded in the absence of divalent cations and in 100 μM EDTA (Fig. 3C). Hence, the block of desensitization in GluR-Bi (R628E) is not caused by divalent cations and is probably caused by the R → E substitution itself.

An electrostatic interaction underlies the disruption of desensitization

To further address the mechanism underlying the nondesensitizing phenotype, we substituted various amino acids at the R position in the M3–S2 linker in GluR-Bi and characterized the effect of these substitutions on desensitization properties (Fig. 4). All of the individual substitutions at the R position, like the R → E substitution, significantly slowed, albeit weakly, the rate of desensitization (Fig. 4A). On the other hand, only oppositely charged substitutions of this position significantly reduced the extent of desensitization (Fig. 4B), with the strongest reduction occurring for glutamate (R628E), followed by the aspartate (R628D) and the cysteine (R628C, which is ~10% charged under physiological pH) side chains. The positively charged histidine (R628H) also reduced %des_{CTZ}, but this effect was quite small (94.0 ± 1.2%; n = 5) compared with wild type (97.7 ± 0.4%; n = 8). Surprisingly, the neutral alanine (R628A) and the polar glutamine (R628Q) had no effect on %des_{CTZ} (Fig. 4B) but did significantly reduce the rate of recovery from desensitization (Fig. 4C). Overall, the results shown in Figure 4 suggest that the effect of R → E substitution on desensitization is at least in part attributable to electrostatic interactions.

Substitutions of a negatively charged glutamate in the M3–S2 linker enhance desensitization

If block of desensitization by the R → E substitution is attributable to electrostatic interactions, there may be other charged residues in the protein, presumably negatively charged, that complement the conserved R. One potential candidate is within the M3–S2 linker itself. Indeed, adjacent to the conserved R in AMPAR and low-affinity KAR subunits is a negatively charged E (Fig. 1B), and these oppositely charged E and R residues could interact either in an intrasubunit or intersubunit state-dependent manner. In AMPAR subunits, either GluR-A_i or -B_i, opposite charge
substitutions of this glutamate (E → R substitution), yielded channels that even in the whole-cell mode had very small glutamate-activated current amplitudes (typically <10 pA compared with 3–5 nA for wild type). Hence, we could not consistently measure desensitization properties in outside-out patches. Notable, however, is that with added CTZ, peak current amplitudes were dramatically increased (Fig. 5A). The increase was maximal (~700%) for the E → R substitution, although comparable effects were observed for other substitutions at this position (Fig. 5B). As discussed below, this dramatic current increase produced by CTZ arises at least in part because the E → R substitution enhances desensitization. Hence, without CTZ, E → R-substituted channels are mainly in the desensitized state in the absence of glutamate or desensitize more rapidly than rates of channel opening, solution exchange, or both.

In contrast to AMPAR subunits, the E → R substitution in GluR-6 [GluR-6 (E631R)] expressed well. Currents through these channels showed a faster rate of desensitization and recovery from it and a greater extent of desensitization than wild type GluR-6 channels (also see Table 1, plateau/peak ratio). In addition, we found that the E → R substitution in a mutant GluR-B, subunit in which an asparagine (N) occupies the Q/R-site [GluR-Bi (N)] also showed reasonable current amplitudes. GluR-Bi (N) channels themselves showed a slower rate of desensitization (τ_{des} = 13 msec) and a reduced extent of desensitization (%des = 92.5%) compared with GluR-B (Q) channels (τ_{des} = 5.6 msec; %des = 96.5%) (Fig. 5C,D). In the GluR-B (N) background, the E → R substitution, like that in GluR-Bi (Q), slowed the rate of desensitization by approximately twofold (Fig. 5C) and dramatically reduced the extent of desensitization (Fig. 5D). In contrast, the E → R substitution produced essentially opposite effects, significantly increasing the rate and extent of desensitization (Fig. 5C,D).

In summary, an E residue in the M3–S2 linker, adjacent to the conserved R, also affects desensitization. The effect of the E → R substitution is essentially opposite to that of the R → E substitution. Accordingly, the apparent glutamate affinity for channels with E → R substitution in the GluR-B (Q) background was significantly reduced (E_{Ca} = 1.3 ± 0.1 mM; n = 5) compared with that in wild type (E_{Ca} = 134 μM) (compare Fig. 3C). These results therefore support the idea that electrostatic interactions involving charged residues in the M3–S2 linker contribute to channel gating. On the other hand, the opposite effects of the E → R and R → E substitutions on desensitization suggest that these electrostatic interactions are not confined to E and R themselves, and some other complementary charge(s) located elsewhere in the protein have to be involved.

Orientiation of the ER dipole is important to channel gating
Although the adjacent opposite charged residues (E and R) in the M3–S2 linker have no net charge, they presumably form an elec-
Subunit-specific and state-dependent reactivity of cysteines substituted in the M3–S2 linker

The subunit-specific differences between AMPAR subtypes appear not to be attributable to a charge difference but rather may reflect structural differences in the M3–S2 linker. To address this idea, we compared glutamate-activated current amplitudes, in mutant GluR-Ai and GluR-B, channels containing cysteines substituted at the E and R positions, before \((I_{\text{pre}})\) and after \((I_{\text{post}})\) the extracellular application of the positively charged methanethiol sulfonate (MTS) reagent MTSET (Fig. 7A). We were also interested in the state dependence of reactivity, so we applied MTSET when channels were mainly (but not exclusively) in the open (top trace), closed (middle trace), and desensitized (bottom trace) states. The maximal occupation of the various states was achieved by applying MTSET in the continuous presence of Glu (1 mM) and saturating concentrations of CTZ (50 μM; open state, O), in the absence of glutamate and in the presence of the competitive AMPAR antagonist CNQX (10 μM; closed state, C), and, finally, in the continuous presence of Glu but in the absence of CTZ (desensitized state, D). However, the distinction between states is by no means absolute. For example, the R → C substitution in GluR-Bi [GluR-Bi (R628C)] disrupts desensitization (Fig. 4B), indicating that even in the absence of CTZ (presence of glutamate), the channels will spend considerable time in the open state.

Figure 7B summarizes the mean percentage change in current amplitude after the application of MTSET for wild-type and cysteine-substituted GluR-Ai and GluR-B channels. For wild-type channels, MTSET had no persistent effects on current amplitudes under any of the conditions. On the other hand, in cysteine-substituted mutant channels, the effect of MTSET on current amplitude was strongly subunit specific and state dependent. At the E position, MTSET altered current amplitudes under all conditions for GluR-Bi, whereas it produced a significant change only when applied in the desensitized state for GluR-Ai. In contrast, at the R position, MTSET did not alter current amplitude under any conditions in GluR-Ai, whereas it produced a significant change in GluR-Bi only when applied in the desensitized state.

The results with the homologous positions in GluR-Ai (E623 and R624) and GluR-B, (E627 and R628) suggest that there are structural differences in the M3–S2 linker between different Glur subtypes. Additionally, the state-dependent reactivity of GluR-Ai (E623C) and GluR-Bi (R626C) with MTSET suggests that conformations of the M3–S2 linker are different in the nonconducting closed and desensitized states.

Discussion

GluR desensitization is an important mechanism shaping excitatory postsynaptic potentials in the brain (Jones and Westbrook, 1996; Glavinovic and Rabie, 1998). As a gating process, desensitization originates in the ligand-binding domain and propagates to the ion channel. Our results represent the first identification of residues outside the ligand-binding domain that can alter desensitization in non-NMDAR channels and suggest that electrostatics involving in part charged residues in the M3–S2 linker contribute to gating in GlurRs.

Charged residues in the M3–S2 linker influence channel gating

Opposite charge substitutions of two adjacent charged residues, a glutamate (E627 in GluR-Bi) and a conserved arginine (R628 in GluR-Bi), strongly affected desensitization in both AMPA and
KA GluR subtypes (Figs. 2, 5; Table 1). These two residues are positioned outside the ligand-binding domain (Armstrong et al., 1998; Armstrong and Gouaux, 2000) being located N-terminal to S2 (see Fig. 1). On the other hand, substitutions of these residues in AMPARs (Jaitke et al., 2003) and of homologous positions in the NMDAR NR1 subunit (Watanebe et al., 2002) affect Ca$^{2+}$ permeation, suggesting that they are associated with the ion conduction pathway. Nevertheless, ER is positioned four amino acid residues external to SYTANLAFF, a motif forming the most extracellular part of the channel (Sobolevsky et al., 2003). We therefore refer to the location of ER between the ligand-binding domain and the ion channel as the M3–S2 linker.

The reduced extent of desensitization in R→E-substituted GluR-B channels (%des = 14%; compared with %des = 97% in wild type) is comparable with that produced by the leucine-to-tyrosine (L→Y) substitution in the ligand-binding domain (%des = 8–10%) in the same background (Stern-Bach et al., 1998; Sun et al., 2002). The L→Y substitution as well as binding of CTZ block desensitization by stabilizing a dimer interface in the ligand-binding domain (Sun et al., 2002), slowing the rate of desensitization by nearly 30-fold (Robert et al., 2002). In contrast, the R→E substitution is located well outside the ligand-binding domain and slows the rate of desensitization by only approximately twofold (Table 1). Therefore, the L→Y and R→E substitutions apparently block desensitization in GluRs by fundamentally different mechanisms.

**Charged residues alter channel gating via electrostatic interactions**

Channels containing an opposite charge substitution of a conserved arginine (R→E substitution, EE channels) in the M3–S2 linker showed a reduced degree of desensitization (Figs. 2, 3; Table 1). Neutralizing substitutions (EA and EQ channels), although not changing significantly the degree of desensitization, strongly slowed down the rate of recovery from desensitization (Fig. 4C). An opposite charge substitution of E, a residue adjacent to the conserved R (E→R substitution, RR channels), produced effects essentially opposite to that of the R→E substitution (Figs. 5, 6), yet mutant channels in which both charges were reversed (ER→RE substitution) still showed gating differences (Fig. 6). Together these results suggest that electrostatics as contributed by the charges or the dipole of ER or both in the M3–S2 linker influence gating in GluRs.

The electrostatic interactions involving E and R imply the existence of closely located charges that interact with each other in a state-dependent manner (Fig. 7). The possibility of intrasubunit or intersubunit interactions involving only E and R themselves seems unlikely because opposite charge substitutions at either position (EE or RR channels) produced opposite effects on channel gating rather than the anticipated comparable effect (Fig. 6).

**Figure 7.** State-dependent effect of a sulphydryl-specific reagent on glutamate-activated currents in cysteine-substituted AMPAR channels. A, Protocols to measure the state dependence of reactivity. Whole-cell currents recorded from Xenopus oocytes expressing GluR-Ai (top trace), GluR-Bi (middle trace), and GluR-Bi (R628C) (bottom trace) subunits are shown. Currents were elicited by applications of glutamate (1 mM; thin lines; typically 15 sec in duration except during the MTSET application). MTSET (2 mM; thick lines) was applied in the presence of glutamate and CTZ (50 μM; open boxes, top trace), in the absence of glutamate and in the presence of the competitive antagonist CNQX (10 μM; shaded box, middle trace), or just in the presence of glutamate (bottom trace). During these different applications, we assume that the predominant channel state is open (O; top trace), closed (C; middle trace), or desensitized (D; bottom trace). Current amplitudes were measured either before (Ipre) or after (Ipst) application of MTSET. B, Changes in the current amplitude measured before (Ipre) and after (Ipst) exposure to MTSET (1–3 min) according to protocols in A for wild-type (first row) and cysteine-substituted (second, third rows) GluR-Ai (left column) and GluR-Bi (right column). Up- and down-pointing bars indicate inhibition and potentiation of glutamate-activated currents, respectively. Filled bars indicate a significant difference between Ipre and Ipst (n = 3–8).

**Contributions of the M3–S2 linker to gating**

In all non-NMDAR subunits, channels containing the R→E substitution showed a reduced degree of desensitization (Fig. 2, Table 1). In terms of the %des, this effect was strongest in the
GluR-Bi background, but the fold increase for plateau/peak ratios was approximately the same in all backgrounds (Table 1). Indeed, the %des for wild-type GluR-6, GluR-A, and GluR-B (Table 1) decreased in parallel with the ratio of the rate of exit from (γ) and entry into (δ) desensitization [γ/δ = 10^{-5} – 10^{-4} for GluR-6 (Bowie et al., 1998), 10^{-3} for GluR-A (Partin et al., 1996; Robert and Howe, 2003), and 10^{-2} – 10^{-1} for GluR-B (Koike et al., 2000)]. In addition, the subunit-specific effect of the R → E substitution can be simulated by the same modification of the wild-type models, specifically increasing the rate of exit from the desensitized state by ~10^2 times as well as decreasing the rate of agonist dissociation (k_{off}) by ~10 times (data not shown). Hence the electrostatic interactions involving charged residues in the M3–S2 linker contribute to gating in all non-NMDAR subunits and subtypes similarly, but the visible effect on desensitization depends on the extent of desensitization for the corresponding wild type.

Structurally, the difference in the maximal degree of desensitization may reflect either a difference in the secondary structure or more likely a difference in the spatial arrangement of the M3–S2 linker relative to other domains, specifically those containing a complementary charge (Fig. 8). The differential reactivity of MTSET with cysteine substituted at E and R positions in GluR-A and GluR-B channels (Fig. 7) supports the latter alternative. In AMPAR subunits, transmembrane domains and associated linkers are highly conserved (nearly 100% sequence identity, with the only exception being in the intracellularly located M1–M2 linker), suggesting that the structural difference in the M3–S2 linker between non-NMDAR subunits and subtypes is probably not attributable to differences in the transmembrane domains but rather to the ligand-binding domain. Supporting this idea is the differential effect of the R → E substitution on desensitization in the flip and flop variants of GluR-B (Table 1). Hence the structure of the ion channel at the level of the M3–S2 linker parallels that of the ligand-binding domain. This conclusion agrees with recent results suggesting that the outer pore of the AMPAR channel has a rotational symmetry similar to that of the ligand-binding domain (Sobolevsky et al., 2004).

**Different structures of the M3–S2 linker in the closed and desensitized states**

The M3–S2 linker is a polypeptide of unknown secondary structure. One end of M3–S2 is connected to the S2 domain, a part of the ligand-binding domain with the structure solved by crystallography (Armstrong et al., 1998; Armstrong and Gouaux, 2000). On the other side, M3–S2 is connected to the M3 segment that, based on sidedness of accessibility to MTS reagents, is α-helical (Sobolevsky et al., 2003). The secondary structure of the M3–S2 linker itself may be difficult to infer from the pattern of accessibility because this region is outside the transmembrane portion of GluR and hence may be water-accessible on all sides. Nevertheless, it seems likely that at least part of the M3–S2 linker is extended, specifically that part around ER.

Previous results have suggested that the M3–S2 linker has a different conformation in the open and closed states (Sobolevsky et al., 2003). The accessibility pattern shown in Figure 7B suggests that the nonconducting closed and desensitized states of the AMPAR channel are also structurally different at the linker level. Of course, this interpretation is limited by the assumptions of the substituted cysteine accessibility method (Karlin and Akabas, 1998), and at present it is unclear how far into the pore this asymmetry between the closed and desensitized conformations exists. Future experiments will be needed to further delineate the structural similarities and differences between the open, closed, and desensitized states.

**References**


