Beta-adrenergic receptor activation rescues theta frequency stimulation-induced LTP deficits in mice expressing C-terminally truncated NMDA receptor GluN2A subunits

Citation for published version:

Digital Object Identifier (DOI):
10.1101/lm.2045311

Link:
Link to publication record in Edinburgh Research Explorer

Document Version:
Publisher's PDF, also known as Version of record

Published In:
Learning & Memory

Publisher Rights Statement:
Copyright © 2011 Cold Spring Harbor Laboratory Press

General rights
Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy
The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.
β-adrenergic receptor activation rescues theta frequency stimulation-induced LTP deficits in mice expressing C-terminally truncated NMDA receptor GluN2A subunits

Teena D. Moody,1 Ayako M. Watabe,2,3 Tim Indersmitten,1 Noboru H. Komiyama,4 Seth G.N. Grant,4 and Thomas J. O’Dell5,6

1Interdepartmental PhD Program for Neuroscience, University of California Los Angeles, Los Angeles, California 90024, USA; 2Laboratory of Neurophysiology, Department of Neuroscience, Jikei University School of Medicine, Tokyo 105-8461, Japan; 3PRESTO, Japan Science and Technology Agency (JST), Saitama 332-0012, Japan; 4Wellcome Trust Sanger Institute, Cambridge CB10 1SA, United Kingdom; 5Department of Physiology, David Geffen School of Medicine, University of California Los Angeles, Los Angeles, California 90024, USA

Through protein interactions mediated by their cytoplasmic C termini the GluN2A and GluN2B subunits of NMDA receptors (NMDARs) have a key role in the formation of NMDAR signaling complexes at excitatory synapses. Although these signaling complexes are thought to have a crucial role in NMDAR-dependent forms of synaptic plasticity such as long-term potentiation (LTP), the role of the C terminus of GluN2A in coupling NMDARs to LTP enhancing and/or suppressing signaling pathways is unclear. To address this issue we examined the induction of LTP in the hippocampal CA1 region in mice lacking the C terminus of endogenous GluN2A subunits (GluN2AΔC/ΔC). Our results show that truncation of GluN2A subunits produces robust, but highly frequency-dependent, deficits in LTP and a reduction in basal levels of extracellular signal regulated kinase 2 (ERK2) activation and phosphorylation of AMPA receptor GluA1 subunits at a protein kinase A site (serine 845). Consistent with the notion that these signaling deficits contribute to the deficits in LTP in GluN2AΔC/ΔC mice, activating ERK2 and increasing GluA1 S845 phosphorylation through activation of β-adrenergic receptors rescued the induction of LTP in these mutants. Together, our results indicate that the capacity of excitatory synapses to undergo plasticity in response to different patterns of activity is dependent on the coupling of specific signaling pathways to the intracellular domains of the NMDARs and that abnormal plasticity resulting from mutations in NMDARs can be reduced by activation of key neuromodulatory transmitter receptors that engage converging signaling pathways.

N-methyl-D-aspartate receptors (NMDARs) are heteromeric ligand-gated ion channels comprised of an obligatory GluN1 subunit along with different combinations of GluN2 or GluN3 subunits (Cull-Candy and Leszkiewicz 2004). In the adult brain, most NMDARs exist as heteromeric combinations of two GluN1 subunits along with different members of the GluN2 family of subunits (GluN2A, -2B, -2C and -2D). Importantly, the presence of different types of GluN2 subunits determines a number of functional characteristics of NMDARs, including agonist affinity, channel kinetics, and membrane localization (for review, see Cull-Candy et al. 2001, Cull-Candy and Leszkiewicz 2004). The cytoplasmic C-terminal tails of the different GluN2 subunits also have a crucial role in coupling NMDARs to various adaptor/scaffolding proteins, such as the membrane associated guanylate kinases or MAGUKs (Kornau et al. 1995; Kim and Sheng 2004). The MAGUKs, which include proteins such as PSD-95, PSD-93, and SAP102, in turn couple NMDARs to a host of postsynaptic signaling pathways, including the Ras GTPase-activating protein SynGAP (Chen et al. 1998; Kim et al. 1998), the A kinase anchoring protein AKAP79/150 (Colledge et al. 2000), and neuronal nitric oxide synthase (nNOS) (Brenman et al. 1996). Moreover, the C termini of GluN2A and GluN2B also directly bind key components of additional signaling pathways, such as α-calcium/calmodulin-activated protein kinase II (CamKII) (Leonard et al. 1999; Bayer et al. 2001) and phospholipase C (Gurd and Bissoon 1997). Thus, GluN2B subunits not only regulate the biophysical properties of NMDARs but also have an essential role in the formation of NMDAR signaling complexes that organize signaling downstream from NMDAR activation. Indeed, disruption of these complexes is associated with pronounced alterations in NMDAR-dependent forms of synaptic plasticity such as long-term potentiation (LTP) and long-term depression (LTD) (Migaud et al. 1999; Sprengel et al. 1998; Sprengel et al. 1998; Barria and Malinow 2005; Cuthbert et al. 2007; Zhou et al. 2007; Carlisle et al. 2008; Xu et al. 2008; Foster et al. 2010).

Although proteomic approaches have revealed much about the molecular composition of NMDAR signaling complexes (Husi et al. 2000; Collins et al. 2006), the role of these complexes in regulating LTP and LTD induction is still poorly understood. For example, previous studies using genetically modified mice where the endogenous GluN2A subunits lack the cytoplasmic C terminus (GluN2AΔC) have found that these mutants exhibit deficits in both LTP (Sprengel et al. 1998; Köhr et al. 2003) and some forms of hippocampus-dependent learning (Sprengel et al. 1998;
Bannerman et al. 2008). This suggests that the C terminus of GluN2A subunits has an important role in recruiting signaling molecules that facilitate LTP induction to NMDAR signaling complexes. In contrast to these studies where the endogenous GluN2A subunit gene was modified, other studies have overexpressed cDNAs encoding truncated GluN2A. For example, in hippocampal slice cultures overexpression of truncated GluN2A subunits can restore LTP in cells where LTP induction is disrupted by knockdown of GluN2B subunits while overexpression of full-length, wild-type GluN2A subunits cannot (Foster et al. 2010). While these data support the conclusion that the C terminus of GluN2A is involved in signaling, it remains unclear if both LTP inhibiting or enhancing pathways act via the C terminus and to what extent these pathways might be recruited by different patterns of synaptic activity.

Here we have reexamined the role of the C terminus of NMDAR GluN2A subunits in LTP using a variety of different LTP induction protocols and biochemical approaches to examine NMDAR signaling in hippocampal slices from wild-type and homozygous mutant mice expressing truncated GluN2A subunits lacking the cytoplasmic C terminus (GluN2AΔC/ΔC) (Sprengel et al. 1998). We find that while high-frequency stimulation (HFS) induced LTP is normal, the induction of LTP by theta-frequency patterns of synaptic activation is strongly disrupted in GluN2AΔC/ΔC mice. This suggests that the C terminus of GluN2A subunits couples NMDARs to downstream signaling pathways that normally facilitate the induction of LTP by low frequency patterns of synaptic activity. Western blot analysis of hippocampal homogenates revealed that basal levels of extracellular signal regulated kinase 2 (ERK2) activation and phosphorylation of AMPA receptor (AMPAR) GluA1 subunits at serine 845 (S845) signal regulated kinase 2 (ERK2) activation and phosphorylation.

**Results**

**Basal synaptic transmission is normal in GluN2AΔC/ΔC mutant mice**

To examine whether truncation of GluN2A subunits alters basal synaptic transmission, we recorded AMPAR-mediated spontaneous miniature excitatory postsynaptic currents (mEPSCs) in CA1 pyramidal cells from wild-type and GluN2AΔC/ΔC mice. Consistent with previous findings suggesting that AMPAR-mediated synaptic transmission is not altered in GluN2A mutant mice (Sprengel et al. 1998; Steigerwald et al. 2000), both the amplitude and frequency of mEPSCs were normal in GluN2AΔC/ΔC mice (Fig. 1A–C). Evoked EPSCs recorded at postsynaptic membrane potentials of −80 mV and +40 mV were also similar in wild-type and GluN2AΔC/ΔC mice (Fig. 1D) and the ratio of NMDAR to AMPAR-mediated currents measured at both membrane potentials in GluN2AΔC/ΔC mice was not significantly different from that seen in wild-type cells (Fig. 1E). While this difference did not reach statistical significance (P = 0.08), it is consistent with previous findings indicating that GluN2AΔC-containing NMDARs may be less tightly localized with respect to presynaptic release sites (Steigerwald et al. 2000).

**GluN2AΔC/ΔC exhibit theta frequency-specific deficits in LTP**

To examine whether truncation of the C terminus GluN2A subunits alters the induction of LTP by different patterns of synaptic activity. Western blot analysis of hippocampal homogenates revealed that basal levels of extracellular signal regulated kinase 2 (ERK2) activation and phosphorylation of AMPA receptor (AMPAR) GluA1 subunits at serine 845 (S845) are reduced in GluN2AΔC/ΔC mice. Consistent with the notion that these signaling deficits contribute to the disruption of LTP in these mutants, enhancing ERK activation and increasing GluA1 phosphorylation at S845 by β-adrenergic receptor activation reduced, and for some stimulation protocols completely rescued, the theta frequency stimulation-induced LTP deficits in GluN2AΔC/ΔC mice.

![Figure 1](image_url) Basal synaptic transmission is normal in GluN2AΔC/ΔC mice. (A) Traces show example mEPSCs recorded in CA1 pyramidal cells from wild-type (GluN2AΔC/ΔC) and GluN2AΔC/ΔC mice. (B,C) Both mEPSC amplitude and frequency are normal in cells from GluN2AΔC/ΔC mice. Results are from 13 cells from three wild-type mice and 11 cells from three GluN2AΔC/ΔC mice. (D) Traces show examples of evoked EPSCs recorded at holding potentials of −80 mV and +40 mV in wild-type (left) and GluN2AΔC/ΔC cells (right). (E) NMDAR to AMPAR ratios at both −80 mV and +40 mV are similar in cells from wild-type (n = 14 cells from three mice) and GluN2AΔC/ΔC mice (n = 12 cells from three mice). (F) Weighted decay time constants (τw) of EPSCs recorded at +40 mV were 84 ± 2 msec in wild-type cells and 101 ± 7 msec in GluN2AΔC/ΔC cells (P = 0.08 compared to wild type).
presynaptic fiber stimulation we first examined the effects of multiple trains of HFS (100 Hz) on synaptic strength in GluN2A<sup>−/−</sup> and wild-type mice. As shown in Figure 2A, the induction of LTP by six trains of 100 Hz in slices from GluN2A<sup>−/−</sup> was similar to that seen in wild-type slices (fEPSPs potentiated to 190 ± 18% of baseline in wild-type slices and to 183 ± 13% of baseline in slices from GluN2A<sup>−/−</sup> mutants, P = 0.75 compared to wild type). A weaker HFS protocol (two trains of 100 Hz stimulation) also induced robust LTP in slices from GluN2A<sup>−/−</sup> that was not significantly different from that seen in slices from wild-type mice (fEPSPs potentiated to 177 ± 13% of baseline in wild-type slices and to 172 ± 15% of baseline in slices from GluN2A<sup>−/−</sup>, P = 0.82 compared to wild type; Fig. 2B). Together, these findings are consistent with previous reports showing that the induction of LTP by multiple trains of high-frequency stimulation is normal in GluN2A<sup>−/−</sup> mutants (Köhr et al. 2003).

Although our results indicate that HFS-induced LTP is normal in GluN2A<sup>−/−</sup> mice, significant LTP deficits were observed in experiments where we examined the induction of LTP by lower frequency patterns of presynaptic fiber stimulation. As shown in Figure 2A, a long train (900 pulses) of 20 Hz stimulation induced wild-type levels of LTP in slices from GluN2A<sup>−/−</sup> mice (Fig. 2C). However, the same number of presynaptic fiber stimulation pulses delivered at 10 Hz, a frequency that falls within the upper range of the hippocampal theta rhythm (Buzsáki 2002), induced significantly less LTP in slices from GluN2A<sup>−/−</sup> mice (fEPSPs potentiated to 157 ± 11% of baseline in wild-type slices and to 122 ± 5% of baseline in slices from GluN2A<sup>−/−</sup> mice, P < 0.01 compared to wild type) (Fig. 2D). Similarly, 900 pulses of presynaptic fiber stimulation at the lower range of the theta rhythm (5 Hz) induced a small (126 ± 10% of baseline), yet significant potentiation of synaptic transmission in slices from wild-type mice (P < 0.05 compared to baseline) but had no lasting effect on synaptic strength in slices from GluN2A<sup>−/−</sup> mutant mice (fEPSPs were 94 ± 7% of baseline, P < 0.05 compared to wild type; Fig. 2E). Thus, as summarized in Figure 2F, truncation of the C terminus

![Figure 2. GluN2A<sup>−/−</sup> mutants exhibit frequency-dependent deficits in LTP. (A) LTP induced by six trains of 100 Hz stimulation (starting at time = 0, intertrain interval = 5 min) in slices from GluN2A<sup>−/−</sup> mice (n = 8 slices from five mice) is not significantly different from that seen in wild-type slices (n = 4 slices from four mice, P = 0.75). (B) Two trains of 100 Hz stimulation (delivered at time = 0, intertrain interval = 10 sec) induces similar levels of LTP in slices from wild-type (n = 6 slices from four mice) and GluN2A<sup>−/−</sup> mice (n = 5 slices from five mice). (C) The induction of LTP by a train of 20 Hz stimulation (900 pulses, delivered at time = 0) in slices from GluN2A<sup>−/−</sup> (n = 9 slices from seven mice) is not significantly different from that seen in wild-type slices (n = 4 slices from four mice, P = 0.63). (D) Ten Hz stimulation (900 pulses delivered at time = 0) induces significantly less LTP in GluN2A<sup>−/−</sup> mutants (n = 14 slices from seven mice) than in wild-type slices (n = 9 slices from five mice, P < 0.01). (E) Five Hz stimulation (900 pulses delivered at time = 0) induced a small amount of potentiation in slices from wild-type mice (n = 8 slices from six mice, P < 0.05 compared to baseline) but had no lasting effect on synaptic strength in slices from GluN2A<sup>−/−</sup> mutants (n = 6 slices from six mice, P < 0.05 compared to wild type). (F) Summary of the effects of different frequencies of presynaptic fiber stimulation on the induction of LTP in wild-type and GluN2A<sup>−/−</sup> mice (⁎P < 0.01; ⁎⁎ P < 0.05). Results for 100 Hz stimulation are from experiments shown in B.)
of GluN2A subunits strongly disrupts the induction of LTP by trains of synaptic stimulation delivered at the theta frequency (5–10 Hz) but has no effect on the induction of LTP by higher frequencies of synaptic stimulation (20 and 100 Hz).

Basal levels of ERK2 and GluA1 S845 phosphorylation are altered in GluN2A<sup>AC/Ac</sup> mutant mice

Unlike HFS-induced LTP, the induction of LTP by theta-frequency stimulation protocols is dependent on ERK activation (Winder et al. 1999; Watabe et al. 2000; Opazo et al. 2003) and strongly inhibited by protein phosphatase activity (Thomas et al. 1996; Brown et al. 2000). We thus examined ERK activation and phosphorylation of AMPAR GluA1 subunits at sites dephosphorylated by protein phosphatases activated downstream from NMDARs to determine whether truncation of GluN2A subunits might lead to deficits in theta-frequency stimulation-induced LTP through alterations in these signaling pathways. As shown in Figure 3A, immunoblotting of homogenates prepared from wild-type and GluN2A<sup>AC/Ac</sup> slices revealed a modest, but significant, reduction in basal levels of the phosphorylated, active form of ERK2 in slices from GluN2A<sup>AC/Ac</sup> mice (levels were reduced to 40% of wild-type levels, P < 0.05) with no difference in total ERK2 levels.

However, the relative increase in ERK2 phosphorylation induced by bath application of NMDA (20 μM NMDA (N) induced similar increases in ERK2 phosphorylation in slices from wild-type (open bars, n = 5) and GluN2A<sup>AC/Ac</sup> mice (filled bars, n = 7). *P < 0.01 compared to untreated (UT) control slices.

Figure 3. Basal levels of ERK2 phosphorylation are significantly reduced in GluN2A<sup>AC/Ac</sup> mice. (A) Levels of phospho-ERK2 in untreated hippocampal slices from wild-type (open bars, n = 5) and GluN2A<sup>AC/Ac</sup> mice (filled bars, n = 7). Total ERK1/2 levels in GluN2A<sup>AC/Ac</sup> mice were not different from wild type (P = 0.91). (B) A 3 min bath application of 20 μM NMDA (N) induced similar increases in ERK2 phosphorylation in slices from wild-type (open bars, n = 5) and GluN2A<sup>AC/Ac</sup> mice (filled bars, n = 7). *P < 0.01 compared to untreated (UT) control slices.

Basal levels of GluA1 phosphorylation at T840 were not altered in slices from GluN2A<sup>AC/Ac</sup> mice (levels were 91 ± 6% of wild-type levels, P = 0.38) while basal levels of GluA1 phosphorylation at S845 were lower in slices from GluN2A<sup>AC/Ac</sup> compared to wild-type slices (levels were 80 ± 5% of wild-type levels, P = 0.05) (Fig. 4A,B). However, dephosphorylation of AMPAR GluA1 subunits at S845 and T840 induced by transient NMDA receptor activation (bath application of 20 μM NMDA for 3 min) was not altered in slices from GluN2A<sup>AC/Ac</sup> mice (Fig. 5A,B). Although NMDAR activation triggers dephosphorylation of AMPAR GluA1 subunits at T840 and S845, phosphorylation of another site in GluA1, S831, is increased (Vanhoose and Winder 2003; Delgado et al. 2007). Thus, as an additional test for potential changes in protein kinase and/or protein phosphatase signaling in GluN2A<sup>AC/Ac</sup> mice we also compared phospho-S831 GluA1 levels in untreated and NMDA-treated slices from wild-type and GluN2A<sup>AC/Ac</sup> mice. As shown in Figures 4C and 5C, both basal levels of GluA1 phosphorylation at S831 and the increase in S831 phosphorylation induced by NMDAR activation were normal in slices from GluN2A<sup>AC/Ac</sup> mice. Together, these findings suggest that truncation of the C terminus of GluN2A subunits does not produce significant deficits in the ability of bath applied NMDA to activate the ERK pathway, protein kinases that regulate S831 phosphorylation (such as PKC and CaMUK), or protein phosphatases that dephosphorylate T840 and S845. It is, however, associated with modest, but
significant, decreases in basal levels of both ERK2 activation and GluA1 S845 phosphorylation.

β-adrenergic receptor activation rescues the induction of LTP in GluN2A\textsuperscript{ΔC/ΔC} mice

The above data shows that the GluN2A C terminus is required for the basal level of phosphorylation of synaptic proteins while leaving intact those kinase and phosphatase pathways that are induced by transient NMDA receptor activation. This led us to ask: Could reduced basal levels of ERK2 and GluA1 S845 phosphorylation contribute to the deficits in theta-frequency stimulation-induced LTP seen in GluN2A\textsuperscript{ΔC/ΔC} mice? To examine this we took advantage of the fact that activation of β-adrenergic receptors in the hippocampal CA1 region activates both the ERK pathway (Roberson et al. 1999; Winder et al. 1999; Opazo et al. 2003) and increases GluA1 phosphorylation at S845 (Opazo et al. 2003; Vanhoose and Winder 2003; Delgado et al. 2007; Joiner et al. 2010) and tested whether activation of β-adrenergic receptors could rescue theta-frequency stimulation-induced LTP in GluN2A\textsuperscript{ΔC/ΔC} mice. As shown in Figure 6, a 10 min bath application of the β-adrenergic receptor agonist isoproterenol (ISO, 1 μM) induced an approximately twofold increase in ERK2 phosphorylation and a more than threefold increase in GluA1 S845 phosphorylation in slices from both wild-type and GluN2A\textsuperscript{ΔC/ΔC} mice. Thus, we next examined whether β-adrenergic receptor activation can rescue theta-frequency stimulation-induced LTP in GluN2A\textsuperscript{ΔC/ΔC} mutant mice. Although long trains of 5 Hz stimulation induce relatively little LTP in slices from wild-type mice (Fig. 2E), shorter trains of 5 Hz stimulation (150 pulses) can induce robust LTP (Thomas et al. 1996, 1998; Carlisle et al. 2008) that is relatively insensitive to the effects of β-adrenergic receptor activation (Thomas et al. 1996). Consistent with this, short trains of 5 Hz stimulation induced robust LTP in slices from wild-type mice (fEPSPs were potentiated to 180 ± 23% of baseline; Fig. 7A) that was not significantly enhanced by bath application of ISO prior to 5 Hz stimulation (fEPSPs were potentiated to 204 ± 16% of baseline; Fig. 6B). A very different pattern of results was observed, however, in slices from GluN2A\textsuperscript{ΔC/ΔC} mice. First, the
induction of LTP by 150 pulses of 5 Hz stimulation alone was significantly reduced compared to that seen in slices from wild-type mice (fEPSPs were 115 ± 5% of baseline, \( P < 0.05 \) compared to wild-type controls; Fig. 7A). Second, activation of \( \beta \)-adrenergic receptors strongly enhanced the induction of LTP by 150 pulses of 5 Hz stimulation in GluN2A\(^{AC/AC} \) mutants (fEPSPs were potentiated to 167 ± 7% of baseline, \( P < 0.001 \) compared to GluN2A\(^{AC/AC} \) control, \( P < 0.05 \) compared to wild-type 5 Hz + ISO; Fig. 7B). Together, these results indicate that \( \beta \)-adrenergic receptor activation can, at least partially, rescue the induction of theta-frequency stimulation-induced LTP in GluN2A\(^{AC/AC} \) mice. Indeed, although the induction of LTP by long trains of 5 Hz stimulation is significantly compromised in slices from GluN2A\(^{AC/AC} \) mice (Fig. 2E), \( \beta \)-adrenergic receptor activation completely rescued the induction of LTP by this pattern of synaptic stimulation in GluN2A\(^{AC/AC} \) mice (Fig. 7C).

Discussion

Theta-frequency stimulation-induced LTP is specifically disrupted by loss of the C terminus of NMDAR GluN2A subunits

Consistent with previous findings (Sprengel et al. 1998; Köhr et al. 2003), our results indicate that the C-terminal domain of GluN2A subunits has a crucial, facilitatory role in the induction of LTP in the hippocampal CA1 region. Our results indicate, however, that the LTP deficits observed in GluN2A\(^{AC/AC} \) mutants are remarkably frequency dependent and are only observed following the induction of LTP by frequencies of presynaptic fiber stimulation that fall within the theta rhythm, a 5–12 Hz oscillation of neural activity that occurs within the hippocampus during certain behaviors (Buzsáki 2002). Like high-frequency stimulation-induced LTP, the induction of LTP by theta frequency patterns of synaptic stimulation is NMDAR-dependent (Thomas et al. 1996, 1998; Winder et al. 1999; Watabe and O’Dell 2003). Thus, a reduction in NMDAR activity that increases the threshold for LTP induction could provide a simple explanation for the theta-frequency LTP deficits in GluN2A\(^{AC/AC} \) mice. We found, however, that both basal AMPA-mediated synaptic transmission and the NMDAR-mediated component of evoked EPSCs were normal in slices from GluN2A\(^{AC/AC} \) mice (Fig. 1). This suggests that the LTP deficits observed in GluN2A\(^{AC/AC} \) mice are not due to alterations in NMDAR currents but instead reflect alterations in signaling downstream from NMDAR activation (also see Köhr et al. 2003).

C-terminal truncation of NMDAR GluN2A subunits leads to alterations in both ERK and AMPA receptor GluA1 subunit phosphorylation

The induction of LTP by theta-frequency stimulation protocols exhibits a number of unique properties that may provide important clues regarding how disruption of signaling pathways dependent on interactions with the C terminus of GluN2A subunits compromises the induction of LTP. For example, unlike high-frequency stimulation-induced LTP, the induction of theta-frequency stimulation-induced LTP is highly dependent on ERK signaling (Winder et al. 1999; Watabe et al. 2000; Opazo et al. 2003). Thus we investigated whether alterations in NMDAR-dependent activation of the ERK pathway might contribute to the theta-frequency stimulation-induced LTP deficits in GluN2A\(^{AC/AC} \) mice. Although activation of the ERK pathway induced by bath application of NMDA was not altered, we did find a modest, but significant, decrease in basal levels of ERK phosphorylation in GluN2A\(^{AC/AC} \) mice. Some studies suggest that activation of GluN2A subunit-containing NMDARs is specifically involved in NMDAR-dependent activation of the ERK pathway through activation of the Ras guanine nucleotide exchange factor Ras-GRF2 (Li et al. 2006; Jin and Feig 2010). Although this suggests that deficits in ERK signaling in GluN2A\(^{AC/AC} \) mutants may be due to alterations in the ability of NMDARs to couple to activators of the ERK pathway, the role of the C terminus of GluN2A subunits in coupling GluN2A-containing NMDARs to Ras-GRF2 is not known. Moreover, other studies have found that activation of either GluN2A or GluN2B-containing NMDARs can produce activation of the ERK pathway in mature neurons (Kim et al. 2005). Importantly, GluN2B-containing receptors, but not GluN2A-containing NMDARs, are coupled to SynGAP (Kim et al. 2005), a synaptic Ras GTase-activating protein that acts as a suppressor of both basal (Komiyama et al. 2002) and NMDAR-dependent increases in ERK1/2 phosphorylation (Rumbaugh et al. 2006). GluN2B-containing NMDARs also appear to be preferentially coupled to the tyrosine phosphatase
striatal-enriched phosphatase (STEP), another negative regulator of ERK activation (Surojit and Connor 2010). Thus, truncation of the C terminus of GluN2A subunits may alter the balance of activators and inhibitors of the ERK pathway coupled to NMDARs in a way that favors inhibition of ERK activation.

The induction of LTP by theta-frequency stimulation is also strongly regulated by activation of protein phosphatases 1 and/or 2A that normally act to inhibit the induction of LTP by these patterns of synaptic stimulation (Thomas et al. 1996; Brown et al. 2000). Thus we also investigated whether alterations in NMDAR-dependent activation of these protein phosphatases might contribute to the theta-frequency stimulation-induced LTP deficits in GluN2A<sup>AC</sup>/<sup>AC</sup> mice. Although NMDA-induced dephosphorylation of AMPAR GluA1 subunit phosphorylation at S845 and T840 was not altered, basal levels GluA1 phosphorylation at S845 were significantly reduced in hippocampal slices from GluN2A<sup>AC</sup>/<sup>AC</sup> mice. Protein kinase A (PKA) mediated phosphorylation of GluA1 at S845 is thought to trigger the insertion of AMPARs into extrasynaptic sites where they are primed for insertion into synaptic sites following the induction of LTP (Esteban et al. 2003; Oh et al. 2006). Thus, lower levels of GluA1 S845 phosphorylation could contribute to the LTP deficits in GluN2A<sup>AC</sup>/<sup>AC</sup> by decreasing the pool of extrasynaptic AMPARs available for incorporation into synapses during LTP induction. Recent studies suggest that ERK also has a role in AMPAR trafficking during LTP (Patterson et al. 2010) and thus alterations in AMPAR trafficking due to reduced ERK activity in GluN2A<sup>AC</sup>/<sup>AC</sup> mice may also contribute to deficits in LTP in these mutants. Notably, basal levels of GluA1 phosphorylation at T840 and S831 were not altered in slices from GluN2A<sup>AC</sup>/<sup>AC</sup> mice. This suggests that the lower levels of S845 phosphorylation are not due to changes in the activity of protein phosphatases 1 and 2A, which regulate GluA1 phosphorylation at all these sites. It thus seems more likely that loss of the C terminus of GluN2A subunits leads to alterations in NMDAR coupling to PKA signaling. One possibility is that truncation of the C terminus of GluN2A subunits compromises coupling of NMDARs to MAGUKs, such as PSD-95, that normally recruit PKA to NMDAR signaling complexes by interacting with a kinase anchoring proteins (Colledge et al. 2000). PSD-95 also couples GluN2A-containing NMDARs to nNOS (Brennan et al. 1996; Christopherson et al. 1999; Al-Hallaq et al. 2007) which, when activated, leads to the production of cGMP and activation of cGMP-dependent protein kinases that can also phosphorylate GluA1 subunits at S845 (Serulle et al. 2007). Thus, deletion of the C terminus of GluN2A subunits may also lead to alterations in GluA1 S845 phosphorylation via changes in NOS and cGMP signaling.

β-adrenergic receptor activation rescues the induction of LTP in GluN2A<sup>AC</sup>/<sup>AC</sup> mice

Consistent with the notion that alterations in ERK signaling and GluA1 S845 phosphorylation contribute to LTP deficits in GluN2A<sup>AC</sup>/<sup>AC</sup> mutants, activation of β-adrenergic receptors, which both activates ERK and increases GluA1 phosphorylation at S845 (Fig. 5), rescued the induction of theta-frequency stimulation-induced LTP in GluN2A<sup>AC</sup>/<sup>AC</sup> mice (Fig. 6). Importantly, β-adrenergic receptor stimulation and activation of PKA can enhance LTP induction via a number of different mechanisms in addition to ERK activation and AMPAR phosphorylation (O'Dell et al. 2010). For example, β-adrenergic receptor-activation mediated inhibition of small conductance, Ca<sup>2+</sup>-activated potassium channels as well as A-type potassium channels have been implicated in the LTP enhancing effects of β-adrenergic receptor activation (Yuan et al. 2002; Faber et al. 2008; for review, see O’Dell et al. 2010). Moreover, β-adrenergic receptor activation can inhibit the activity of protein phosphatases that oppose LTP induction (Thomas et al. 1996; Brown et al. 2000) as well as enhance the translation of proteins needed for the long-term maintenance of LTP (Gelin et al. 2007). The ability of β-adrenergic receptor activation to rescue theta-frequency stimulation-induced LTP in GluN2A<sup>AC</sup>/<sup>AC</sup> mutants may thus also reflect compensation for deficits in NMDAR modulation of these processes caused by loss of the C terminus of GluN2A subunits. Our results suggest, however, that protein phosphorylation downstream from NMDARs is not altered in GluN2A<sup>AC</sup>/<sup>AC</sup> mice. Because the deficits in theta-frequency stimulation-induced LTP in GluN2A<sup>AC</sup>/<sup>AC</sup> mutants are evident at very early time points post-LTP induction (5–10 min) our results also suggest that alterations in protein synthesis are unlikely to be involved. They do not, however, rule out the possibility that deficits in basal translation rates, perhaps due to alterations in basal ERK activity, might contribute to LTP deficits in GluN2A<sup>AC</sup>/<sup>AC</sup> mutants. Our results also do not rule out the possibility that deletion of the C terminus of GluN2A subunits leads to alterations in NMDAR modulation of dendritic potassium channels. Indeed, ERK activation inhibits A-type potassium channels in hippocampal dendrites (Yuan et al. 2002) and thus the lower basal levels of ERK activation in GluN2A<sup>AC</sup>/<sup>AC</sup> mice might lead to changes in dendritic excitability that contribute to the LTP deficits in these mutants. Although we did not observe any obvious alterations in postsynaptic complex spike bursting during theta frequency stimulation (Thomas et al. 1998) in GluN2A<sup>AC</sup>/<sup>AC</sup> mice (data not shown), future experiments that examine this possibility in more detail could provide important insights into the mechanisms underlying LTP deficits produced by truncation of the C terminus of GluN2A subunits.

Comparison of results between gene targeting and overexpression approaches

Our LTP findings are in general agreement with previous studies of GluN2A<sup>AC</sup>/<sup>AC</sup> mutants generated by gene targeting of the endogenous exon encoding the GluN2A C terminus (Sprengel et al. 1998; Köhr et al. 2003). In contrast to these studies that found a reduction in LTP with one stimulation protocol we have examined a wide range of frequencies in identical experimental conditions, which uncovered frequency-specific effects. A completely different experimental approach was reported in a recent study using RNA interference approaches in hippocampal organotypic slice cultures from wild-type rats came to just the opposite conclusion—that the C terminus of GluN2A subunits recruits negative regulators of LTP induction to NMDA receptor signaling complexes (Foster et al. 2010). Their conclusion derived from experiments that combined a knockdown of GluN2B with the overexpression of a cDNA encoding a truncated GluN2A in the presence of expression of the endogenous full-length GluN2A (full-length and truncated GluN2A are coexpressed). In part, the conclusions of Foster et al. (2010) were based on the finding that deficits in pairing-induced LTP caused by knockdown of GluN2B subunits could be overcome by overexpression of GluN2A subunits lacking the C terminus but not by overexpression of full-length GluN2A subunits. There are a number of methodological differences that may contribute to this discrepancy, but there are three that we believe are potentially interesting. First, GluN2B-containing NMDARs are not altered in GluN2A<sup>AC</sup>/<sup>AC</sup> mutants (Köhr et al. 2003) while the experiments of Foster et al. (2010) examined the role of the C terminus of GluN2A subunits in LTP in the context of GluN2B knockdown. The different effects of C-terminal truncation of GluN2A subunits on LTP induction thus appears to depend on the presence of GluN2B subunits. This suggests that interactions between signaling molecules recruited to NMDAR signaling complexes through
the C termini of both GluN2A and GluN2B subunits may have an important role in regulating LTP induction. Second, pairing protocols induce an ERK-independent form of LTP in the mouse hippocampus (Ozawa et al. 2003) while the induction of LTP by theta-frequency stimulation is highly ERK-dependent. Thus, another possibility is that the C terminus of GluN2A subunits couples NMDARs to both LTP enhancing and suppressing signaling pathways that have distinct roles in the induction of LTP by different patterns of synaptic activation. Finally, in our experiments we examined the role of the C terminus of GluN2A subunits in LTP induction in the hippocampus of adult mice while organotypic slices prepared from the hippocampus of 7-d-old rats were used in the experiments of Foster et al. (2010). Thus, there may be important changes in the molecular composition of NMDAR signaling complexes that occur during development that are dependent on the C terminus of NMDAR GluN2A subunits. Although additional experiments will be needed to address these possibilities, our results along with the findings of previous studies (Sprengel et al. 1998; Körn et al. 2003; Foster et al. 2010) suggest that a better understanding of the role of the C terminus of GluN2A subunits in organizing NMDAR signaling complexes is likely to reveal important new insights into the mechanisms underlying the induction of NMDAR-dependent forms of LTP.

Future strategies for rescue of mutations affecting NMDA receptor complexes

Human mutations in proteins within the signaling complexes of NMDA receptor complexes (Husi et al. 2000; Collins et al. 2006; Fernandez et al. 2009) are known to be involved in many brain diseases (Grant et al. 2005). For example, recently it was reported that mutations in GluN2A and GluN2B result in epilepsy and mental retardation (Endele et al. 2010). In the current study of GluN2A mutations, we have found that the LTP deficits can be significantly reduced using β-adrenergic stimulation based on the insight that phosphorylation of signaling proteins downstream from β-adrenergic receptors are shared with those impacted by the glutamate receptor mutation. This demonstrates that mutation phenotypes affecting plasticity mechanisms are amenable to pharmacological manipulation and suggests an interesting avenue for further research that may lead to development of new therapies for brain diseases.

Materials and Methods

Slice preparation and extracellular recording

Hippocampal slices (400 μm thick) from adult (three to eight months of age) wild-type and GluN2AΔC/ΔC mice (Sprengel et al. 1998) were prepared using standard techniques approved by the UCLA Institutional Animal Care and Use Committee. Slices were allowed to recover for at least 1 h in an interface-slice type recording chamber (at 37°C) perfused with oxygenated (95% O2/5% CO2) artificial cerebrospinal fluid (ACSF) containing 124 mM NaCl, 4.4 mM KCl, 25 mM NaHCO3, 1 mM NaH2PO4, 2 mM CaCl2, 1.2 mM MgSO4, and 10 mM glucose. For extracellular recordings a bipolar, nichrome wire stimulating electrode placed in stratum radiatum of the CA1 region was used to activate Schaffer collateral/commissural fiber synapses (basal stimulation rate = 0.02 Hz) and the evoked field excitatory postsynaptic potentials (fEPSPs) were recorded using a glass microelectrode filled with ACSF (resistance = 5–10 MΩ). At the start of each experiment we first identified the maximal fEPSP amplitude that could be elicited in each slice and then adjusted the intensity of presynaptic fiber stimulation to evoke fEPSPs with amplitude approximately 50% of the maximal amplitude. LTP was elicited using either conventional HFS protocols (multiple trains of 100 Hz stimulation, each 1 sec in duration, with inter-train intervals of 10 sec or 5 min) or different numbers of single pulses of presynaptic fiber stimulation delivered at 5–20 Hz. For each protocol the average fEPSP slope measured over the last 5 min of the experiment was used for statistical comparisons (two-tailed Student t-tests). In cases where the same stimulation protocol was delivered to multiple slices from the same animal the results were averaged and the N used for statistical tests corresponds to the number of animals in each group. All results are reported as mean ± SEM.

Whole-cell recordings

Low resistance (2–6 MΩ) patch electrodes filled with a solution containing 102 mM cesium gluconate, 17.5 mM CsCl, 10 mM TEA-Cl, 5 mM QX314, 4.0 mM Mg-ATP, 0.3 mM Tris-GTP, and 20 mM HEPES (pH = 7.2) were used to record excitatory postsynaptic currents (EPSCs). In these experiments slices with CA3 region removed were maintained in a submerged-slice recording chamber and bathed in a modified ACSF containing picrotoxin (100 μM), elevated concentrations of CaCl2 and MgSO4 (4 mM each) and a lower concentration of KCl (2.4 mM). To examine the AMPAR and NMDAR-mediated components of the evoked EPSCs we measured the amplitude of EPSCs recorded at postsynaptic membrane potentials (Vhold) of −80 or +40 mV. The AMPAR-mediated component of the EPSC was measured at the time after EPSC onset corresponding to the peak amplitude of the EPSC at −80 mV while the amplitude of the EPSC 30 msec after onset was used to estimate the NMDAR-mediated component. The decay time constant of NMDAR-mediated EPSCs was obtained from double exponential fits to the decay of synaptic currents measured at +40 mV and calculated as a weighted mean decay time constant (Rumbaugh and Vicini 1999). The intensity of presynaptic fiber stimulation was adjusted to elicit EPSCs with a peak amplitude of 200–300 pA at Vhold = −80 mV. Miniature EPSCs (mEPSCs) were recorded at −80 mV in cells from slices bathed in a modified ACSF containing 1.0 μM tetrodotoxin and 100 μM picrotoxin. A template-based event detection routine in pClamp 10 (Molecular Devices) was used to analyze mEPSCs. Detected events smaller than 6 pA were excluded from the analysis. Statistical comparisons of evoked EPSCs were performed using two-tailed Student t-tests and Kolmogorov-Smirnov tests were used for comparisons of mEPSC amplitude and inter-event interval distributions. The results from multiple cells from the same animal were averaged and the N used for statistical tests equals the number of animals in each group.

Western immunoblotting

Hippocampal slices from wild-type and GluN2AΔC/ΔC mice were maintained in interface chambers perfused at 2 mL/min with warm (30°C), oxygenated ACSF and allowed to recover for 2–2.5 h. Slices were either left untreated or exposed to bath applied receptor agonists and then collected by rapidly transferring them into pre-frozen microcentrifuge tubes (Kontes Glass Company) kept on crushed dry ice. Treated slices were collected for analysis after a 10 min application of ISO (1 μM) or a 3 min application of NMDA (20 μM). Stock solutions of ISO (1 mM in H2O) were prepared before each experiment and diluted to 1 μM in ACSF prior to application. In all experiments three slices were pooled for each treatment condition.

The techniques used for Western blot analysis are described in detail elsewhere (Delgado et al. 2007). Briefly, slices were homogenized in an ice-cold homogenization buffer that contained both protease inhibitors (Protease Inhibitors Complete, Roche Molecular Biochemicals) and a mixture of different protein phosphatase inhibitors (Phosphatase Inhibitors (PhosphoStop) 1 and II, Sigma-Aldrich). 20 μg of total protein from each sample was resolved on 12% SDS-PAGE gels and then transferred onto nitrocellulose membranes. Primary antibody incubations were done overnight (at 4°C) and, after three washes in Tris-buffered saline containing 0.05% Tween-20, blots were incubated for two or more hours at room temperature in HRP-conjugated 1 and II antibodies. Images of immunoreactive bands visualized using enhanced chemiluminescence were acquired using a cooled CCD
camera-based image acquisition and analysis system (Quantity One, Bio-Rad). All blots were re-probed with an anti-tubulin antibody and the optical density values for each band of interest was normalized to the density values obtained for tubulin in the same lane to control for potential variations in loading. Mann-Whitney tests and two-tailed Student t-tests were used to determine statistical significance.

Antibodies to dually phosphorylated ERK1/2 and total ERK1/2 were obtained from Cell Signaling Technology. Antibodies to GluA1 and phospho-T840 GluA1 were obtained from Abcam and antibodies for phospho-S845 GluA1 and the βIIl isofrom of tubulin were obtained from Upstate Biotechnology (Millipore). Anti-phospho-S831 antibodies were obtained from Invitrogen.

Acknowledgments

This work was supported by National Institute of Mental Health grant number MH609197 (T.J.O.) and the Wellcome Trust and Wellcome Trust Genes to Cognition Programme (S.G.N.G.). T.J.O. is a member of the UCLA Brain Research Institute. We are grateful to Rolf Sprengel and Peter Seeburg for GluN2A mouse; to Maksym Kopański, Tomás Ryan, and Erin Gray for helpful comments on the manuscript; and to Kathryn Elsegood, Carole Froiz, and Dierk Biggs for help with animal shipments.

References


Patterson MA, Szatmari EM, Yasuda R. 2010. AMPA receptors are exocytosed in stimulated spines and adjacent dendrites in a Ras-ERK-dependent manner during long-term potentiation. *Proc Natl Acad Sci* **107**: 15951–15956.


Received August 10, 2010; accepted in revised form November 19, 2010.