Two serine residues on GluN2A C-terminal tails control NMDA receptor current decay times

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NMDA receptors are glutamate-activated, Ca2+-permeable ion channels with critical roles in synaptic transmission and plasticity. The shape and size of their current is modulated by several kinase/phosphatase systems, and numerous residues located on the receptors’ intracellular C-termini are phosphorylated in vivo. To investigate the mechanisms by which phosphorylation may control channel gating, we examined the single-channel behaviors of receptors carrying the S900A or S929A substitution in their GluN2A subunits and thus were rendered resistant to phosphorylation at those sites. We found that the mutations reduced channel open probability primarily by increasing the frequency of desensitized events. The kinetic models we developed revealed complex but similar changes in mechanism for the two mutants, leading to the view that dephosphorylation at either site may cause receptors to activate slower, deactivate faster and desensitize more frequently. This modulatory mechanism is consistent with the proposed roles for these residues in Ca2+-dependent desensitization and calcineurin-mediated reduction of current during brain development.

Introduction

NMDA receptors (NRs) are prominent members of the ionotropic glutamate receptor family whose activation initiates synaptic plasticity at a majority of synapses in the central nervous system and also mediates neurotoxic actions of glutamate. They have high Ca2+ permeability and characteristically slow gating kinetics. Consistent with their central roles in health and disease, NR activity is controlled by numerous cellular mechanisms including reversible covalent modification by kinase/phosphatase pathways.

Functional receptors are tetramers of two GluN1 (N1) and two GluN2 (N2) subunits.1 Each subunit spans the synaptic membrane and projects significantly into both the extra- and the intra-cellular fields. The large N-termini are organized into eight distinct globular domains; whereas C-terminal domains (CTDs), which represent about one third of the receptor’s mass, are believed to be intrinsically disordered.1,2 Substantial functional diversity arises from differential splicing of the N1 transcript combined with regulated expression of four separate N2 subunits, denoted A through D.3 In addition, both the level and the kinetics of the NR-generated current are controlled by mechanisms with remarkable diversity. Numerous diffusible ligands bind at specific extracellular sites and control receptor kinetics with allosteric mechanisms, while several inorganic divalent cations and organic small molecules bind within the membrane-spanning pore to affect the level and composition of ionic flux.1 Further, covalent modifications, mainly by phosphorylation of intracellular residues, have been proposed to control receptor permeation and gating.4-6

We have recently discovered that, relative to wild-type receptors, NRs engineered to lack the CTDs of N1 subunits (N1ΔC838/N2A) produce currents with larger unitary conductance but
similar kinetics, whereas receptors engineered to lack the CTD of N2A subunits (N1/N2AΔCTD) produce currents of similar unitary conductance but dwell significantly longer in open and desensitized states. In addition, we found that the gating effect of N2A CTD truncation is mediated solely by the membrane-proximal region, residues 844 to 944, of the C-terminal tail. Based on these results, we hypothesized that the C-termini of N2A subunits may exert their effects on receptor gating through specific sites that may be subject to phosphorylation by cellular kinases. Based on available literature, two residues in this region, Ser-900 and Ser-929, stand out in functional significance. These were identified by alanine scanning mutagenesis to be responsible for endogenous phosphorylation at Ser-900 or Ser-929 of N2A, that cannot be phosphorylated at these positions: N1/N2A900A and N1/N2A929A. We used the cell-attached patch-clamp technique, which preserves the integrity of the intracellular milieu, and omitted adding Ca\(^{2+}\) to the extracellular recording solution, such that the inward current passed by the channel consisted of only Na\(^{+}\). We reasoned that in these conditions, and for the entire duration of the observation window, the trapped receptor will be in the high-activity form that exists basally before Ca\(^{2+}\) entry initiates desensitization, inactivation or rundown events.

We obtained several current traces, each lasting between 10 and 90 min, from patches containing one N1/N2A900A (n = 6) or one N1/N2A929A channel (n = 8). We compared these to previously reported data sets for wild-type receptors (N1/N2A, n = 14) and receptors with truncated N2A CTDs (N1/N2AΔCTD, n = 7), which were obtained and analyzed in similar conditions (Fig. 1A). All channels examined opened to a single amplitude level that was not different in value across the four data sets (p > 0.05, one-way ANOVA) (Table 1). Relative to N1/N2A receptors, the open probability (P\(_{o}\)) of N1/N2A900A and N1/N2A929A, as calculated from entire records, decreased 2- and 3-fold, respectively (p < 0.01). For both mutants, this drop in activity originated mainly from a -4-fold increase in mean closed time (MCT), with only N1/N2A929A also having shorter (-1.5-fold) mean open time (MOT). Neither mutation reproduced the full phenotype of low activity we observed previously for N1/N2AΔCTD, which originated from a combination of much longer MCT (-50-fold) and longer MOT (-3-fold). These results support the interpretation that perturbations to either of the examined serine residues, including by dephosphorylation, will contribute substantially to the low activity observed for the full CTD deletion; however, because the CTD truncations have a much more severe loss of function phenotype, we surmise that yet undetermined sites within the N2A C-termini module channel activity by lengthening desensitized and/or open events.

In saturating concentrations of agonists and the absence of divalent cations, as employed in this study (1 mM Glu, 0.1 mM Gly and 1 mM EDTA), traces recorded from wild-type NRs have event durations with complex multi-exponential distributions. The closed event distributions of the mutants we examined had five exponential components, E\(_1\)–E\(_5\), as described previously for wild-type NRs (Fig. 1B). The three shorter components (E\(_1\)–E\(_3\)) occur largely within bursts and reflect receptor states visited along the activation pathway, whereas the two longest components, E\(_4\) and E\(_5\), represent largely closures that terminate bursts and inform about desensitized receptor states. The closed event durations measured from N1/N2A900A and N1/N2A929A revealed that the longer MCT values could be attributed to slightly (< 2-fold) longer intra-burst, E\(_4\)–E\(_5\), events, 10-fold longer E\(_4\)-type closures and a 6-fold larger relative area for the E\(_5\) component (Table 2). Notably, the time constant of E\(_5\) was not significantly different from wild-type for either mutant (p > 0.05), and was ~3 sec for each. This was a conspicuous difference from N1/N2AΔCTD, whose longest closures (E\(_5\)) were significantly longer (p < 0.05) with a time constant of ~8 sec. These results support the hypothesis that dephosphorylation of either serine would lengthen specific time components in a similar manner.

Although the lower P\(_{o}\) exhibited by the S900A and S929A mutations was only partially attributed to a reduction in mean open time, we went on to examine the open duration distributions for the two mutants to learn about possible changes in modal gating. Aside from gating and desensitization transitions, NRs also experience modal transitions, which change the receptor’s activity pattern for seconds or minutes at a time. For N1/N2A receptors, three modes can be queried experimentally by determining how many types of openings occur in a record generated by an individual channel. In any mode, defined as a period of uniform kinetics, N1/N2A receptors generate two types of openings, one short and one long. Shifts in modes are characterized by changes in duration of the long openings, while length of the short openings remains constant across all modes. Thus, the open duration distribution will reveal two, three or four open components for records that capture one, two or three modes, respectively. The molecular determinants of NR modal gating are not yet known; however phosphorylation of intracellular residues...
has been mentioned as a possible mechanism for modal shifts. We found that the open interval distributions of both N1/N2A<sup>900A</sup> and N1/N2A<sup>929A</sup> receptors had four components and that their time constants and relative areas were not statistically different from those calculated for N1/N2A receptors (data not shown). This result demonstrates that neither Ser-900 nor Ser-929 influences modal gating of N1/N2A receptors.

Next, we summarized the kinetic changes we measured for the two mutants using a commonly used state model for NR gating (Fig. 2). In this scheme, all states represent receptors that are fully occupied with glutamate and glycine; C<sub>3</sub>, C<sub>2</sub>, and C<sub>1</sub> represent a sequence of non-conductive conformations en route to channel open states; C<sub>4</sub> and C<sub>5</sub> are non-conductive states that represent functionally desensitized receptors; and last, O stands for all active receptor conformations. The rates estimated by fitting this model to entire recordings were largely similar for N1/N2A<sup>900A</sup> and N1/N2A<sup>929A</sup> (Fig. 2), as expected from their similar kinetic parameters (Table 1) and duration distributions (Table 2). Relative to wild-type channels, N1/N2A<sup>900A</sup> and N1/N2A<sup>929A</sup> had slower activation rates (C<sub>3</sub> → C<sub>2</sub>), slower recovery from the desensitized C<sub>4</sub> state (C<sub>2</sub> ← C<sub>4</sub>) and faster access into the most stable desensitized state, C<sub>5</sub> (C<sub>3</sub> → C<sub>5</sub>). Relative to the full N2A CTD truncation, the changes in the rate constants for activation (C<sub>3</sub> → C<sub>2</sub>), desensitization (C<sub>4</sub> ← C<sub>3</sub>) and resensitization (C<sub>4</sub> ← C<sub>2</sub>) were less pronounced. This
if the mechanisms we propose here are correct, they should also account for the macroscopic results reported previously for S900A and S929A in recombinant systems,8 as well as the changes in the NR synaptic response upon dephosphorylation at Ser-900 by calcineurin.9 To determine how well our models account for these known effects, we used them to simulate macroscopic currents (100 channels, 10 pA each) in response to desensitizing (5 sec) and synaptic-like (1 ms) pulses of glutamate (1 mM).

Results of simulations with long stimuli show that both N1/N2A900A and N1/N2A929A receptors produce macroscopic currents that desensitize faster (Fig. 3A). This result is consistent with measurements done by the Westbrook group in that both constructs have pronounced effects on macroscopic desensitization; even though in the previous study the receptors used also lacked the C-termini of N1 subunits (N1Δ838/N2A900A and N1Δ838/N2A929A).8 In addition, our model predicted that both N1/N2A900A and N1/N2A929A will reduce the peak macroscopic response (Fig. 3B), and overall will reduce charge transfer by 40% and 70%, respectively. This information cannot be obtained experimentally because in whole-cell current measurements the total number of channels is unknown; however it is valuable in estimating how the specific serine residues may control the actual charge transfer that generates the activation pathway C2 ↔ C1. This result may have structural implications, but these are insufficiently determined at this time.19

State models are valuable in that they summarize and organize all observed kinetic changes into a reaction mechanism. More importantly, they have the power to predict channel responses over a broad spectrum of stimulation protocols. This is useful in testing the model experimentally but also in controlling variables that cannot be determined empirically, such as the number of receptors contributing to a macroscopic response. Thus, if the mechanisms we propose here are correct, they should also account for the macroscopic results reported previously for S900A and S929A in recombinant systems,8 as well as the changes in the NR synaptic response upon dephosphorylation at Ser-900 by calcineurin.9 To determine how well our models account for these known effects, we used them to simulate macroscopic currents (100 channels, 10 pA each) in response to desensitizing (5 sec) and synaptic-like (1 ms) pulses of glutamate (1 mM).

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Table 1. Single-channel properties of N1/N2A900A and N1/N2A929A

| Receptor       | Amplitude (pA) | Po       | MCT (ms) | MOT (ms) | N    | Events |
|----------------|----------------|----------|----------|----------|------|--------|
| N1/N2A         | 8.7 ± 0.3      | 0.54 ± 0.04 | 4.7 ± 0.5 | 5.4 ± 0.5 | 14   | 4.9 × 10⁶ |
| N1/N2A900A     | 8.3 ± 1.6      | 0.28 ± 0.10† | 18 ± 6†  | 3.9 ± 0.7† | 6    | 1.7 × 10⁶ |
| N1/N2A929A     | 8.9 ± 1.6      | 0.18 ± 0.05* | 21 ± 6*  | 3.1 ± 0.4*† | 8    | 1.6 × 10⁶ |
| N1/N2AΔ844     | 9.7 ± 0.3      | 0.22 ± 0.04* | 50 ± 10* | 13 ± 2*  | 7    | 5.0 × 10³ |

Symbols mark values that are statistically different compared with N1/N2A (*) or N1/N2AΔ844 (†), (p < 0.05, Student’s t-test).

Table 2. Closed components of N1/N2A900A and N1/N2A929A

| Receptor       | τE1 (ms) | aE1 (%) | τE2 (ms) | aE2 (%) | τE3 (ms) | aE3 (%) | τE4 (ms) | aE4 (%) | τE5 (ms) | aE5 (%) |
|----------------|----------|---------|----------|---------|----------|---------|----------|---------|----------|---------|
| N1/N2A         | 0.14 ± 0.01 | 45 ± 3  | 1.4 ± 0.1 | 24 ± 2  | 4.4 ± 0.2 | 30 ± 3  | 22 ± 4  | 1.5 ± 0.3 | 3,000 ± 400 | 0.08 ± 0.01 |
| N1/N2A900A     | 0.17 ± 0.01* | 31 ± 4*  | 2.1 ± 0.2* | 38 ± 4*† | 5.6 ± 0.6 | 29 ± 6 | 100 ± 50* | 1.2 ± 0.7 | 3300 ± 500† | 0.5 ± 0.1* |
| N1/N2A929A     | 0.18 ± 0.01* | 32 ± 6*  | 2.1 ± 0.2* | 30 ± 3*  | 6.3 ± 0.7* | 36 ± 6 | 230 ± 80* | 1.3 ± 0.3* | 3400 ± 700* | 0.5 ± 0.1* |
| N1/N2AΔ844     | 0.15 ± 0.02 | 46 ± 4  | 2.0 ± 0.3 | 46 ± 4* | 7 ± 1   | 11 ± 2* | 900 ± 400* | 0.5 ± 0.1* | 8,000 ± 1,000* | 0.7 ± 0.1* |

Symbols mark values that are statistically different compared with N1/N2A (*) or N1/N2AΔ844 (†), (p < 0.05, Student’s t-test).

Figure 2. Reaction mechanisms of N1/N2A900A and N1/N2A929A. Rates (s⁻¹) were optimized by fitting the illustrated model directly to idealized single-channel data in each record and are presented as mean values for each data set rounded to the first significant figure. Symbols (*) mark values that are significantly different, larger (red) or smaller (blue) relative to those obtained for N1/N2A receptor (p < 0.05, Student’s t-test).
Ca\textsuperscript{2+} transient. Our results with brief synaptic-like stimuli predicted that both N1/N2A\textsuperscript{900A} and N1/N2A\textsuperscript{929A} receptors will generate macroscopic currents with lower peaks and shorter decay times (Fig. 3C). Notably, these results match closely previous reports of calcineurin-dependent reduction in synaptic NR current by direct dephosphorylation of Ser-900.\textsuperscript{9}

**Discussion**

Based on our previous findings that the N2A CTD has substantial influence over NR gating,\textsuperscript{7} we investigated how phosphorylation/dephosphorylation cycles may control NR output. Our approach was to examine the gating properties of NRs that were rendered resistant to phosphorylation at two specific sites by serine-to-alanine single-residue substitutions. We recorded currents from individual channels with the cell-attached patch-clamp technique; examined the kinetic distributions of the recorded open and closed intervals; and used kinetic modeling to summarize and interpret the observed changes in terms of a previously proposed reaction scheme. We found that relative to wild-type N1/N2A receptors, N1/N2A\textsuperscript{900A} and N1/N2A\textsuperscript{929A} receptors were substantially less active, and their P_o values were low primarily because these channels remained closed for longer periods (longer MCT). Of the five components present in the closed duration distributions, all were longer, except for E_c. However, this longest time component, which reflects desensitized intervals, represented a substantially higher fraction of all closures. For N1/N2A\textsuperscript{929A}, but not for N1/N2A\textsuperscript{900A}, we also observed a slight decrease in mean open time; however, the open duration distributions for both mutants were very similar to those of wild-type receptors, an indication that perturbations to Ser-900 or Ser-929 of N2A subunits neither causes nor modulates modal gating. Instead, kinetic modeling of our single channel data indicated changes at several activation and desensitization steps, and predicted that the phosphorylation status at these serine residues may control the kinetics and extent of macroscopic desensitization and also the decay of the synaptic response.

Because in our records we tracked channel activity as pure Na\textsuperscript{+} currents, and because all our measurements were done with the cell-attached technique, we presume that the receptor trapped within the patch pipette displayed the high level of basal activity that precedes Ca\textsuperscript{2+}-dependent desensitization, inactivation or run down events.\textsuperscript{8,10-15} This premise was supported by our observation that in the records we obtained, which lasted between 10 and 90 min, channel activity was independent of recording time. For these reasons we assume that the reduction in P_o we observed for N1/N2A\textsuperscript{900A} and N1/N2A\textsuperscript{929A} receptors was caused directly by the site-specific perturbations we introduced, and also that the behaviors we characterized for each mutant may inform about the gating mechanism of wild-type N1/N2A receptors that are dephosphorylated at these sites.

Our simulations based on the kinetic models developed from single-channel data of N1/N2A\textsuperscript{900A} and N1/N2A\textsuperscript{929A} agree well with the conclusions by the Westbrook group that these two residues control macroscopic desensitization.\textsuperscript{8} The shape of our simulated currents, however, was distinct from those reported by the Westbrook group. Namely, they found that N1\textsuperscript{838}/N2A\textsuperscript{900A} had a rapid initial desensitization, reminiscent of wild type receptors that were repeatedly exposed to glutamate in the presence of Ca\textsuperscript{2+}, while N1\textsuperscript{838}/N2A\textsuperscript{929A} were resistant to this time-dependent form of desensitization. Our simulations predicted that N1/N2A\textsuperscript{900A} will have deeper desensitization than wild type receptors, whereas N1/N2A\textsuperscript{929A} will desensitize even more. However, these quantitative discrepancies may originate from the different constructs used (with or without the N1 CTD) and also permeant ions present (with or without Ca\textsuperscript{2+}), and thus may point to the likelihood that in addition to kinase/phosphatase systems acting at the two serine residues examined here, separate biochemical pathways may target sites located elsewhere on NR C-termini.

Overall, our simulations with brief pulses of glutamate predicted that substituting an alanine residue at Ser-900 or Ser-929 will cause NR-mediated synaptic currents to become smaller and to decay faster (Fig. 3C). These constructs were developed with the intent of replicating a WT receptor that cannot be phosphorylated at these sites. Our results are consistent with a series of literature reports that

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**Figure 3.** Predicted effects of N2A S900A and S929A substitutions on NR macroscopic responses. Responses to 5 sec (A and B) or 1 ms (C) applications of glutamate (1 mM) were simulated with the models in Figure 2. Current traces are normalized to peak in (A).
show kinetic effects of dephosphorylation at these sites. Tong and Jahr reported that repeated stimulation of synaptic NRs in the presence of physiologic Ca\textsuperscript{2+} reduces the NR-mediated synaptic current by a mechanism that involves Ca\textsuperscript{2+} activation of calcineurin. At least part of this effect is mediated by dephosphorylation at Ser-900 of N2A subunits. Further, inhibiting protein kinase A (PKA) prevents the calcineurin-mediated reduction of NR currents and conversely, stimulating PKA overcomes the ability of calcineurin to reduce NR currents; however, PKA stimulation alone does not elevate NR currents above basal levels. The latter observations are salient to our report, in light of biochemical evidence that PKA directly phosphorylates NRs in vivo at multiple sites on both N1 and N2A subunits.

Together, the available data on NR modulation by kinase/phosphatase systems paint the view that during quiescent periods, synaptic NRs are basally phosphorylated by (at least) PKA action and that the Ca\textsuperscript{2+} influx that accompanies synaptic stimulation of NRs results in calcineurin-mediated dephosphorylation of specific sites on NR. During development of the visual system, this latter effect causes NR currents to become smaller and to decay faster.

We should note that in addition to preventing phosphorylation at the examined sites, our serine-to-alanine substitutions may have also perturbed CTD structure in other ways, such as by altering gating directly or by abolishing protein-protein interactions aside from kinases. Despite this, the CTDs of NRs are believed to lack ordered structures, and to our knowledge, there is not yet evidence of protein-protein interactions at the examined sites. Given that the CTD of NRs make as much as one third of the receptor’s mass, it is likely that multiple mechanisms exist which can modulate gating through this domain. The role of CTD in NR gating is just beginning to be explored, and additional investigations are necessary to delineate these.

Overall, the state models we developed in this study extend on previous reports regarding changes in macroscopic NR responses caused by dephosphorylation of N2A Ser-900 or Ser-929 by illuminating the kinetic mechanisms responsible for the observed changes. These models will therefore be instrumental for quantitative investigations into the structural determinants of channel gating, as well as for studies that aim to reveal the mechanisms of NR regulation by kinase/phosphatase systems and their functions in health and disease.

Materials and Methods

DNA constructs and receptor expression. Rat GluN1-1a (U8261) and GluN2A (M91561) were subcloned into pcDNA 3.1(+). Alanine substitutions were introduced into the GluN2A sequence at positions Ser-900 or Ser-929 using standard molecular biology methods. Wild-type or mutated GluN2A subunits were co-expressed with GluN1-1a and GFP after the encoding plasmids were introduced into HEK293 cells with the Ca\textsuperscript{2+} precipitate method previously described in more detail. Transfected cells were cultured in standard medium supplemented with 2 mM Mg\textsuperscript{2+} and were used for experiments 24 to 48 h post transfection.

Electrophysiology. Steady-state activity was recorded as inward Na\textsuperscript{+}-only currents from cell-attached patches that contained one active channel after applying 100 mV through the recording pipette, which was filled with (extra-cellular) solution containing (in mM): 150 NaCl, 2.5 KCl, 10 HEPES, 1 EDTA, 1 glutamate and 0.1 glycine, adjusted to pH 8.0 (NaOH). In all experiments, currents were amplified and low-pass filtered at 10 kHz using Axopatch 200B, were digitally sampled at 40 kHz using National Instruments PCI-6229 A/D board, and were stored into digital files using QuB software (www.qub.buffalo.edu).

Kinetic analysis and modeling. Current amplitude levels and event durations were measured directly from the single-channel record using QuB software with methods described in detail elsewhere. Briefly, records were first minimally preprocessed to correct for baseline drifts and noise artifacts and were idealized with the SKM algorithm after imposing a 12 kHz Gaussian digital filter. Kinetic modeling was done with the MIL algorithm by fitting the illustrated models directly to the idealized data after imposing a dead time of 0.075 ms. This procedure also calculated time constants and areas for the exponential components present in the closed and open event distributions.

Simulations. To simulate glutamate-elicited currents we appended the illustrated kinetic models, which represent saturation gating, with glutamate association steps leading into the closed fully liganded state C3. We used association and dissociation rate constants previously determined for N1/N2A receptors in conditions similar to those described here (1.7 × 10\textsuperscript{3} M\textsuperscript{-1}s\textsuperscript{-1} and 60 sec\textsuperscript{-1}, respectively). Macroscopic responses were calculated as the time-dependent occupancies of the open state after exposing 100 resting channels (10 pA/each) to pulses (5 sec or 1 ms) of glutamate (1 mM).

Statistical tests. Results are reported as means ± SEM, unless otherwise indicated. Statistical significance of differences was evaluated with ANOVA or Student’s t-test, as indicated, and was considered significant for p values < 0.05.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

Acknowledgments
We thank Eileen Kasperek for help with molecular biology and tissue culture. This work was supported by the National Institutes of Health National Institute of Neurological Disorders and Stroke (grant 052669) to G.K.P.

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