

# Doc2 Proteins Are Not Required for the Increased Spontaneous Release Rate in Synaptotagmin-1-Deficient Neurons

Rocío Díez-Arazola,<sup>1</sup>  Marieke Meijer,<sup>2</sup> Quentin Bourgeois-Jaarsma,<sup>1</sup>  L. Niels Cornelisse,<sup>2</sup>  Matthijs Verhage,<sup>1,2</sup> and  Alexander J. Groffen<sup>2</sup>

<sup>1</sup>Department of Functional Genomics, Center for Neurogenomics and Cognitive Research, Neuroscience Campus Amsterdam, Vrije Universiteit Amsterdam, 1081 HV Amsterdam, The Netherlands, and <sup>2</sup>Department of Clinical Genetics, Center for Neurogenomics and Cognitive Research, Amsterdam UMC, Vrije Universiteit Amsterdam, 1081 HV Amsterdam, The Netherlands

Regulated secretion is controlled by  $\text{Ca}^{2+}$  sensors with different affinities and subcellular distributions. Inactivation of *Syt1* (synaptotagmin-1), the main  $\text{Ca}^{2+}$  sensor for synchronous neurotransmission in many neurons, enhances asynchronous and spontaneous release rates, suggesting that *Syt1* inhibits other sensors with higher  $\text{Ca}^{2+}$  affinities and/or lower cooperativities. Such sensors could include *Doc2a* and *Doc2b*, which have been implicated in spontaneous and asynchronous neurotransmitter release and compete with *Syt1* for binding SNARE complexes. Here, we tested this hypothesis using triple-knock-out mice. Inactivation of *Doc2a* and *Doc2b* in *Syt1*-deficient neurons did not reduce the high spontaneous release rate. Overexpression of *Doc2b* variants in triple-knock-out neurons reduced spontaneous release but did not rescue synchronous release. A chimeric construct in which the C2AB domain of *Syt1* was substituted by that of *Doc2b* did not support synchronous release either. Conversely, the soluble C2AB domain of *Syt1* did not affect spontaneous release. We conclude that the high spontaneous release rate in synaptotagmin-deficient neurons does not involve the binding of *Doc2* proteins to *Syt1* binding sites in the SNARE complex. Instead, our results suggest that the C2AB domains of *Syt1* and *Doc2b* specifically support synchronous and spontaneous release by separate mechanisms. (Both male and female neurons were studied without sex determination.)

## Significance Statement

Neurotransmission in the brain is regulated by presynaptic  $\text{Ca}^{2+}$  concentrations. Multiple  $\text{Ca}^{2+}$  sensor proteins contribute to synchronous (*Syt1*, *Syt2*), asynchronous (*Syt7*), and spontaneous (*Doc2a/Doc2b*) phases of neurotransmitter release. Genetic ablation of synchronous release was previously shown to affect other release phases, suggesting that multiple sensors may compete for similar release sites, together encoding stimulus–secretion coupling over a large range of synaptic  $\text{Ca}^{2+}$  concentrations. Here, we investigated the extent of functional overlap between *Syt1*, *Doc2a*, and *Doc2b* by reintroducing wild-type and mutant proteins in triple-knock-out neurons, and conclude that the sensors are highly specialized for different phases of release.

## Introduction

Neurotransmitter secretion is induced by action potentials (APs) which activate voltage-dependent  $\text{Ca}^{2+}$  channels and, subsequently,  $\text{Ca}^{2+}$  sensor proteins in synapses (Südhof, 2013). The

canonical  $\text{Ca}^{2+}$  sensor synaptotagmin-1 (*Syt1*) is anchored in the synaptic vesicle membrane. It contains two C2 domains which bind  $\text{Ca}^{2+}$  ions in a cooperative manner. Ablation of *Syt1* selectively impairs synchronous neurotransmission, whereas asynchronous (slower release triggered by action potentials) and

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Correspondence should be addressed to Alexander J. Groffen at a.j.a.groffen@vu.nl.

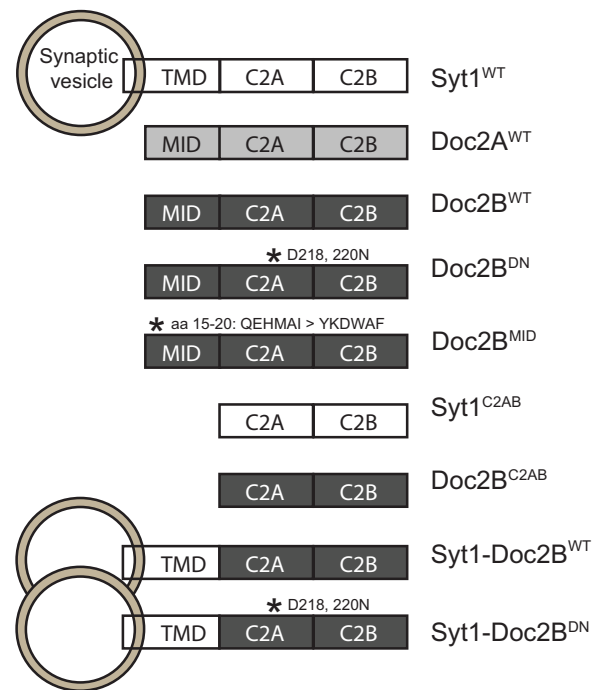
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spontaneous release (quantal events occurring at resting membrane potential) are unaffected (Littleton et al., 1993; Broadie et al., 1994; Geppert et al., 1994). In adrenal chromaffin cells, Syt1 is important for fast-burst exocytosis but not sustained release (Voets et al., 2001). These findings prompted a “two release component” hypothesis (Broadie et al., 1994; Geppert et al., 1994; Goda and Stevens, 1994), stating that other than the canonical sensor, a second sensor can occupy release sites and codetermine the  $\text{Ca}^{2+}$  dependence of secretion. In Syt1-deficient neurons, a higher apparent  $\text{Ca}^{2+}$  sensitivity and lower cooperativity was observed (Xu et al., 2009). Similarly, Syt2 deficiency in the calyx of Held synapse strongly reduces  $\text{Ca}^{2+}$  cooperativity (Sun et al., 2007; Kochubey and Schneggenburger, 2011), suggesting involvement of additional release mechanisms, such as alternative sensors with relatively low cooperativities and high  $\text{Ca}^{2+}$  affinities.

Following Syt1, a number of additional  $\text{Ca}^{2+}$  sensors were identified (reviewed by Walter et al., 2011). Syt2 and Syt9 rescue synchronous release in Syt1-deficient neurons (Xu et al., 2007). Syt7 and Syt10 do not rescue fast release but contribute to other  $\text{Ca}^{2+}$ -secretion coupling mechanisms in neurons and other cell types (Gustavsson et al., 2008, 2009; Cao et al., 2011). Acute Syt7 knock-down studies in Syt1-null neurons show that Syt7 contributes to asynchronous release (Bacaj et al., 2013). Syt7 enhances neurotransmission during high-frequency trains, thereby compensating for synaptic depression (Turecek et al., 2017). In chromaffin cells, Syt7 is required for sustained secretion (Schonn et al., 2008). In both neurons and chromaffin cells, the combined removal of Syt1 and Syt7 leaves a remaining release component. In particular, Syt1-deficient neurons exhibit an increased spontaneous release rate (Littleton et al., 1993; Broadie et al., 1994; Xu et al., 2009). Spontaneous release is unaffected by Syt7 knockdown and inhibited by Syt7 overexpression (Bacaj et al., 2013), again suggesting that additional release mechanisms exist.

Spontaneous neurotransmission is mediated by Doc2 proteins, which contain C2AB domains similar to those of Syt1 (Orita et al., 1995; Sakaguchi et al., 1995). The C2AB domains of Doc2b and Syt1 share several properties:  $\text{Ca}^{2+}$ -dependent phosphatidylserine binding, phosphatidylinositol (4,5)-bisphosphate and SNARE complex binding, membrane curvature induction, and fusion of SNARE-containing liposomes (Brose et al., 1992; Rickman et al., 2004; Martens et al., 2007; Groffen et al., 2010). Doc2a and Doc2b translocate from the cytosol to the plasma membrane with high  $\text{Ca}^{2+}$  sensitivity (Groffen et al., 2004, 2006). Doc2b may function as a high-affinity  $\text{Ca}^{2+}$  sensor (Groffen et al., 2006, 2010; Courtney et al., 2018) or by another mechanism (Pang et al., 2011). The N-terminal domain contains a Munc13-interacting domain (MID) which mediates plasma membrane recruitment in response to diacylglycerol (Orita et al., 1997; Duncan et al., 1999). Doc2 proteins also contribute to asynchronous neurotransmission (Yao et al., 2011a). In chromaffin cells, Doc2b enhances burst exocytosis and inhibits sustained release (Friedrich et al., 2008; Pinheiro et al., 2013; Houy et al., 2017). Doc2 proteins also mediate endocrine processes in pancreatic  $\beta$  cells (Miyazaki et al., 2009; Ramalingam et al., 2012), adipocytes, muscle (Fukuda et al., 2009; Ramalingam et al., 2012), and mast cells (Higashio et al., 2008). Because Syt1 and Doc2 compete for binding to SNARE complexes (Groffen et al., 2010), Doc2 proteins might be responsible for the high spontaneous release frequency observed in the absence of Syt1.



**Figure 1.** Schematic overview of the constructs investigated in this study, expressed from lentiviral vectors. The N-terminal domain of rat Syt1 (amino acids 1–141) is abbreviated as TMD to emphasize that it includes a transmembrane domain. The N-terminal domains of rat Doc2a (amino acids 1–92) and Doc2b (amino acids 1–126) are designated MID to emphasize that they contain a Munc13-binding domain. C2A–C2B domains comprise residues 142–421 (Syt1), 93–403 (Doc2a), and 127–412 (Doc2b). Asterisks indicate targeted mutations as specified.

Here, we generated mice lacking Syt1, Doc2a, and Doc2b and found that Doc2a/Doc2b do not account for the high spontaneous release rate of Syt1-deficient neurons. Our data demonstrate that the C2AB domains of Syt1 and Doc2b are functionally distinct, suggesting separate mechanisms for evoked and spontaneous neurotransmitter release.

## Materials and Methods

**Generation of triple-knock-out mice.** Animals were housed, bred, and handled in accordance with Dutch and European Union (EU) governmental guidelines. Protocols were approved by the Vrije Universiteit Animal Ethics and Welfare Committee (approval number FGA 11-06). Wild-type C57BL/6 mice were obtained from Charles River Laboratories. Null alleles for *Syt1* (Geppert et al., 1994), *Doc2a* (Sakaguchi et al., 1999), and *Doc2b* (Groffen et al., 2010) were maintained by extensively backcrossing heterozygous mice with C57BL/6. To produce Syt1 single null mice, heterozygotes were intercrossed. To produce triple-knock-out (TKO) mice, a *Doc2a*<sup>-/-</sup> *Doc2b*<sup>-/-</sup> *Syt1*<sup>+/-</sup> mouse line was first established and then intercrossed.

**Construction of lentiviral overexpression vectors.** Expression vectors encoding wild-type, mutated, or hybrid variants of Syt1 and Doc2b were constructed as illustrated in Figure 1. NCBI reference sequences are NP\_001028852.2 (rat Syt1), NP\_112404.1 (rat Doc2b), and NP\_075226.1 (rat Doc2a).

**Autapse culture electrophysiology.** Autapse cultures were prepared as previously described (Groffen et al., 2010). In brief, neurons were dissociated from WT, Syt1 knock-out, and Syt1/Doc2a/Doc2b TKO mouse hippocampi at embryonic stage E18 and plated on glial microislands. Whole-cell recordings were performed at room temperature between 14 and 18 d *in vitro*. The intracellular solution was free of EGTA; it contained the following: 125 mM K<sup>+</sup>-gluconate, 10 mM NaCl, 4.6 mM MgCl<sub>2</sub>, 4 mM K<sub>2</sub>-ATP, 15 mM creatine phosphate, 10 U/ml phosphocreatine kinase, pH 7.30, 300 mOsm. The external medium contained the follow-

ing (in mM): 140 NaCl, 2.4 KCl, 4 CaCl<sub>2</sub>, 4 MgCl<sub>2</sub>, 10 glucose, and 10 HEPES, pH 7.30, 300 mOsm. Clampfit v10.2 (Molecular Devices) was used for offline analysis of evoked EPSCs. Data acquisition rate was 10 kHz. Miniature EPSCs were detected using Mini Analysis v6.0.3 (Synaptosoft, Inc.) using a threshold amplitude of 8 pA. Each trace was visually inspected to confirm accuracy of event detection. To calculate synchronous and asynchronous components of the EPSC charge as described, a homemade routine was written in the MATLAB environment (MathWorks; for details, see Bourgeois-Jaarsma et al., 2019). For overexpression of full-length rat Doc2a, Doc2b, and Syt1, cells were infected with lentiviral vectors encoding Doc2b<sup>WT</sup>, Syt1<sup>WT</sup>, Doc2b<sup>DN</sup>, Doc2b<sup>MID</sup>, Syt1<sup>C2AB</sup>, Syt1-Doc2b<sup>WT</sup>, or Syt1-Doc2b<sup>DN</sup> on DIV2. Throughout this paper, DN designates the D218,220N mutation in Doc2b. To control for effects caused by the lentiviral infection, we used particles encoding GFP alone. To facilitate a side-by-side comparison between all constructs, mEPSC frequencies were normalized to that of a control group included in all experiments. Recordings and analyses were performed while the identity of experimental groups was unknown to the experimenter.

**Network culture and electrophysiology.** Hippocampal neurons from WT, Syt1 knock-out, and Syt1/Doc2a/Doc2b TKO mice were prepared as previously described (Groffen et al., 2010). Shortly, neurons were plated at 50,000 per well on a confluent layer of glia in 12-well plates. The cells were infected with lentiviral vectors encoding GFP, Doc2b<sup>WT</sup>, Doc2b<sup>C2AB</sup>, Doc2a<sup>WT</sup>, Syt1<sup>WT</sup>, Syt1-Doc2b<sup>WT</sup>, or Syt1-Doc2b<sup>DN</sup> on DIV2. For whole-cell recordings at DIV10–17, the neurons were analyzed with the same internal and external media as in autapses, except that the bath contained 1  $\mu$ M tetrodotoxin (TTX) to block Na<sup>+</sup> currents (Ascent) and 20  $\mu$ M gabazine (GZ) to block GABA<sub>A</sub> receptor-mediated currents (Sigma). The experimenters were blind to the expressed constructs during the recordings and analyses.

**Immunocytochemistry.** To verify the expression and localization of the different constructs, mouse hippocampal neurons were cultured in networks at a density of 25,000/well and transduced at DIV2 with lentiviral particles (3  $\mu$ l/well). At DIV14, the cells were fixed with 4% PFA and permeabilized with 0.1% Triton X-100 in PBS for 5 min. Constructs containing the N-terminal domain of Syt1 were stained with a mouse monoclonal antibody raised against amino acids 1–12 of rat Syt1 (1:2000 in PBS supplemented with 10% normal goat serum; #105221, Synaptic Systems). This antibody does not react with endogenous mouse Syt1. Synapses were stained with rabbit polyclonal E028 against Synapsin 1/2 (1:1000; Hosaka and Südhof, 1999). All constructs containing the N-terminal domain of Doc2a/Doc2b were stained with rabbit polyclonal antibody (pAb) 13.2 (fraction 5, diluted 1:200), which was cross absorbed against Doc2a/Doc2b DKO mouse brain lysate to remove aspecific activity (Groffen et al., 2010). A mouse monoclonal against synaptobrevin-2 was used to identify synapses (1:2000, clone 69.1; Synaptic Systems). To stain the Doc2b<sup>C2AB</sup> construct, which lacks the immunogen used to raise pAb 13.2, a mouse pAb anti-Doc2b (H00008447, Abnova) was used at 1:200, combined with rabbit anti-Synapsin. After labeling by secondary antibodies conjugated to Alexa 546 and Alexa 647, coverslips were mounted in Mowiol 4–88 (Aldrich) and images were acquired on an A1R confocal laser microscope (Nikon) controlled by NIS-elements AR version 4.30 (Laboratory Imaging).

**Experimental design and statistical analysis.** Neurons were cultured from E18-stage embryos without sex determination, thus using both male and female mice. In experiments aimed at testing the effect of overexpressed proteins, untransfected cells from the same culture batch were used as an internal control. This is important to minimize confounding effects caused by weekly experimental variations: each week uses a different culture batch from a different mouse embryo, which is reflected by weekly variations in the average mEPSC frequency of the untransfected control cells. Details of all statistical tests are listed in Table 1.

## Results

To investigate whether Doc2 proteins underlie the high rate of spontaneous release in Syt1-deficient neurons, we generated TKO mice lacking Syt1 (Geppert et al., 1994), Doc2a (Sakaguchi et al., 1999), and Doc2b (Groffen et al., 2010). TKO mice

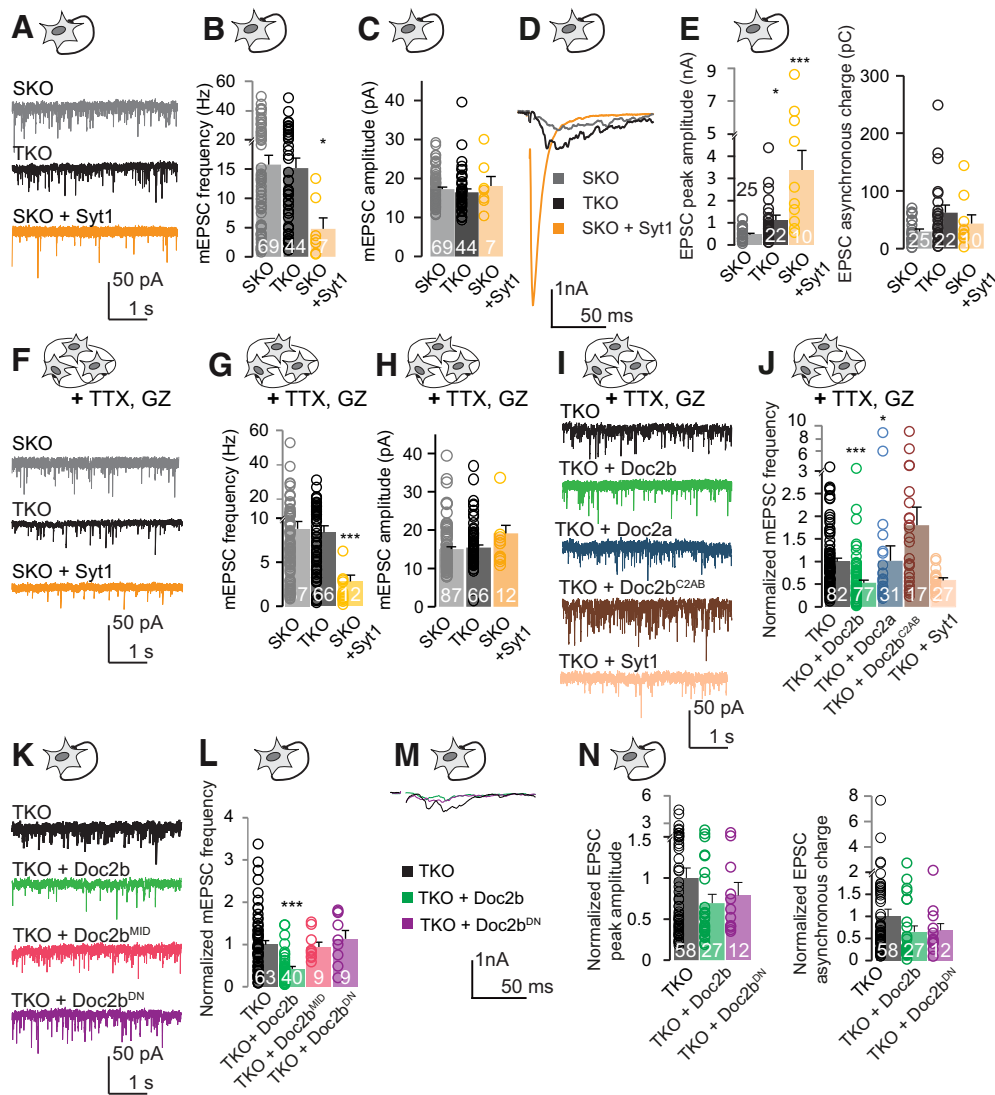
showed a similar severe phenotype as Syt1-null mice, characterized by normal embryonic development and death immediately following birth, associated with a lack of movement and respiration.

### Doc2a/Doc2b are not responsible for the high spontaneous release rate in Syt1-deficient neurons

TKO hippocampal neurons were cultured either on glial microislands to induce self-innervation (autapses; Fig. 2A–E) or in network cultures (Fig. 2F–H). As expected due to the absence of Syt1 (Geppert et al., 1994), TKO neurons showed no synchronous neurotransmission upon induction of a single action potential (Fig. 2D,E). In a direct comparison between neurons lacking Syt1 alone (SKO; *Syt1*<sup>-/-</sup>) or lacking all three sensors (TKO; *Syt1*<sup>-/-</sup>, *Doc2a*<sup>-/-</sup>, *Doc2b*<sup>-/-</sup>), the peak EPSC amplitude in TKO neurons was slightly higher than that of SKO neurons (1.1  $\pm$  0.22 and 0.47  $\pm$  0.06 nA, respectively;  $p$  = 0.0322, Kruskal–Wallis and Dunn's test; Fig. 2E, left). The deletion of Doc2 proteins did not reduce the asynchronous charge (SKO, 30  $\pm$  4.4 pC; TKO, 63  $\pm$  13 pC; SKO + Syt1 neurons, 44  $\pm$  15 pC;  $p$  = 0.0922, Kruskal–Wallis and Dunn's tests; Fig. 2E, right). Expression of Syt1 in SKO neurons caused much higher EPSC amplitudes (3.4  $\pm$  0.9 nA;  $p$  = 0.000063, Kruskal–Wallis and Dunn's tests). The mean spontaneous release frequency was not significantly reduced in the absence of Doc2 proteins (Fig. 2B). In previous studies, autaptic and continental Syt1-deficient neurons were shown to have different spontaneous release phenotypes (Liu et al., 2009). Therefore, we also used network cultures and recorded spontaneous mEPSC events in the presence of tetrodotoxin (TTX) and gabazine (GZ). Again, the genetic ablation of Doc2 proteins did not reduce the mini frequency (Fig. 2F,G). In both islands and networks, the expression of Syt1 in SKO neurons reduced the mEPSC frequency to normal levels, as expected (Fig. 2B,G;  $p$  = 0.019 and  $p$  = 0.000035, respectively, Dunn's tests). These results confirm earlier observations in an shRNA-based study (Pang et al., 2011), together indicating that the high mini frequency in Syt1 knock-out neurons is not attributable to Doc2a/Doc2b. Unexpectedly, we observed a decrease in the mEPSC 10–90% rise time upon expression of Syt1 in null neurons in networks ( $p$  = 0.01), but this observation was not made in autapses.

### Expression and subcellular distribution of recombinant proteins

In view of the similar SNARE-binding and phospholipid-binding activities of Syt1 and Doc2, we also investigated the effect of Doc2 overexpression on spontaneous release. In addition to wild-type Syt1, Doc2a, and Doc2b, we designed mutated and hybrid variants of these proteins (see Fig. 1 for an overview). To test the expression and localization of these constructs, neurons expressing them were stained with antibodies against Doc2 and synaptic markers (Fig. 3). Most Doc2 constructs were homogeneously distributed throughout the soma and neurite extensions. In addition, Doc2b<sup>DN</sup> was found at the plasma membrane. Despite the lack of a synaptic targeting signal, immunoreactivity was also detectable in synapses (Fig. 3, arrowheads). Likewise, all constructs containing the N-terminal domain of Syt1 were readily detectable by immunocytochemistry. As predicted by the trafficking signals in the N-terminal domain (Kang et al., 2004), and consistent with previous experiments (Xue et al., 2015), these constructs were effectively targeted to synapses (Fig. 4).



**Figure 2.** Doc2 proteins are not responsible for the high frequency of spontaneous release in SKO neurons, and the MID is necessary for an inhibitory role of Doc2b overexpressed in TKO neurons. **A–C**, Representative traces (**A**), average frequency (**B**), and amplitude (**C**) of spontaneous mEPSCs recorded from autaptic cultures of SKO, TKO, or SKO neurons overexpressing Syt1. **D, E**, Representative traces (**D**) and average peak amplitude (**E**, left) and asynchronous charge (**E**, right) of evoked EPSCs recorded from autaptic neurons with the same genotypes. **F–H**, Representative traces (**F**), average frequency (**G**), and amplitude (**H**) of spontaneous mEPSCs recorded from network cultures of SKO, TKO, or SKO neurons overexpressing Syt1. TTX (1  $\mu$ M) and GZ (20  $\mu$ M) were added to the extracellular solution to prevent network activity and isolate excitatory events. **I, J**, Representative traces (**I**) and average frequency (**J**) of spontaneous mEPSCs recorded from network cultures of TKO neurons overexpressing Doc2b, Doc2a, Doc2b<sup>C2AB</sup>, or Syt1 in the presence of 1  $\mu$ M TTX and 20  $\mu$ M GZ. **K, L**, Representative traces (**K**) and average frequency (**L**) of spontaneous mEPSCs recorded from autaptic cultures of TKO neurons overexpressing Doc2b, Doc2b<sup>MID</sup>, Doc2b<sup>DN</sup>, or Syt1. **M, N**, Representative traces recorded from TKO neurons overexpressing Doc2b or Doc2b<sup>DN</sup> (**M**) and average peak amplitude (**N**, left) and asynchronous charge (**N**, right) of evoked EPSCs. **J, L, N**, Values of cells expressing various constructs were normalized to the mean value of untransfected cells. Error bars indicate SEs. Cell numbers are indicated within bars. \* $p < 0.05$ , \*\*\* $p < 0.001$ .

### Doc2b overexpression inhibits spontaneous release in TKO neurons

Interestingly, Doc2b overexpression caused a twofold reduction of the mEPSC frequency in TKO islands (from 100% in untransfected cells to  $46 \pm 5\%$  in Doc2b<sup>WT</sup>-expressing cells;  $p = 0.000006$ , Kruskal–Wallis and Dunn’s tests; Fig. 2*K, L*, compare green and black). The data are also provided in Table 1. A similar reduction was observed in Doc2b-overexpressing networks (to  $50 \pm 5\%$  of controls;  $p = 0.000001$ , Kruskal–Wallis and Dunn’s tests; Fig. 2*I, J*, green versus black). This inhibitory effect was specific for Doc2b (Fig. 2*J*), even though Doc2a was present in synapses (Fig. 3). Although Dunn’s test reported a different data distribution for Doc2a ( $p = 0.015$ ), the mean

mEPSC frequency was identical to that of untransfected neurons, so we conclude that Doc2a did not inhibit spontaneous release. The inhibitory effect of Doc2b was accompanied by a slightly reduced asynchronous release component, but this effect was not significant (Fig. 2*M, N*, right, green versus black). Synchronous release remained absent (Fig. 2*M, N*, green versus black). Thus, in TKO neurons, overexpression of Doc2b inhibits spontaneous release.

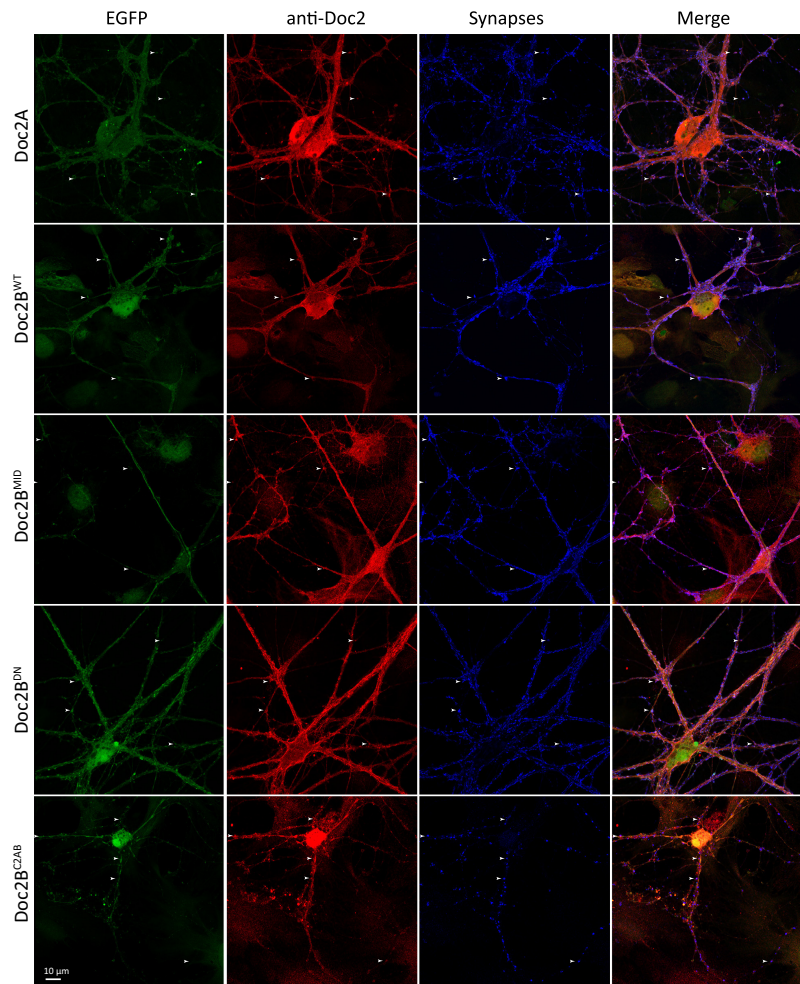
### Spontaneous release inhibition by Doc2b depends on its Munc13-interacting domain

In wild-type neurons, Doc2b overexpression did not inhibit spontaneous release, as reported previously (Groffen et al., 2010).

We hypothesized that Doc2, which is soluble in the cytoplasm, cannot compete efficiently with Syt1, which is anchored to the vesicle membrane of wild-type cells. In addition to  $\text{Ca}^{2+}$ , Munc13 can also target Doc2b to the plasma membrane by its N-terminal MID (Orita et al., 1997; Duncan et al., 1999; Friedrich et al., 2013; Houy et al., 2017). To test whether Doc2b-mediated inhibition of spontaneous release requires Munc13 binding, the N-terminal domain was either deleted or mutated. Upon deletion of the N-terminal domain, the C2AB fragment of Doc2b (Doc2b<sup>C2AB</sup>) lost its inhibitory effect on spontaneous release (Fig. 2J). Likewise, when the Munc13 interaction was blocked by a previously characterized mutation at amino acids 15–20 (Orita et al., 1997; Mochida et al., 1998; Groffen et al., 2004), full-length Doc2b lost its ability to inhibit spontaneous release (Fig. 2L, Doc2b<sup>MID</sup>). The subcellular distribution of the Doc2b<sup>MID</sup> mutant was not discernible from Doc2b<sup>WT</sup>, including immunoreactivity in synapses (Fig. 3). Thus, Munc13 interaction is not required for synaptic delivery of Doc2b. Instead, the mutation may affect the distribution of Doc2 or Munc13 within synaptic microdomains, or alter the functional activity of either protein. Finally, a mutation that mimics the effect of  $\text{Ca}^{2+}$  binding to the C2A domain (Groffen et al., 2004; Friedrich et al., 2008) did not inhibit spontaneous release (Fig. 2L, Doc2b<sup>DN</sup>). The asynchronous release component was not reduced by overexpression of this mutant either (Fig. 2N, right). Consistent with other studies, Doc2b<sup>DN</sup> was enriched at the plasma membrane of transfected neurons (Groffen et al., 2006; Xue et al., 2015; Houy et al., 2017; Bourgeois-Jaarsma et al., 2019). Together, we conclude that the inhibitory activity of Doc2b in TKO cells requires its Munc13 binding activity and is lost after neutralization of the  $\text{Ca}^{2+}$ -binding aspartates D218 and D220 in the C2A domain.

### Doc2b overexpression in SKO neurons

Similar to the aforementioned experiments in TKO cells, full-length Doc2a and Doc2b were also expressed in SKO neurons lacking only Syt1. As previously reported (Geppert et al., 1994), SKO autapses showed extremely small EPSC amplitudes (Fig. 5D,E), which were effectively rescued upon reintroduction of Syt1 (the amplitude increased 10-fold after Syt1 expression in SKO neurons, gray versus orange;  $p = 0.0008$ , Kruskal–Wallis and Dunn's tests). Overexpression of Doc2b failed to rescue the EPSC amplitude, which was almost identical to that of SKO neurons (gray versus yellow). Overexpression of Doc2b reduced the mini frequency in SKO networks to  $65 \pm 10\%$  (Fig. 5F,G;  $p = 0.0387$ , Kruskal–Wallis and Dunn's tests), but this effect was not observed in autapses, where Doc2b-overexpressing cells had a mini frequency of  $90 \pm 20\%$  relative

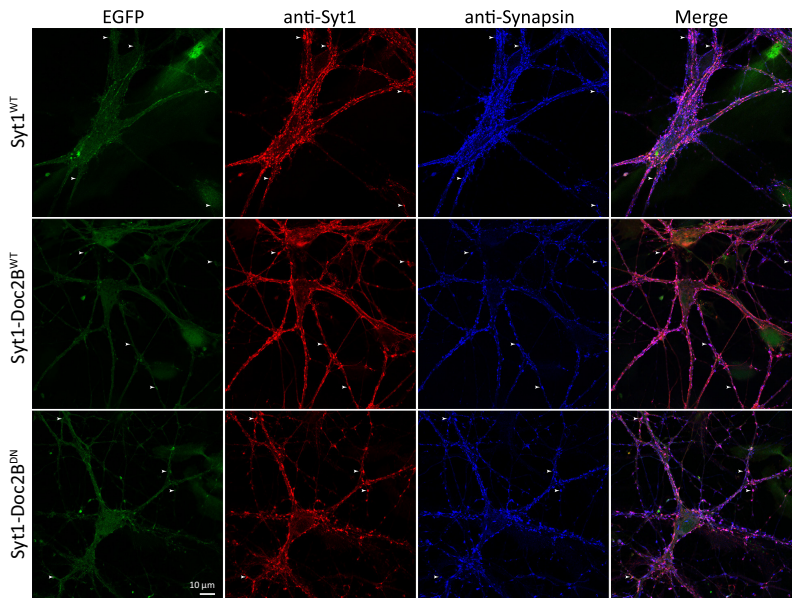


**Figure 3.** Overexpressed Doc2 constructs are widely distributed in mouse hippocampal neurons with detectable expression in synapses (arrowheads). Doc2a<sup>WT</sup>, Doc2b<sup>WT</sup>, Doc2b<sup>MID</sup>, Doc2b<sup>DN</sup>, or Doc2b<sup>C2AB</sup> was coexpressed with EGFP from an internal ribosomal entry site. Doc2a/Doc2b immunoreactivity was stained with rabbit polyclonal 13.2 combined with mouse-anti-synaptobrevin as a synaptic marker. Alternatively, Doc2b<sup>C2AB</sup>, which lacks the 13.2 epitope, was stained with mouse-anti-Doc2b (Abnova) combined with rabbit anti-synapsin-1/2 as the synaptic marker. Most Doc2 constructs are homogeneously distributed in the soma and neurites of transfected neurons, except Doc2b<sup>DN</sup>, which was enriched on the plasma membrane. When available, mouse neurons were derived from Doc2a/Doc2b DKO mice (Doc2b<sup>WT</sup>, Doc2b<sup>MID</sup>), but in some cases (other constructs) wild-type cells were used. Arrowheads indicate selected synapses to illustrate that Doc2 immunoreactivity is also detectable in synapses.

to untransfected controls (Fig. 5A,B, gray versus yellow;  $p = 0.7289$ , Kruskal–Wallis and Dunn's tests). Doc2a overexpression did not affect the spontaneous release frequency (Fig. 5F), although an increase in mEPSC amplitude was observed (Fig. 5H;  $p = 0.0475$ , Kruskal–Wallis and Dunn's tests). Finally, as expected, overexpression of wild-type Syt1 readily decreased the mini frequency in both networks (to  $30 \pm 8\%$  of controls; Fig. 5F;  $p = 0.0016$ , Kruskal–Wallis and Dunn's tests) and autaptic neurons (to  $22 \pm 7\%$  of controls; Fig. 5B;  $p = 0.0027$ , Kruskal–Wallis and Dunn's tests). Together, the results suggest that Doc2b does not support secretion at Syt1-dependent release sites: neither during the synchronous phase (measured as EPSCs) nor during rest (mEPSCs).

### The Syt1 N-terminal domain is essential for spontaneous release inhibition

Previous studies suggested that the transmembrane domain (TMD) is not essential for the fusogenic activity of Syt1 (Mar-



**Figure 4.** Overexpressed Syt1 and Syt1-Doc2b are targeted to synapses. Rat Syt1<sup>WT</sup>, Syt1-Doc2b<sup>WT</sup>, or Syt1-Doc2b<sup>DN</sup> was expressed in wild-type mouse hippocampal neurons. EGFP was expressed as a separate protein from an internal ribosomal entry site. After fixing at DIV14, the cells were stained with a monoclonal antibody specific for amino acids 1–12 of rat Syt1. This antibody does not react with mouse Syt1. Synapsin-1/2 was stained to mark synapses. Many synapses express recombinant Syt1; a few examples are highlighted by arrowheads.

tens et al., 2007; Hui et al., 2009; Wang et al., 2011), although it may contribute to vesicle docking (de Wit et al., 2009) and recycling (Yao et al., 2011b). Interestingly, overexpression of the C2AB fragment of Syt1 (Syt1<sup>C2AB</sup>) in autaptic neurons had no significant effect on the spontaneous release rate ( $80 \pm 10\%$  of control; Fig. 5B;  $p > 0.9999$ , Kruskal–Wallis and Dunn’s tests). Thus, the TMD is essential for spontaneous release inhibition by Syt1.

#### The C2AB fragment of Syt1 synchronizes fusion during sustained activity

Overexpression of Syt1<sup>C2AB</sup> in Syt1-null neurons had a minor effect on the EPSC amplitude (1.7-fold increase in SKO + Syt1<sup>C2AB</sup>; Fig. 5E;  $p = 0.1640$ , Kruskal–Wallis and Dunn’s test). During trains of 100 action potentials at 5 Hz, 50% of the neurons overexpressing Syt1<sup>C2AB</sup> showed a remarkable facilitation of the EPSC amplitude, increasing from  $0.8 \pm 0.2$  nA (first AP) to  $2.5 \pm 0.4$  nA (10th AP; Fig. 5I–K;  $p = 0.000244$ , Wilcoxon matched-pairs signed rank test), thereby reaching similar amplitudes as SKO neurons overexpressing full-length Syt1 (increasing from  $2.7 \pm 0.6$  nA after the first AP to  $3.1 \pm 0.4$  nA after the 10th AP;  $p = 0.9860$ , Kruskal–Wallis and Dunn’s test). A small increase was observed in Syt1-null neurons (first pulse,  $0.66 \pm 0.18$  nA; 10th pulse,  $0.9 \pm 0.18$  nA; Fig. 5J, left;  $p = 0.021545$ , Wilcoxon matched-pairs signed rank test); however, the amplitude of Syt1-null neurons did not reach similar levels as Syt1-null neurons overexpressing full-length Syt1 ( $p = 0.0006$ , Kruskal–Wallis and Dunn’s test). The effect of Syt1<sup>C2AB</sup> was accompanied by a reduction of the 10–90% rise time of the EPSCs that matched the 10–90% rise time of the 10th pulse of Syt1-null neurons overexpressing full-length Syt1 (SKO + Syt1<sup>C2AB</sup>, first pulse:  $9.12 \pm 3.12$  ms, 10th pulse:  $2.43 \pm 0.16$  ms; SKO + Syt1<sup>WT</sup>, first pulse:  $2.14 \pm 0.21$  ms, 10th pulse:  $2.62 \pm 0.3$  ms; Fig. 5J, right;  $p = 0.390805$ , Kruskal–Wallis and Dunn’s test). In contrast, SKO neurons did not show a reduction of the 10–90% EPSC rise time

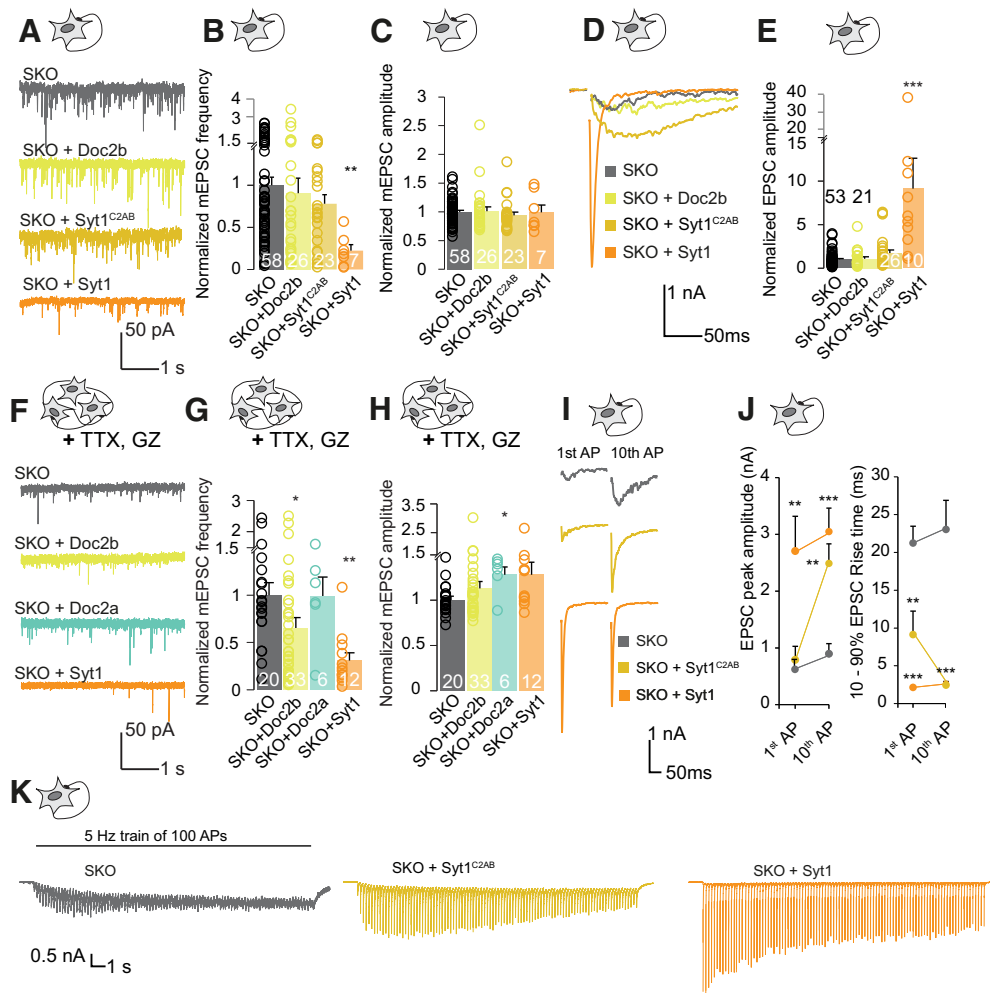
upon synaptic activity (first pulse,  $20.9 \pm 2.6$  ms; 10th pulse,  $21.2 \pm 3.54$  ms;  $p > 0.9999$ , Wilcoxon matched-pairs signed rank test), indicating that the release remained asynchronous. We conclude that overexpression of the C2AB fragment of Syt1 is sufficient for driving synchronous release in a subset of neurons, and that this effect depends on synaptic activity.

#### Intrinsic differences in fusion enhancement by Doc2b<sup>C2AB</sup> and Syt1<sup>C2AB</sup>

Given the high degree of biochemical similarity between the Syt1<sup>C2AB</sup> and Doc2b<sup>C2AB</sup> fragment, we next investigated whether a hybrid construct, composed of the N-terminal domain of Syt1 and the C2AB domain of Doc2b, substitutes for wild-type Syt1 in synchronous neurotransmitter release. In hippocampal autapses, the high spontaneous release rates of Syt1-null neurons remained high after expression of the hybrid construct, and the mEPSC rise time was reduced compared with that of Syt1-null neurons (Fig. 6A, B;  $p = 0.01$ , Kruskal–Wallis and Dunn’s test). A 1.7-fold increase in mEPSC frequency was observed in cells expressing the Syt1-Doc2b<sup>DN</sup> hybrid ( $p = 0.01054$ , Kruskal–Wallis and Dunn’s test). Moreover, an increase in the mEPSC amplitude was observed in this group ( $p = 0.0035$ , Kruskal–Wallis and Dunn’s test). Synchronous neurotransmission was not rescued by the Syt1-Doc2b hybrid, nor by the mutant variant Syt1-Doc2b<sup>DN</sup> (Fig. 6D, E), even though the hybrid constructs were targeted to synapses (Fig. 4). In network cultures, Syt1-Doc2b<sup>DN</sup> expression did not significantly change the mEPSC frequency and amplitude (Fig. 6G;  $p = 0.2614$  and  $0.7487$ , respectively, Kruskal–Wallis and Dunn’s test). Together, we conclude that the C2AB domains of Syt1 and Doc2b are not functionally interchangeable.

#### Discussion

Previous studies reported a strong increase in the frequency of spontaneous neurotransmitter release events in neurons lacking Syt1 (Littleton et al., 1993; Broadie et al., 1994; Xu et al., 2009). To explain this phenomenon, it has been hypothesized that Doc2 proteins occupy vacant Syt1-dependent release sites, increasing their apparent  $\text{Ca}^{2+}$  sensitivity and thereby causing a higher release probability at low resting  $\text{Ca}^{2+}$  concentrations (Walter et al., 2011). Here, we tested this hypothesis by triple homologous gene targeting of Syt1, Doc2a, and Doc2b in mouse neurons. We conclude that the high spontaneous release rate in Syt1 cells does not require Doc2a/Doc2b activity. This finding is consistent with earlier knock-down experiments (Pang et al., 2011). Our data show that the functional differences between Doc2b and Syt1 are not limited to their different N-terminal domains but are also encoded in the C2AB domain. In particular, the C2AB domain of Doc2b cannot functionally replace that of Syt1, as demonstrated by the inability of hybrid Syt1-Doc2b to support synchronous fusion. In a previous study where pre-synaptic neurons were locally stimulated in a network configuration, similar Syt1-Doc2b chimeras also did not support synchronous fusion (Xue et al., 2015). Interestingly, this study



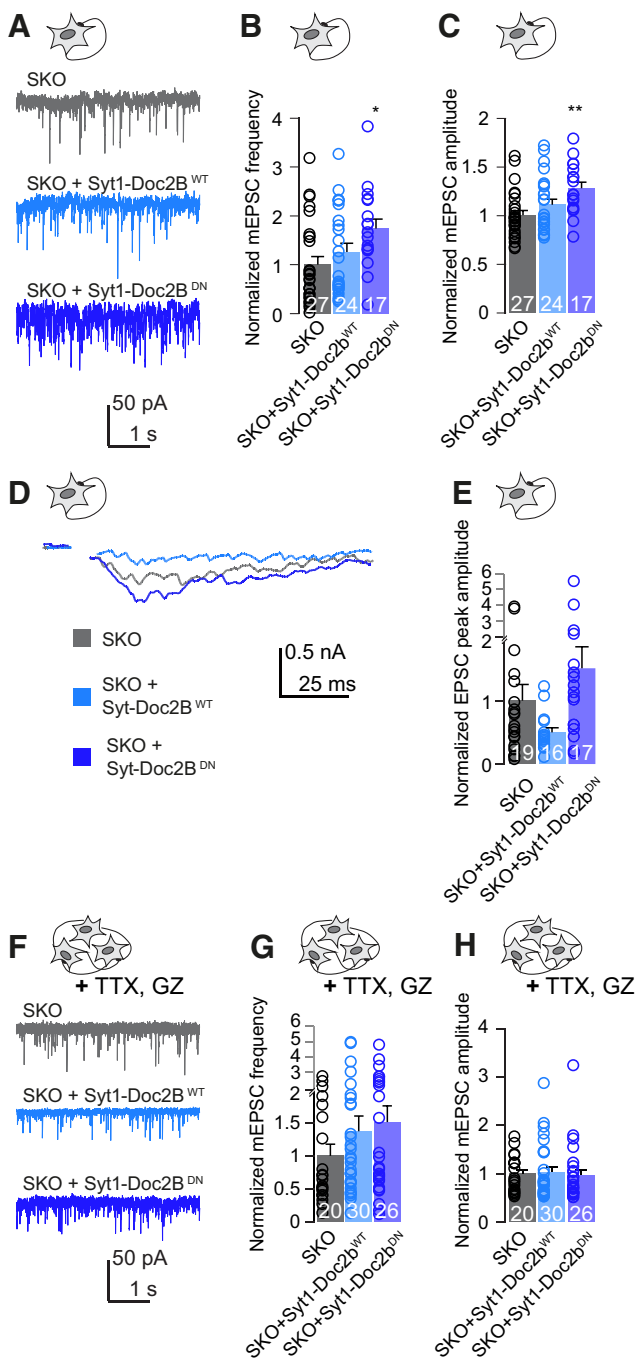
**Figure 5.** The transmembrane domain of Syt1 is necessary for its inhibitory role on mEPSCs in hippocampal neurons. To control for variation between independent experiments, mEPSC frequencies and amplitudes were normalized to the mean value of untransfected neurons measured in the same experiment. **A–C**, Representative traces (**A**), average frequency (**B**), and amplitude (**C**) of spontaneous mEPSCs recorded from Syt1 SKO neurons in autaptic culture, untransfected or overexpressing Doc2b, Syt1<sup>C2AB</sup>, or Syt1. **D, E**, Representative traces (**D**) and average peak amplitude (**E**) of evoked EPSCs recorded from autaptic SKO neurons, untransfected or overexpressing Doc2b, Syt1<sup>C2AB</sup>, or Syt1. **F–H**, Representative traces (**F**), average frequency (**G**), and amplitude (**H**) of spontaneous mEPSCs recorded from network cultures of SKO neurons, untransfected or overexpressing Doc2b, Doc2a, or Syt1. To prevent network activity and isolate excitatory events, TTX (1  $\mu$ M) and GZ (20  $\mu$ M) were added to the extracellular solution. **I**, Representative traces of the first and the 10th evoked EPSCs from a train of 100 APs elicited at 5 Hz, recorded from autaptic SKO neurons, untransfected or overexpressing Syt1<sup>C2AB</sup> or Syt1. **J**, Average peak amplitude (left) and 10–90% rise time (right) of the first and 10th evoked EPSC from the train ( $n = 15$  untransfected, 13 Syt1<sup>C2AB</sup>-expressing and 7 Syt1-expressing cells). **K**, Representative traces showing the full EPSC recording throughout the 5 Hz train. Error bars indicate SEs. Cell numbers are indicated within bars. \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ .

also demonstrated that the C2B domain of Doc2b exhibits slower kinetics than Syt1. Thus, differences in both the kinetics and affinity of  $\text{Ca}^{2+}$  and phospholipid binding likely contribute to the divergent C2AB domain activity.

It is conceivable that different sensors have evolved to mediate specialized  $\text{Ca}^{2+}$ -secretion coupling processes in different subcellular compartments. Moreover, different types of neurons may bear different sets of presynaptic proteins to regulate vesicle release. In line with this, a recent study has shown that Doc2a and Doc2b are differentially expressed in the brain, facilitating glutamatergic and GABAergic spontaneous release, respectively (Courtney et al., 2018). Upon overexpression, however, Doc2b restored the spontaneous release frequency in Doc2a-deficient glutamatergic cells and vice versa, suggesting functional redundancy (Courtney et al., 2018). Interestingly, our data now indicate a functional difference between Doc2a and Doc2b: Doc2b inhibited the spontaneous release rate, whereas Doc2a did not. This observation was made in both TKO and SKO networks (Figs. 2I, J, 5F, G).

The inhibitory effect of Doc2b was evident in SKO network cultures, but not autapses. As a key difference, networks are mixed cultures that contain GABAergic cells. The long-term spontaneous release of GABA can regulate mEPSC frequencies of glutamatergic cells in a mixed culture (Wierda and Sørensen, 2014). In several studies, this effect has presumably masked the high mEPSC frequency of Syt1-deficient neurons (Liu et al., 2009; Wierda and Sørensen, 2014). In a similar manner, it is possible that the inhibitory effect of Doc2b is an indirect consequence of increased spontaneous GABA release. Because we included gabazine to block GABA<sub>A</sub> receptors, this effect would have to be mediated by metabotropic GABA<sub>B</sub> receptors. Interestingly, mixed culture effects may also help explain why experiments in locally stimulated networks suggest a role for Doc2b in asynchronous release (Yao et al., 2011a), whereas autapse experiments do not (Groffen et al., 2010).

At physiological extracellular  $\text{Ca}^{2+}$  concentrations ( $[\text{Ca}^{2+}]_0$ ), both Syt1 and Doc2b are required for driving spontaneous release in inhibitory neurons, whereas Syt1 is dispensable for facilitating



**Figure 6.** The C2AB domain of Doc2b is not capable of replacing that of Syt1 in synchronous neurotransmitter secretion. To control for variation between independent experiments, mEPSC frequencies and amplitudes were normalized to the mean value of untransfected neurons measured in the same experiment. **A–C**, Representative traces (**A**), average frequency (**B**), and amplitude (**C**) of spontaneous mEPSCs recorded from autaptic cultures of SKO neurons overexpressing Syt1-Doc2b<sup>WT</sup> or Syt1-Doc2b<sup>DN</sup>. **D, E**, Representative traces (**D**) and average peak amplitude (**E**) of evoked EPSCs recorded from autaptic SKO neurons overexpressing Syt1-Doc2b<sup>WT</sup> or Syt1-Doc2b<sup>DN</sup>. **F–H**, Representative traces (**F**), average frequency (**G**), and amplitude (**H**) of spontaneous mEPSCs recorded from network hippocampal cultures of SKO overexpressing Syt1-Doc2b<sup>WT</sup> or Syt1-Doc2b<sup>DN</sup>. TTX (1  $\mu$ M) and GZ (20  $\mu$ M) were added to the extracellular solution to prevent network activity and isolate excitatory events. Error bars indicate SE. Cell numbers are indicated in bar graphs. \* $p < 0.05$ , \*\* $p < 0.01$ .

minis in glutamatergic neurons, an action that depended on Doc2a alone (Courtney et al., 2018). Thus, it is possible that the promoting role of Syt1 on spontaneous release described by Xu et al. (2009) is restricted to higher  $[Ca^{2+}]_0$ . Our results support a

multisensor model where specialized  $Ca^{2+}$  sensors act together to determine the  $Ca^{2+}$ -dependent synaptic release rate, each contributing to a distinct kinetic component of the overall release in hippocampal synapses. This model is compatible with the previously described two-sensor model (Goda and Stevens, 1994), which already noted that the “second sensor” could represent a collection of molecules. We suggest that at least three classes of sensors underlie synaptic secretion. The first class of sensors (Syt1, Syt2, or Syt9) promotes fast membrane fusion during action potential-evoked  $Ca^{2+}$  transients but does not support secretory events at resting  $Ca^{2+}$ , at least not in excitatory neurons (Courtney et al., 2018). A second class of sensors is responsible for the “asynchronous” release of neurotransmitters evoked by action potentials. Syt7 was shown to belong to this class (Bacaj et al., 2013; Luo and Südhof, 2017); however, it is not responsible for increased spontaneous release in Syt1-deficient neurons. A third class, including Doc2b, does not support synchronous fusion but promotes secretory events in the absence of action potentials. Doc2 proteins have also been proposed to play a role in asynchronous release (Yao et al., 2011a), although this finding was not reproducible in our hands, possibly due to different experimental conditions. A recent study also showed a novel role for Doc2 proteins as part of the  $Ca^{2+}$  sensor machinery that drives augmentation (Xue et al., 2018). Finally, the mechanism of spontaneous release in TKO neurons remains to be identified. It is possible that this class comprises a fourth class of  $Ca^{2+}$ -sensing molecules, but alternatively, sensorless fusion may occur. The likelihood of spontaneous release events is regulated by extracellular  $Ca^{2+}$  concentrations (Xu et al., 2009; Smith et al., 2012), the stochastic opening of voltage-dependent channels (Ermolyuk et al., 2013), or intracellular  $Ca^{2+}$  channels (Llano et al., 2000; Emptage et al., 2001).

In wild-type neurons, Doc2b positively contributes to spontaneous release (Groffen et al., 2010), and its overexpression does not inhibit the mini frequency, suggesting that endogenous Doc2b is not rate limiting. Overexpression of Doc2b in the absence of Syt1, however, revealed a negative contribution to spontaneous release. The inhibitory activity of Doc2b required an intact N-terminal domain and was blocked by mutations in the Munc13-interacting domain. In chromaffin cells, Doc2b also has mixed positive and inhibitory effects on regulated secretion, both of which are dependent on the MID domain (Houy et al., 2017). The domain likely contributes to local enrichment of Doc2b at release sites, even though the Doc2b<sup>MID</sup> mutant is expressed in synaptic structures. The MID domains of Doc2a and Doc2b are highly conserved, and both isoforms bind to Munc13 (Orita et al., 1997; Duncan et al., 1999). Local accumulation also appears to be relevant for Syt1-dependent inhibition of spontaneous release, because this activity is abolished by deletion of the transmembrane domain. Thus, in general,  $Ca^{2+}$  sensors may inhibit spontaneous release by binding to release sites under conditions in which the C2AB domain is not activated. These conditions may include low intracellular  $Ca^{2+}$  concentrations, sensor protein mistargeting, or mutations that block fusogenic activity. In line with this model, minimal fragments comprising the C2AB domain of Syt1 and Doc2b are sufficient to drive membrane fusion in reconstituted SNARE liposomes (Martens et al., 2007; Hui et al., 2009; Groffen et al., 2010; Wang et al., 2011). The C2AB fragment of Doc2b has also been shown to interact with opposing membranes, directly participating in  $Ca^{2+}$ -dependent and phosphatidylserine-dependent stalk formation and hemifusion in a SNARE-free assay (Brouwer et al., 2015).



**Table 1. Summary of data and statistical analyses**

	Parameter	df	Test/H		df	Test/H		df	Test/H
<b>Figure 2</b>									
Genotype autaptic cells ( <i>n</i> )	mEPSC frequency (Hz)		Kruskal–Wallis	mEPSC amplitude (pA)		Kruskal–Wallis	mEPSC rise time (ms)		Kruskal–Wallis
SKO (control; 69)	15.7 ± 1.6			17.3 ± 0.5			1.8 ± 0.05		
TKO (44)	15.2 ± 1.7 ( <i>p</i> > 0.9999)	2	7.032	16.5 ± 0.8 ( <i>p</i> = 0.3836)	2	1.920	1.9 ± 0.1 ( <i>p</i> > 0.9999)	2	7.750
SKO + Syt1 (7)	4.8 ± 1.9* ( <i>p</i> = 0.0190)	2	7.032	18.1 ± 2.4 ( <i>p</i> > 0.9999)	2	1.920	1.4 ± 0.2 ( <i>p</i> = 0.0232)	2	7.750
Genotype continental cells ( <i>n</i> )	mEPSC frequency (Hz)		Kruskal–Wallis	mEPSC amplitude (pA)		Kruskal–Wallis	mEPSC rise time (ms)		Kruskal–Wallis
SKO (control; 87)	8.8 ± 0.8			15 ± 0.6			1.9 ± 0.1		
TKO (66)	8.4 ± 0.7 ( <i>p</i> = 0.9584)	2	21.48	15.4 ± 0.7 ( <i>p</i> = 0.2187)	2	3.150	1.7 ± 0.1 ( <i>p</i> = 0.3322)	2	8.457
SKO + Syt (12)	2.8 ± 0.7*** ( <i>p</i> = 0.000035)	2	21.48	19.2 ± 2.1 ( <i>p</i> = 0.5328)	2	3.150	1.6 ± 0.1* ( <i>p</i> = 0.0100)	2	8.457
Genotype autaptic cells ( <i>n</i> )	EPSC size (pA)		Kruskal–Wallis	EPSC async. charge (pC)		Kruskal–Wallis			
SKO (control; 25)	466.1 ± 64.3			30 ± 4					
TKO (22)	1128 ± 222.6* ( <i>p</i> = 0.0322)	2	18.22	63 ± 13 ( <i>p</i> = 0.0922)	2	4.053			
SKO + Syt1 (10)	3376 ± 887*** ( <i>p</i> = 0.000063)	2	18.22	44 ± 15 ( <i>p</i> > 0.9999)	2	4.053			
Genotype autaptic cells ( <i>n</i> )	Normalized mEPSC frequency		Kruskal–Wallis	Normalized mEPSC amplitude		Kruskal–Wallis	Normalized mEPSC rise time		Kruskal–Wallis
TKO (control) (63)	1 ± 0.1			1 ± 0.03			1 ± 0.03		
TKO + Doc2b (40)	0.46 ± 0.07*** ( <i>p</i> = 0.000006)	3	28.41	0.8 ± 0.03 ( <i>p</i> = 0.167)	3	8.527	1 ± 0.05 ( <i>p</i> > 0.9999)	3	2.485
TKO + Doc2b <sup>MD</sup> (9)	0.9 ± 0.1 ( <i>p</i> > 0.9999)	3	28.41	0.9 ± 0.1 ( <i>p</i> = 0.4114)	3	8.527	0.9 ± 0.06 ( <i>p</i> > 0.9999)	3	2.485
TKO + Doc2b <sup>DN</sup> (9)	1.1 ± 0.2 ( <i>p</i> > 0.9999)	3	28.41	0.9 ± 0.05 ( <i>p</i> > 0.9999)	3	8.527	1.2 ± 0.2 ( <i>p</i> = 0.5603)	3	2.485
Genotype continental cells ( <i>n</i> )	Normalized mEPSC frequency		Kruskal–Wallis						
TKO (control; 82)	1 ± 0.07								
TKO + Doc2b (77)	0.5 ± 0.06*** ( <i>p</i> < 0.000001)	4	44.96						
TKO + Doc2a (31)	1 ± 0.3* ( <i>p</i> = 0.0153)	4	44.96						
TKO + Doc2b <sup>2AB</sup> (17)	1.8 ± 0.4 ( <i>p</i> = 0.9818)	4	44.96						
TKO + Syt1 (27)	0.6 ± 0.06 ( <i>p</i> = 0.1566)	4	44.96						
Genotype autaptic cells ( <i>n</i> )	Normalized EPSC peak amplitude		Kruskal–Wallis	Normalized EPSC asynchronous charge		Kruskal–Wallis			
TKO (control; 58)	1 ± 0.1			1 ± 0.16					
TKO + Doc2b (27)	0.6 ± 0.1 ( <i>p</i> = 0.1462)	2	3.3323	0.65 ± 0.13 ( <i>p</i> = 0.0684)	2	4.487			
TKO + Doc2b <sup>DN</sup> (12)	0.8 ± 0.1 ( <i>p</i> > 0.9999)	2	3.3323	0.7 ± 0.15 ( <i>p</i> > 0.9999)	2	4.487			
<b>Figure 5</b>									
Genotype autaptic cells ( <i>n</i> )	Normalized mEPSC frequency		Kruskal–Wallis	Normalized mEPSC amplitude		Kruskal–Wallis	Normalized mEPSC rise time		Kruskal–Wallis
SKO (control; 58)	1 ± 0.1			1 ± 0.03			1 ± 0.03		
SKO + Doc2b (26)	0.9 ± 0.2 ( <i>p</i> = 0.7289)	3	11.35	1 ± 0.07 ( <i>p</i> > 0.9999)	3	3.788	0.9 ± 0.05 ( <i>p</i> = 0.2735)	3	3.776
SKO + Syt <sup>2AB</sup> (26)	0.8 ± 0.1 ( <i>p</i> > 0.9999)	3	11.35	0.9 ± 0.05 ( <i>p</i> = 0.1824)	3	3.788	0.9 ± 0.04 ( <i>p</i> > 0.9999)	3	3.776
SKO + Syt1 (7)	0.22 ± 0.07** ( <i>p</i> = 0.0027)	3	11.35	0.9 ± 0.12 ( <i>p</i> > 0.9999)	3	3.788	0.9 ± 0.13 ( <i>p</i> = 0.6479)	3	3.776
Genotype continental cells ( <i>n</i> )	Normalized mEPSC frequency		Kruskal–Wallis	Normalized mEPSC amplitude		Kruskal–Wallis	Normalized mEPSC rise time		Kruskal–Wallis
SKO (control; 20)	1 ± 0.13			1 ± 0.04			1 ± 0.05		
SKO + Doc2b (33)	0.65 ± 0.1* ( <i>p</i> = 0.0387)	3	15.21	1.1 ± 0.06 ( <i>p</i> > 0.9999)	3	8.058	1.13 ± 0.05 ( <i>p</i> = 0.4392)	3	14.09
SKO + Doc2a (6)	0.1 ± 0.2 ( <i>p</i> > 0.9999)	3	15.21	1.3 ± 0.08* ( <i>p</i> = 0.0475)	3	8.058	0.8 ± 0.06 ( <i>p</i> = 0.5051)	3	14.09
SKO + Syt1 (12)	0.3 ± 0.08** ( <i>p</i> = 0.0016)	3	15.21	1.3 ± 0.13 ( <i>p</i> = 0.1388)	3	8.058	0.7 ± 0.06 ( <i>p</i> = 0.1324)	3	14.09
Genotype autaptic cells ( <i>n</i> )	Normalized EPSC size		Kruskal–Wallis						
SKO (control; 53)	1 ± 0.11								
SKO + Doc2b (21)	1.1 ± 0.22 ( <i>p</i> > 0.9999)	3	15.57						
SKO + Syt <sup>2AB</sup> (26)	1.74 ± 0.35 ( <i>p</i> = 0.1640)	3	15.57						
SKO + Syt1 (10)	10.8 ± 5.7*** ( <i>p</i> = 0.0008)	3	15.57						
Genotype autaptic cells ( <i>n</i> )	EPSC size first pulse (pA)		Kruskal–Wallis	EPSC size 10th pulse (pA)		Kruskal–Wallis	EPSC size first versus 10th (paired measurements)		
SKO (control) (15)	628.2 ± 180.7			909.3 ± 183			SKO* ( <i>p</i> = 0.021545)		Wilcoxon matched-pairs signed rank test <sup>a</sup>
SKO + Syt <sup>2AB</sup> (13)	790.2 ± 237.3 ( <i>p</i> > 0.9999)	2	11.79	2490.4 ± 373.4** ( <i>p</i> = 0.0032)	2	16.69	SKO + Syt <sup>2AB</sup> *** ( <i>p</i> = 0.000244)		Wilcoxon matched-pairs signed rank test <sup>a</sup>

(Table continues)

Table 1. Continued

	Parameter	df	Test/H		df	Test/H		df	Test/H
SKO + Syt1 (7)	2707 ± 610** ( <i>p</i> = 0.0022)	2	11.79	3098 ± 449.4*** (vs control) ( <i>p</i> = 0.0006); n.s. (vs SKO + Syt <sup>C2AB</sup> ) ( <i>p</i> = 0.9860)	2	16.69	SKO + Syt ( <i>p</i> = 0.2188)		Wilcoxon matched-pairs signed rank test <sup>a</sup>
Genotype autaptic cells ( <i>n</i> )	EPSC first pulse 10–90% rise time (ms)		Kruskal–Wallis	EPSC 10th pulse 10–90% rise time (ms)		Kruskal–Wallis	Rise time first vs 10th pulse (paired measurements)		
SKO (control) (15)	21.2 ± 2.2			23 ± 3.8			SKO ( <i>p</i> = 0.9999)		Wilcoxon matched-pairs signed rank test <sup>a</sup>
SKO + Syt <sup>C2AB</sup> (13)	9.1 ± 3.1** ( <i>p</i> = 0.00483)	2	23.65	2.4 ± 0.2*** ( <i>p</i> = 0.000002)	2	31.17	SKO + Syt <sup>C2AB</sup> ( <i>p</i> = 0.108154)		Wilcoxon matched-pairs signed rank test <sup>a</sup>
SKO + Syt1 (7)	2.1 ± 0.2*** vs control ( <i>p</i> = 0.000011); n.s. vs SKO + Syt <sup>C2AB</sup> ( <i>p</i> = 0.390805)	2	23.65	2.6 ± 0.3*** vs control ( <i>p</i> = 0.000026); n.s. vs SKO + Syt <sup>C2AB</sup> ( <i>p</i> > 0.999999)	2	31.17	SKO + Syt1* ( <i>p</i> = 0.0420)		Wilcoxon matched-pairs signed rank test <sup>a</sup>
Figure 6									
Genotype autaptic cells ( <i>n</i> )	Normalized mEPSC frequency		Kruskal–Wallis	Normalized mEPSC amplitude		Kruskal–Wallis	Normalized mEPSC rise time		Kruskal–Wallis
SKO (control; 27)	1 ± 0.16			1 ± 0.04			1 ± 0.03		1 ± 0.04
SKO + Syt1-Doc2b <sup>WT</sup> (24)	1.2 ± 0.17 ( <i>p</i> = 0.5254)	2	7.790	1.1 ± 0.05 ( <i>p</i> = 0.4791)	2	9.855	0.8 ± 0.04** ( <i>p</i> = 0.01)	2	12.99
SKO + Syt1-Doc2b <sup>DN</sup> (17)	1.7 ± 0.2* ( <i>p</i> = 0.01054)	2	7.790	1.2 ± 0.06** ( <i>p</i> = 0.0035)	2	9.855	1.1 ± 0.05 ( <i>p</i> = 0.7952)	2	12.99
Genotype continental cells ( <i>n</i> )	Normalized mEPSC frequency		Kruskal–Wallis	Normalized mEPSC amplitude		Kruskal–Wallis	Normalized mEPSC rise time		Kruskal–Wallis
SKO (control; 20)	1 ± 0.17			1 ± 0.08			1 ± 0.08		1 ± 0.07
SKO + Syt1-Doc2b <sup>WT</sup> (30)	1.4 ± 0.2 ( <i>p</i> = 0.4809)	2	2.415	1 ± 0.1 ( <i>p</i> > 0.9999)	2	0.7947	1.18 ± 0.07 ( <i>p</i> = 0.2751)	5.509	1.2 ± 0.1 ( <i>p</i> = 0.1080)
SKO + Syt1-Doc2b <sup>DN</sup> (26)	1.5 ± 0.2 ( <i>p</i> = 0.2614)	2	2.415	0.9 ± 0.1 ( <i>p</i> = 0.7487)	2	0.7947	1.24 ± 0.07 ( <i>p</i> = 0.0383)	5.509	1.1 ± 0.05 ( <i>p</i> = 0.3633)
Genotype autaptic cells ( <i>n</i> )	Normalized EPSC size		Kruskal–Wallis						
SKO (control) (19)	1 ± 0.25								
SKO + Syt1-Doc2b <sup>WT</sup> (16)	0.5 ± 0.07 ( <i>p</i> = 0.3221)	2	9.055						
SKO + Syt1-Doc2b <sup>DN</sup> (15)	1.5 ± 0.34 ( <i>p</i> = 0.1758)	2	9.055						

Values of *p* correspond to Dunn's adjusted *p* values, except for paired tests where the Wilcoxon matched-pairs signed rank test was applied. \**p* < 0.05, \*\**p* < 0.01, \*\*\**p* < 0.001, n.s., not significant.

<sup>a</sup>Wilcoxon matched-pairs signed rank test was applied.

In most cases, the overexpressed constructs did not alter the mEPSC amplitude, which is primarily determined by postsynaptic processes. However, in SKO neurons expressing Doc2a, Syt1, or Syt1-Doc2b<sup>DN</sup>, a small but significant increase of the mEPSC amplitude was observed. In another study, Doc2a expressed in wild-type neurons did not affect the mEPSC amplitude (Yao et al., 2011a). Syt1 expressed in SKO neurons also did not affect it (Tagliatti et al., 2020). Consistent with our data, mEPSC amplitude was unchanged by Syt1-Doc2b<sup>WT</sup> expressed in SKO neurons (Xue et al., 2015). Overall, we are reluctant to draw strong conclusions from the small effects on mEPSC amplitude.

Previous work also indicated that the C2AB domain of Syt1 is sufficient for eliciting synchronous release when targeted to the plasma membrane using GAP-43 (Yao et al., 2011b). In this work, we show that the soluble C2AB fragment of Syt1 is capable of driving synchronous release in a subset of Syt1-null neurons, and that this action appears to depend on synaptic activity and concurrent Ca<sup>2+</sup> accumulation at the synaptic terminal. Recent work has characterized a role for Syt1 in tethering synaptic vesicles close to the active zone at rest (Chang et al., 2018). This action does not involve its Ca<sup>2+</sup>-binding regions but requires the polybasic side (K324–K327) and bottom residues (R398, R399) of the C2B domain, as well as an (at least partially) assembled SNARE complex. Upon AP induction and Ca<sup>2+</sup> entry, the authors observed an increase in synaptic vesicle density at the active zone, an effect that depended on Ca<sup>2+</sup> sensing by Syt1. We hypothesize that local accumulation of the soluble C2AB fragment of Syt1 at the active zone by a classical C2 activation mechanism could explain the partial rescue of synchronous release. The levels of soluble C2AB fragments present at the active zone are likely to vary among synapses, giving rise to differences in the

extent to which synchronous release is restored. In line with the aforementioned study, we observed a progressive increase in EPSC size as well as resynchronization of release upon synaptic activity, consistent with a Ca<sup>2+</sup>-dependent action of the C2AB fragment in rapidly recruiting synaptic vesicles to the active zone and catalyzing fast release by bringing the synaptic vesicle membrane and plasma membrane closer together. Thus, we conclude that the transmembrane domain is crucial for reliable triggering of synchronous release, whereas the C2AB fragment of Syt1 alone requires membrane enrichment to carry out its Ca<sup>2+</sup>-dependent docking and bridging actions that trigger synchronous release.

In summary, we conclude that, despite their similarity, the C2AB fragments of Doc2b and Syt1 have divergent functions in driving spontaneous and synchronous neurotransmitter release, and that the high quantal release frequency in Syt1-null neurons relies on mechanisms independent of Doc2a/Doc2b.

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