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Genetic Analysis of Synaptotagmin C2 Domain Specificity in Regulating Spontaneous and Evoked Neurotransmitter Release

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Synaptic vesicle fusion mediates communication between neurons and is triggered by rapid influx of Ca^{2+} . The Ca^{2+} -triggering step for fusion is regulated by the synaptic vesicle transmembrane protein Synaptotagmin 1 (Syt1). Syt1 contains two cytoplasmic C2 domains, termed C2A and C2B, which coordinate Ca^{2+} binding. Although C2A and C2B share similar topology, binding of Ca^{2+} ions to the C2B domain has been suggested as the only critical trigger for evoked vesicle release. If and how C2A domain function is coordinated with C2B remain unclear. In this study, we generated a panel of Syt1 chimeric constructs in *Drosophila* to delineate the unique and shared functions of each C2 domain in regulation of synaptic vesicle fusion. Expression of Syt1 transgenes containing only individual C2 domains, or dual C2A–C2A or C2B–C2B chimeras, failed to restore Syt1 function in a *sytl*^{−/−} null mutant background, indicating both C2A and C2B are specifically required to support fast synchronous release. Mutations that disrupted Ca^{2+} binding to both C2 domains failed to rescue evoked release, but supported synaptic vesicle docking and endocytosis, indicating that these functions of Syt1 are Ca^{2+} -independent. The dual C2 domain Ca^{2+} -binding mutant also enhanced spontaneous fusion while dramatically increasing evoked release when coexpressed with native Syt1. Together, these data indicate that synaptic transmission can be regulated by Syt1 multimerization and that both C2 domains of Syt1 are uniquely required for modulating Ca^{2+} -independent spontaneous fusion and Ca^{2+} -dependent synchronous release.

Introduction

Neurotransmitter release at synapses occurs within milliseconds following Ca^{2+} influx through voltage-gated channels (Llinás et al., 1981; Neher and Zucker, 1993; Sabatini and Regehr, 1996; Sun and Wu, 2001; Wojcik and Brose, 2007). Although the full complement of Ca^{2+} binding proteins that coordinate synaptic vesicle fusion is unknown, the synaptic vesicle protein Synaptotagmin 1 (Syt1) has emerged as a key Ca^{2+} sensor that regulates fast synchronous release. Syt1 consists of a short intraluminal N-terminal region, a single transmembrane domain, and two cytoplasmic PKC-homologous repeats (C2 domains) that bind Ca^{2+} via negatively charged aspartate residues (Perin et al., 1990; Perin et al., 1991; Sutton et al., 1995; Ubach et al., 1998; Desai et al., 2000). *In vitro* studies have demonstrated Ca^{2+} -dependent interactions between Syt1 and plasma membrane phospholipids (Brose et al., 1992; Chapman and Jahn,

1994; Sutton et al., 1995; Fernandez et al., 2001), as well as the neuronal SNARE complex (Chapman et al., 1995; Zhang et al., 2002). The loss of Syt1 severely impairs Ca^{2+} -dependent synchronous vesicle release (Geppert et al., 1994; Geppert et al., 1997; Voets et al., 2001; Yoshihara and Littleton, 2002; Nishiki and Augustine, 2004b; Liu et al., 2009).

To determine how Syt1 regulates fusion, several studies have focused on the function of its two C2 domains, termed C2A and C2B. Despite the robust interaction between the C2A domain and membrane phospholipids (Chapman and Davis, 1998; Bai et al., 2000; Bai et al., 2002; Stevens and Sullivan, 2003; Paddock et al., 2008), neutralization of negatively charged aspartate residues (D to N) in C2A does not disrupt synchronous neurotransmitter release (Fernández-Chacón et al., 2002; Robinson et al., 2002; Stevens and Sullivan, 2003). In contrast, a D229E substitution in the *Drosophila* C2A domain reduced vesicle fusion (Striegel et al., 2012), whereas neutralization of D232 (D232N) in mice enhanced synaptic transmission (Fernández-Chacón et al., 2002; Stevens and Sullivan, 2003; Pang et al., 2006a). Together, these results suggest that the C2A domain may regulate exocytosis, but its specific function remains unclear. In contrast to C2A, Ca^{2+} binding to the C2B domain is required for synchronous release (Littleton et al., 1994; Littleton et al., 2001; Mackler et al., 2002; Nishiki and Augustine, 2004a; Nishiki and Augustine, 2004b; Shin et al., 2009; Yoshihara et al., 2010).

Although these studies emphasize the importance of Ca^{2+} -binding to C2B, there is no experimental evidence to indicate that

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C2B can function without C2A. Indeed, membrane penetration by Syt1 requires cooperativity between its tandem C2 domains (Bai et al., 2002; Herrick et al., 2006). Here, we tested the function of each C2 domain and identified cooperative actions of C2A and C2B that regulate release at *Drosophila* neuromuscular junctions (NMJs). We generated transgenic Syt1 constructs bearing Ca^{2+} -binding mutations in each C2 domain or that expressed proteins with chimeric C2 domain structure. Our results indicate that the essential function for the C2B domain in fusion requires C2A. In addition, similar interactions are required for regulation of spontaneous vesicle release, suggesting that Syt1 C2 domain cooperativity regulates multiple synaptic vesicle release pathways.

Materials and Methods

Drosophila stocks and genetics. *Drosophila melanogaster* were cultured on standard medium at 22°C. Female larvae and adult flies were used for analyses described below unless indicated. DNA constructs for *UAS-synaptotagmin 1 (syt1)^{C2B-D3/4N}* encoding Syt1^{D416N, D418N} (C2A-C2B*) were obtained from N.E. Reist (Colorado State University, Fort Collins, CO). DNA for *UAS-syt1^{C2A-D3/4N}* encoding Syt1^{D282N, D284N} (C2A*-C2B) was generated using the QuikChange multiSite-Directed Mutagenesis Kit (Stratagene) with the primer ctcgtgttggccatttcAacttcAatcgc. A similar strategy was used to generate *UAS-syt1^{C2A-D3/4N-C2B-D3/4N}* (C2A*-C2B*) encoding Syt1^{D282N, D284N, D416N, D418N}. Isolated C2 domain constructs with or without D3/4N mutations (C2A, C2A*, C2B, and C2B*) were generated by PCR reactions with the following primer sets: cgGAATTCatgccgcaaatgcaaatcgg (5'-EcoRI) and gCTCTAGAtatcttcaacgtgaccaggtc (3'-XbaI) for C2A and C2A*; cgGAATTCatgccgcaaatgcaaatcgg (5'-EcoRI), ccgCTCGAGctgttctctctgcacccctc (3'-XhoI for a cytoplasmic linker), gCGCTCGAGagcggtgaaggagggcgagc (5'-XhoI for C2B), and cgTCTAGAttacttcatgttcttcaggatc (3'-XbaI) for C2B and C2B*. In addition, constructs composed of two homologous C2 domains tagged with hemagglutinin (C2A-C2A and C2B-C2B) were generated by the *Drosophila* Gateway vector system with the following primer sets: cgGAATTCatgccgcaaatgcaaatcgg (5'-EcoRI), gaGATCTtttctctgcccctctcttc (3'-BglII for the first C2A), GAagatctCAGAGCGAGCAGAAgctggggc (5'-BglII for the second C2A), tccTCTAGActttctgtccgcccctctcttc (3'-XbaI for the second C2A), CACCATggcgcaaatgcaaatcgg (5'-Gateway), and tccTCTAGActttctctgtc (3'-Gateway) for C2A-C2A; cgGAATTCatgccgcaaatgcaaatcgg (5'-EcoRI), GAagatctCTTCATGTTCTTCAGGATCTCGTC (3'-BglII for the first C2B), GAagatctAGCGTTGAAGGAGAGGGCGGACAG (5'-BglII for the second C2B), tccTCTAGActtcatgttcttcaggatc (3'-XbaI for the second C2B), CACCATggcgcaaatgcaaatcgg (5'-Gateway), and tccTCTAGActtcatgttcttc (3'-Gateway) for C2B-C2B. Transgenic strains were generated using standard microinjection into *white*⁻ embryos performed by Duke University Model System Genomics (Durham, NC) and Genetics Services (Cambridge, MA). *UAS-syt1* transgenes were expressed using a *GAL4* driver under the control of the pan-neuronal *elav* promoter (Campos et al., 1987) in the *syt1* null (*syt1*^{-/-}) background. Null mutants lacking endogenous Syt1 were generated by crossing *syt1*^{N13}, an intragenic *syt1* deficiency (Littleton et al., 1994), with *syt1*^{AD4}, which truncates Syt1 before the transmembrane domain (DiAntonio and Schwarz, 1994).

Western blot analysis. Western blots were performed using standard laboratory procedures. Nitrocellulose membranes were probed with mouse anti-Discs-Large (Dlg) (4F3, 1:2000) and rabbit anti-Syt1 antibodies (1:200). The 4F3 antibody was obtained from the Developmental Studies Hybridoma Bank (University of Iowa, Iowa City, IA). Blocking was performed in a solution containing four parts PBS (4.3 mM Na_2HPO_4 , 1.5 mM KH_2PO_4 , 137 mM NaCl, pH 7.4) to one part Odyssey Blocking Buffer (LI-COR Biosciences). Antibody incubations were performed in a solution containing four parts PBST (1× PBS with 1% Tween 20) to one part Odyssey Blocking Buffer. The probes were detected using AlexaFluor-680-conjugated goat anti-rabbit IgG at a dilution of 1:5000 (Invitrogen) and IR Dye 800-conjugated goat anti-mouse IgG at a dilution of 1:5000 (Invitrogen). Visualization was done using the LI-COR Odyssey Imaging System (LI-COR Biosciences).

Immunohistochemistry. Third instar larvae were reared at 22°C and dissected in HL3.1 saline (70 mM NaCl, 5 mM KCl, 10 mM NaHCO_3 , 4 mM MgCl_2 , 5 mM trehalose, 115 mM sucrose, 5 mM HEPES, pH 7.2). Larvae were fixed for 20 min in HL3.1 containing 4% formaldehyde. Following washes, larvae were incubated with primary antibody overnight at 4°C, incubated with secondary antibodies for 2 h at room temperature, and mounted in 80% glycerol for imaging. The dilutions of primary antibodies were as follows: nc82 (1:50) and Syt1 (1:500). The mouse nc82 antibody against Bruchpilot, developed by Erich Buchner, was obtained from the Developmental Studies Hybridoma Bank (University of Iowa, Iowa City, IA). Secondary antibodies from Jackson ImmunoResearch Laboratories, used at a dilution of 1:250, were as follows: FITC-conjugated goat anti-rabbit, FITC-conjugated goat anti-mouse, and rhodamine red/FITC-conjugated goat anti-HRP. Visualization was performed with confocal microscopy (Axioplan 2; Carl Zeiss Microscopy) using PASCAL software (Carl Zeiss Microscopy).

Protein expression and protein binding assays. Syt1 was amplified by PCR and subcloned into pGEX-2T. Recombinant Syt1 fused with GST was expressed in *Escherichia coli* (BL21) and purified using glutathione-Sepharose beads (GE Healthcare). A second version of recombinant Syt1 fused with His₆ was generated by PCR and subcloning into pTrcHisA, expressed in *E. coli* (BL21), and purified using the ProBond purification system (Invitrogen). The concentration of purified proteins was determined by SDS-PAGE separation, Coomassie Blue staining, and comparison with BSA standards.

GFP-fused Syt1 fraction was made by freezing 500–1000 male flies in liquid nitrogen. Heads were obtained by sieving and then homogenized on ice in TS buffer supplied with protease inhibitors (cOmplete Mini Protease Inhibitor Cocktail Tablets, Roche Applied Sciences). The homogenate was subsequently solubilized with 0.5% Triton X-100 for 30 min at 4°C. The resulting supernatant (after removing cell debris by centrifugation at 15,000 × g for 20 min) was used in binding assays. Protein concentration of head supernatants was measured with Pierce BCA reagents (Thermo Fisher Scientific). GST-fused Syt1 immobilized on glutathione-Sepharose beads was incubated with head supernatants or His₆-tagged Syt1 at 4°C for 2 h in TS buffer containing either 2 mM EGTA or 1 (or 10) mM Ca^{2+} . Equal fractions were then subjected to SDS-PAGE and immunoblotting. Polyclonal antiserum against GFP (1:2000; Invitrogen) and Syt1 (1:200) was used for detection of protein binding.

Electron microscopy. Wandering third instar larvae expressing Syt1 transgenic constructs in the wild-type or *syt1*^{-/-} mutant backgrounds were dissected, fixed, and processed as previously described (Rodal et al., 2011), and subjected to a poststaining step with uranyl acetate (2%; Electron Microscopy Sciences) for electron microscopy. Thin sections (50 nm) were imaged at a 49,000 × magnification at 80 kV on an electron microscope (Tecnai G2 Spirit, FEI) equipped with a charge-coupled device camera (Advanced Microscopy Techniques). Type 1b boutons with small clear vesicles were selected for measurements for the number and diameter of synaptic vesicles.

Electrophysiology. Preparation of wandering third instar larvae and intracellular recordings of excitatory junctional potentials (EJPs) and miniature EJPs (mEJPs) were performed as described previously (Lee et al., 2008) at the indicated extracellular calcium concentrations in HL3.1 saline at muscle fiber 6 of segments A3–A5. All recordings were done using an Axoclamp-2B amplifier (Molecular Devices) and digitized with Digidata 1330 (Molecular Devices). The data were acquired using Axoscope (version 9.0, Molecular Devices) and analyzed using CLAMPFIT (version 9.0, Molecular Devices) and Origin software (version 8.5, Origin Lab). The correction procedure for nonlinear summation of synaptic potential (Martin, 1955) was applied for EJP amplitude comparison when indicated. mEJPs, detected at resting membrane potentials more negative than −60 mV, were analyzed for their frequency and amplitude (MiniAnalysis, Synaptosoft).

Results

Generation of Syt1 transgenic constructs defective in Ca^{2+} binding

It has been well established that five conserved aspartate residues (termed D1–D5) form a Ca^{2+} -binding pocket within each C2 domain (Fig. 1A, asterisks; Shao et al., 1996; Sutton et al., 1999; Fernandez et al., 2001). Neutralization of these residues leads to a

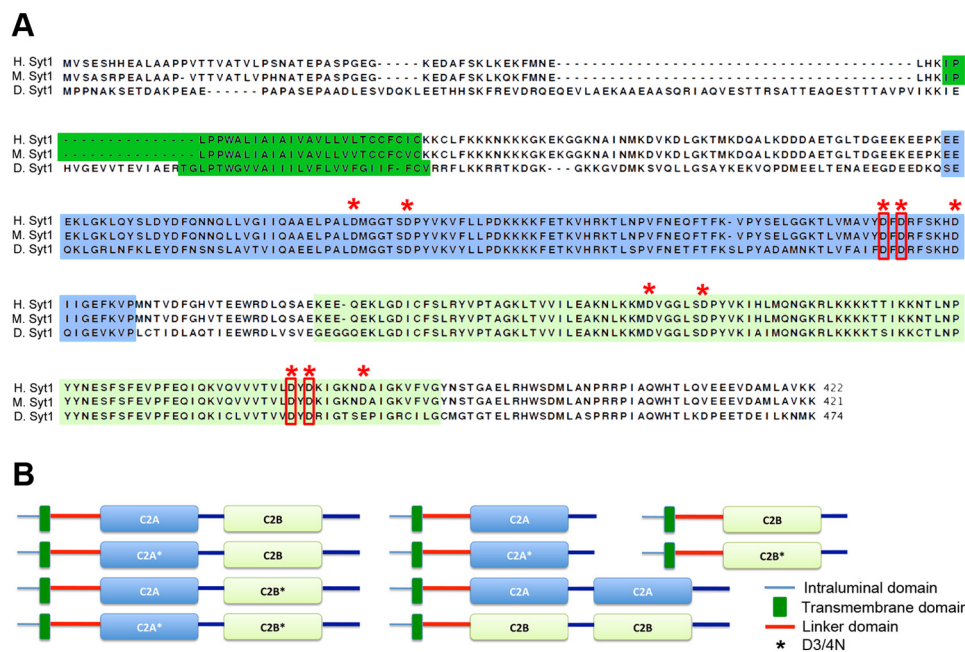


Figure 1. Syt1 transgenic constructs and sequence similarity. **A**, Amino acid sequences of human, mouse, and *Drosophila* Syt1 are compared. A single transmembrane domain and two C2 domains, C2A and C2B, are indicated as dark green, blue, and light green blocks, respectively. The five aspartate residues (D) involved in binding Ca^{2+} ions in each C2 domain are indicated with red asterisks. The third and fourth of these five aspartate residues were mutated to asparagines (N) to disrupt Ca^{2+} -binding ability of each C2 domain (red boxes). **B**, The design of transgenic UAS-Syt1 constructs used for the analysis is shown. Syt1 consists of a short intraluminal region (blue line), a single transmembrane domain (dark green box), a cytoplasmic linker (red), two C2 domains (C2A, blue; C2B, green) with a short linker between them, and a C-terminal tail. *C2 domains containing mutations (described in **A**) (red boxes).

loss of Ca^{2+} binding and subsequent disruption in Syt1 function (Littleton et al., 2001; Fernández-Chacón et al., 2002; Mackler et al., 2002; Stevens and Sullivan, 2003). Although the C2B domain is critical for activating synchronous vesicle release (Littleton et al., 1994; Littleton et al., 2001; Mackler et al., 2002; Nishiki and Augustine, 2004a; Nishiki and Augustine, 2004b; Shin et al., 2009; Yoshihara et al., 2010), there are no experimental data to indicate the C2B domain can function as the sole Ca^{2+} -binding module independent of C2A. To delineate isolated functions conveyed by each C2 domain versus cooperative interactions requiring both C2 domains, we generated transgenic Syt1 constructs bearing mutations in the key aspartate residues in each C2 domain in *Drosophila* (Fig. 1B). The five conserved aspartate residues involved in binding Ca^{2+} are highly conserved across the phyla (Fig. 1A, asterisks). Similar to prior approaches to eliminate Ca^{2+} binding (Littleton et al., 2001; Fernández-Chacón et al., 2002; Mackler et al., 2002; Stevens and Sullivan, 2003), two of the five residues in each C2 domain were mutated to asparagine (Fig. 1A, red boxes; D282/284N and in C2A and D416/418N in C2B). This allowed generation of *syt1* mutants disrupting Ca^{2+} binding to C2A, C2B, or both C2 domains. These mutations (indicated hereafter as C2A*, C2B*, or C2A*-C2B*) were introduced into transgenic constructs (Fig. 1B, left) and expressed in the *syt1* null mutant (*syt1*^{-/-}) background using the GAL4-UAS system (Brand and Perrimon, 1993).

To begin analyzing these transgenic animals, we first examined expression of the mutant protein in the *syt1*^{-/-} background to determine whether they localized properly at NMJs, similar to endogenous Syt1. Immunoreactivity against Syt1 indicated that the mutant proteins were distributed at NMJs in a pattern similar to that of endogenous Syt1 (Fig. 2A), suggesting that Ca^{2+} binding to the C2 domains is not required for synaptic targeting of Syt1. To avoid complications in functional analysis that might originate from differential expression of each construct, trans-

genic lines with similar expression levels by Western analysis (Fig. 2B) were chosen for comparison in subsequent experiments.

Essential role of Ca^{2+} binding to the Syt1 C2B domain for synchronous neurotransmitter release

To functionally characterize the role of Ca^{2+} binding to each C2 domain in neurotransmitter release, we measured nerve-evoked excitatory junction potentials (eEJPs) at third instar NMJs in the presence of low (0.2 mM) and high (1.0 mM) external $[\text{Ca}^{2+}]$ (Fig. 3A, upper and lower traces, respectively). We found that synchronous release at lower Ca^{2+} levels (0.2 mM) was restored more effectively in null mutants rescued with Syt1 lacking C2A Ca^{2+} binding (C2A*-C2B) than those with full-length wild-type Syt1 (C2A-C2B) (Fig. 3B, left, $p < 0.01$), similar to observations made at *Drosophila* embryonic NMJs (Yoshihara et al., 2010). Despite differences in the degree of rescue at a lower Ca^{2+} level, both C2A-C2B and C2A*-C2B Syt1 constructs restored synchronous release to near wild-type levels at more physiological Ca^{2+} levels (Fig. 3A, bottom traces in each pair; Fig. 3B, right). However, as previously observed (Littleton et al., 2001; Fernández-Chacón et al., 2002; Mackler et al., 2002; Stevens and Sullivan, 2003), Syt1 with defective C2B Ca^{2+} binding (C2A-C2B*) failed to rescue synchronous release, regardless of the presence of a wild-type or mutated C2A domain (C2A-C2B* and C2A*-C2B*) or external Ca^{2+} concentration (0.2 and 1.0 mM) (Fig. 3A,B; $p < 0.01$ at 0.2 mM $[\text{Ca}^{2+}]$ and $p < 0.001$ at 1.0 mM). As a result, the ratio of synaptic failure, defined as no detectable eEJP following nerve stimulation, was comparable between *syt1*^{-/-} and C2A-C2B* or C2A*-C2B* rescued lines (Fig. 3C; $p < 0.05$ or $p < 0.001$), whereas failures were absent in C2A*-C2B or wild-type Syt1 rescued larvae at a higher Ca^{2+} level (1.0 mM) (Fig. 3C). Together, these data support the model that Ca^{2+} binding to C2B, but not C2A, is essential for fast synchronous neurotransmitter release.

Cooperativity between Syt1 C2 domains in synchronous neurotransmitter release

Rescue data indicate that the C2B domain is the major Ca^{2+} -binding module involved in regulation of synchronous fusion. However, these results do not exclude a contribution from the C2A domain in C2B-driven Ca^{2+} -dependent exocytosis. Given that mutations in the C2B domain (C2A-C2B*) led to a complete failure to rescue, regardless of the presence of a wild-type or Ca^{2+} -binding defective C2A domain (Fig. 3, C2A-C2B* and C2A*-C2B*), it is possible that C2B functions as a major Ca^{2+} -binding module without a C2A domain present. To assay the function of each C2 domain separately, we created transgenic constructs that deleted the entire C2A or C2B domain. In addition, we created chimeric transgenic lines that expressed a dual C2A-C2A or C2B-C2B Syt1 protein where the corresponding endogenous C2 domain was replaced to generate a homologous C2 domain architecture (Fig. 1B, right). A construct lacking the entire C2B domain (C2A only) was properly localized at NMJs, albeit at a lower level, when detected with anti-Syt1 antibodies. However, the counterpart lacking the C2A domain (C2B only) was not recognized by immunocytochemistry (data not shown), possibly because of a lack of antigenicity to C2B epitopes with our anti-Syt1 antisera. To circumvent this problem, the dual C2A-C2A and C2B-C2B constructs were tagged with hemagglutinin. These constructs were detected at NMJs at similar patterns to endogenous Syt1 when visualized with an anti-hemagglutinin antibody (data not shown).

We first examined whether a functional C2A domain in the absence of C2B could restore release in *syt1*^{-/-}. Syt1 transgenes expressing C2A in the absence of C2B failed to rescue fast synchronous release, even in high extracellular $[\text{Ca}^{2+}]$ (1.0 mM, Fig. 4A, B, $p < 0.001$). A similar failure to rescue was also observed in the case of Syt1 transgenes expressing C2A that lacks Ca^{2+} -binding ability (C2A*, data not shown). To determine whether the inability of C2A to restore release was due to the lack of a C2B domain or simply due to the requirement for the dual C2 domain structure found in all synaptotagmin isoforms (Sudhof and Rizo, 1996), we examined transgenic strains that replaced the C2B domain with a second C2A to retain the characteristic twin C2 domain structure (Figs. 1B, right, 4A, C2A-C2A). We observed that C2A-C2A transgenic rescue lines had no ability to restore release, as the amplitude of eEPs remained similar to that in *syt1*^{-/-} and far below control levels (Fig. 4B, $p < 0.001$). These data indicate that the C2A domain alone, or as a dual C2A-C2A module, cannot support fast synchronous release.

We next assayed whether C2B alone was sufficient to support synchronous release when expressed in *syt1*^{-/-} mutants. Constructs expressing a Syt1 construct lacking the C2A domain were

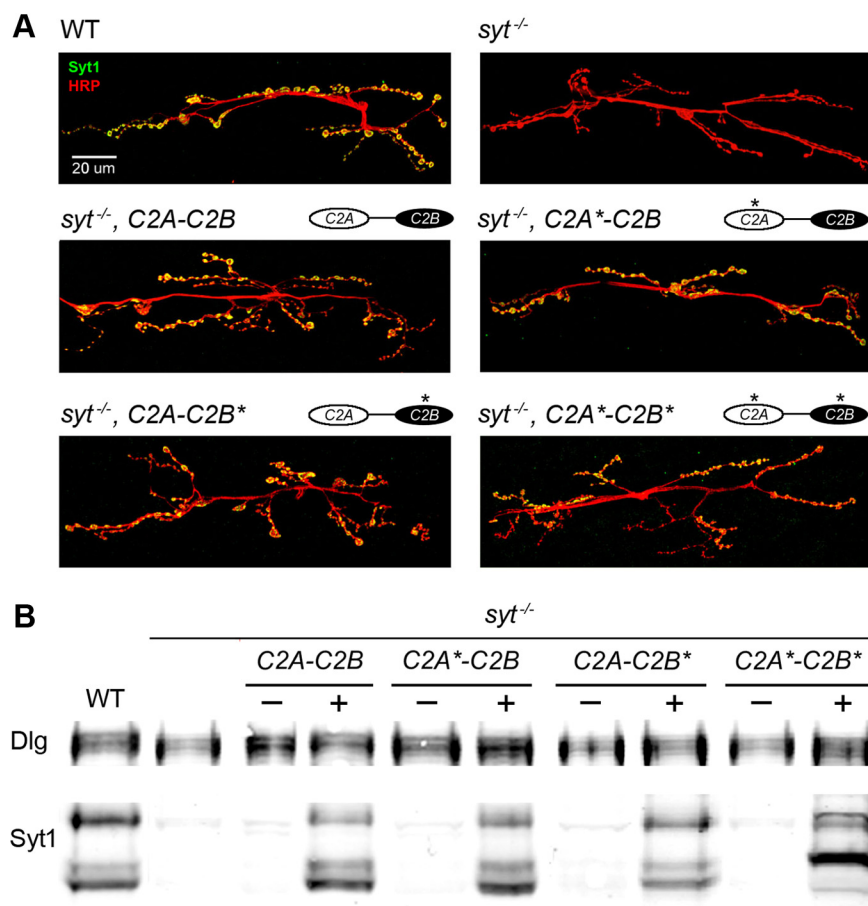


Figure 2. Expression of Syt1 transgenic constructs. **A**, Presynaptic distribution of Syt1 at the third instar larval NMJ is visualized by Syt1 immunoreactivity (green) in wild-type, *syt1*^{-/-} (null), and *syt1*^{-/-} animals rescued with the indicated transgenic constructs. The presynaptic nerve terminal is visualized with anti-HRP immunoreactivity (red) in a collapsed Z-stack confocal image stack. Scale bar, 20 μm . **B**, Western blot analysis is shown for the expression level of Syt1 in *syt1*^{-/-} mutants rescued with the indicated transgenic constructs compared with endogenous Syt1 levels in controls (WT-left lane). The — lanes are extracts from animals lacking the neuronal GAL4 driver, while the + lanes are from extracts of animals containing the GAL4 driver. The levels of the Discs-large (Dlg) protein are compared as loading controls. Note that the C2A*-C2B* construct results in a change in the size of the protease-sensitive breakdown product previously described (Littleton et al., 1993).

not capable of restoring synchronous release, regardless of the Ca^{2+} -binding ability of C2B (Fig. 4A, B, $p < 0.001$; data not shown for C2B*). We next examined whether a transgenic protein expressing dual C2B domains (C2B-C2B) was able to function in vesicle fusion. Synchronous release in *syt1*^{-/-} larvae expressing this construct was similar to that observed in *syt1*^{-/-} alone (Fig. 4A), indicating that C2B cannot function in the absence of C2A (Fig. 4B, $p < 0.001$). In summary, these data demonstrate that Ca^{2+} -dependent exocytosis mediated by the C2B domain requires the presence of C2A, indicating that the activity of both C2 domains is specifically required to regulate synchronous synaptic vesicle fusion.

C2A and C2B function cooperatively to regulate spontaneous neurotransmitter release

In addition to evoked neurotransmitter release, Syt1 has been shown to regulate spontaneous fusion, as evidenced by the elevated rates of spontaneous release in *syt1*^{-/-} null mutants (DiAntonio and Schwarz, 1994; Geppert et al., 1994; Littleton et al., 1994; Mackler et al., 2002; Yoshihara and Littleton, 2002; Pang et al., 2006b; Xu et al., 2009). However, the role of the C2A and C2B domains in regulation of spontaneous fusion has just begun to be explored (Xu et al., 2009) and remains poorly characterized. We

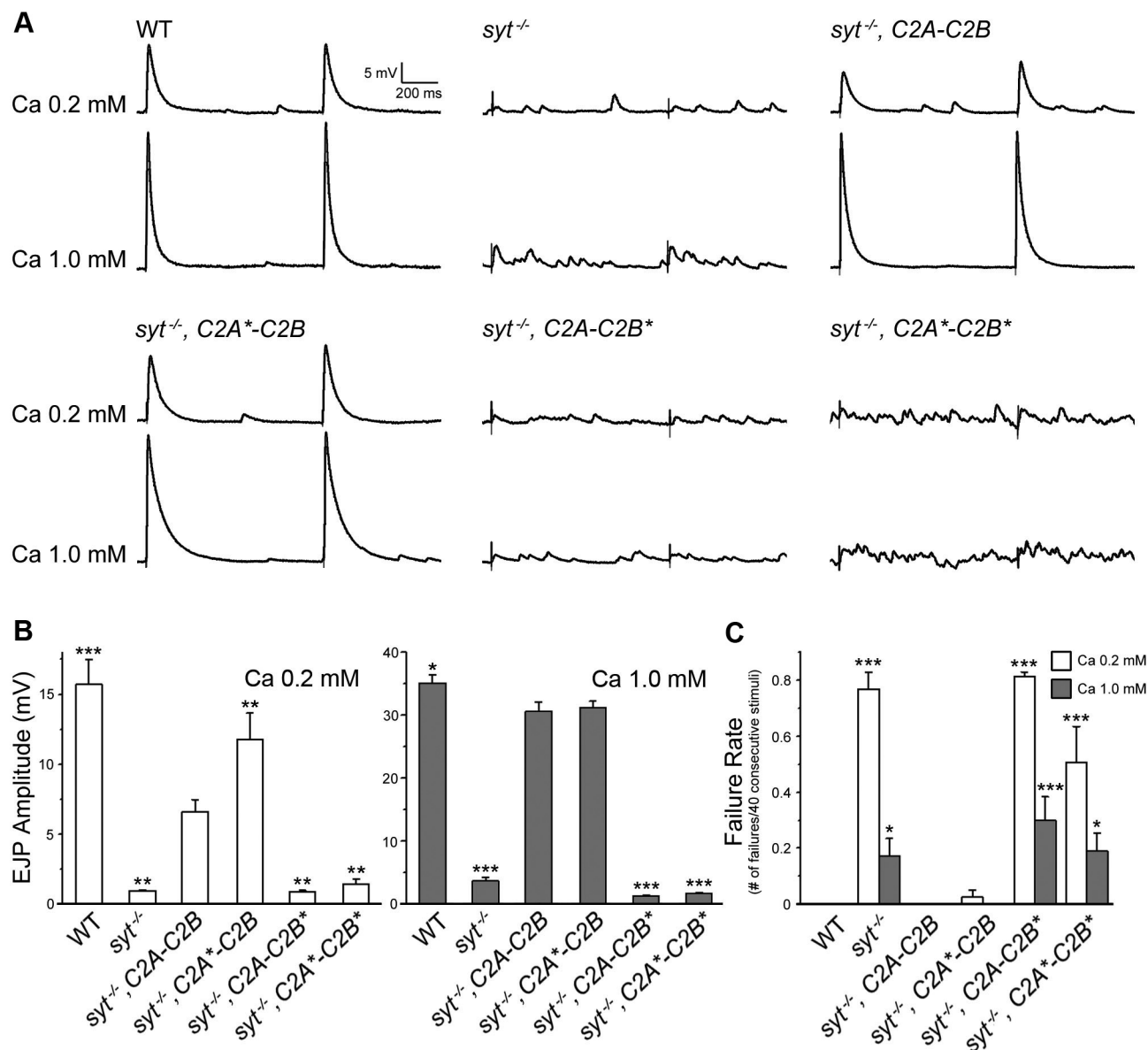


Figure 3. Effects of Ca^{2+} -binding mutations in each Syt1 C2 domain on evoked synchronous release. **A**, Representative traces of two consecutive eEPJ responses are shown for wild-type, *syt1*^{-/-} (null), and *syt1*^{-/-} rescued with the indicated transgenic constructs. Example traces are shown for recordings at low (0.2 mM, top) and high (1.0 mM, bottom) $[\text{Ca}^{2+}]$ in HL3.1 saline. Calibration: 5 mV, 200 ms. **B**, The mean amplitude of eEPJ responses are summarized for each genotype indicated. **C**, Failure rates, calculated by counting trials with no detectable eEPJ for 40 consecutive stimuli, are shown for each genotype. **B**, **C**, Data are mean \pm SEM. *** $p < 0.001$, ** $p < 0.01$, and * $p < 0.05$, one-way ANOVA with multiple comparisons using the Fisher's LSD test between *syt1*^{-/-} rescued with the C2A-C2B and the indicated genotypes. Number of NMJs examined (0.2 and 1.0 mM $[\text{Ca}^{2+}]_o$): WT, 7 and 7; *syt1*^{-/-}, 6 and 16; *syt1*^{-/-}, C2A-C2B, 10 and 11; *syt1*^{-/-}, C2A*-C2B, 9 and 7; *syt1*^{-/-}, C2A-C2B*, 6 and 10; and *syt1*^{-/-}, C2A*-C2B*, 6 and 12.

examined whether C2A and C2B regulate spontaneous release by measuring mEPJ frequency and amplitude in *syt1*^{-/-} larvae rescued with control and mutated transgenic Syt1 constructs (Fig. 5). Consistent with previous reports, *syt1*^{-/-} mutants displayed a slight but significant increase in the frequency of spontaneous release (Fig. 5A, B, bottom, $p < 0.05$). Expression of the wild-type Syt1 transgene (C2A-C2B) in *syt1*^{-/-} larvae restored spontaneous release to control levels (Fig. 5A, B, bottom). Syt1 transgenes containing neutralization of two aspartate residues (D3/4N) in C2A (C2A*-C2B) appeared to be more effective in clamping spontaneous release compared with those rescued with wild-type Syt1, although the effect did not reach statistical significance (Fig. 5B, bottom, $p = 0.32$). In contrast to C2A mutations, mEPJ frequency appeared elevated in larvae expressing Ca^{2+} -binding mutations in C2B (C2A-C2B*) (Fig. 5B, bottom, $p = 0.14$), com-

parable to *syt1*^{-/-} (Fig. 5A, B, bottom), suggesting that spontaneous release may be differentially regulated by Ca^{2+} -binding to the C2A and C2B domains.

To determine whether the ability of C2B to regulate spontaneous fusion requires wild-type C2A, we assayed Syt1 transgenes expressing Ca^{2+} binding mutants in both C2 domains (C2A*-C2B*). Surprisingly, expression of the C2A*-C2B* transgenic rescue in the *syt1*^{-/-} null background induced a dramatic increase in mEPJ frequency regardless of external $[\text{Ca}^{2+}]$ (Fig. 5A, B, bottom, $p < 0.001$). The elevated mini-frequency was far beyond the levels observed in *syt1*^{-/-} mutants alone or rescue strains expressing C2A-C2B* (Fig. 5B, $p < 0.001$, C2A*-C2B* vs *syt1*^{-/-} or C2A-C2B*).

The increase in spontaneous release induced by expression of a Ca^{2+} -binding defective form of Syt1 could be secondary to

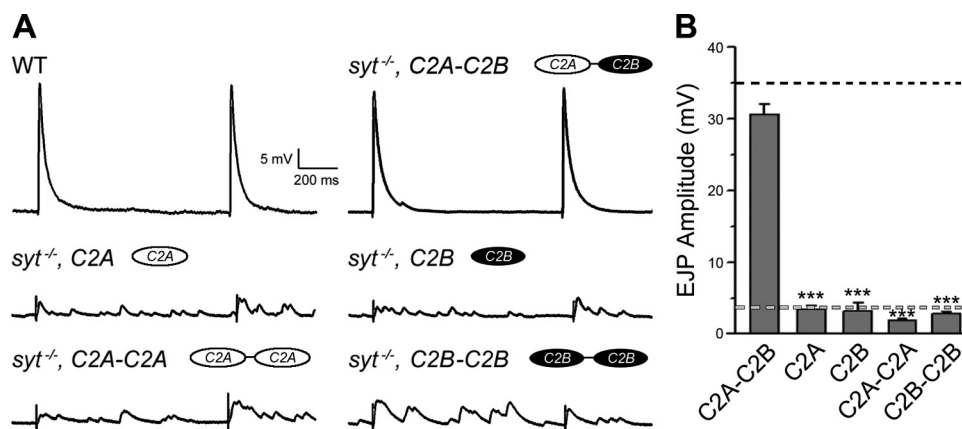


Figure 4. Functional characterization of isolated or dual Syt1 C2 domains on synchronous release. **A**, Representative eEPs recorded in the presence of high $[Ca^{2+}]_o$ (1.0 mM) are shown for wild-type and *syt1*^{-/-} rescued with the indicated transgenic constructs. **B**, Mean amplitude of eEP responses are indicated for each genotype. The mean eEP amplitude for wild-type and *syt1* null mutants (compare Fig. 3B) are indicated with black and light gray dotted lines, respectively. Data are mean \pm SEM. *** p < 0.001, one-way ANOVA with multiple comparisons using the Fisher's LSD test between *syt1*^{-/-} rescued with the C2A-C2B (Fig. 3) and the indicated genotypes. Number of NMJs examined: *syt1*^{-/-}, C2A-C2B, 11; *syt1*^{-/-}, C2A, 6; *syt1*^{-/-}, C2B, 5; *syt1*^{-/-}, C2A-C2A, 7; and *syt1*^{-/-}, C2B-C2B, 13.

Ca^{2+} -independent roles for the protein in synaptic vesicle docking, priming, or endocytosis that enhance the number of docked synaptic vesicles. To test this possibility, we examined the distribution of synaptic vesicles in *syt1*^{-/-} null mutants expressing C2A*-C2B* using transmission electron microscopy and compared them with null mutants alone or rescue lines expressing the wild-type Syt1 protein (C2A-C2B). As previous reports have indicated (Reist et al., 1998; Liu et al., 2009; Young and Neher, 2009), *syt1*^{-/-} null mutants displayed a reduction in docked synaptic vesicles (Fig. 6A,D, black column, p < 0.001), as well as a reduced vesicle density, measured as the number of vesicles per unit bouton area (Fig. 6E, p < 0.001). Expression of C2A*-C2B* resulted in an increase in docked synaptic vesicles adjacent to active zones, similar to that in strains expressing the wild-type transgene (Fig. 6A,D, black columns). Likewise, vesicle density in *syt1*^{-/-} mutants expressing C2A*-C2B* was comparable to *syt1*^{-/-} animals rescued with the wild-type protein and significantly greater than animals lacking Syt1 altogether (Fig. 6E, p < 0.001). We conclude that a Syt1 transgene with no Ca^{2+} -binding ability (C2A*-C2B*) can partially rescue defects in synaptic vesicle docking and total vesicle number. Given that docked vesicle number is similar in *syt1*^{-/-} mutants expressing the wild-type or C2A*-C2B* Syt1, it is unlikely that structural changes in synaptic vesicle distribution underlie the enhanced spontaneous release induced by C2A*-C2B* (Fig. 5A,B). Rather, Ca^{2+} -binding defective Syt1 appears to increase the fusogenicity of synaptic vesicles, resulting in a dramatic elevation in mEJP frequency.

Because the effects on evoked release suggested cooperative actions of the C2 domains, we investigated whether such cooperativity between C2 domains exists for the regulation of spontaneous release. We tested the role of the C2 domains by measuring mEJP frequency and amplitude in *syt1*^{-/-} mutants expressing single or dual C2 domains in either wild-type or mutated form (C2A, C2A*, and C2A-C2A; C2B, C2B*, and C2B-C2B). mEJP frequency in larvae expressing transgenic constructs that lack C2B (C2A, C2A*, and C2A-C2A) remained comparable with *syt1*^{-/-} mutants (Fig. 5C,D, gray dashed line) but was still far below the level of those expressing C2A*-C2B* (Fig. 5D, black dashed line, p < 0.001). Similarly, transgenic constructs consisting solely of the C2B domain (C2B, C2B*, and C2B-C2B) failed to

rescue the increased mini-frequency in null mutants (Fig. 5C,D) but remained far below the level observed in *syt1*^{-/-} rescued with C2A*-C2B* (Fig. 5D, p < 0.001). These results indicate that cooperativity between C2A and C2B is required not only for synchronous neurotransmitter release but also for Syt1 function as a partial clamp of spontaneous release. It should be noted that there are no observed Ca^{2+} -dependent changes in mEJP frequency within a range (0.2–2.0 mM) of external $[Ca^{2+}]$ at *Drosophila* NMJs (data not shown), in contrast to the Ca^{2+} -dependent nature of spontaneous release at mammalian synapses (Xu et al., 2009; Kochubey and Schneggenburger, 2011). Thus, it is likely that Syt1 exerts its function as a partial clamp in a Ca^{2+} -independent manner. Given that C2A*-C2B* lacks Ca^{2+} -binding ability and Ca^{2+} -dependent interaction with SNAREs (Earles et al., 2001; Shin et al., 2009), the mutated protein may alter Ca^{2+} -independent interactions of Syt1 with other partners regulating vesicle release, such as SNAREs and membrane phospholipids. Unlike mEJP frequency, mEJP amplitude remained similar among the different transgenic expressions, except for *syt1*^{-/-} larvae expressing a C2B-C2B chimeric protein, which significantly enhanced mEJP amplitude (Figs. 5C, 6B, p < 0.001). The increase in mEJP amplitude in C2B-C2B-expressing animals was associated with the appearance of significantly larger synaptic vesicles by transmission electron microscopy (Fig. 6A, arrowheads) and with an increase in diameter of synaptic vesicles near active zones, compared with *syt1*^{-/-} rescued with the wild-type Syt1 (C2A-C2B) (Fig. 6C). Furthermore, we also found a significant reduction in the number of vesicles distant from active zones in *syt1*^{-/-} larvae expressing a C2B-C2B protein (Fig. 6D), resulting in a reduced total vesicle density similar to *syt1*^{-/-} (Fig. 6E, p < 0.001). The Syt1 C2B domain is known to interact with several proteins involved in synaptic vesicle endocytosis (Jorgensen et al., 1995; Haucke and De Camilli, 1999; Littleton et al., 2001; Poskanzer et al., 2003; Nicholson-Tomishima and Ryan, 2004; Yao et al., 2012a; Yao et al., 2012b). Considering that the distribution of vesicles near active zones in *syt1*^{-/-} rescued with C2B-C2B was restored to wild-type levels (Fig. 6D, black column), our findings argue that the C2B-C2B protein is capable of promoting vesicle docking similar to wild-type Syt1. In contrast, the reduction in total vesicle density, together with the increased vesicle diameter and enlarged mEJP amplitude, indicates that the

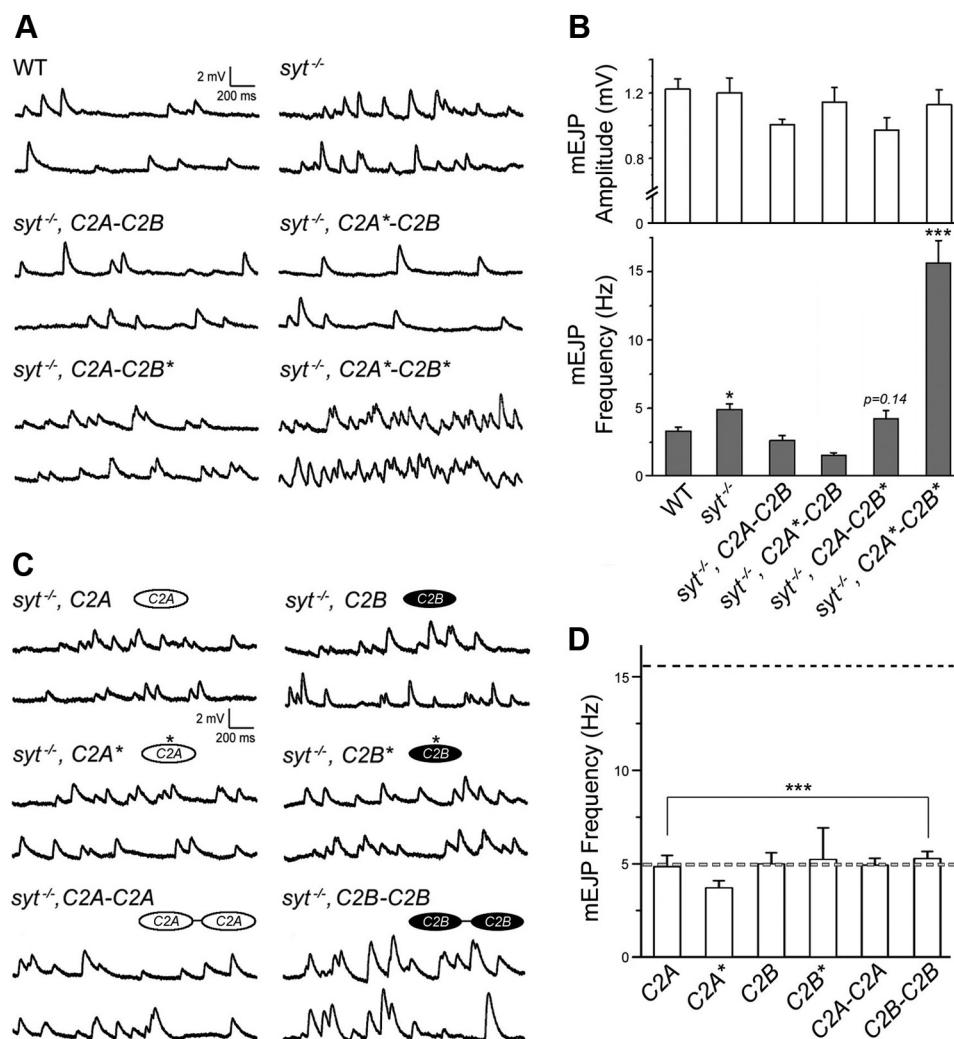


Figure 5. Contributions of Syt1 C2 domains to spontaneous synaptic release. **A**, Representative mEJPs recorded in the presence of low $[Ca^{2+}]_o$ (0.2 mM) are shown for wild-type, *syt*^{-/-} (null), and *syt*^{-/-} rescued with the indicated transgenic constructs. Calibration: 2 mV, 200 ms. **B**, Summary data for mean mEJP amplitude (top) and frequency (bottom) are shown for the indicated genotypes. Data are mean \pm SEM. *** $p < 0.001$ and * $p < 0.05$, one-way ANOVA with multiple comparison using the Fisher's LSD test between *syt*^{-/-} rescued with the C2A-C2B and the indicated genotypes. Number of NMJs examined: WT, 5; *syt*^{-/-}, 19; *syt*^{-/-}, C2A-C2B, 15; *syt*^{-/-}, C2A*-C2B, 8; *syt*^{-/-}, C2A*-C2B*, 11; *syt*^{-/-}, C2A-C2B*, 11. **C**, Representative mEJPs are shown for *syt*^{-/-} rescued with isolated C2 domains (C2A or C2B), with or without mutations in Ca^{2+} -binding residues, or with dual C2A (C2A-C2A) or C2B (C2B-C2B) domains. **D**, Summary of mean mEJP frequency is shown for the indicated genotypes. The levels measured in *syt*^{-/-} null mutants (light gray) and *syt*^{-/-} rescued with Ca^{2+} -binding defective C2A*-C2B* (black) are indicated with dotted lines. *** $p < 0.001$, one-way ANOVA analysis with multiple comparisons using the Fisher's LSD test between C2A*-C2B* (black dashed line) and the indicated genotypes. Number of NMJs examined: *syt*^{-/-}, C2A, 5; *syt*^{-/-}, C2A*, 6; *syt*^{-/-}, C2B, 5; *syt*^{-/-}, C2B*, 5; *syt*^{-/-}, C2A-C2A, 7; and *syt*^{-/-}, C2B-C2B, 18.

dual C2B-C2B protein likely alters the association of Syt1 with the endocytotic machinery, resulting in defects in vesicle number and size.

Enhanced neurotransmitter release induced by coexpression of native and Ca^{2+} -insensitive Syt1

Although expression of transgenic constructs in the *syt*^{-/-} null background allows characterization of their function in the absence of endogenous Syt1, overexpression of these constructs in the wild-type background can assay any dominant-negative or gain-of-function properties of the mutant proteins. Prior work demonstrated that overexpression of the Ca^{2+} binding C2B mutant (C2A-C2B*) induces a decrease in evoked neurotransmitter release at *Drosophila* larval NMJs (Mackler et al., 2002). We observed a similar effect caused by overexpression of C2A-C2B* using *elav*^{C155}-GAL4, which resulted in a significant decrease in eEJP amplitude compared with overexpression of wild-type Syt1 (C2A-C2B) (Fig. 7A, $p < 0.001$). Such dominant-negative effects

on synchronous release were accompanied by an increase in asynchronous release (Fig. 7A, arrows) and enhanced spontaneous fusion (Fig. 7B, bottom), presumably because of the altered nature of the mutated Syt1 C2B domain interaction with membrane phospholipids (Paddock et al., 2011).

Whereas overexpression of wild-type or C2A Ca^{2+} binding Syt1 mutants (C2A*-C2B) did not alter EJP amplitude at 0.2 mM extracellular $[Ca^{2+}]_o$, overexpression of the C2A*-C2B* protein that completely lacks Ca^{2+} -binding ability induced a dramatic increase in EJP amplitude (Fig. 7A, $p < 0.001$) without affecting asynchronous release or spontaneous fusion (Fig. 7B). Such enhancement in synchronous release could reflect an increase in the number of synaptic vesicles available for immediate release. To test this hypothesis, we examined the synaptic ultrastructure of larvae expressing wild-type or C2A*-C2B* Syt1 (Fig. 8A, B). We found no discernible abnormalities associated with overexpression of C2A*-C2B*, including no significant increase in the number of synaptic vesicles near active zones (Fig. 8B). Indeed, the

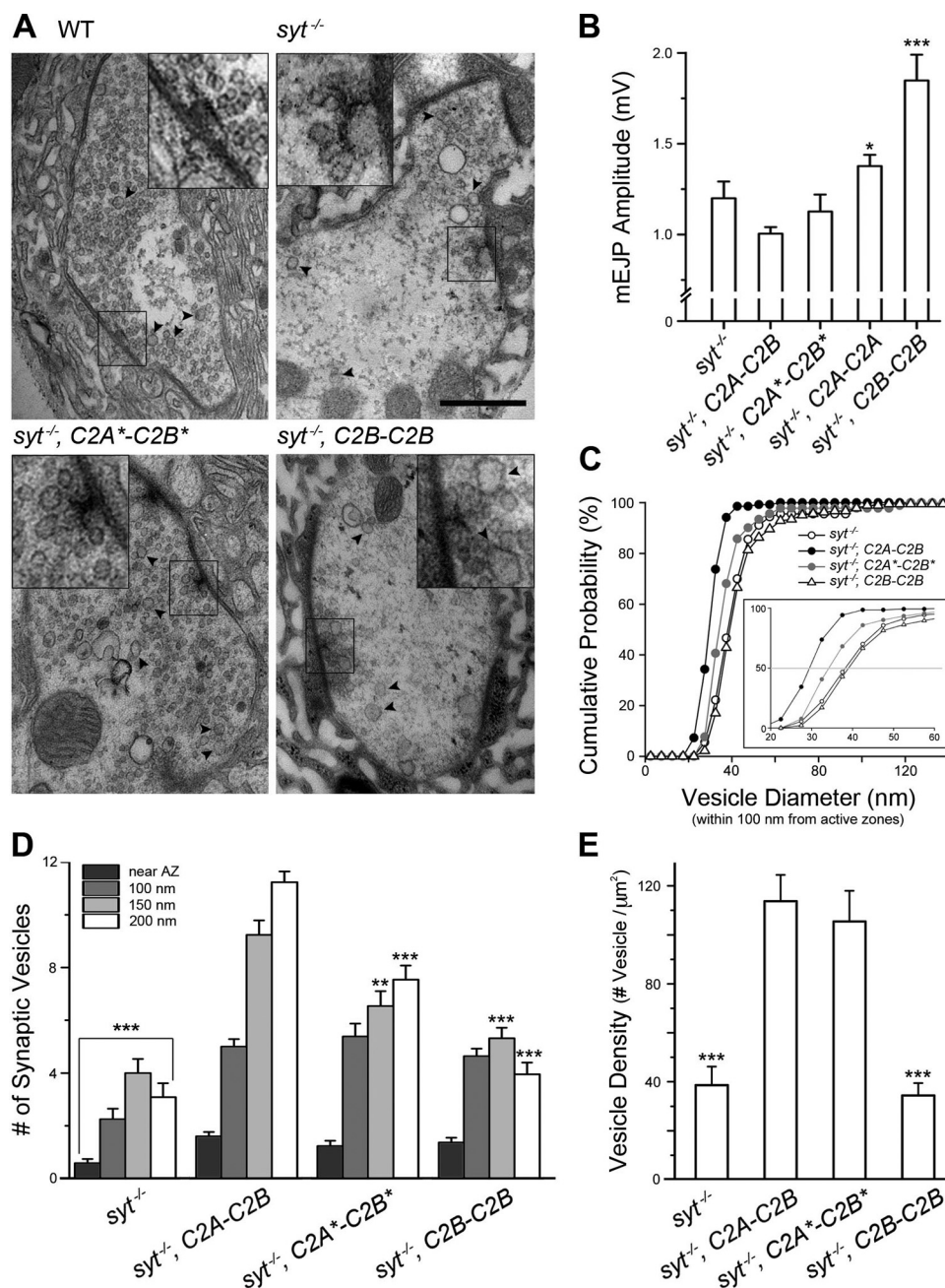


Figure 6. Role of the Syt1 C2B domain in regulation of synaptic vesicle size and density. **A**, Electron micrographs are shown for wild-type, *syt1*^{-/-} (null), and *syt1*^{-/-} rescued with Ca²⁺-binding defective C2 domains (C2A*-C2B*) or with dual C2B domains (C2B-C2B). Boxed regions in each micrograph are magnified 2.5 × in insets. Synaptic vesicles with abnormally large diameters are indicated with arrowheads. Scale bar, 500 nm. **B**, Summary of the mean mEJP amplitude is shown for *syt1*^{-/-} as well as *syt1*^{-/-} rescued with wild-type Syt1 (C2A-C2B), Ca²⁺-binding defective C2A*-C2B*, or dual C2A/C2B domains. Data for *syt1*^{-/-} as well as *syt1*^{-/-}, C2A-C2B and *syt1*^{-/-}, C2A*-C2B* from Figure 5 are presented for comparison. Data are mean ± SEM. ****p* < 0.001 and **p* < 0.05, one-way ANOVA with multiple comparisons using the Fisher's LSD test between *syt1*^{-/-} rescued with the C2A-C2B and the indicated genotypes. The number of NMJs examined for each genotype is listed in Figure 5. **C**, Cumulative diameter distributions of synaptic vesicles residing within a 100 nm radius of active zones are shown for *syt1*^{-/-} as well as those rescued with wild-type, Ca²⁺-binding defective C2 domains, or dual C2B domains. A boxed inset is provided for a detailed comparison. **D**, **E**, The number of synaptic vesicles near or in the vicinity of active zones (**D**) and total synaptic vesicle density (**E**) are summarized for each genotype indicated. Data are mean ± SEM. ****p* < 0.001 and ***p* < 0.01, one-way ANOVA with multiple comparisons using the Fisher's LSD test between *syt1*^{-/-} rescued with the C2A-C2B and the indicated genotypes. Number of active zones analyzed for **D** and **E**: *syt1*^{-/-}, 24 and 14; *syt1*^{-/-}, C2A-C2B, 17 and 14; *syt1*^{-/-}, C2A*-C2B*, 13 and 5; and *syt1*^{-/-}, C2B-C2B, 22 and 14.

total number of synaptic vesicles residing within a 200 nm radius of active zones was mildly reduced in larvae overexpressing C2A*-C2B* compared with wild-type Syt1 (Fig. 8B, *p* = 0.10), suggesting that the enhanced synchronous release is not secondary to increases in docked vesicles. Another possibility for the enhanced evoked release induced by C2A*-C2B* would be an increase in the number of release sites. However, there was no

significant change in the number of release sites when visualized by immunoreactivity against the active zone protein Bruchpilot (Fig. 8C,D), indicating that structural changes are unlikely to contribute to the increased synchronous release upon overexpression of C2A*-C2B*.

We next investigated which functional aspects of the release process were modified by overexpression of C2A*-C2B*. For in-

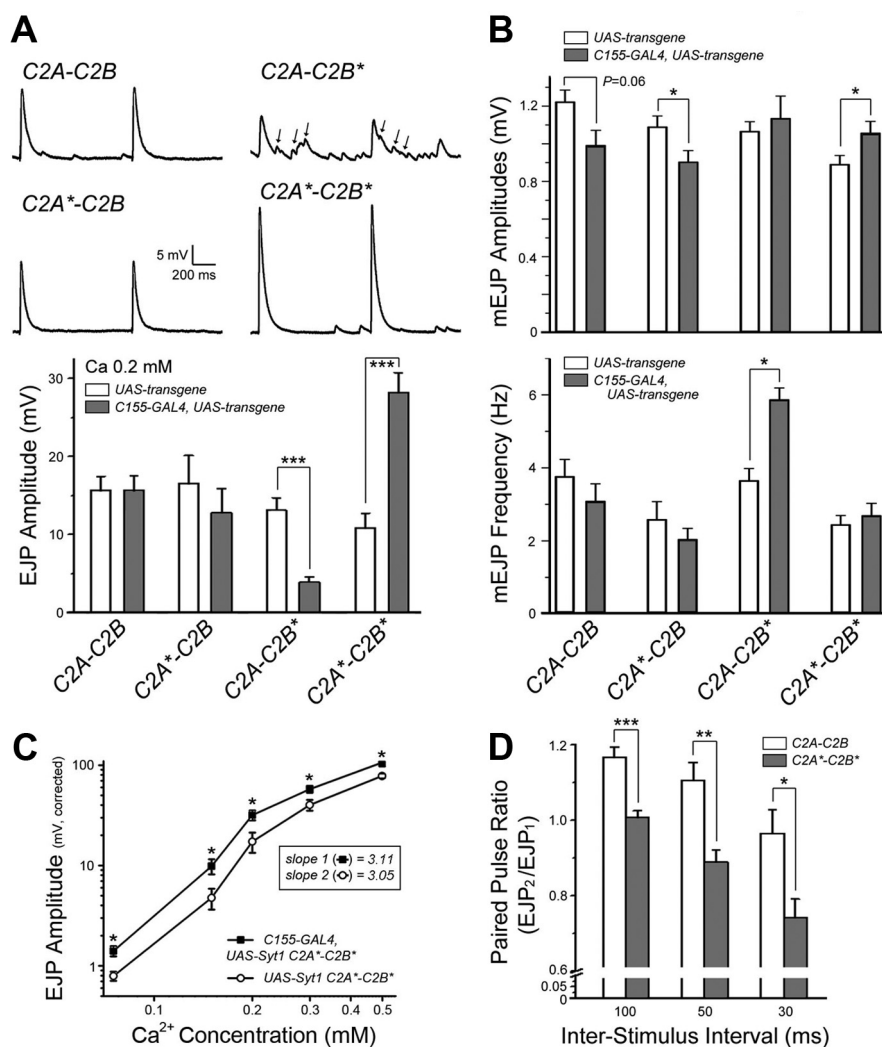


Figure 7. Interplay between endogenous and transgenic Syt1 constructs on evoked synchronous release. **A**, Top, Representative traces of two consecutive EJPs recorded in the presence of low $[Ca^{2+}]_o$ (0.2 mM) are shown for wild-type larvae overexpressing the indicated transgenic Syt1 constructs. Calibration: 5 mV, 200 ms. Asynchronous release events during stimulation are indicated with arrows. (Bottom) Mean eEJP amplitude is summarized for the indicated genotypes. Animals carrying each transgenic construct without a GAL4-driver (white) served as controls for comparison with the same transgenic constructs driven by *elav*^{C155}-GAL4 (gray). **B**, Mean mEJP amplitude (top) and frequency (bottom) are summarized for the indicated genotypes. **A**, **B**, Number of NMJs examined (control and transgene expression): C2A-C2B, 7 and 8; C2A*-C2B, 8 and 5; C2A-C2B*, 11 and 8; and C2A*-C2B*, 10 and 11. **C**, Log-log plot for eEJP amplitudes at varying $[Ca^{2+}]_o$ is shown for animals overexpressing the C2A*-C2B* construct (■) and its transgenic control without a GAL4-driver (○). The slope values calculated from a linear fit of the first three data points (0.075–0.2 mM $[Ca^{2+}]_o$) are indicated in the box. Number of NMJs examined (control and transgene expression): 7 and 11 at 0.075 mM $[Ca^{2+}]_o$; 7 and 11 at 0.15 mM; 6 and 11 at 0.2 mM; 6 and 10 at 0.3 mM; and 6 and 9 at 0.5 mM. **D**, The ratios of eEJP responses in a paired-pulse stimulation paradigm are displayed for wild-type animals overexpressing the C2A-C2B (white) or C2A*-C2B* (gray) constructs. Number of NMJs examined (C2A-C2B and C2A*-C2B*): 8 and 6 at 30 ms interval; 8 and 7 at 50 ms; and 8 and 7 at 100 ms. **A–D**, Data are mean \pm SEM. *** $p < 0.001$, ** $p < 0.01$, and * $p < 0.05$, Student's *t* test for control (UAS-transgene) versus neuronal expression (C155-GAL4, UAS-transgene) (**A–C**) or for neuronal overexpression of C2A-C2B versus C2A*-C2B* (**D**).

stance, increases in release probability or Ca^{2+} affinity could result in enhanced synchronous release. We compared these properties in larvae overexpressing C2A*-C2B* with either a transgenic control lacking a GAL4-driver or to larvae overexpressing wild-type Syt1 (C2A-C2B). When eEJP amplitudes were measured at varying external $[Ca^{2+}]$, the log-log plot for Ca^{2+} -dependency of release in larvae overexpressing C2A*-C2B* was shifted to the left compared with a transgenic control, without a change in the slope (Fig. 7C, $p < 0.05$; slope for the control and C2A*-C2B*, 3.05 and 3.11, respectively). These data indicate an increase in Ca^{2+} affinity, but not cooperativity, favoring lower

Ca^{2+} conditions to support synchronous release. Furthermore, the paired-pulse ratio was significantly reduced in C2A*-C2B*-overexpressing larvae compared with animals overexpressing wild-type Syt1 (Fig. 7D). Given that the paired-pulse ratio reflects release probability during the first stimulation, these results indicate that overexpression of C2A*-C2B* enhances synchronous release by increasing release probability and lowering the $[Ca^{2+}]$ required to trigger fusion.

Multimerization of native and Ca^{2+} -binding defective Syt1

Our results demonstrate that expression of a Ca^{2+} -binding defective Syt1 (C2A*-C2B*) protein in the absence of native Syt1 robustly increases the rate of spontaneous fusion (Fig. 5), without restoring fast synchronous release (Fig. 3). In contrast, coexpression of C2A*-C2B* and native Syt1 resulted in a large enhancement in synchronous release (Fig. 7A), with no increase in spontaneous fusion (Fig. 7B). If Syt1 regulates synaptic release as a monomer, we predicted that a mixed population of Ca^{2+} -binding defective and native Syt1 monomers would yield one of several possibilities: (1) both synchronous and spontaneous release would remain unchanged compared with control if native wild-type monomers in a mixed population are sufficient to perform the full repertoire of Syt1 functions; (2) synchronous release will be partially disrupted with a slight elevation of spontaneous release if wild-type and Ca^{2+} -binding defective Syt1 monomers participate independently in regulating both synchronous and spontaneous release; or (3) synchronous and spontaneous release will be differentially disrupted if native and mutated Syt1 monomers contribute unequally to different modes of synaptic release, resulting in either relatively normal synchronous release with a slightly enhanced spontaneous release or impaired synchronous release with a wild-type level of spontaneous fusion. However, our results are not consistent with these predictions, given the normal rate of spontaneous release

with enhanced synchronous fusion caused by coexpression of Ca^{2+} -binding defective and native Syt1 (Fig. 7A,B). These results suggest the possibility of a Syt1 hetero-multimeric complex where native Syt1 clamps spontaneous release, whereas C2A*-C2B* Syt1 promotes evoked fusion. Indeed, the synaptotagmin family has been biochemically shown to multimerize in Ca^{2+} -independent (Fukuda et al., 1999; Bai et al., 2000; Fukuda et al., 2001) and Ca^{2+} -dependent manners (Chapman et al., 1996; Damer and Creutz, 1996; Sugita et al., 1996; Osborne et al., 1999; Desai et al., 2000). Thus, we examined whether dimerization could occur between native and Ca^{2+} -insensitive Syt1,

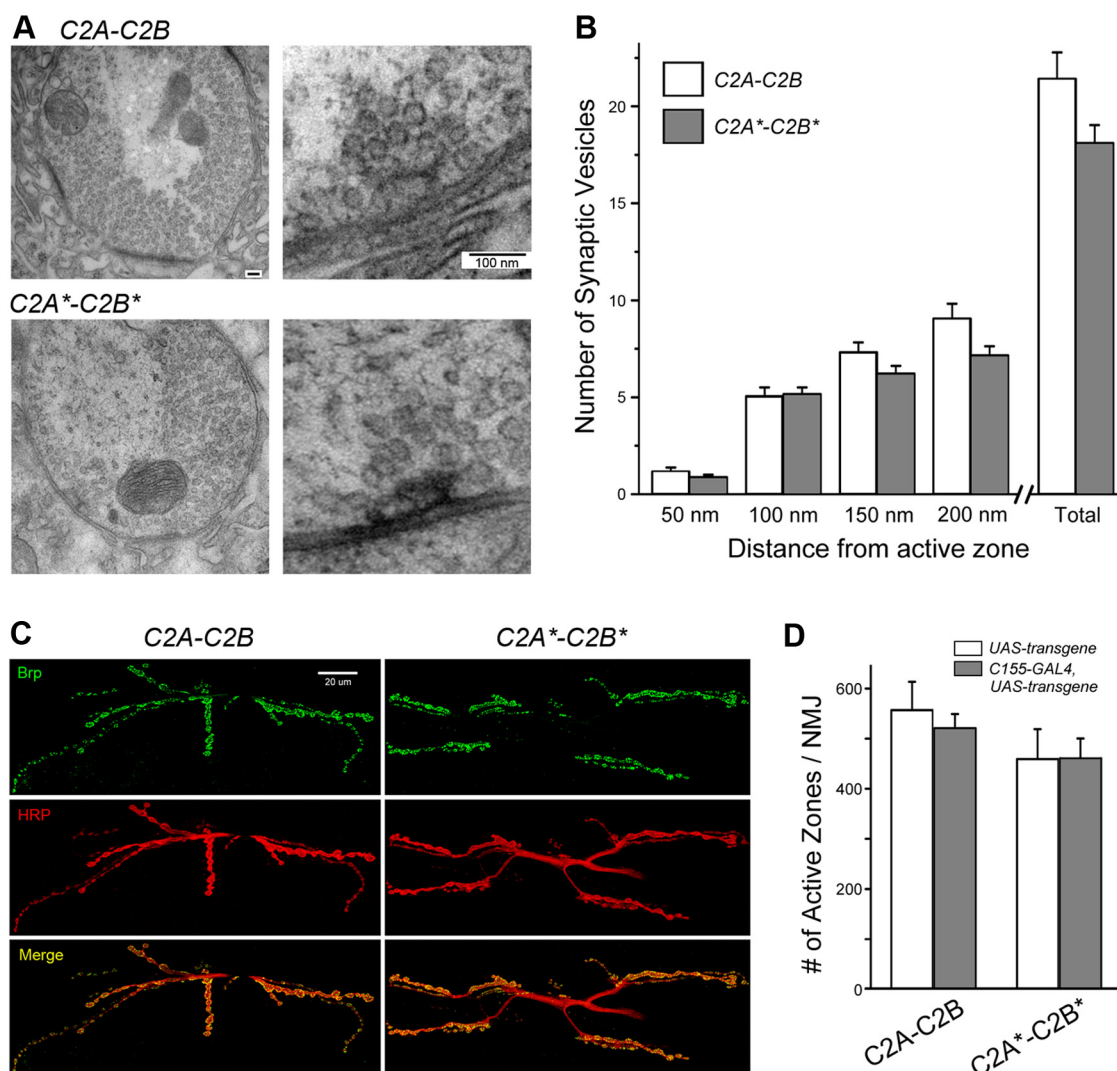


Figure 8. Similar distributions of synaptic vesicles and release sites between animals overexpressing wild-type and Ca^{2+} -binding defective Syt1. **A**, Electron micrographs are shown for larvae overexpressing the wild-type (C2A-C2B, top) or Ca^{2+} -binding defective (C2A*-C2B*, bottom) Syt1 constructs. Detailed view near a single active zone is shown on the right panels. Scale bar, 100 nm. **B**, Mean number of synaptic vesicles at varying distances from active zones is shown for larvae overexpressing the wild-type or Ca^{2+} -binding defective Syt1 constructs. Number of active zones analyzed (C2A-C2B and C2A*-C2B*): 13 and 25 at 50 nm; 19 and 32 at 100–200 nm; 15 and 16 for total number of synaptic vesicles. **C**, Representative confocal images depicting distributions of release sites (active zones) in muscle 6/7 NMJs are shown for larvae overexpressing wild-type or Ca^{2+} -binding defective Syt1. Active zones are identified by immunoreactivity against Brp (top panels, green). The overall structure of NMJs is detected with HRP immunoreactivity (middle panels, red). The merged images of Brp and HRP channels are shown in the bottom panels. Scale bar, 20 μm . **D**, The mean number of active zones per NMJ is summarized for larvae containing each transgenic construct without a GAL4 driver (white, control) and those with transgenic constructs driven by *elav*^{C155}-GAL4 (gray). Number of NMJs examined (control and transgene expression): 12 and 11 for C2A-C2B; 11 and 10 for C2A*-C2B*. **B**, **D**, Data are mean \pm SEM.

which might underlie the enhanced synchronous release observed in larvae overexpressing C2A*-C2B*.

We purified GFP-tagged full-length Syt1 expressed in *syt1*^{-/-} fly heads and performed pull-down assays with GST-fused wild-type (GST-C2A-C2B) or mutated (GST-C2A*-C2B*) recombinant Syt1 (Fig. 9A). Purified GFP-tagged Syt1 bound to the cytoplasmic C2 domains of Syt1 immobilized on glutathione-Sepharose beads in the absence of Ca^{2+} , as visualized by GFP immunoreactivity (Fig. 9A, EGTA, 2 mM). Further multimerization of Syt1 was observed by increasing $[\text{Ca}^{2+}]$ to 1 or 10 mM (Fig. 9A). These Ca^{2+} -independent and Ca^{2+} -dependent interactions did not differ between wild-type and C2A*-C2B* Syt1 (Fig. 9A). Similar results were obtained when we performed pull-down assays with His-tagged wild-type cytoplasmic Syt1 C2 domains to bind GST-fused wild-type or mutated recombinant Syt1. Ca^{2+} -independent binding between His-Syt1 and GST-fused Syt1 (wild-type and C2A*-C2B*) was consistently detected

(Fig. 9B). As seen with endogenous Syt1 pull-down, the interaction between His-Syt1 and GST-tagged Syt1 was enhanced in response to Ca^{2+} (Fig. 9B). Together, these findings indicate that endogenous Syt1 and Ca^{2+} -binding defective Syt1 likely form hetero-multimers in the absence of Ca^{2+} , which enhances the fusogenicity of synaptic vesicles in response to Ca^{2+} (Fig. 7).

Discussion

Requirement of tandem Syt1 C2A-C2B domains in regulation of neurotransmitter release

Using transgenic rescue approaches with chimeric Syt1 proteins, we demonstrate that both the C2A and C2B domains are individually required for Syt1 function in regulating vesicle release at the *Drosophila* NMJ. Prior studies have established that the C2B domain of Syt1 is a critical module for activating synchronous vesicle release in *Drosophila* (Littleton et al., 1994; Littleton et al., 2001; Mackler et al., 2002; Yoshihara et al., 2010). In contrast, the

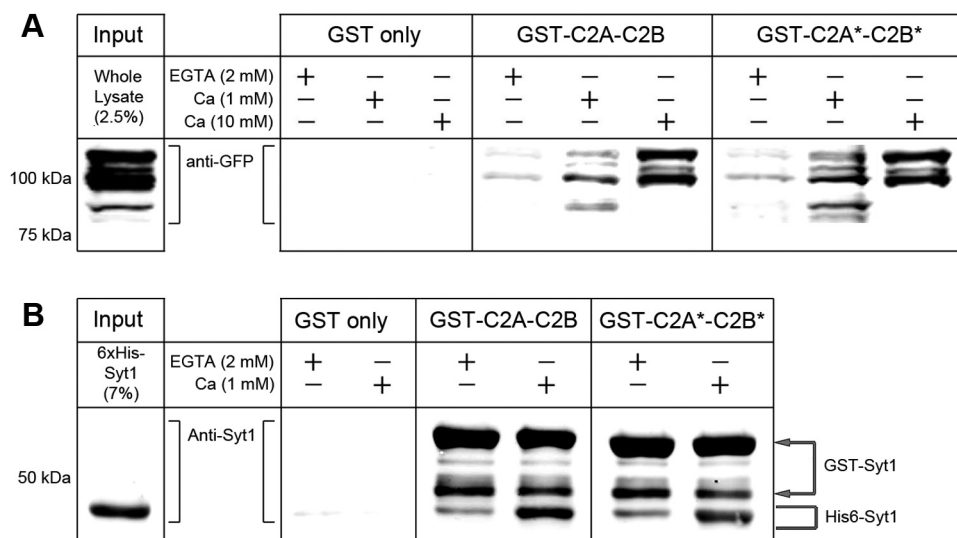


Figure 9. Multimerization of wild-type and C2A*-C2B* Syt1. **A**, Western blot with anti-GFP antibodies is shown to demonstrate Ca^{2+} -independent and Ca^{2+} -dependent binding of endogenous GFP-tagged Syt1, purified from adult head lysates, to GST-fused wild-type (C2A-C2B) or Ca^{2+} -binding defective (C2A*-C2B*) Syt1. Fly head extracts are incubated with GST-fused Syt1 in the absence (EGTA, 2 mM) or presence of Ca^{2+} (1 or 10 mM). **B**, Binding of purified wild-type Syt1-His₆ to wild-type or Ca^{2+} -binding defective GST-Syt1. GST-Syt1 (C2A-C2B and C2A*-C2B*) and interacting wild-type Syt1-His₆ products are indicated with arrows (top two bands) and a bracket (a single bottom band), respectively.

contribution of the C2A domain is still being elucidated (Yoshihara et al., 2010; Striegel et al., 2012). Although the activities of isolated C2 domains in their interaction with SNAREs and membrane phospholipids remain controversial in *in vitro* studies, growing evidence suggests synergistic actions of both C2 domains. Isolated C2 domains, either C2A or C2B, bind syntaxin and SNAP-25 less effectively than C2A-C2B constructs in response to Ca^{2+} in *in vitro* binding assays (Chapman et al., 1996; Rickman and Davletov, 2003) or reconstituted vesicle fusion assays (Tucker et al., 2004). Synergistic actions of the C2 domains also occur during lipid binding, as isolated C2 domains bind poorly to liposomes made from chromaffin granule lipid extracts compared with dual C2A-C2B proteins (Damer and Creutz, 1994). Likewise, the interaction between the C2B domain and membrane phospholipids requires an intact C2A domain, even in the absence of its Ca^{2+} - and lipid-binding abilities (Bai et al., 2002). Furthermore, Ca^{2+} -dependent penetration of both C2 domains into reconstituted lipid bilayers appears to occur simultaneously (Hui et al., 2006). These *in vitro* studies indicate that cooperative actions of C2 domains regulate SNARE and lipid interactions and match our *in vivo* observations that evoked neurotransmitter release requires both C2 domains of Syt1.

In addition to a dual C2 domain structure, our findings indicate that C2A and C2B are uniquely required. Double C2 domain constructs containing only C2A-C2A fail to restore Syt1 function (Fig. 4), consistent with a lack of Ca^{2+} -dependent binding to t-SNAREs in the absence of C2B (Earles et al., 2001). Similarly, Syt2 constructs lacking the C2B domain at the calyx of Held failed to support evoked fusion in the null mutant background (Kochubey and Schneggenburger, 2011). Although the C2B domain rather than C2A plays a more prominent role in evoked fusion (Fig. 3), our results also indicate that the C2B domain is not solely responsible for Syt1 function in synaptic vesicle fusion, as an isolated or dual C2B domain cannot support synaptic vesicle fusion (Fig. 4). In addition, dual C2B-C2B constructs induce additional defects in vesicle endocytosis pathways that increase vesicle size and

mEJP amplitude (Fig. 6). This contrasts with the ability of purified Syt1 dual C2B domains to induce liposome aggregation and SNARE-dependent vesicle fusion comparable with the wild-type Syt1 (C2A-C2B) *in vitro* (Hui et al., 2011), indicating a limitation of *in vitro* fusion assays in recapitulating *in vivo* requirements for synaptic vesicle release.

Regulation of spontaneous fusion by Syt1 C2 domains

In addition to triggering synchronous fusion, Syt1 also partially clamps spontaneous fusion in the absence of a Ca^{2+} signal. Transgenic constructs with Ca^{2+} binding mutations in C2A fully rescued the elevated spontaneous fusion rate in *sytl*^{-/-} null mutants, whereas those with Ca^{2+} binding mutations in C2B did not (Fig. 5). Prior *in vitro* studies have demonstrated that C2A*-C2B is indistinguishable from wild-type C2A-C2B in its binding to membrane phospholipids (Bai et al., 2002; Bai et al., 2004), whereas mutations in the C2B domain (C2A-C2B*) facilitated Ca^{2+} -independent binding of Syt1 to phospholipids (Mackler et al., 2002). Such differences in Ca^{2+} -independent lipid binding may underlie the differential regulation of spontaneous release in *sytl*^{-/-} mutants rescued with C2A*-C2B versus C2A-C2B*. Alternatively, neutralizations of these Ca^{2+} -binding residues may induce differential changes in C2 domain structure that influence Ca^{2+} -independent interactions between Syt1 and SNAREs. Syt1 has been shown to form a Ca^{2+} -independent complex with SNARE proteins *in vitro* (Bennett et al., 1992; Rickman and Davletov, 2003; Shin et al., 2003) and to stabilize partially assembled trans-SNARE complexes to function as a fusion clamp (Chicka et al., 2008).

Unexpectedly, we found a dramatic enhancement in the rate of spontaneous vesicle release far beyond that observed in *sytl*^{-/-} null mutants when we rescued with transgenic proteins carrying mutations of Ca^{2+} -binding residues in both C2 domains (C2A*-C2B*, Fig. 5). Unlike the Ca^{2+} -dependent nature of spontaneous release at mammalian synapses (Xu et al., 2009; Kochubey and Schneggenburger, 2011), Ca^{2+} -dependent changes in mEJP frequency within a compatible

range (0.2–2.0 mM) of external $[Ca^{2+}]$ at *Drosophila* NMJs has not been observed. As such, how can the dual Ca^{2+} -binding mutations of Syt 1 dramatically alter spontaneous release at *Drosophila* NMJs? Recently, conformational changes in C2 domain structure associated with Ca^{2+} binding have been suggested based on crystallization studies (Fuson et al., 2007). Indeed, we observed a change in the size of a protease-sensitive breakdown product previously described for Syt1 (Littleton et al., 1993) in *syt1*^{-/-} null mutants rescued with C2A*-C2B* (Fig. 2), suggesting subtle conformational changes in Syt1 structure. Ca^{2+} -binding mutations neutralize negatively charged aspartate residues (i.e., C2A*-C2B*), somewhat similar to the effect of bound Ca^{2+} ions. If these Ca^{2+} -binding mutations in the C2 domains induce structural changes, it might alter Ca^{2+} -independent interactions between Syt1 and SNARE complexes to increase the frequency of spontaneous vesicle release. It is unlikely that the effect of C2A*-C2B* on mEJP frequency would be due solely to the rescue of the docking and endocytosis defects observed in *syt1*^{-/-} mutants, as wild-type C2A-C2B transgenes also rescue these defects but did not induce a dramatic elevation in spontaneous fusion.

Physiological significance of Syt1 multimerization

Although the Syt family can form multimers *in vitro*, the relevance of such multimeric complexes on synaptic transmission *in vivo* has remained unclear. Different Syt family members can form homodimers via their N-terminal cysteine residues in a Ca^{2+} -independent manner (Fukuda et al., 1999; Bai et al., 2000; Fukuda et al., 2001), whereas Ca^{2+} -dependent self-association and formation of heterodimers between different Syt family members are linked to activity of the C2B domain (Chapman et al., 1996; Damer and Creutz, 1996; Sugita et al., 1996; Osborne et al., 1999; Desai et al., 2000). Genetic analyses in *Drosophila* have suggested Syt oligomerization may regulate synaptic transmission. Mutations in the C2B domain that alter Syt1 oligomerization (*Syt*^{AD3}) (DiAntonio and Schwarz, 1994; Littleton et al., 1994; Fukuda et al., 2000) disrupt synaptic vesicle release (Littleton et al., 2001), although the effects of this mutation on phosphoinositide and SNARE binding, as well as defects in interactions with endocytosis regulators, might also contribute to the inability of the protein to control fusion.

The effects of the C2A*-C2B* transgene on release in the null versus wild-type backgrounds argue that the mutant form of Syt1 may form multimers with native Syt1 *in vivo* to generate a complex with unique properties. In the absence of endogenous Syt1, the C2A*-C2B* mutant failed to rescue any aspect of evoked release but strongly activated spontaneous fusion, suggesting that Ca^{2+} -binding defective Syt1 at rest was fusion-prone (Fig. 5). In contrast, when overexpressed with wild-type Syt1, there was no change in mEJP frequency, indicating that a multimer of native Syt1 and C2A*-C2B* Syt 1 may be able to effectively clamp spontaneous release (Fig. 7B). In addition, this complex would be able to dramatically enhance evoked release (Fig. 7A). We hypothesize that overexpression of C2A*-C2B* allows this fusogenic version of Syt1 to multimerize with endogenous Syt1 in both Ca^{2+} -independent and Ca^{2+} -dependent manners *in vivo*, as suggested by binding assays (Fig. 9). These heteromultimers exhibit characteristics of both Syt1 molecules: they are more effective in Ca^{2+} -dependent vesicle fusion, as evidenced by increased release probability (Fig. 7D) and Ca^{2+} sensitivity (Fig. 7C) while maintaining a clamping function provided by endogenous Syt1 to reduce Ca^{2+} -independent spontaneous fusion (compare Figs. 7B

5). Although additional studies will be required to confirm that the multimeric Syt1 complex exists *in vivo*, our data indicate that the different modes of synaptic release, Ca^{2+} -independent spontaneous fusion and Ca^{2+} -dependent synchronous vesicle release, are likely to be regulated by multimerization of Syt1. In addition, our findings provide evidence that both C2 domains of Syt1 are uniquely required for the two forms of synaptic neurotransmitter release.

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