Synaptically Released Glutamate Activates Extrasynaptic NMDA Receptors on Cells in the Ganglion Cell Layer of Rat Retina

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NMDA and AMPA receptors (NMDARs and AMPARs) are colocalized at most excitatory synapses in the CNS. Consequently, both receptor types are activated by a single quantum of transmitter and contribute to miniature and evoked EPSCs. However, in amphibian retina, miniature EPSCs in ganglion cell layer neurons are mediated solely by AMPARs, although both NMDARs and AMPARs are activated during evoked EPSCs. One explanation for this discrepancy is that NMDARs are located outside of the synaptic cleft and are activated only when extrasynaptic glutamate levels increase during coincident release from multiple synapses. Alternatively, NMDARs may be segregated at synapses that either are not spontaneously active or yield miniature EPSCs that are too small to detect. In this study, we examined excitatory, glutamatergic synaptic inputs to neurons in the ganglion cell layer of acute slices of rat retina. EPSCs, elicited by electrically stimulating presynaptic bipolar cells, exhibited both NMDAR- and AMPAR-mediated components. However, spontaneous EPSCs exhibited only an AMPAR-mediated component. The effects of low-affinity, competitive receptor antagonists indicated that NMDARs encounter less glutamate than AMPARs during an evoked synaptic response. Reducing glutamate uptake or changing the probability of release preferentially affected the NMDAR component in evoked EPSCs; reducing uptake revealed an NMDAR component in spontaneous EPSCs. These results indicate that NMDARs are located extrasynaptically and that glutamate transporters prevent NMDAR activation by a transmitter released from a single vesicle and limit their activation during evoked responses.

Key words: rat; retina; ganglion cell; low-affinity antagonist; glutamate transporter; spillover

At most excitatory synapses in the CNS, AMPA receptors (AMPARs) and NMDA receptors (NMDARs) are colocalized in the postsynaptic membrane (Bekkers and Stevens, 1989; McBain and Dingledine, 1992; Silver et al., 1992). Consequently, both receptor types usually contribute to evoked EPSCs and to miniature EPSCs, the postsynaptic response to a single quantum of transmitter.

In the retina, ganglion cells receive excitatory and inhibitory synaptic input from bipolar cells and amacrine cells at synapses in the inner plexiform layer (IPL). Ganglion cells integrate and translate this input into patterns of action potentials that are propagated along their axons in the optic nerve to targets in the lateral geniculate nucleus and superior colliculus. Cells in the ganglion cell layer (GLCs) (Matsui et al., 1998) of numerous species express both NMDARs and AMPARs (Aizenman et al., 1988; Mittman et al., 1990; Cohen et al., 1994). Accordingly, EPSCs evoked by either light or electrical stimulation exhibit NMDAR and AMPAR components (Mittman et al., 1990; Diamond and Copenhagen, 1993; Lukasiewicz and Roeder, 1995; Matsui et al., 1998; Higgs and Lukasiewicz, 1999; Matsui et al., 1999). However, in acute slices of amphibian retina spontaneous EPSCs (SEPSCs) lack an NMDAR-mediated component (Taylor et al., 1995; Matsui et al., 1998). Two mechanisms have been proposed to explain this result: first, that NMDARs are located extrasynaptically and are activated only by the concomitant release of many vesicles (Matsui et al., 1998; Higgs and Lukasiewicz, 1999); second, that NMDA and AMPA receptors are expressed separately at different synapses and that only the AMPAR-mediated SEPSCs are detectable (Taylor et al., 1995). Physiological data from amphibian retina supports the first possibility (Matsui et al., 1998), but the punctate expression and colocalization of NMDARs and the postsynaptic density protein PSD-95 in the IPL of rat retina (Fletcher et al., 2000) support the second idea, although the identity of the postsynaptic, immunopositive neurons was not established in the latter study.

We examined this issue in the rat retina by recording evoked and spontaneous EPSCs from GLCs in acute slices. Consistent with reports in amphibian retina, we find that electrically evoked EPSCs are mediated by both AMPARs and NMDARs, whereas SEPSCs are mediated solely by AMPARs. Using low-affinity competitive antigens of either receptor type, we show that AMPARs are exposed to more synaptically released glutamate than are NMDARs during an evoked response. In addition, reducing glutamate uptake or changing release probability affects the amplitudes of NMDAR EPSCs to a greater extent than those of AMPAR EPSCs. Finally, reducing glutamate uptake causes an NMDAR-mediated component to emerge in the SEPSCs. Taken together, these results suggest that NMDARs on GLCs are located outside of excitatory synapses and are activated only when multiple release events increase extrasynaptic glutamate levels sufficiently. These results suggest that glutamate transporters regulate NMDAR activation and, subsequently, the manner in which GLCs integrate synaptic input.
**MATERIALS AND METHODS**

**Slice preparation and solutions.** Retinal slices were prepared from Sprague Dawley rats (17–22 d) in accordance with the National Institute of Neurological Disorders and Stroke Animal Care and Use Committee guidelines. Both eyes were removed and immersed in oxygenated extracellular solution at room temperature. Extracellular solution contained (in mM): 119 NaCl, 2.5 KCl, 1.3 MgCl$_2$, 2.5 CaCl$_2$, 26.2 NaHCO$_3$, 1 NaH$_2$PO$_4$, 2 glucose, 2 Na pyruvate, and 4 Na lactate, bubbled with 95% O$_2$ and 5% CO$_2$. The cornea, iris, lens, and vitreous were removed from one eye with scissors. The retina was mechanically detached from the eyecup and immersed in 2% agarose (low-gelling temperature, type VII; Sigma, St. Louis, MO) and cut into 200-μm-thick slices on a vibratome (Leica, Nussloch, Germany). Slices were prepared and stored in oxygenated extracellular solution; they were transferred one at a time to the recording chamber, in which picrotoxin (100 μM) and strychnine (10 μM) were added to oxygenated extracellular solution to block inhibitory synaptic transmission. For outside-out patch experiments, the NaHCO$_3$ in the extracellular solution was replaced with 20 mM HEPES. In magnesium-free solutions, MgCl$_2$ was replaced with CaCl$_2$. The patch pipette solution contained (in mM): 120 Cs methanesulfonate, 10 EGTA, 20 HEPES, 2 MgATP, and 0.2 NaGTP. All solutions were adjusted to pH 7.4 with NaOH or CsOH and adjusted to 290–300 mOsm with sucrose. Reagents were obtained from Sigma, except for L-glutamate, L-2-amino-5-phosphonopentanoic acid (L-AP-5), and d,-threo-β-benzyoxysapparatate (TBOA), which were obtained from Cookson (Ballwin, MO).

**Solution delivery.** The recording chamber was superheated constantly at a low rate (1 ml/min) with control extracellular solution. During outside-out patch recordings, control and test solutions were delivered simultaneously through theta glass tubing (Warner Instruments, Hamden, CT) pulled to a tip width of 100 μm per barrel. The solution flow created a sharp interface between solutions delivered through neighboring barrels. Solution changes were made by moving the tubing rapidly with a piezo-electric bimorph (Piezo Systems, Cambridge, MA), such that the solution interface traversed the width of the patch pipette tip, enabling brief (1–2 msec) applications of L-glutamate.

**Electrophysiology.** All recordings were made from GLCs with an Axopatch 200B amplifiers (Axon Instruments, Foster City, CA) in voltage-clamp mode. Patch electrodes (#0010 glass; World Precision Instruments, Sarasota, FL) had tip resistances of 4–5 MΩ when filled with internal solution. Access resistance was 10–20 MΩ; it was monitored continuously and not compensated. Data acquisition and analysis were performed with custom macros written in IgorPro (WaveMetrics, Lake Oswego, OR). Data were filtered at 5 kHz and sampled at 10 kHz. Responses were elicited with a stainless steel bipolar electrode (Frederick Haer, Bowdoinham, ME), positioned in the outer plexiform layer or the distal part of the inner nuclear layer. Because large stimulus currents often elicited longer-lasting, multiphasic responses, the stimulus strength was adjusted such that the AMPAR EPSC decayed in a relatively rapid, monotonic manner (Fig. 1).

Outside-out patches were obtained by slowly withdrawing the patch pipette after establishing a whole-cell recording. Because L-glutamate-evoked currents were very small in conventional outside-out patches, gentle suction was applied during withdrawal to obtain nucleated patches.

All experiments were performed at room temperature (21–23°C). Unless otherwise indicated, all data are expressed as means ± SD; p values indicate paired t tests, and p < 0.05 was considered significant.

**RESULTS**

**Electrically evoked synaptic responses in GLC neurons**

Whole-cell voltage clamp recordings were made from GLCs in acute slices of rat retina (Fig. 1A). Electrical stimuli (10–20 μA, 600 μsec) were delivered through a stainless steel bipolar stimulating electrode positioned in the outer plexiform layer (Fig. 1A). When the cell was voltage clamped at −80 mV, stimulation elicited inward currents that were reduced either by the Ca$^{2+}$ channel blocker Cd$^{2+}$ (20 μM; 87 ± 10% block; n = 7) (Fig. 1B), 6,7-dinitroquinoxaline-2,3-dione (DNQX, 10 μM; 96 ± 2% block; n = 6) (Fig. 1C) or 1-[4-amino phenyl]-4-methyl-7,8-methylene dioxy-5H-2,3-benzodiazepine (GYKI 52466, 25 μM; 94 ± 6% block; n = 4) (Fig. 1D), indicating that the responses reflected primarily synaptic activation of AMPARs.

In a subset of cells, the charge transfer (Q) during both evoked EPSCs and sEPSCs was measured to estimate the quantal content of the evoked response. The quantal content (Q$_{evoked}$/Q$_{sEPSC}$) was quite variable across cells (16 ± 13; n = 7). This approach requires that sEPSCs reflect the postsynaptic response to a single quantum, rather than multivesicular release, which has been demonstrated at some synapses (Tong and Jahr, 1994; Auger et al., 1998; Wadiche and Jahr, 2001). Q$_{sEPSC}$ was unaffected by changes in p$_f$ (see Fig. 6A,B), consistent with the sEPSCs being unquantal. However, it remains possible that multivesicular release occurs during an evoked response, which could lead to an underestimate of the quantal content with this method.

Electrically evoked EPSCs exhibited faster time courses than light-evoked EPSCs that were recorded in mammalian ganglion cells previously (Cohen, 2000), presumably because the bipolar cell depolarization elicited by electrical stimulation has a faster time course than that elicited by light (cf. Higgs and Lukasiewicz, 1999; Berntson and Taylor, 2000). Although the GCL in rat retina contains both ganglion cells and displaced amacrine cells (Perry, 1981), no systematic differences in EPSC characteristics distinguished the cell types. Consequently, the data from all cells have been pooled; the postsynaptic neurons are referred to as GLCs (Matsui et al., 1998). Generally, larger cells that were not imme-
and the events are referred to as sEPSCs.

The sodium channel blocker tetrodotoxin (TTX; 1 μM) had little effect on the EPSC at –8 mV. Similar results were observed in six cells. When magnesium was removed from the extracellular solution, the charge transferred during an EPSC (Q_{evoked}) clamped at +40 mV was significantly greater than at –80 mV, despite the reduced driving force (Q_{evoked} at +40 mV = 711 ± 656% of Q_{evoked} at –80 mV; n = 7; p = 0.003) (Fig. 3B). However, the average charge transferred during sEPSCs (Q_{spont}) recorded at +50 mV, when scaled to account for the difference in driving force, was not significantly different from that during the Q_{spont} recorded at –80 mV (Q_{spont} (+50, scaled) = 89 ± 27% of Q_{spont} at –80 mV; n = 5; p = 0.49) (Fig. 3D).

When magnesium was removed from the extracellular solution, an NMDAR component was detected in the evoked response at –80 mV (Fig. 3E), which led to a significant increase in Q_{evoked} (411 ± 253% of control; n = 6; p = 0.01) (Fig. 3F). This did not appear to result from any change in the probability of release, because no effects were observed in the

**Evoked responses exhibit AMPAR and NMDAR components**

Synaptic, glutamatergic excitation of GLCs in amphibian retinal slices is mediated by both AMPARs and NMDARs (Mittman et al., 1990; Diamond and Copenhagen, 1993; Lukasiewicz and Roeder, 1995; Lukasiewicz et al., 1997; Matsui et al., 1998). However, in rat ganglion cells grown in culture, evoked synaptic responses are mediated solely by AMPARs, even though the cells express functional NMDARs (Taschenberger et al., 1995). To determine whether this discrepancy is attributable to a difference in species or in preparation, we looked in rat retinal slices to see whether evoked EPSCs in GLCs exhibit an NMDAR component. NMDARs are mostly blocked at negative potentials by external magnesium ions, but this blockade is primarily relieved at positive potentials (Mayer et al., 1984; Nowak et al., 1984). Accordingly, when the postsynaptic membrane was clamped at positive potentials, the evoked EPSC decayed much more slowly (Fig. 2A), indicating the presence of an NMDAR component (Hestrin et al., 1990a; Mittman et al., 1990). The kinetic differences between the two components allowed them to be examined simultaneously: the early component of the EPSC exhibited a linear, ohmic conductance, typical of AMPARs, and the late component exhibited a J-shaped current–voltage relationship, indicative of NMDARs (Mayer et al., 1984; Nowak et al., 1984) (Fig. 2B).

The AMPAR and NMDAR components could also be distinguished pharmacologically. As shown in Figure 1, at –80 mV the response was abolished by AMPAR antagonists, including DNQX or 2,3-dioxo-6-nitro-1,2,3,4-tetrahydrobenzof[1]quinoline-7-sulfonamide (NBQX; 5 μM) (Fig. 2C). In contrast, the NMDAR antagonist (RS)-3-(2-carboxypiperazin-4-yl)-propyl-1-phosphonic acid (CPP; 5 μM) had little effect on the EPSC at –80 mV (Fig. 2D). At +40 mV, NBQX blocked a fast component of the EPSC (Fig. 2C), whereas CPP blocked the slow component (Fig. 2D).

**Spontaneous EPSCs exhibit only an AMPAR component**

At hippocampal and cerebellar synapses, evoked EPSCs are also mediated by NMDARs and AMPARs (Hestrin et al., 1990a; Silver et al., 1992). Moreover, miniature EPSCs also exhibit NMDAR and AMPAR components at these synapses, indicating that the two receptor types are colocalized in the postsynaptic membrane (Bekkers and Stevens, 1989; McBain and Dingledine, 1992; Silver et al., 1992). In contrast, sEPSCs in amphibian (Taylor et al., 1995; Matsui et al., 1998) and mammalian (Taschenberger et al., 1995; Tian et al., 1998) GLCs do not exhibit an NMDAR component, suggesting that NMDARs may be expressed either extrasynaptically or at synapses different from those at which AMPARs are expressed. We addressed this question in rat retinal slices by recording sEPSCs in GLCs (Fig. 3).

The sodium channel blocker tetrodotoxin (TTX; 1 μM) did not affect the frequency, amplitude, or waveform of spontaneous events recorded in GLCs (data not shown); therefore, the experiments described here were performed in the absence of TTX, and the events are referred to as sEPSCs.

As shown above, evoked EPSCs exhibited a prominent NMDAR component at positive potentials (Figs. 2, 3A). Because of the prolonged time course of the NMDAR conductance, the charge transferred during an EPSC (Q_{evoked}) clamped at +40 mV was significantly greater than at –80 mV, despite the reduced driving force (Q_{evoked} at +40 mV = 711 ± 656% of Q_{evoked} at –80 mV; n = 7; p = 0.003) (Fig. 3B). However, the average charge transferred during sEPSCs (Q_{spont}) recorded at +50 mV, when scaled to account for the difference in driving force, was not significantly different from that during the Q_{spont} recorded at –80 mV (Q_{spont} (+50, scaled) = 89 ± 27% of Q_{spont} at –80 mV; n = 5; p = 0.49) (Fig. 3D).
evoked EPSC at +40 mV or in the AMPAR component of the EPSC at −80 mV (Fig. 3E). At −80 mV, removing external magnesium did not affect $Q_{\text{spont}}$ (104 ± 5% of control; $n = 7$; $p = 0.22$) (Fig. 3G,H), and sEPSCs were abolished by 10 μM DNQX (Fig. 3G). The lack of an NMDAR component in the sEPSCs may reflect the depletion of endogenous glycine, a coagonist of the NMDAR (Johnson and Ascher, 1987), from the slice. To control for this possibility, sEPSCs were also recorded in the absence of magnesium and the presence of 10 μM d-serine, a nontransported NMDAR glycine site agonist, with similar results [$Q_{\text{spont}}$ (d-serine, 0 Mg$^{2+}$) = 115 ± 26% of control (0 Mg$^{2+}$); $n = 5$; $p = 0.27$] (data not shown). These results indicate that sEPSCs in rat GLCs are mediated solely by AMPARs, as has been reported for other species (Taylor et al., 1995; Matsui et al., 1998; Tian et al., 1998).

**NMDARs encounter a lower transmitter concentration during a synaptic event**

The experiments illustrated in Figure 3 suggest that NMDARs on rat GLCs may be located either extrasynaptically or in synapses separate from AMPARs. If NMDARs were located extrasynaptically, they would be likely to encounter a lower glutamate...
concentration during a synaptic event than AMPARs located within the synaptic cleft, closer to the site of glutamate release. To test this prediction, we examined the actions of low-affinity competitive antagonists on synaptic responses and on receptor-mediated currents in outside-out, nucleated patches excised from GLC somata. The efficacy of γ-D-glutamylglycine (γ-DGG; 500 μM), a low-affinity AMPAR antagonist (Watkins and Olverman, 1987), and l-AP-5 (200 μM), a low-affinity NMDAR antagonist (Watkins and Olverman, 1987), was calibrated by measuring their effects on patch currents elicited by brief (2 msec) applications of 1 mM L-glutamate (see Materials and Methods) (Fig. 4A,C). In one group of patches, AMPARs were pharmacologically isolated by including 5 μM CPP in all solutions, whereas in another group NMDARs were isolated by including 5 μM NBQX. The duration of glutamate application was monitored by measuring open-tip currents at the end of each experiment (Fig. 4A,C) and was similar for both groups.

γ-DGG (500 μM) reversibly reduced AMPAR responses in excised patches (peak amplitude in γ-DGG was 22 ± 10% of control; n = 6) (Fig. 4A,E). This degree of blockade was comparable to the antagonism of γ-DGG against AMPAR EPSCs (peak amplitude in γ-DGG was 21 ± 9% of control; n = 6; t = 0.83, unpaired t test between patch and EPSC data) (Fig. 4B,E), suggesting that AMPARs encountered comparable amounts of glutamate in the patch and synaptic responses. In contrast, the effect of l-AP-5 on NMDAR patch currents (peak amplitude in l-AP-5 was 67 ± 8% of control; n = 6) (Fig. 4C,F) was significantly weaker than its effect on NMDAR EPSCs (peak amplitude in l-AP-5 was 33 ± 14% of control; n = 6; p = 0.001, unpaired t test between patch and EPSC data) (Fig. 4D,F), suggesting that NMDARs encountered less glutamate during the synaptic responses than during the patch responses. Although the fragility of the patches pulled from GLCs precluded a direct comparison of the two antagonists in the same patch, these results indicate that AMPARs encounter more glutamate than NMDARs during a synaptic response.

**Blocking glutamate uptake preferentially enhances the NMDAR component of the EPSC**

The results presented thus far are consistent with a scenario in which NMDARs on GLC dendrites are located outside the synaptic cleft. Glutamate uptake plays a crucial role in clearing synaptically released glutamate at many synapses (Otis et al., 1996; Diamond and Jahr, 1997; Higgs and Lukasiewicz, 1999; Carter and Regehr, 2000). Despite the presence of glutamate transporters in neuronal membranes (Rothstein et al., 1994), including those of rat bipolar and ganglion cells (Rauen et al., 1996), most glutamate uptake in the inner retina appears to occur at glial (Müller cell) membranes (Rauen et al., 1998), beyond the immediate vicinity of the postsynaptic density. Thus, uptake may limit activation of receptors outside the synaptic cleft (or in neighboring, inactive clefts) more than it would affect receptor activation within an active synapse. Therefore, one would predict that the activation of extrasynaptic NMDARs would be limited more than that of synaptic AMPARs by glutamate uptake.

To test this prediction, we examined the effects of TBOA (Shimamoto et al., 1998) on evoked EPSCs. TBOA is a competitive, nonsubstrate antagonist of glutamate transporters that does not interact with NMDARs (Jabaudon et al., 1999). TBOA (10 μM) did not affect the charge transferred during AMPAR EPSCs (Q_{evoked} = 107 ± 12% of control; n = 5; p = 0.2) (Fig. 5A,C). In contrast, TBOA potentiated and prolonged NMDAR EPSCs (Q_{evoked} = 365 ± 145% of control; n = 5; p = 0.02) (Fig. 5B,C). Therefore, blocking transporters preferentially enhanced the NMDAR component of the EPSC, which is consistent with there being an extrasynaptic location for the NMDARs.

![Figure 4](image-url)
Changing release probability preferentially affects NMDAR activation

NMDAR activation in GLCs appears to require coincident release of multiple quanta. During an evoked response, glutamate released from multiple vesicles at different synapses, or from within the same synapse, may accumulate in the extrasynaptic space to levels sufficient to activate extrasynaptic NMDARs. NMDARs could even be activated by transmitter released at a synapse made on a different postsynaptic cell. One might predict from this model that changing the probability of release (\( p_r \)) would change the extrasynaptic accumulation of glutamate and affect the NMDAR EPSC, perhaps more than the AMPAR EPSC. This would stand in contrast to other central synapses, where changing \( p_r \) affects NMDAR and AMPAR EPSCs to a similar extent (Perkel and Nicoll, 1993; Tong and Jahr, 1994).
solution in the continued presence of TBOA a slow component emerged, significantly increasing $Q_{\text{spont}}$ (210 ± 66% of TBOA alone; $n = 5; p = 0.02$) (Fig. 7A, top left). This slow component was mediated by NMDARs because it was blocked by 5 μM CPP ($Q_{\text{spont}}$ reduced to 116 ± 21% of TBOA alone; $n = 5; p = 0.2$) (Fig. 7A, top left). Subtraction of the average sEPSC waveform in the presence of CPP from that in TBOA/0 Mg$^{2+}$ revealed a CPP-sensitive component with a slow rise and decay (Fig. 7A, bottom left).

Interestingly, the NMDAR component in the TBOA–0 Mg$^{2+}$ condition was most apparent in the largest sEPSCs. No significant effect of removing Mg$^{2+}$ was observed in the smallest 20% of events ($Q_{\text{spont}}$ = 108 ± 30% of TBOA alone; $n = 5; p = 0.6$) (Fig. 7A, center), whereas a marked effect was observed in the largest 20% of events ($Q_{\text{spont}}$ = 183 ± 63% of TBOA alone; $n = 5; p = 0.04$) (Fig. 7A, right). These results suggest that extrasynaptic NMDARs would be activated by glutamate released from a single vesicle were it not for high-affinity glutamate uptake, particularly during the quantal events that generate the largest postsynaptic responses.

**DISCUSSION**

The experiments presented here indicate that NMDARs on GLCs are activated only during evoked responses, when multiple release events occur. Glutamate released from a single quantum is not, by itself, sufficient to activate NMDARs, apparently because glutamate uptake limits transmitter access to the NMDARs. Thus, in contrast to most central synapses, in which glutamate transporters regulate the synaptic activation of NMDARs only moderately (Hestrin et al., 1990b; Sarantis et al., 1993; Asztely et al., 1997; cf. Overstreet et al., 1999), transporters in the inner retina appear to play a critical role in limiting NMDAR activation by synaptically released glutamate.

**Extrasynaptic NMDA receptors**

These results in rat retinas are consistent with previous work in amphibian GLCs that showed that sEPSCs are mediated solely by AMPARs, despite the presence of an NMDAR component in the evoked EPSC (Taylor et al., 1995; Matsui et al., 1998). Taylor et al. (1995) suggested that NMDARs may be segregated from AMPARs at different synapses; Matsui et al. (1998) proposed that NMDARs are located extrasynaptically. The data presented here are interpreted most easily in the context of the second scenario: NMDARs were shown to encounter less glutamate during a synaptic response (Fig. 4); reducing transport with TBOA or increasing $p_r$ preferentially enhanced the NMDAR component of the EPSC (Figs. 5, 6); and TBOA caused an NMDAR component to emerge in sEPSCs (Fig. 7). Taken together, these results suggest that NMDARs are located at some distance from the site of release.

However, most of our results can also be interpreted in terms of the segregated-receptor hypothesis (Taylor et al., 1995). This scenario is potentially consistent with immunohistochemical evidence from rat retina that NMDARs colocalize with PSD-95 at postsynaptic densities in the IPL (Fletcher et al., 2000), although the NMDAR-immunopositive dendrites were not identified in that study and could have originated from amacrine cells in the inner nuclear layer. If NMDARs were segregated at synapses into which less glutamate was released during a synaptic response, perhaps attributable to differences in release machinery (Choi et al., 2000), NMDAR sEPSCs could be difficult to detect, and low-affinity antagonists would block the evoked NMDAR EPSC.
to a relatively greater extent than the AMPAR EPSC (Fig. 4). Blocking transporters could encourage spillover between neighboring synapses, preferentially enhancing the NMDAR component of the evoked EPSC (Fig. 5) (Asztely et al., 1997). However, TBOA would cause an NMDAR component to emerge in the sEPSCs (Fig. 7) only if substantial spillover consistently occurred between synaptic contacts made on the same postsynaptic neuron. At “dyad” synapses in the IPL, bipolar cell synaptic terminals are apposed to two postsynaptic elements that only very rarely, if ever, arise from the same ganglion cell (Dowling, 1987). Even if they did, it seems unlikely, given the small dimensions of dyad synapses (~200 nm diameter) (Koulen et al., 1998; Fletcher et al., 2000), that even a high density of glutamate transporters could so sharply partition the transmitter concentration within the synapse. Given these considerations, the results presented here are most consistent with an extrasynaptic location of NMDARs on GLC dendrites.

In rat retinal ganglion cells grown in cell culture, neither sEPSCs nor EPSCs exhibit an NMDAR component, even though the cells exhibit functional NMDARs (Taschenberger et al., 1995). Whereas it is possible that culture conditions would, for some reason, favor AMPAR synapses over NMDAR synapses [although this does not appear to be the case in the hippocampus (Bekkers and Stevens, 1989; Gomperts et al., 1998)], it seems more likely that synthetically released glutamate, on reaching the perimeter of the cleft, would be diluted by the large extracellular volume of the culture media before reaching extrasynaptic NMDARs.

**Transporters limit glutamate receptor activation**

Decreasing glutamate transport with TBOA enhanced the NMDAR EPSC but did not affect the AMPAR EPSC. This is in contrast to previous reports in amphibian retinas that showed that the glutamate transport inhibitor 1-trans-pyrrolidine-2,4-dicarboxylic acid (PDC) prolonged AMPAR EPSCs evoked by electrical or light stimulation (Higgs and Lukasiewicz, 1999; Matsui et al., 1999). In both of these previous studies, blocking transporters enhanced primarily a slow component of the evoked EPSC that was not evident in the responses reported here. The discrepancy is likely attributable to differences in stimulus strength; when we used stronger stimuli, we also observed a slow component in the EPSC, similar to that reported in amphibians (Higgs and Lukasiewicz, 1999; Matsui et al., 1999), that appeared to be attributable to the spillover of glutamate onto AMPARs in other synapses. It was enhanced by TBOA and blocked to a greater extent than the fast component by γ-DGG (data not shown). We purposely limited stimulus intensity to examine the differences between the NMDAR and AMPAR components of the EPSC, although the effect of PDC on the light-evoked AMPAR EPSC (Higgs and Lukasiewicz, 1999; Matsui et al., 1999) strongly suggests that glutamate transporters limit AMPAR activation in a physiologically meaningful manner.

When glutamate uptake was reduced with TBOA, an NMDAR component emerged in sEPSCs, but only in larger events (Fig. 7). One possible explanation for this result is that larger sEPSCs may reflect activity at larger synapses that express more AMPARs within the synaptic cleft and NMDARs extrasynaptically. In contrast, smaller synapses would express fewer AMPARs and perhaps no extrasynaptic NMDARs. Alternatively, larger sEPSCs may reflect the release of more glutamate, which could activate a larger fraction of synaptic AMPARs and, with transporters inhibited, might succeed in activating extrasynaptic NMDARs. A wide range of transmitter concentrations would be achieved if larger sEPSCs reflected spontaneous multivesicular release. However, sEPSC amplitude was insensitive to changes in [Ca$^{2+}$]o (Fig. 6A,B), a manipulation that affects the incidence of multivesicular release (Tong and Jahr, 1994; Auger et al., 1998; Wadiche and Jahr, 2001), suggesting that sEPSCs reflect postsynaptic responses to single quanta. Such a range of transmitter concentrations, then, would have to arise from variations in vesicular transmitter content (Ferking et al., 1995; Liu et al., 1999). Additional experiments are required to distinguish between these presynaptic and postsynaptic possibilities.

**Possible physiological roles for extrasynaptic NMDARs**

Functional NMDARs are expressed by retinal ganglion cells in numerous species, but a specific role for NMDARs in ganglion cell synaptic processing remains unclear. Changing [Ca$^{2+}$]o from 2.5:1.3 to 3.8:0 caused little change in the EPSC amplitude (Fig. 3E), suggesting that $p_\text{r}$ at bipolar cell terminals may be maximal under control conditions (2.5 mM [Ca$^{2+}$]o, 1.3 mM [Mg$^{2+}$]o). Increasing [Ca$^{2+}$]o from 1.3:1.0 to 3:1.3 enhanced NMDAR EPSCs (indicating a modulation of $p_\text{r}$) but had relatively little effect on the AMPAR EPSC (Fig. 6). This may mean that $p_\text{r}$ in 1 mM [Ca$^{2+}$]o is nearly 1; if synaptic receptors were significantly occupied under these conditions, then increasing $p_\text{r}$ further would cause a subproportional change in receptor activation. Perhaps their extrasynaptic location allows NMDARs to avoid saturation and accurately reflect increases in $p_\text{r}$, a range over which synaptic receptors may be relatively insensitive. It is possible that NMDARs may not be activated under low-light conditions but may play a role in boosting the synaptic response to stronger light stimuli. Although NMDARs on salamander ganglion cells appear to mediate a similar fraction of the response to weak and strong light stimulation (Diamond and Copenhagen, 1995), recent work in mouse amacrine cells suggests that postsynaptic NMDAR activation during a light response may depend on bipolar cell terminal $p_\text{r}$, which is regulated by feedback inhibition from amacrine cells (Matsui et al., 2001).

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