Development/Plasticity/Repair

Exocytosis of Vesicular Zinc Reveals Persistent Depression of Neurotransmitter Release during Metabotropic Glutamate Receptor Long-Term Depression at the Hippocampal CA3–CA1 Synapse

Jing Qian and Jeffrey L. Noebels
Department of Neurology, Baylor College of Medicine, Houston, Texas 77030

Exocytosis can be directly measured in mammalian brain slices by fluorescence detection of vesicular zinc release. Detection of the low-level evoked zinc signal [Zn]t was first demonstrated at the zinc-rich hippocampal mossy fiber pathway and required the use of high-frequency presynaptic stimulation. Here, we show that release after individual action potentials can be reliably detected even at non-mossy fiber, zinc-poor synapses in the hippocampus, a major enhancement in the temporal resolution of the technique. Short-term facilitation of release properties of zinc-positive CA3–CA1 Schaffer collateral/commissural synapses in the stratum radiatum differ from those at mossy fibers but are similar to those measured for the EPSP [field EPSP (fEPSP)]. The N-type Ca2+ channel toxin α-conotoxin GVIA inhibited both the [Zn]t and fEPSP equally, and the modulation of neurotransmitter release by neuropeptide Y, baclofen, and adenosine as revealed by [Zn]t closely resembles that measured for the fEPSP. A long-standing controversy in hippocampal synaptic plasticity involves the site of long-term depression (LTD) at these synapses. Using zinc release as a direct marker for exocytotic events and a surrogate marker for glutamate release, we demonstrate that persistent depression of presynaptic release occurs in the late expression of DHPG ([S]-3,5-dihydroxyphenylglycine]-induced LTD at this synapse. The ability to examine release dynamics with zinc fluorescence detection will facilitate exploration of the molecular pharmacology and plasticity of exocytosis at many CNS synapses.

Key words: vesicular zinc release; FluoZin-3; plasticity; LTD; DHPG; hippocampus

Introduction
Metabotropic glutamate receptors (mGluRs) play a critical role in synaptic plasticity in the CNS (Fagni et al., 2000). In the hippocampal Schaffer collateral/commissural pathway, long-lasting depression of synaptic transmission [long-term depression (LTD)] can be induced by activating mGluRs at a low frequency with synaptically released glutamate (Bolshakov and Siegelbaum, 1994; Oliet et al., 1997) or with exogenously applied (S)-3,5-dihydroxyphenylglycine (DHPG), an agonist of mGluRs. Although the induction of mGluR-LTD is widely thought to be postsynaptic, a definitive site for LTD expression has not been confirmed. Evidence for internalization or redistribution of ionotropic glutamate receptors and a requirement for postsynaptic protein synthesis in cultured neurons after application of DHPG supports a postsynaptic mechanism underlying mGluR-LTD (Snyder et al., 2001; Xiao et al., 2001); however, more recent evidence, including altered paired-pulse facilitation (PPF), decreased miniature EPSC (mEPSC) frequency, and the absence of a change in postsynaptic AMPA receptor sensitivity by DHPG all favor presynaptic expression of DHPG-LTD (Fitzjohn et al., 2001; Faas et al., 2002; Rammes et al., 2003; Rouach and Nicoll, 2003; Nosyreva and Huber, 2005). Recently, a developmental switch in the mechanism of mGluR-LTD expression has been proposed, based on evidence that mGluR-LTD at neonatal synapses is mediated by presynaptic mechanisms, whereas mGluR-LTD at mature synapses may result from postsynaptic modifications (Nosyreva and Huber, 2005). Although DHPG is widely used to study mGluR-LTD and DHPG-LTD shares many features with synaptically induced mGluR-LTD, there are some unusual properties of DHPG-LTD. The DHPG-LTD expression requires sustained activation of mGluR (Watabe et al., 2002; Rouach and Nicoll, 2003; Huang and Hsu, 2006) and the induction of DHPG-LTD is independent of postsynaptic Ca2+ entry (Fitzjohn et al., 2001).

The ability to directly examine presynaptic neurotransmitter release rather than inferring changes in evoked quantal output by analysis of postsynaptic responses can provide critical information in localizing the site of plasticity at synapses. We recently introduced such a method based on fluorescence detection of vesicular zinc by an extracellular zinc indicator dye (Qian and Noebels, 2005). Zinc has been shown to coexist with glutamate within synaptic vesicles in glutamatergic synapses in the many parts of brain including the hippocampus (Frederickson et al.,...
2000). Fluorescence detection of exocytosis by monitoring evoked vesicular zinc signals provides a novel way to assess neurotransmitter release at the zinc-containing hippocampal mossy fiber synapse (Qian and Noebels, 2005).

In the present study, we adapted the techniques developed for the mossy fiber synapse to measure vesicular zinc release evoked by individual action potentials at CA3–CA1 synapses within the hippocampal stratum radiatum. Zinc histochemistry detects a low level of vesicular zinc-containing terminals within this region. Recent analysis reveals that these synapses can be divided into two populations, those containing vesicular zinc and those without (Sindreu et al., 2003). The release properties of these populations have never been selectively characterized. We now demonstrate that fluorescence detection of zinc release can selectively resolve neurotransmitter release dynamics in the Schaffer collateral/commissural pathway, and apply this direct measure to examine DHPG-LTD at CA3–CA1 synapses. Our experimental results demonstrate a persistent depression of neurotransmitter release during DHPG-LTD at the zinc-containing mouse hippocampal CA3–CA1 synapse.

Materials and Methods

Preparation of brain slices. All procedures were performed in accordance with the guidelines of the National Institutes of Health, as approved by the Animal Care and Use Committee of Baylor College of Medicine. Transverse brain slices (200–250 μm thickness: 200 μm, optical recording; 250 μm, extracellular recording of field EPSP (fEPSP)) were prepared from hippocampi of wild-type (C57BL/6) and Znt3+/− (C57BL/6 × 129/SvEv) mice. For the adult mouse group, the animal age ranges between 6 and 10 weeks. For the young mouse group in the LTD experiments, the animal age was between 15 and 18 d. Znt3+/− mutant mice were generously provided by R. D. Palmer (University of Washington, Seattle, WA). Brain slices were incubated in artificial CSF (ACSF) at 32°C for 1 h and then transferred into a submerged recording chamber mounted on an inverted microscope (Axiovert 100; Zeiss, Oberkochen, Germany). The ACSF contained the following (in mM): 119 NaCl, 2.5 KCl, 2.5 CaCl2, 26 NaHCO3, 2 MgSO4, 1 NaH2PO4, 11 n-glucose, 0.2 Ca-EDTA gassed with 95% O2/5% CO2 to maintain a constant pH of 7.4. The temperature of the recording chamber was controlled at 30°C.

Zinc fluorescence detection in stratum radiatum of the CA1 region. A bipolar tungsten electrode was positioned in the stratum radiatum to stimulate the CA3–CA1 synapse. The stimulation pulse intensity was adjusted to 20 μA/0.1 ms to evoke a submaximal response for optical recording and to 10 μA/0.1 ms for extracellular recording of the fEPSP. A train of five pulses at 5 Hz was used to evoke the transient zinc fluorescence signal (ΔF or [Zn]), and the fluorescence response of FluoZin-3 after correction for photobleaching. Be-

Figure 1 is a schematic diagram detailing the measurement of extracellular zinc accumulation ([Zn]o) evoked by stimulating the CA3–CA1 synapse in the stratum radiatum of the hippocampal CA1 region of mouse brain slices. The top inset (Fig. 1Aa) shows the fluorescence response of the high affinity zinc indicator FluoZin-3 ([Zn]o) evoked by a stimulation protocol consisting of five pulses at 5 Hz. Over the sampling period of 2.5 s, ~1% of the signal diminished because of photobleaching of the zinc indicator. To obtain a more accurate time course of the [Zn], decay, dye bleaching was corrected by linear extrapolation of the baseline before stimulation as shown by the dashed line. The bottom inset (Fig. 1Ab) shows sample traces of autofluorescence, autofluorescence in the presence of CNQX/d-APV, and the fluorescence response of FluoZin-3 after correction for photobleaching. Because there was a significant amount of postsynaptic activity-dependent change of autofluorescence in the CA1 region, the glutamate receptor antagonists CNQX (10 μM) and d-APV (25 μM) were routinely added to the perfusate to eliminate the interference of autofluorescence in all zinc measurements. Figure 1B summarizes the experimental results from wild-type (+/+) and vesicular zinc transporter Znt3-deficient mutant mice (Znt3−/−). As demonstrated here, CNQX and d-APV abolished the autofluorescence transient but not the [Zn]. The average peak amplitude of the [Zn] was 2.1 ± 0.5% (ΔF/F) in wild-type mice (n = 24). However, no [Zn] was observed in slices from Znt3−/− mice (n = 5). These data confirm that the [Zn] is a result of the interaction between vesicular zinc and FluoZin-3 after action potential-induced exocytosis of synaptic vesicles. The absence of an evoked FluoZin-3 signal in Znt3−/− mice also excludes the possibility of irrelevant artifacts such as light scattering contributing to the FluoZin-3 signal.

Zinc release and glutamate release share the same short-term facilitation and Ca2+ channel reliance at the CA3–CA1 synapse

Vesicular zinc coreleased with glutamate has been demonstrated to serve as a surrogate marker for presynaptic neurotransmitter release at the intensely zinc-rich hippocampal mossy fiber synapse (Qian and Noebels, 2005). In contrast to the mossy fiber synapse, where zinc coexists with glutamate in synaptic vesicles
within every mossy fiber bouton, only ~45% of the boutons making axospinous glutamatergic synaptic contacts in the stratum radiatum of the CA1 region are zinc containing (Sindreu et al., 2003). We reasoned, however, that if zinc-positive synapses possess similar release properties as zinc-negative synapses in this pathway (see below), then \([Zn]_t\) measured at the zinc-positive synapse and compared it with fEPSPs obtained from the CA3–CA1 synapses. Therefore, in the course of the fluorescence intensity (\(\Delta F\)) and the peak amplitude of the transient fluorescence evoked by individual action potentials at the CA3–CA1 synapse.

Figure 2 summarizes the short-term facilitation of zinc release and glutamate release. Zinc release exhibits the same pattern of short-term facilitation as glutamate release. To quantify \(\Delta F\) for each action potential, 10 traces of the \([Zn]_t\) during steady state were first averaged and then smoothed with an average time window of 10 ms, and then the \(\Delta F\) corresponding to each action potential (AP) was measured. The smoothing algorithm does not alter the kinetics of the \([Zn]_t\). The N-type \(Ca^{2+}\) channel blocker \(\omega\text{-CgTX GVIA}\) equally inhibited the \([Zn]_t\) and fEPSP. The time course of the \([Zn]_t\) was quantified by measuring the total \(\Delta F\) evoked by a 5 Hz stimulation protocol. To be better compared with the time course of the \([Zn]_t\), a summation of all five fEPSPs within the stimulation train was quantified as the time course of fEPSP. Insets show sample traces of the \([Zn]_t\) (smoothed), first and fifth fEPSP before and after application of \(\omega\text{-CgTX GVIA}\). We did not examine the contribution of P/Q-type \(Ca^{2+}\) channels to the \([Zn]_t\), because blockade of P/Q-type \(Ca^{2+}\) channels, which

Figure 1. Vesicular zinc release evoked by individual action potentials at the CA3–CA1 synapse. A. Schematic diagram of a hippocampal slice showing stimulating and recording sites, DG, Dentate gyrus. Inset, Sample traces of autofluorescence and fluorescence signal of FluoZin-3 in response to 5 Hz stimulation. a, FluoZin-3 fluorescence before correction for photobleaching. b, Auto-fluorescence in the presence of the glutamate receptor antagonist CNQX/D-APV and FluoZin-3 fluorescence after correction of photobleaching. The time course of the photobleaching was corrected by linear extrapolation of the time course before stimulation. F, Fluorescence transient but not the FluoZin-3 signal. Right, Time course of the fluorescence intensity (\(\Delta F\)) and the peak amplitude of the transient fluorescence evoked by individual action potentials at the CA3–CA1 synapse.

Figure 2. Short-term facilitation and \(Ca^{2+}\) channel reliance of zinc release are similar to glutamate release. A. Left, Superimposed fEPSPs evoked by the 5 Hz stimulation protocol. Right, Summary data of short-term facilitation of zinc release and glutamate release. Zn release exhibits the same pattern of short-term facilitation as glutamate release. To quantify \(\Delta F\) for each action potential, 10 traces of the \([Zn]_t\) during steady state were first averaged and then smoothed with an average time window of 10 ms, and then the \(\Delta F\) corresponding to each action potential (AP) was measured. The smoothing algorithm does not alter the kinetics of the \([Zn]_t\). The N-type \(Ca^{2+}\) channel blocker \(\omega\text{-CgTX GVIA}\) equally inhibited the \([Zn]_t\) and fEPSP. The time course of the \([Zn]_t\) was quantified by measuring the total \(\Delta F\) evoked by a 5 Hz stimulation protocol. To be better compared with the time course of the \([Zn]_t\), a summation of all five fEPSPs within the stimulation train was quantified as the time course of fEPSP. Insets show sample traces of the \([Zn]_t\) (smoothed), first and fifth fEPSP before and after application of \(\omega\text{-CgTX GVIA}\).
contribute predominantly to the remainder of the presynaptic Ca\(^{2+}\) required for exocytosis at this synapse (Wheeler et al., 1994), should yield a similar result on the [Zn]\(_i\) and fEPSP (see Discussion).

**Zinc release and glutamate release are modulated identically at CA3–CA1 synapses**

Next, we characterized the modulation of zinc release and compared it with that measured for the fEPSP. Neurotransmitter release at CA3–CA1 synapses is regulated by presynaptic receptors for NPY, GABAB\(_\text{R}\), and adenosine acting via presynaptic Ca\(^{2+}\) channels (Qian et al., 1997; Wu and Saggu, 1997). Figure 3, A and B, shows the effect of NPY on the [Zn]\(_i\) and fEPSP. Application of 1 \(\mu\)M NPY led to a similar effect on both [Zn]\(_i\) and fEPSP. NPY inhibited the [Zn]\(_i\) by 78 \(\pm\) 1% and 79 \(\pm\) 1% \((n = 4)\) for the first and fifth action potentials, respectively. The average reduction of the corresponding fEPSPs was 80 \(\pm\) 4 and 78 \(\pm\) 1% \((n = 4)\). Similarly, activation of presynaptic GABAB\(_\text{R}\) and adenosine receptors resulted in the same amount of inhibition of both the [Zn]\(_i\) and fEPSP. As shown in Figure 3, C and D, baclofen (10 \(\mu\)M), the GABAB\(_\text{R}\) receptor agonist, reduced the [Zn]\(_i\) by 71 \(\pm\) 4 and 70 \(\pm\) 5% \((n = 4)\) for the first and fifth action potentials, respectively. The mean reduction of the corresponding fEPSPs was 76 \(\pm\) 4 and 68 \(\pm\) 4% \((n = 4)\). Not surprisingly, application of 5 \(\mu\)M adenosine also reversibly inhibited the [Zn]\(_i\) and fEPSPs to a similar extent (Fig. 3 E, F). The [Zn]\(_i\) during the peak effect of adenosine was 54 \(\pm\) 11 and 51 \(\pm\) 8% \((n = 4)\) of baseline for the first and fifth action potentials, respectively, and the corresponding fEPSPs were 50 \(\pm\) 7 and 53 \(\pm\) 8% \((n = 4)\) of baseline. These results, combined with similar short-term facilitation and Ca\(^{2+}\) channel reliance of zinc release and glutamate release confirm that the [Zn]\(_i\) is a valid surrogate marker for presynaptic neurotransmitter release at the CA3–CA1 synapse.

**Persistent depression of neurotransmitter release after activation of mGluR**

Using fluorescence detection of zinc release, we then explored the expression site of mGluR-LTD with the mGluR agonist DHPG at this synapse. To compare results with previous studies, a 10 min application of 100 \(\mu\)M NPY led to persistent depression of synaptic transmission at the CA3–CA1 synapse \((n = 5)\) as shown in Figure 4 A. The action of DHPG was completely blocked with the broad mGluR antagonist LY341495 (20 \(\mu\)M; \(n = 3\)), indicating the requirement for mGluR activation in DHPG-LTD. DHPG-LTD was independent of the protocols used for testing. As shown in Figure 4 B, a short application of DHPG in slices receiving the same stimulation protocol used in optical recordings induced an equivalent amount of LTD. n-APV (25 \(\mu\)M) was routinely applied in this set of LTD experiments to eliminate a possible contribution from NMDA receptor-mediated LTD. The average fEPSP 40 min after DHPG application was 67 \(\pm\) 11 and 72 \(\pm\) 9% of baseline for first and fifth fEPSP, respectively \((n = 10)\).

We then examined presynaptic neurotransmitter release during DHPG-LTD by measuring the [Zn]\(_i\) in response to DHPG. As shown in Figure 4 C, after an initial decrease, the [Zn]\(_i\) only partially recovered 40 min after washout of DHPG and exhibited a pattern of long-term depression. The first and fifth [Zn]\(_i\) were 70 \(\pm\) 7 and 69 \(\pm\) 7% of baseline, respectively \((n = 5)\). The persistent reduction of the [Zn]\(_i\) was not attributable to a possible deletion of vesicular zinc, because the [Zn]\(_i\) in control experiments remained at a stable level throughout the entire measurement. The first and fifth [Zn]\(_i\) were 97 \(\pm\) 2 and 95 \(\pm\) 3% \((n = 4)\) of baseline by the end of the experiments. Our results here clearly indicate a presynaptic expression of LTD after activation of mGluR at the CA3–CA1 synapse.

Recently, a developmental switch in the mechanism of
mGluR-LTD expression has been proposed, based on evidence that mGluR-LTD at neonatal synapses is mediated by presynaptic mechanisms, whereas mGluR-LTD at mature synapses may arise from postsynaptic modifications (Nosyreva and Huber, 2005). To learn whether presynaptic mechanisms of DHPG-LTD fade from postsynaptic modifications (Nosyreva and Huber, 2005). As shown in Figure 2, inhibition of the $\Delta Zn^{\text{t}}$ by the N-type Ca$^{2+}$ channels was identical to that measured electrophysiologically for the global fEPSP, implying a similar degree of persistent depression of synaptic transmission at CA3–CA1 synapses after activation of mGluR with DHPG. Our results are congruent with previous conclusions based on electrophysiological and vesicle dye loading methods of the presynaptic site for LTD expression at CA3–CA1 synapses, and reveal additional properties of this heterogeneous presynaptic terminal population.

Release properties of zinc-containing synapses at the CA3–CA1 synapses

In contrast to the mossy fiber pathway, where zinc coexists with glutamate in synaptic vesicles within every mossy fiber bouton, only about one-half of the CA3–CA1 synapses in the stratum radiatum are zinc/Znt3 positive (Sindreu et al., 2003). This raises several key questions. First, where are the parent neurons of zinc containing synapses in this region? The vesicular zinc transporter gene Znt3 is strongly expressed in CA3 pyramidal neurons (Palmiter et al., 1996), consistent with its presence in fiber terminals in the Schaffer collateral pathway. The origin of zinc-negative synapses, however, is less clear, and these terminals may represent a functional subgroup of neurons residing in CA3 or elsewhere (Sindreu et al., 2003). Second, do the zinc-positive synapses in this lamina possess different release properties and plasticity compared with zinc-negative synapses? Using only conventional electrophysiological recording techniques, it is impossible to separate the zinc-positive synaptic component of the fEPSP from the one that is zinc negative; however, fluorescence detection of zinc release provides a direct way to selectively assess exocytosis at the zinc-positive subpopulation. In this study, we found that neurotransmitter release at zinc-containing synapses relies on the same presynaptic Ca$^{2+}$ channel subtype ratio as zinc-negative synapses. As shown in Figure 2, inhibition of the $\Delta Zn^{\text{t}}$, by the N-type Ca$^{2+}$ blocker 6-CgTx GV1A was identical to that measured electrophysiologically for the global fEPSP, implying a similar degree of N-type channel contribution to neurotransmitter release at both zinc-positive and zinc-negative synapses. In addition, presynaptic N- and P/Q-type Ca$^{2+}$ channels are sensitive to various neuromodulators (Qian et al., 1997; Wu and Saggau, 1997). Consistent with its origin of zinc/Znt3 positive CA1 synapses at the CA3–CA1 synapses after activation of mGluR with DHPG. Our results are congruent with previous conclusions based on electrophysiological and vesicle dye loading methods of the presynaptic site for LTD expression at CA3–CA1 synapses, and reveal additional properties of this heterogeneous presynaptic terminal population.

Discussion

The ability to detect exocytosis at zinc-containing synapses provides a novel method to analyze plasticity of the release process in hippocampal slices. Refinements in the method allow exploration of release dynamics in non-mossy fiber terminals, and here we have used the zinc fluorescence detection technique to explore the site of persistent depression of synaptic transmission at CA3–CA1 synapses after activation of mGluR with DHPG. Our results are congruent with previous conclusions based on electrophysiological and vesicle dye loading methods of the presynaptic site for LTD expression at CA3–CA1 synapses, and reveal additional properties of this heterogeneous presynaptic terminal population.

**Figure 4.** Persistent depression of neurotransmitter release after mGluR activation. A, Time course of fEPSPs evoked by a pair of stimuli separated by an interval of 50 ms after a 10 min application of 100 $\mu$M DHPG. The brief application of DHPG led to a persistent depression of synaptic transmission ($n = 5$) that was blocked by the nonspecific mGluR antagonist LY341495 ($n = 3$). B, Time course of summation of fEPSPs evoked by 5 Hz stimulation protocol in response to the brief application of DHPG. C, Time course of the $\Delta Zn^{\text{t}}$, in response to DHPG in adult mouse hippocampus (6–10 weeks). After DHPG application, there was a persistent depression of the $\Delta Zn^{\text{t}}$. The first and fifth $\Delta Zn^{\text{t}}$ responses in the same age control group were $7 \pm 7$ and $69 \pm 7\%$ ($n = 5$) of baseline when measured 40 min after washout of DHPG. D, Time course of summation of fEPSPs evoked by 5 Hz stimulation protocol in response to the brief application of DHPG. The first and fifth $\Delta Zn^{\text{t}}$ results indicate that the same presynaptic mechanism controls level during the entire experimental period in the control group ($n = 4$). The persistent depression of the $\Delta Zn^{\text{t}}$, clearly indicates a presynaptic expression of DHPG-LTD. E, Time course of the $\Delta Zn^{\text{t}}$, in response to DHPG in developing brain (15–18 d). In slices from immature hippocampus, DHPG-induced initial depression of neurotransmitter release than in the adult group (C). However, the persistent depression of neurotransmitter release during DHPG-LTD was nearly identical in the two age groups. The first and fifth $\Delta Zn^{\text{t}}$ responses were $7 \pm 7$ and $66 \pm 3\%$ ($n = 5$) of baseline when measured 40 min after washout of DHPG.
NMDA receptors were blocked with CNQX and D-APV to eliminate the interference of autofluorescence in our zinc measurement, the generation of such a retrograde messenger must be independent of a postsynaptic Ca\(^{2+}\) triggered signal contributed via Ca\(^{2+}\) entry through either NMDA receptors or voltage-gated Ca\(^{2+}\) channels. In the Fitzjohn et al. (2001) study, DHPG was shown to induce LTD in the absence of extracellular Ca\(^{2+}\). This result implies that postsynaptic Ca\(^{2+}\) entry via NMDA or voltage-dependent calcium channels is not required for DHPG-LTD, consistent with our conclusion. Because postsynaptic Ca\(^{2+}\) entry is required for synthetically induced mGluR-LTD (Bolshakov and Siegelbaum, 1994; Oliet et al., 1997; Otani and Connor, 1998), our result raises the possibility that DHPG-LTD may have a different mechanism. Because of the significant autofluorescence, it is not currently possible to precisely monitor zinc release by action potentials without blocking postsynaptic activity, and we were therefore unable to verify the presynaptic expression of mGluR-LTD induced solely by synaptic activity. Furthermore, as shown in Figure 4, C and D, we found that the persistent depression of neurotransmitter release after application of DHPG was nearly identical in developing and mature mouse brain. Therefore, we were unable to confirm a developmental switch of the mGluR-LTD mechanism as reported in adolescent rats (Nosyreva and Huber, 2005). It is unclear whether the discrepancy between these two studies might be attributable to the species difference of experimental animals.

Vesicular zinc exocytosis at non-mossy fiber terminals
Zinc, as an essential divalent metal in the brain, exists in various forms throughout many subcellular compartments of the CNS (Frederickson et al., 2005). Previous studies of vesicular zinc exocytosis at synapses have naturally focused on the zinc-rich mossy fiber pathway in the hippocampus (Thompson et al., 2000; Li et al., 2001; Ueno et al., 2002; Kay, 2003; Qian and Noebels, 2005). Little attention has been paid to neuronal circuits with weak histochemical vesicular zinc staining, although the widespread presence of mRNA for Znt3, the gene underlying zinc transport into synaptic vesicles suggests that many glutamate synapses other than the mossy fiber synapse may also have zinc available for release (Palmiter et al., 1996). Because of differences in dye concentration and stimulus strength required to resolve vesicular zinc exocytosis in the CA1 region, the amplitudes of ΔF/ΔF are not directly comparable with those reported at mossy fiber synapses (Qian and Noebels, 2005). Our zinc signal neither reflects absolute levels of zinc, nor distinguishes between free zinc and the externalization of membrane-bound zinc in the synaptic cleft as discussed by Kay (2006). In summary, the enhanced ability to resolve exocytosis of vesicular zinc evoked by action potentials opens the exploration of presynaptic release dynamics at many other zinc-containing neuronal circuits in the CNS.

References


