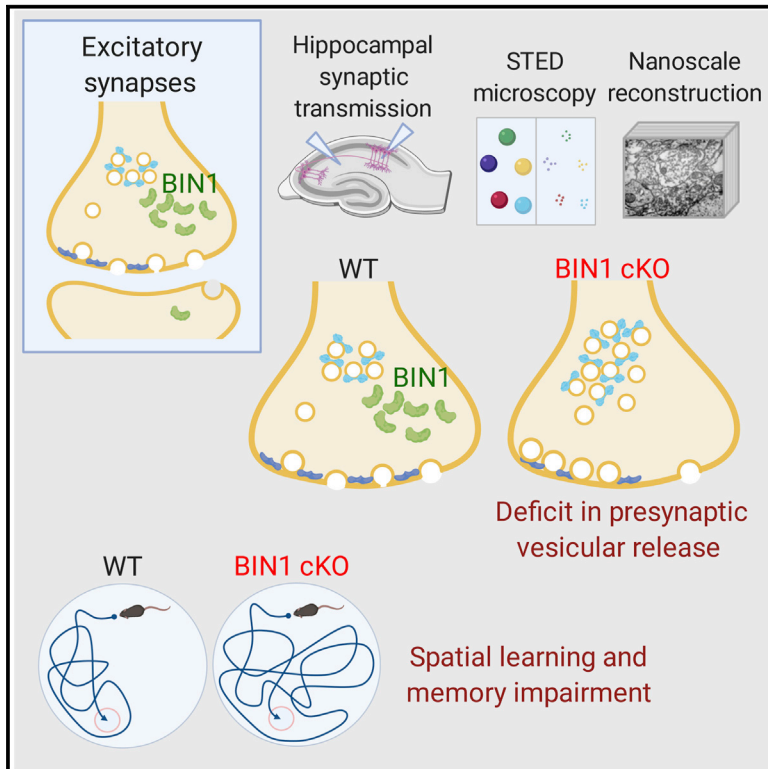


## Neuronal BIN1 Regulates Presynaptic Neurotransmitter Release and Memory Consolidation

### Graphical Abstract



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### In Brief

*BIN1* is a significant risk factor for late-onset Alzheimer disease. *BIN1* has a general role in endocytosis and membrane dynamics in non-neuronal cells. De Rossi et al. show that *BIN1* localizes to presynaptic terminals and plays an indispensable role in excitatory synaptic transmission by regulating neurotransmitter vesicle dynamics.

### Highlights

- The loss of *BIN1* in neurons leads to impaired spatial memory consolidation
- Neuronal *Bin1* cKO mice have deficits in excitatory synaptic transmission
- *BIN1* regulates presynaptic vesicular release in hippocampal excitatory synapses
- The results highlight a non-redundant role for *BIN1* in presynaptic regulation



# Neuronal BIN1 Regulates Presynaptic Neurotransmitter Release and Memory Consolidation

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## SUMMARY

**BIN1, a member of the BAR adaptor protein family, is a significant late-onset Alzheimer disease risk factor. Here, we investigate BIN1 function in the brain using conditional knockout (cKO) models. Loss of neuronal *Bin1* expression results in the select impairment of spatial learning and memory. Examination of hippocampal CA1 excitatory synapses reveals a deficit in presynaptic release probability and slower depletion of neurotransmitters during repetitive stimulation, suggesting altered vesicle dynamics in *Bin1* cKO mice. Super-resolution and immunoelectron microscopy localizes BIN1 to presynaptic sites in excitatory synapses. *Bin1* cKO significantly reduces synapse density and alters presynaptic active zone protein cluster formation. Finally, 3D electron microscopy reconstruction analysis uncovers a significant increase in docked and reserve pools of synaptic vesicles at hippocampal synapses in *Bin1* cKO mice. Our results demonstrate a non-redundant role for BIN1 in presynaptic regulation, thus providing significant insights into the fundamental function of BIN1 in synaptic physiology relevant to Alzheimer disease.**

## INTRODUCTION

Genome-wide association studies (GWAS) have identified *BIN1* as a major susceptibility locus for late-onset Alzheimer disease (LOAD). BIN1, also known as Amphiphysin 2 (Amph2), is an

adaptor protein that, among other roles, regulates membrane dynamics in the context of endocytosis and membrane remodeling (Prokic et al., 2014). Two alternative mechanisms have been proposed to link BIN1 with AD Tau pathology from work in cultured neurons: first, BIN1 can bind to Tau in the cytosol, and second, the function of BIN1 in endocytosis limits extracellular Tau uptake and pathology propagation (Calafate et al., 2016; Chapuis et al., 2013). Microglial BIN1 also has been recently found to influence the release of Tau in extracellular vesicles (Crotti et al., 2019). The loss of BIN1 expression influences AD  $\beta$ -amyloid pathogenesis in cultured neurons, but this does not appear to be the case *in vivo* (Andrew et al., 2019; Miyagawa et al., 2016). Among older individuals without dementia, carriers of the BIN1 rs744373 risk allele were found to have similar amyloid pathology but increased Tau pathology and significantly impaired memory performance (Franzmeier et al., 2019). The *Bin1* gene undergoes complex alternative splicing to generate >10 tissue-specific and ubiquitous isoforms (De Rossi et al., 2016a; Prokic et al., 2014). Independent studies have reported a decrease in the longest BIN1 isoform 1, which contains a central clathrin-associated protein-binding region (CLAP domain), in the brains of individuals with LOAD (De Rossi et al., 2016a; Holler et al., 2014; McKenzie et al., 2017). Despite the burgeoning interest in deciphering the involvement of *BIN1* in AD pathophysiology, fundamental information on its physiological function in the brain, especially in neurons and synapses, is still lacking.

BIN1 and its mammalian paralog, Amph1, are members of the BAR (Bin/Amphiphysin/Rversus) domain protein superfamily. Amph1 was discovered as a synaptic vesicle-associated protein enriched in nerve terminals (Lichte et al., 1992) and suggested to play a role in synaptic vesicle endocytosis based on interactions with the AP-2 adaptor protein, dynamin 1, and synaptojanin (Micheva et al., 1997). Amph1 knockout (KO) mice have reduced



viability and cognitive impairments. Neurons cultured from *Amph1* KO mice revealed defects in synaptic vesicle recycling (Di Paolo et al., 2002). Furthermore, stiff person syndrome-associated autoantibodies against *Amph1* compromised endocytosis preferentially at inhibitory GABAergic synapses by altering synaptic vesicle dynamics (Geis et al., 2010; Werner et al., 2016).

The high sequence similarity between *Amph1* and BIN1 and the ability of BIN1 to interact with *Amph1*, dynamin1, AP-2, clathrin, synaptotagmin, and endophilin suggested a role for BIN1 at the presynapse (Prokic et al., 2014). A recent study reported altered AMPA receptor-mediated synaptic transmission in cultured cortical neurons following acute knockdown of BIN1 expression (Schürmann et al., 2019). Aside from this single report of an *in vitro* study, the BIN1 function in synaptic transmission has not been characterized. Here, we investigated the role of BIN1 at the synapse and in learning and memory using neuronal conditional KO (cKO) mice. We report the impairment of spatial memory in association with the loss of BIN1 expression in the hippocampus. Functional analysis of synaptic transmission in the CA1 of the hippocampus demonstrated impairment in neurotransmitter vesicle release at excitatory synapses of the cKO mice. Super-resolution direct stochastic optical reconstruction microscopy (dSTORM) and immunoelectron microscopy (immuno-EM) localized BIN1 to presynaptic sites in glutamatergic synapses. Confocal and stimulated emission depletion (STED) microscopy analysis of presynaptic morphology in cKO mice revealed a decrease in the density of presynaptic sites and the size of presynaptic protein clusters. Finally, analysis of 3D EM image stacks from hippocampal CA3-CA1 synapses revealed a significant increase in docked and reserve pools of synaptic vesicles in cKO mice. These results demonstrate the presynaptic localization and functional role for BIN1 that are highly relevant to AD pathophysiology.

## RESULTS

### Characterization of BIN1 Neuronal Expression in the Brain

We generated neuronal *Bin1* cKO mice by crossing the *Bin1<sup>fl/fl</sup>* strain with the *Syn1-Cre* and *Emx1-IRES-Cre* driver lines to explore the function of BIN1 in neurons. The *Syn1-Cre* transgene is specific to neurons with high expression in the hippocampus (Zhu et al., 2001), and *Emx1-Cre* is expressed in excitatory neurons and glia in the cortex and hippocampus (Gorski et al., 2002). The loss of BIN1 in the *Bin1<sup>fl/fl</sup>;Syn1-Cre* (henceforth referred to as *Syn* mice) and *Bin1<sup>fl/fl</sup>;Emx1-IRES-Cre* (henceforth referred to as *Emx* mice) cKO models were confirmed by immunoblots and immunostaining. Both BIN1:H and BIN1:L isoforms were significantly decreased in different areas of the brain in the *Syn* cKO and the *Emx* models (Figures 1A and S1). The loss of BIN1 expression in the cKO mice did not affect the expression of the paralog of BIN1, *Amph1*, and several other proteins in the brain (Figure S2; *Emx* data not shown). Immunostaining of BIN1 using monoclonal antibody (mAb) 13463 under conditions that preferentially stain neuronal BIN1 (Andrew et al., 2019) revealed an almost complete loss of BIN1 in the hippocampus (Figure 1B). Microdissection immunoblot analysis confirmed a significant decrease in both the BIN1:H and BIN1:L isoforms

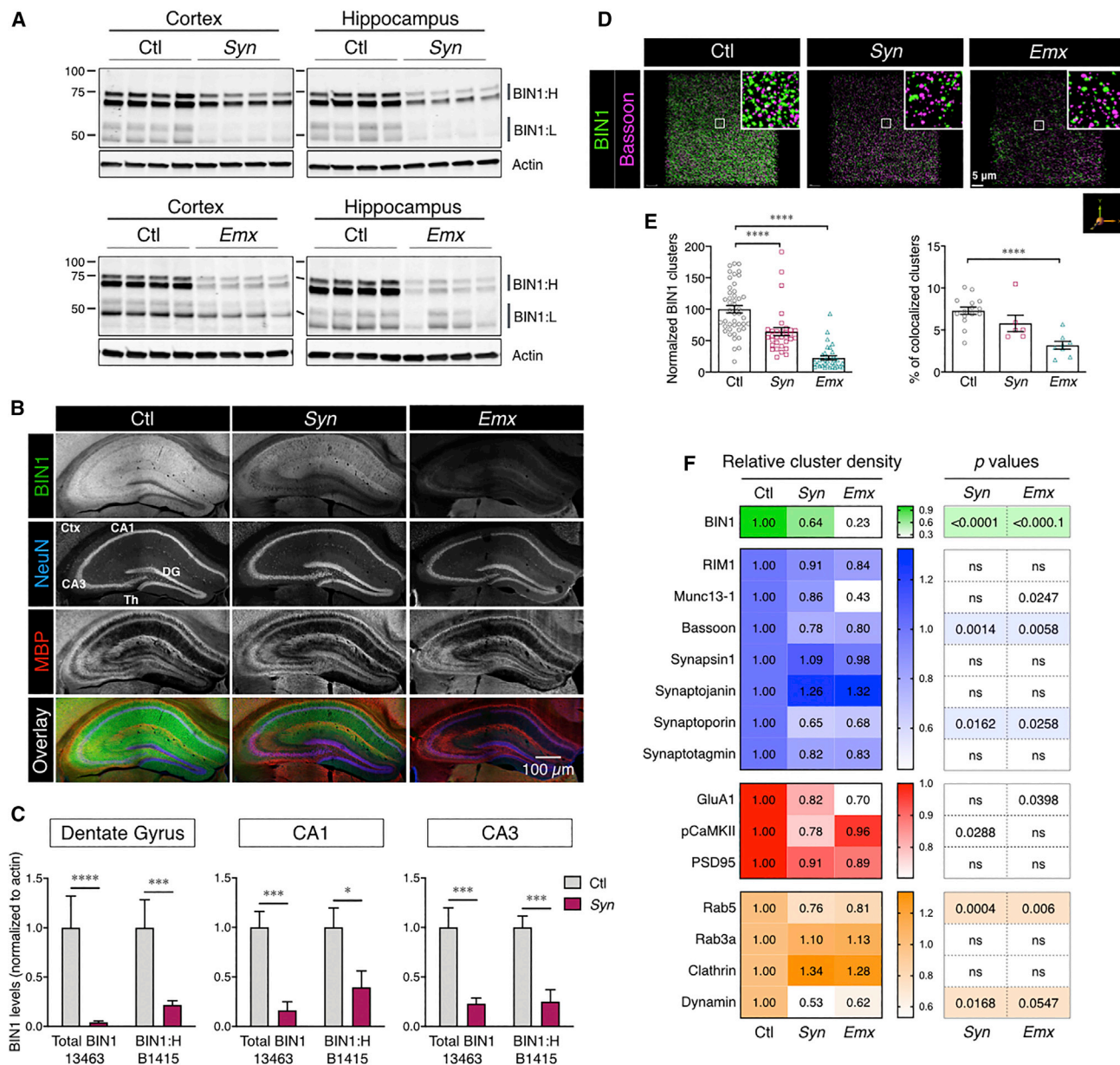
(Figures 1C and S2), suggesting that hippocampal neurons express multiple isoforms of BIN1, including those lacking the brain-specific CLAP domain, as confirmed by immunoblotting using pAb B1415, an antibody that is specific for the longest BIN1 brain isoforms 1 and 2 (BIN1:H).

### Impaired Recognition Memory and Spatial Learning in Mice Lacking Neuronal BIN1 Expression

We used multiple behavior tests to assess memory acuity, in which the hippocampus plays a major role. In the Y-maze test, a working memory task that engages neural circuits in the prefrontal cortex, hippocampus, basal forebrain, and septum (Lalonde, 2002), the loss of neuronal BIN1 expression did not affect total arm entries or spontaneous alternations (Figure 2A). In the novel object recognition test (NOR), a task that examines the communication between the hippocampus and the cortex in recognition memory (Winters et al., 2008), no difference was observed during the learning phase. Quantification of the interaction index during the memory recollection phase revealed a significantly impaired NOR performance in the *Emx* model but not in the *Syn* model (Ctl versus *Syn*  $p = 0.6411$ ; Ctl versus *Emx*  $p = 0.0014$ ). (Figure 2B). These data suggest that the loss of *Bin1* in excitatory neurons in the forebrain led to an impairment in a task that assesses recognition memory performance.

We then tested whether BIN1 is required for behavioral tasks that engage the amygdala and hippocampal circuits using a fear-conditioning paradigm (Kim and Jung, 2006). No difference in the cumulative freezing time was observed in *Bin1* cKO mice as compared to littermate Ctl (Figure 2C). These results suggested that mice are able to associate a specific environment with the aversive stimulus in the absence of BIN1 expression. Moreover, both Ctl and *Syn* mice showed the same fear extinction pattern when mice were returned once per week in the contextual chamber without shock (Figure 2D), confirming normal amygdala-dependent learning.

Finally, we tested spatial learning memory using the Morris water maze and observed a clear difference between Ctl mice and both neuronal *Bin1* cKO models. Over the 7-day training period, Ctl mice significantly reduced the time to find the platform (>80% success of finding the platform in 3 consecutive days), indicative of a successful learning process. However, the two cKO models failed to show a definite improvement in the latency to find the platform during the acquisition phase (the latency in seconds to find platform day 1 versus day 7: Ctl  $74.57 \pm 3.9$  versus  $45.86 \pm 4.65$ ,  $p < 0.0001$ ; *Syn*  $80.51 \pm 5.26$  versus  $73.86 \pm 7.10$ ,  $p = 0.449$ ; *Emx*  $68.33 \pm 6.3$  versus  $55.23 \pm 7.12$ ,  $p = 0.1439$ , two-way ANOVA with Fisher's least significant difference [LSD] post hoc test). The *Emx* cohort was observed to have faster mobility and longer swim distance. When the swim distance was included as a covariate in the linear mixed effects model, there was a significant effect of genotype on the rate of learning (latency to find the platform;  $F_{(2, 107)} = 19.501$ ,  $p < 0.0005$ ) (Figure 2E). We also examined the explorative strategies that mice used to find the platform after 3 to 4 days of training (Graziano et al., 2003). The Ctl mice scanned the arena using a random search or circling strategy over the initial days and then switched to self-orienting and approaching target strategy (5–20 s escape latency) or direct finding (1–10 s



**Figure 1. Loss of Neuronal BIN1 Expression in cKO Mice**

(A) Immunoblot analysis of BIN1 levels in the cortex and hippocampus of BIN1 cKO mice (polyclonal antibody [pAb] 14647).

(B) Confocal microscopy analysis of neuronal BIN1 expression in control (Ctl) and cKO (*Syn* and *Emx*) mice (mAb 13463; no epitope retrieval). Adjacent image panels acquired using a 10× objective were stitched together for the visualization of the entire hippocampus. Scale bar: 200 μm.

(C) Analysis of the diminution of BIN1 expression in the micro-dissected areas of the hippocampus of cKO mice. The means ± SEMs are plotted in each graph. \**p* < 0.05, \*\**p* < 0.01, \*\*\**p* < 0.001, and \*\*\*\**p* < 0.0001.

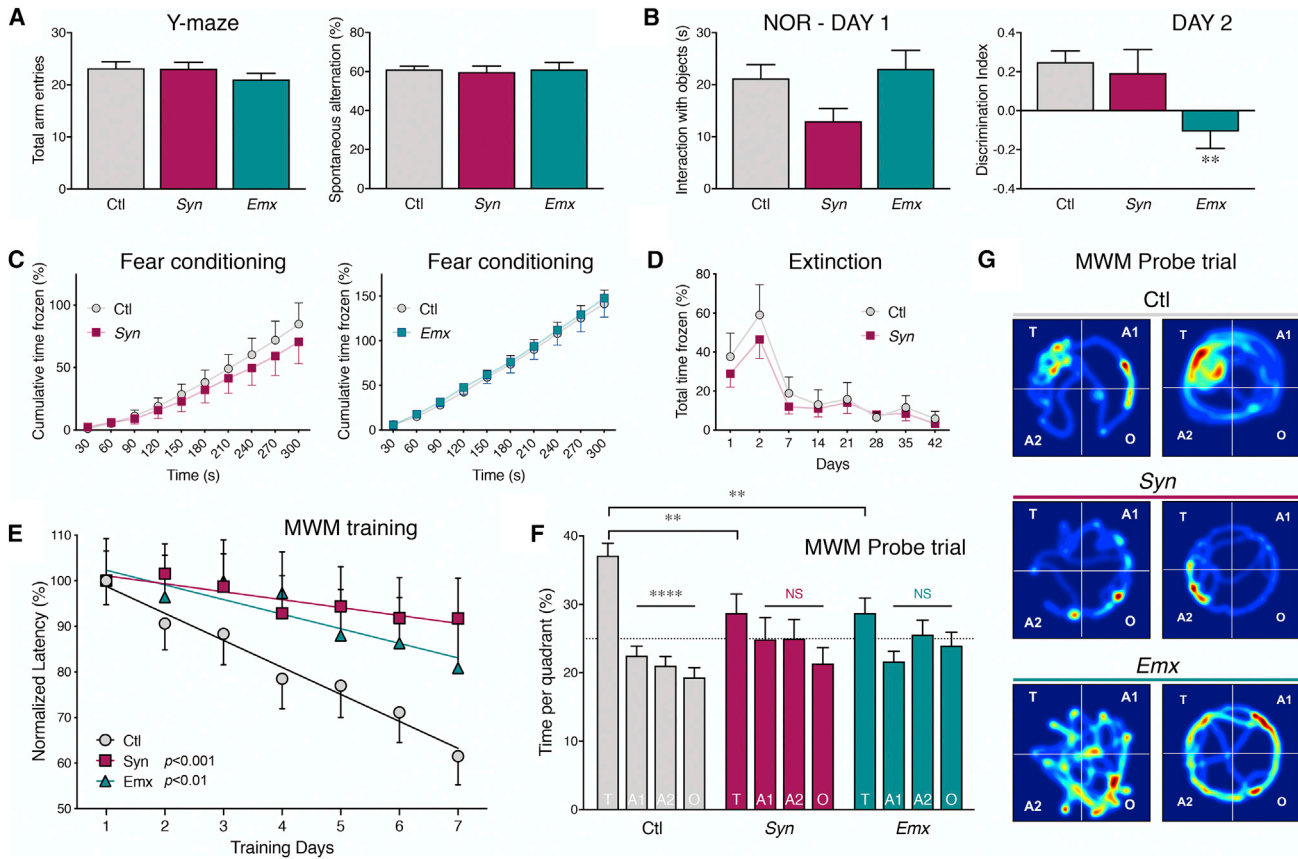
(D) Decreased presynaptic marker density in neuronal *Bin1* cKO mice. Masks of BIN1 and Bassoon immunostaining in confocal images (see [Method Details](#)) were used to quantify synaptic marker densities from a block of 50 z stack images representing a depth of ~25 μm (96,193 μm<sup>3</sup> volume).

(E) Decrease in BIN1 cluster density (left) in cKO mice (ANOVA:  $F_{(2, 107)} = 50.7$ , *p* < 0.0001; Fisher's LSD post hoc test: Ctl versus *Syn* and Ctl versus *Emx* *p* < 0.0001). Decrease of and BIN1-Bassoon colocalization, normalized to Bassoon cluster density (right) in *Emx* mice (ANOVA:  $F_{(2, 26)} = 14.18$ , *p* < 0.0001; Fisher's LSD post hoc test: Ctl versus *Syn* *p* = 0.0907 and Ctl versus *Emx* *p* < 0.0001).

(F) Heatmap representing mean changes in the density of different presynaptic markers (blue), postsynaptic markers (red), and presynaptic endocytic vesicle markers (orange).

See also [Figures S1](#), [S2](#), and [S7](#) and [Tables S1](#) and [S2](#).





**Figure 2. Loss of Neuronal BIN1 Expression Impairs Spatial Learning and Memory**

Neuronal cKO mice, 6 to 10 weeks old, and their littermate Ctl were subject to multiple behavioral tests.

(A) Y-maze test shows no significant difference between the groups in the number of arm entries (left) or spontaneous alternation behavior (ANOVA  $F_{(2, 78)} = 0.07933$ ,  $p = 0.9238$ ;  $n = 44$  Ctl, 18 *Syn*, and 19 *Emx* mice) (right).

(B) In the novel object recognition test, no significant difference between groups was observed during the interaction with the objects on day 1 (ANOVA:  $F_{(2, 101)} = 2.419$ ,  $p = 0.0942$ ;  $n = 56$  Ctl, 23 *Syn*, and 25 *Emx* mice). The discrimination index is plotted at right. ANOVA ( $F_{(2, 97)} = 5.560$ ,  $p = 0.0052$ ;  $n = 56$  Ctl, 19 *Syn*, and 25 *Emx* mice) showed no significant difference between Ctl and *Syn* (Fisher's LSD post hoc test,  $p = 0.6411$ ) and a significant difference between Ctl and *Emx* (Fisher's LSD post hoc test,  $p = 0.0014$ ).

(C) Contextual fear conditioning test shows no difference between cKO mice and their respective littermate Ctl in the cumulative time frozen (Ctl versus *Syn*:  $t_{27} = 0.5876$ ,  $p = 0.5617$ ,  $n = 15$  Ctl and 14 *Syn* mice; Ctl versus *Emx*:  $t_{32} = 0.2483$ ,  $p = 0.8055$ ,  $n = 21$  Ctl and 13 *Emx* mice; two-tailed Student's t test).

(D) No statistical difference was observed in fear conditioning extinction between *Syn* mice and their littermate Ctl (Ctl versus *Syn*  $p > 0.24$  at all times;  $n = 19$  Ctl and 18 *Syn* mice; multiple t test analysis).

(E) Morris water maze test shows spatial learning defects in *Bin1* cKO mice performance over the 7 days of training. There was a significant effect of genotype on the rate of learning (raw latency to find the platform; [ $F_{(2, 107)} = 19.501$ ,  $p < 0.0005$ ];  $n = 54$  Ctl, 28 *Syn*, and 27 *Emx* mice). The performance of Ctl mice significantly improved during training as compared with *Syn* ( $p < 0.0005$ ) and *Emx* mice ( $p = 0.004$ ).

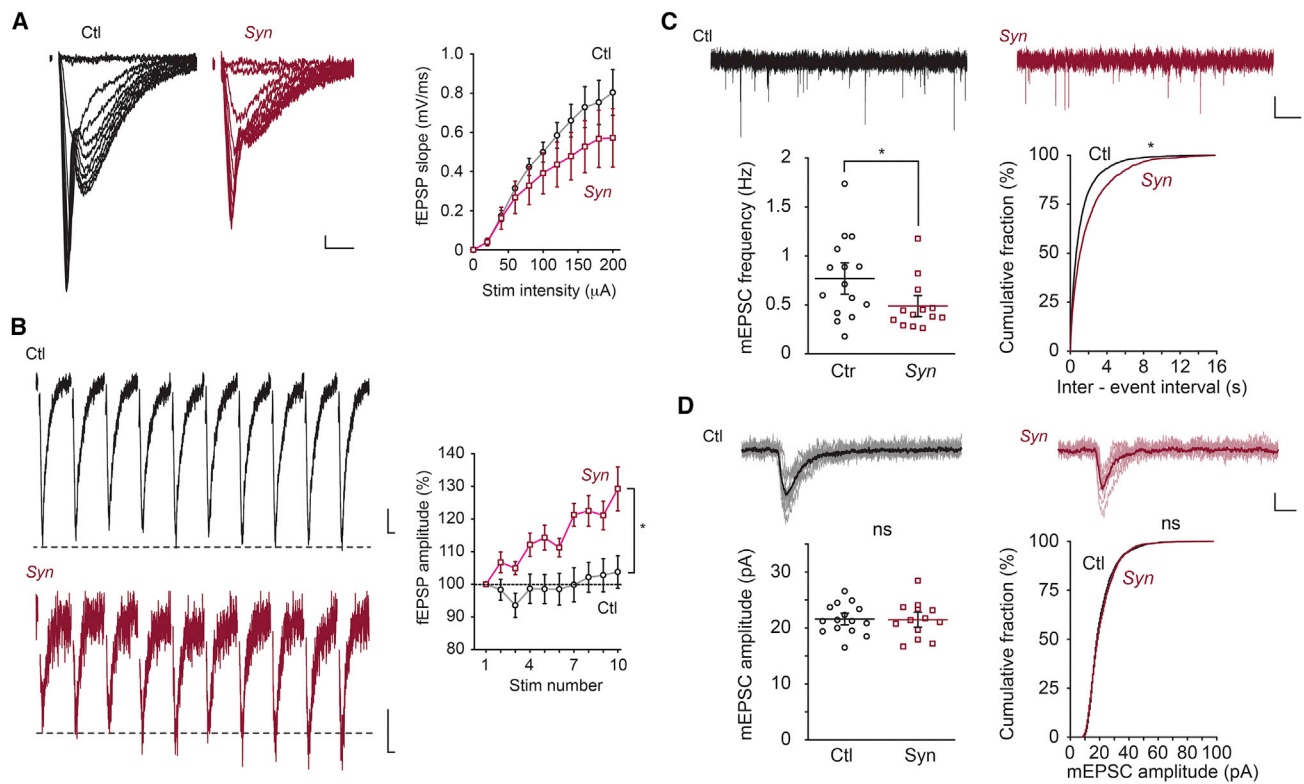
(F) In the probe trial, *Bin1* cKO mice spent significantly less time in the target quadrant as compared with Ctl mice ( $F_{(2, 105)} = 5.497$ ,  $p = 0.0054$ ). The percentage of time spent in each quadrant is plotted. T, target; A1 and A2, adjacent; O, opposite quadrant. ANOVA analysis ( $F_{(6, 424)} = 2.930$ ,  $p = 0.0082$ ); post hoc Tukey's multiple comparisons for Ctl mice: T versus A1,  $p < 0.0001$ ; T versus A2,  $p < 0.0001$ ; T versus O,  $p < 0.0001$ ; no significant difference between the quadrants was recorded for *Syn* and *Emx* mice.

(G) Representative heatmaps of average search during the probe trial for Ctl and cKO groups.

See also Figure S3.

in the following days. However, most of the cKO mice did not change their strategy from random searching (40–120 s) or scanning (15–60 s), indicating that the capacity for spatial learning is limited in the absence of BIN1 expression. In the probe trial, Ctl mice spent more time in the target area compared to the other quadrants. In contrast, the neuronal *Bin1* cKO mice spent the same amount of time in each quadrant (the percentage of time spent in target versus opposite: Ctl  $37.11 \pm 1.81$  versus

$19.33 \pm 1.43$ ,  $p < 0.0001$ ; *Syn*  $28.76 \pm 2.75$  versus  $21.36 \pm 2.31$ ,  $p > 0.05$ ; *Emx*  $28.75 \pm 2.18$  versus  $23.99 \pm 1.94$ ,  $p > 0.05$ , two-way ANOVA with Tukey's multiple comparisons test) (Figure 1F). These results suggest a significant deficit in spatial learning memory in *Bin1* cKO mice. All of the mice performed equally in the visible platform test, and the swimming parameters were identical for all of the genotypes (Figures S3A and S3B). Thus, the failure to complete the task was not related to a



**Figure 3. *Bin1* Deletion Causes an Altered Presynaptic Function**

(A) I/O curve of fEPSPs. Representative fEPSP traces (left) and grouped data for average slope (right) in WT (Ctl) and *Bin1* cKO (*Syn*) mice. There is a significant interaction between stimulation intensity and genotype;  $p < 0.0005$ . Calibration: 10 ms, 0.1 mV.

(B) Short-term plasticity (1 Hz). Representative fEPSP traces (left) and grouped data (right) in Ctl and *Syn* mice. Calibration: 10 ms, 0.1 mV.

(C) mEPSC frequency. Representative mEPSC traces (top) and group data for average frequency (bottom left) and cumulative plot of the interspike interval (bottom right) in Ctl and *Syn* mice. Calibration: 2 s, 10 pA.

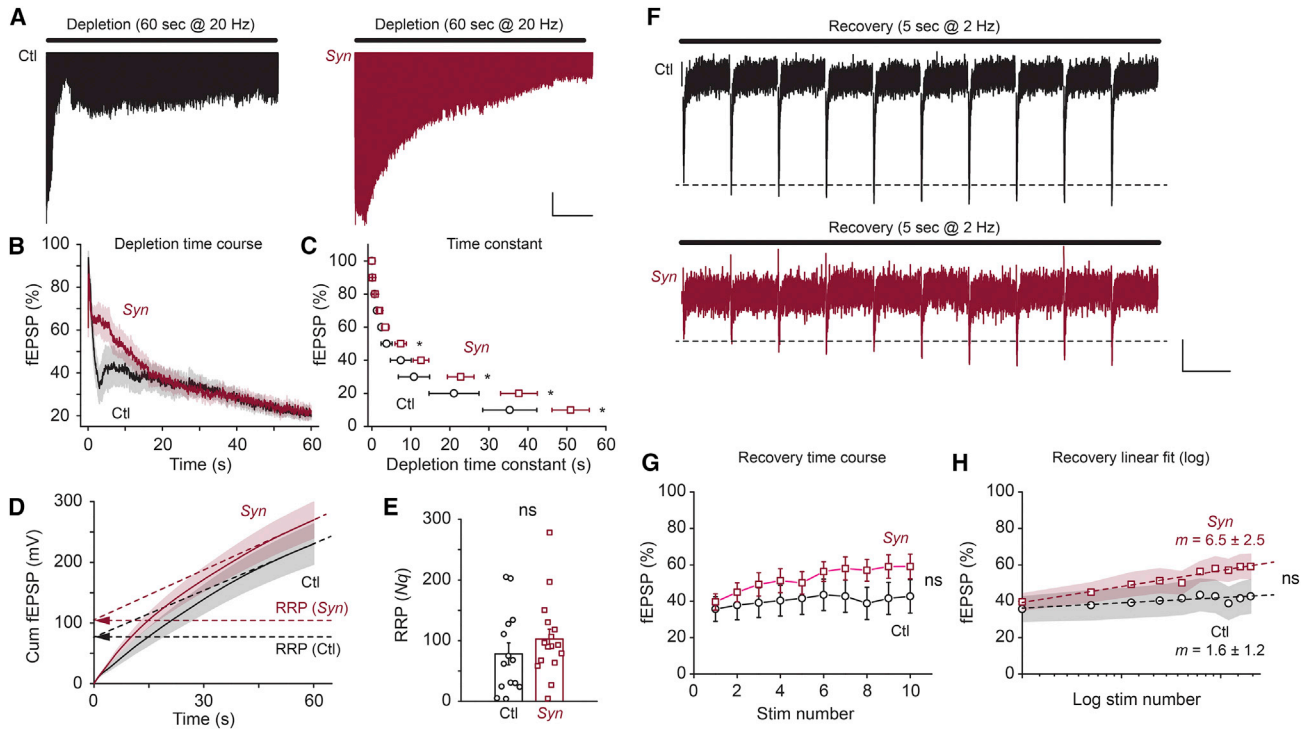
(D) mEPSC amplitude. Representative individual mEPSC traces. Ten representative events and the averaged trace were presented as lighter and darker lines, respectively (top). Grouped data for average amplitude (bottom left) and cumulative plot of amplitude (bottom right) in Ctl and *Syn* mice. Calibration: 10 ms, 10 pA; 15–18 slices from 4 Ctl mice and 13–18 slices from 4 *Syn* mice were used; ns  $p \geq 0.05$ .

physical problem but was due to a spatial memory deficit. We tested motor impairment in *Syn* and *Emx* models using rotarod tests and DigiGait analyses (Figures S3C–S3G). All of the measured parameters showed no difference between *Bin1* cKO and Ctl mice. These data suggest a role for BIN1 in memory formation and consolidation.

### Absence of *Bin1* Causes a Deficit in the Presynaptic Vesicular Release

Given the effect of *Bin1* deletion on learning and memory, we hypothesized that the loss of BIN1 in neurons results in an alteration of synaptic function. First, we compared the input/output (I/O) curve of synaptic responses in the CA1 of Ctl and *Syn* mice to examine whether there are any gross alterations in synaptic transmission. Analysis of the data using a linear mixed effects model showed that there was a significant interaction between stimulation intensity and genotype, suggesting that synaptic responses in slices from *Syn* mice at different stimulation intensities were different from those of Ctl mice ( $F_{(9, 283)} = 3.435$ ,  $p < 0.0005$ ;  $n = 16$  Ctl and 18 *Syn*). However, the overall effect of the genotype was not significant ( $F_{(1, 32)} = 0.671$ ,  $p = 0.42$ ) (Fig-

ure 3A). Second, we examined short-term plasticity induced by short trains of stimuli to determine whether there was any alteration in presynaptic release probability (Klemmer et al., 2011; Lou et al., 2012; Zucker and Regehr, 2002). During a train of 10 synaptic stimuli given at 1 Hz, field excitatory postsynaptic potential (fEPSP) responses were potentiated to a significantly higher magnitude in *Syn* mice than in the Ctl group, suggesting that the release probability of CA1 synapses was lower in *Syn* mice ( $F_{(1, 27)} = 8.74$ ,  $p < 0.01$ ;  $n = 15$  Ctl and 14 *Syn*, two-way ANOVA) (Figure 3B). In separate voltage clamp recordings of miniature excitatory postsynaptic currents (mEPSCs) we observed a significantly reduced frequency of synaptic events in *Syn* mice, consistent with reduced release at individual terminals ( $0.77 \pm 0.11$  versus  $0.49 \pm 0.07$  Hz,  $p = 0.038$ , Mann-Whitney test) (Figure 3C). There was no significant difference in the amplitude of the quantal events between Ctl and *Syn* mice ( $21.6 \pm 0.7$  versus  $21.5 \pm 0.9$  pA,  $p = 0.93$ , Mann-Whitney test) (Figure 3D). This latter finding in CA1 neuronal postsynaptic response in intact slices is in contrast to the effect of acute knockdown of BIN1 expression in cultured cortical neurons, in which a reduction in mEPSC amplitudes was observed



**Figure 4. *Bin1* Deletion Causes Altered Depression of the EPSP during Sustained Synaptic Trains**

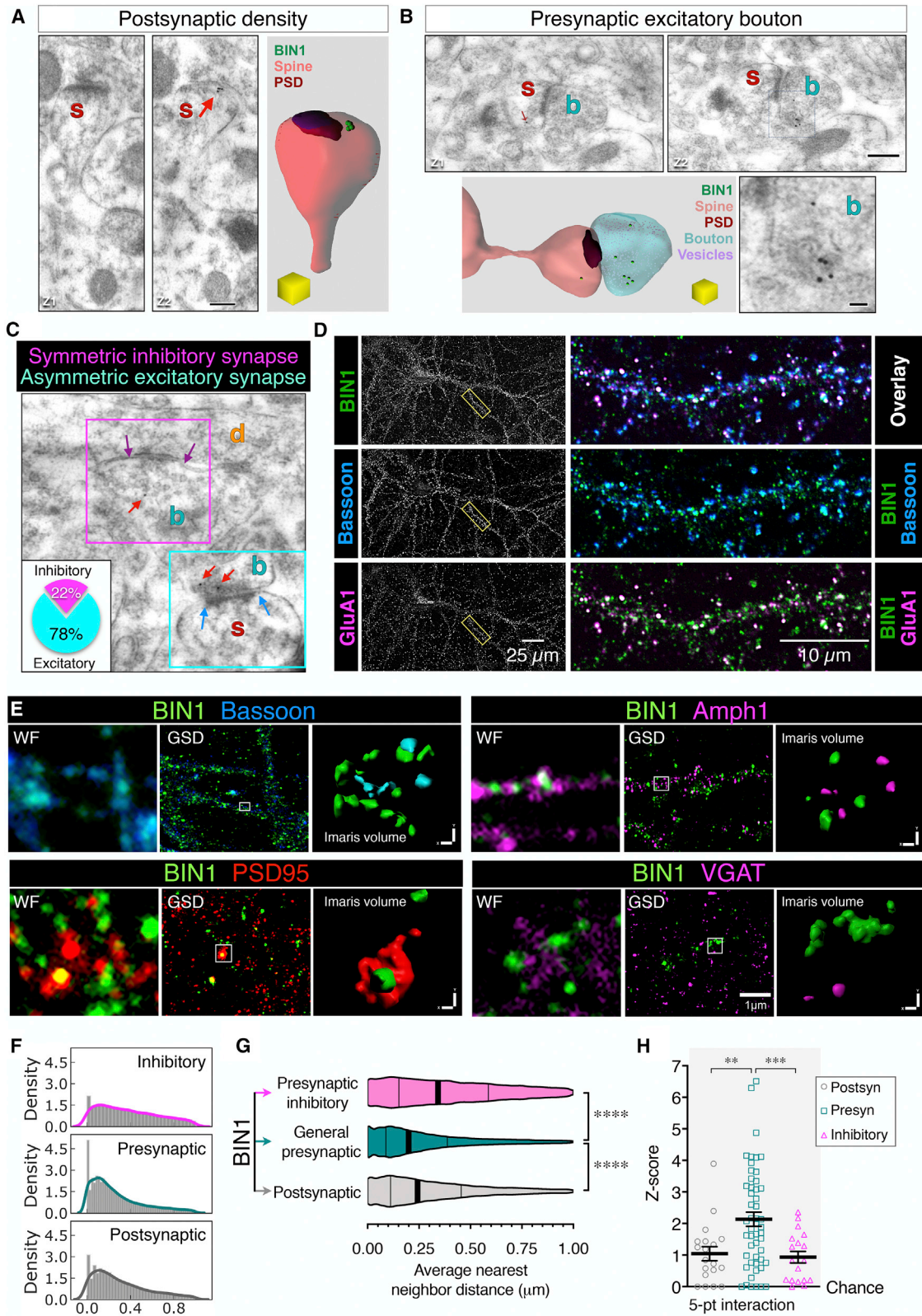
(A) Representative fEPSP traces during the depletion paradigm (20 Hz stimuli for 60 s) in *Bin1* WT (Ctl) and cKO (*Syn*) mice. Calibration: 10 s, 0.1 mV.  
 (B) Grouped data for fEPSP depletion time course in Ctl and cKO mice.  
 (C) Grouped data for fEPSP depletion time constants for 90%–10% of the peak in Ctl and cKO mice.  
 (D) Grouped data for cumulative fEPSP amplitude during the depletion paradigm in Ctl and cKO mice. RRP is estimated from the y axis intercept of the linear regression fit.  
 (E) Grouped data for estimated RRP in Ctl and cKO mice.  
 (F) Representative fEPSP traces during the recovery paradigm (2 Hz stimuli for 5 s) in Ctl and cKO mice. Calibration: 500 ms, 0.1 mV.  
 (G) Grouped data for fEPSP recovery time course in Ctl and cKO mice.  
 (H) Grouped data for fEPSP recovery time course in logarithm fit in Ctl and cKO mice. The slope ( $m$ ) of the linear fit is presented in each genotype; 14 slices from 3 Ctl mice and 16 slices from 4 *Syn* mice were used; ns,  $p \geq 0.05$ .

(Schürmann et al., 2019). To further examine whether there is an alteration in the presynaptic vesicular release probability in *Bin1* cKO mice, we analyzed other proxies of the release probability in the whole-cell recording configuration. Paired pulse facilitation (PPF) with various interstimulus intervals (ISIs) was not significantly different between Ctl and *Syn* mice (PPF with ISI = 20 ms: Ctl,  $1.63 \pm 0.07$  versus *Syn*,  $1.90 \pm 0.11$ ; 40 ms: Ctl,  $1.55 \pm 0.06$  versus *Syn*,  $1.66 \pm 0.07$ ; 60 ms: Ctl,  $1.40 \pm 0.06$  versus *Syn*,  $1.48 \pm 0.07$ ; 80 ms: Ctl,  $1.35 \pm 0.04$  versus *Syn*,  $1.42 \pm 0.06$ ;  $F_{(1, 26)} = 2.851$ ,  $p = 0.103$ ;  $n = 16$  Ctl and 12 *Syn*, two-way ANOVA). In contrast to this, analysis of the coefficient of variation (CV) of EPSCs determined a significantly larger CV in *Syn* mice than in Ctl mice (Ctl,  $0.21 \pm 0.02$  versus *Syn*,  $0.27 \pm 0.02$ ,  $p = 0.013$ , Mann-Whitney test), which would indicate a lower synaptic release probability in *Syn* mice. Despite the lack of difference in PPF between the genotypes, our results from recording in the CA1 of the hippocampus by three measures of release probability indicate that there is a significant reduction in presynaptic vesicular release probability, whereas postsynaptic properties of CA1 synapses are not perturbed in *Syn* mice.

### Deletion of *Bin1* Causes Slower Vesicle Depletion

The results from the hippocampal slices described above contrast with recently reported data from cultured cortical neurons in which the knockdown of BIN1 expression altered AMPA receptor (AMPA) expression and surface localization at the postsynaptic site (Schürmann et al., 2019). Amphiphysins are thought to be involved in synaptic vesicle recycling by interacting with presynaptic endocytic proteins (Calafate et al., 2016; Prokic et al., 2014; Wigge et al., 1997). Therefore, we examined whether the loss of BIN1 expression affects synaptic vesicle depletion during sustained presynaptic stimulation, as reported in mutant mice for other presynaptic proteins (Chung et al., 2010; Cremona et al., 1999; Di Paolo et al., 2002; Ferguson et al., 2007; Lou et al., 2012; Milosevic et al., 2011; Vargas et al., 2014). Repetitive trains of stimuli were applied at a frequency of 20 Hz for 60 s, and fEPSP synaptic responses were recorded. The amplitude of the fEPSP depressed more slowly during the train in *Syn* mice than in Ctl mice ( $p < 0.05$  for depletion constants for 50%, 30%, 20%, and 10%,  $n = 14$  Ctl and 16 *Syn*, Mann-Whitney test) (Figures 4A–C). One possible explanation for the slower rate of depression of synaptic responses is by





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an increased size of the vesicular pool to maintain synaptic strength. Therefore, we calculated the size of the readily releasable pool (RRP) in *Syn* mice by measuring the cumulative fEPSPs during the stimulus trains (Fernandes et al., 2015; Fioravante and Regehr, 2011; Zucker and Regehr, 2002) (Figure 4D). This analysis assumes that the depression of the synaptic response reflects a depletion of the RRP, allowing an estimate to be made of the product of  $Nq$  (where  $N$  is the total number of releasable vesicles and  $q$  is the quantal size) from the cumulative fEPSP plot (Fernandes et al., 2015; Schneggenburger et al., 1999). Given that we had observed no difference in the mEPSC amplitude in Ctl and *Syn* mice (Figure 3D), it can be assumed that the quantal size  $q$  is constant between the groups. With this analysis, we did not find a statistically significant difference in  $Nq$ , suggesting that the RRP ( $N$ ) is not grossly affected by *Bin1* loss ( $Nq$ ,  $78.2 \pm 18.1$  versus  $102.6 \pm 16.3$ ,  $p = 0.29$ ,  $n = 14$  Ctl mice and 16 *Syn* mice, Mann-Whitney test) (Figure 4E). We also examined the recovery time course of the synaptic responses after depletion. Low-frequency stimuli (2 Hz) were applied immediately after the depletion paradigms and the fEPSP response was measured. Again, we did not observe a significant difference in the recovery time constant between Ctl and *Syn* mice ( $1.6 \pm 1.2$  versus  $6.5 \pm 2.5$  (slope of percentage of recovery per stimulus number),  $p = 0.19$ ,  $n = 14$  Ctl mice and 16 *Syn* mice, Mann-Whitney test) (Figures 4F–4H). The physiological characterization of synapses revealed a role for BIN1 in regulating presynaptic release probability, but *Bin1* loss had no measurable effect on the RRP.

### BIN1 Localizes to Excitatory Presynaptic Sites

We performed immunogold EM analyses to determine the localization of BIN1 in CA3–CA1 synapses in the mouse brain using post-embedding immunogold EM on freeze-substituted tissue. Figure 5 shows representative images of BIN1 localization in a dendritic spine, a presynaptic excitatory bouton, and symmetric and asymmetric synapses (Figures 5A–5C). In the volume of tissue examined (578 excitatory pre- and postsynapses and 67 inhibitory boutons), 122 excitatory synapses were found to contain BIN1-immunogold particles as compared with 35 inhibitory presynaptic boutons. Only 35 of 578 postsynaptic sites were immunopositive. Thus, although BIN1 could be localized

by immunogold particles in the postsynaptic compartment of axospinous synapses and symmetric and inhibitory presynaptic terminals, the frequency of labeling was highest in the presynaptic compartment of asymmetric, axospinous synapses. BIN1 was frequently found to be associated with large clusters of synaptic vesicles at axon terminals, but only occasionally located within the docking area of the active zone (Figures 5A–5C; see also Figure S4).

We further characterized BIN1 localization at the synapse in mature rat hippocampal primary neuronal cultures (Figures 5D and S5A). To precisely discriminate between presynaptic and postsynaptic localization of BIN1, we acquired super-resolution images at 20 nm pixel-size resolution using dSTORM (Figures S5B–S5D). Sub-diffraction resolution dSTORM images revealed a lack of enrichment of BIN1 in the dendritic spines (labeled with  $\text{Ca}^{2+}$ /calmodulin-dependent protein kinase II [CaMKII]) or within the dendritic shaft (labeled with MAP2). We observed BIN1 in slender MAP2<sup>−</sup> processes, indicative of axonal localization of BIN1 (Figure S5C, white arrows). Moreover, BIN1 showed considerable overlap with Bassoon as compared with GluA1 or PSD95, suggesting a preponderance of presynaptic BIN1 localization (Figures 5E and S5). Our extensive data from immuno-EM and super-resolution microscopy analysis of hippocampal neurons in mouse brain slices and culture contrast with a recent study that focused on BIN1 postsynaptic localization (Schürmann et al., 2019). However, it should be noted that the authors of the previous report observed an obvious localization of BIN1 at the presynaptic site, visualized as a strong colocalization between BIN1 and synapsin1 (Schürmann et al., 2019).

For unambiguous quantification, BIN1 localization was compared with general presynaptic markers (Bassoon, synapsin1, and synaptophysin), and inhibitory presynaptic (VGAT) or postsynaptic (AMPA-GluA1, PSD95, and CaMKII) markers. Recent studies showed an enrichment and a role for Amph1, the paralog of BIN1, in GABAergic synapses (Geis et al., 2010; Werner et al., 2016). For this reason, we included Amph1 as a marker of inhibitory synapses in addition to VGAT. To precisely determine BIN1 localization in dSTORM images, we mapped the cluster locations of BIN1 and the synaptic markers and calculated the distances between the cluster centers using the Density-Based Spatial Clustering of Applications with Noise

### Figure 5. Investigation of BIN1 Synaptic Localization Using Immuno-EM and Ground-State Depletion (GSD) dSTORM Microscopy

(A–C) Representative images of BIN1 immuno-EM localization in the mouse hippocampal CA1 region, showing BIN1-associated gold particles in the post-synaptic (A), general presynaptic (B), and inhibitory synapses (C). The inset in (C) indicates that of all of the synapses with BIN1-immunogold particles, 78% were excitatory terminals. Scale bars: 250 nm; enlarged inset, 50 nm.

(D) Representative confocal images of BIN1, Bassoon, and GluA1. Image overlays of the boxed region are shown at a higher magnification at right.

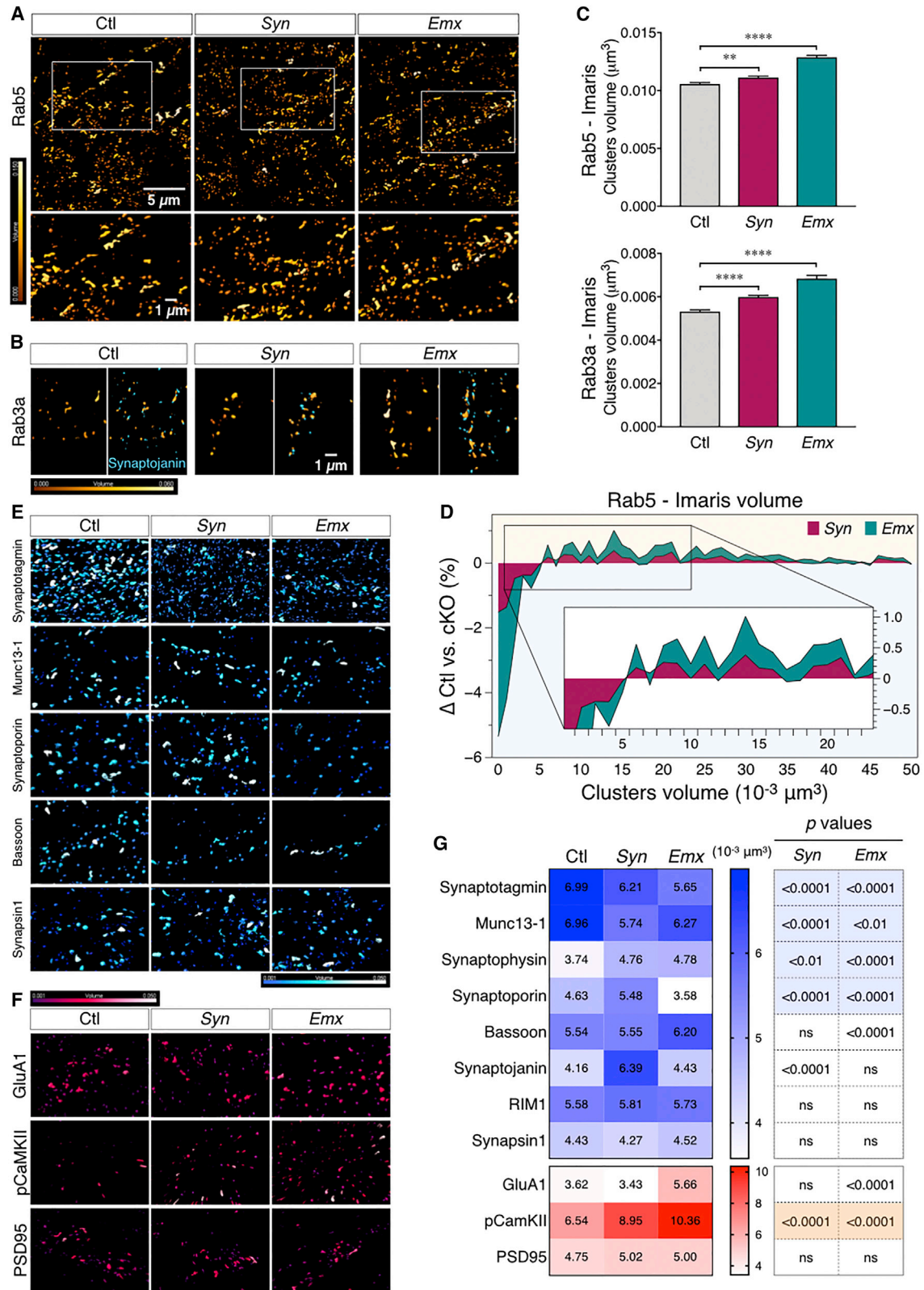
(E) Representative images of BIN1 (green) and markers for postsynaptic sites (PSD95; red), presynaptic sites (Bassoon; blue), and inhibitory presynaptic sites (Amph1 and VGAT; magenta). For each marker, the widefield image (WF), the super-resolution image (GSD), and the Imaris volume view (used for nearest-neighbor calculation) of the identical region are displayed.

(F) The density plot depicts the frequency of the distance between the BIN1 clusters and synaptic markers. The x axis indicates the binned DBSCAN average nearest-neighbor distance (microns) between BIN1 and synaptic markers.

(G) Violin plot of the average distance between BIN1 and its nearest neighbor. ANOVA analysis ( $F_{(2, 80700)} = 1,332$ ,  $p < 0.0001$ ); BIN1-to-marker mean distance: general presynaptic markers =  $0.2708 \pm 0.001 \mu\text{m}$ ; postsynaptic markers =  $0.308 \pm 0.002 \mu\text{m}$ ; and presynaptic inhibitory markers =  $0.3849 \pm 0.002 \mu\text{m}$ . The vertical lines indicate the median and quartiles.

(H) DBSCAN analysis of randomization score to assess the specificity of BIN1 approximation to synaptic markers (5-point interaction; 50 points per cluster at 50-nm distance). The Z-score analysis showed a specific association of BIN1 with the synaptic markers; ANOVA analysis ( $F_{(8, 255)} = 5.446$ ,  $p < 0.0001$ ). When the Z score equals 0, the association between BIN1 and the markers is not greater than a chance occurrence.

See also Figures S4–S6 and Table S1.



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(DBSCAN) (see [Method Details](#)). We found that a significantly higher proportion of BIN1 clusters were within 30, 40, and 50 nm of presynaptic clusters as compared with the proximity of BIN1 to inhibitory presynaptic or postsynaptic clusters (ANOVA, 30-nm threshold:  $F_{(2, 84)} = 30.09$ ,  $p < 0.0001$ ; 40-nm threshold:  $F_{(2, 84)} = 27.05$ ,  $p < 0.0001$ , 50-nm threshold:  $F_{(2, 84)} = 38.73$ ,  $p < 0.0001$ ) ([Figure S6A](#)). However, we note that there still were a sizeable number of BIN1 clusters that were located in nearby postsynaptic markers. We confirmed these results using Imaris software 3D reconstruction of dSTORM images and calculating the nearest-neighbor distance between all BIN1 and the synaptic marker-stained structures. For statistical relevance and to not bias the data, the distance analyses were performed using a 1,000-nm threshold. A frequency distribution analysis revealed that a large number of BIN1 clusters were localized within 250 nm from synapses ([Figures 5F and 5G](#)). The average distance comparison confirmed significant differences in the proximity of BIN1 with presynaptic markers as compared with the nearest-neighbor distance between BIN1 and postsynaptic or inhibitory presynaptic markers ( $p < 0.0001$ ). Next, we calculated how the proportion of estimated synaptic and BIN1 clusters differed from the chance occurrence, by shifting all of the coordinates belonging to a single synaptic cluster by a common vector (see [Method Details](#)). This randomization analysis confirmed a significant association between BIN1 and general presynaptic markers ([Figures 5H and S6C](#)). The compelling results from immuno-EM and dSTORM analyses indicate that BIN1 is preferentially but not exclusively enriched at the excitatory presynaptic terminals in hippocampal neurons.

### Reduced Synaptic Density in *Bin1* cKO Mice

To determine whether the loss of BIN1 alters the levels of synaptic proteins, we analyzed synaptosome fractions prepared from hippocampal samples of cKO mice and their littermate Ctl. Higher levels of the brain-specific BIN1:H were recovered in the non-PSD fractions relative to PSD fractions ( $p < 0.0001$ ), and there was a significant decrease in both fractions of cKO mice as compared with wild-type (WT) animals (t test, non-PSD fraction, BIN1:H  $p < 0.0001$ , PSD fraction, BIN1:H  $p = 0.0161$ ;  $n = 3$  Ctl and 3 *Syn*) ([Figure S7A](#)). BIN1:L isoforms were also recovered in the non-PSD synaptic fractions ( $p = 0.0205$ ) and were significantly reduced in cKO mice ( $p = 0.033$ ). The levels of postsynaptic marker proteins in hippocampal PSD fractions showed no difference in AMPAR phos-

phorylation (p831 or p845) or CaMKII activation (4-month sample:  $F_{(4, 55)} = 0.739$ ,  $p = 0.5694$ ;  $n = 8$ –12 Ctl, 3–7 *Syn*, and 5 *Emx*) ([Figure S7](#)). We also found no statistical differences between the genotypes in the abundance of 10 presynaptic and vesicle-associated proteins (4-month samples:  $F_{(18, 150)} = 0.4135$ ,  $p = 0.9835$ ;  $n = 9$  Ctl, 4 *Syn*, and 5 *Emx*) ([Figure S7C](#)). These results indicate that the loss of neuronal BIN1 expression has little effect on the overall abundance of several synaptic proteins.

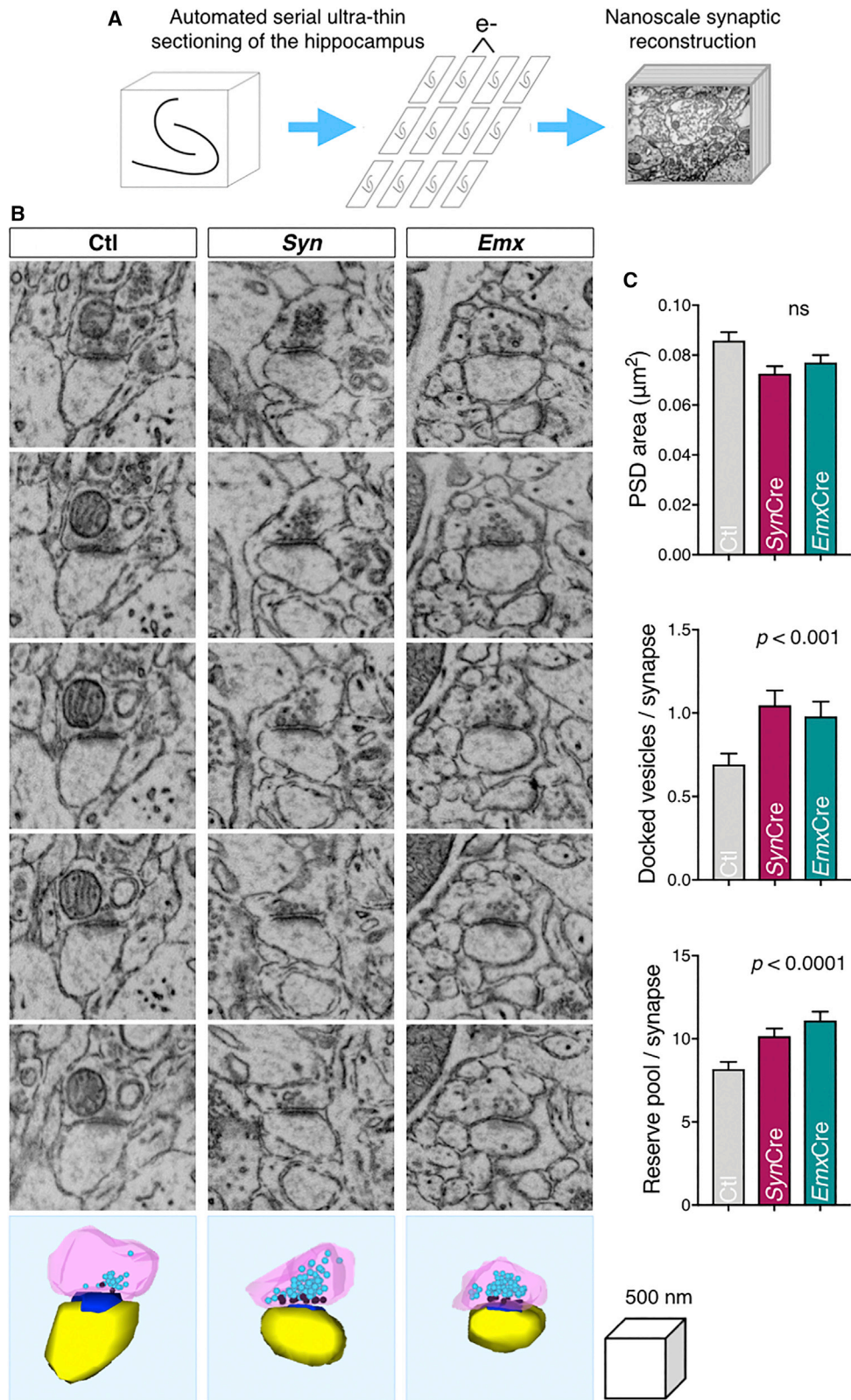
To detect more subtle changes in the localization of synaptic proteins, we performed confocal and super-resolution microscopy, focusing on the CA1 region of the hippocampus, to examine whether the number or the size of presynaptic and postsynaptic clusters were altered in cKO mice. As expected, there was a significant reduction in the number of BIN1 clusters in cKO mice as compared with Ctl mice ([Figures 1D and 1E](#)). In addition, we observed a significant reduction in Bassoon clusters that colocalized with BIN1 in *Syn* and *Emx* mice as compared to Ctl mice. Furthermore, our results showed a significant reduction in the cluster density of SNAP receptor (SNARE)-associated proteins Bassoon and Synaptoporin ([Figure 1F](#); [Table S2](#)). The cluster density of Munc13-1 was also significantly decreased in *Emx* mice, but the smaller decrease observed in *Syn* mice did not reach statistical significance. In addition, we observed a significant decrease in the cluster density of proteins associated with internalization at the presynaptic site: dynamin and Rab5. We also examined the postsynaptic proteins and observed no change in the cluster density of PSD95 ([Figure 1F](#)). The levels of GluA1 and pCamKII (Thr286) were also reduced, but the decrease in GluA1 was statistically significant only in *Emx* mice, and the reduction of pCamKII was significant only in *Syn* mice ([Figure 1F](#); [Table S2](#)). Synaptoporin and Munc13-1 are involved in presynaptic vesicle regulation and SNARE complex association ([He et al., 2017](#); [Li et al., 2017](#); [Singec et al., 2002](#)), and Bassoon is an organizer of the docking area at the presynaptic site ([Annamneedi et al., 2018](#); [Gundelfinger et al., 2016](#)). These results suggested that the loss of BIN1 leads to a potential defect in synaptic density and the organization of synaptic proteins, including SNARE proteins at the presynaptic sites.

We, therefore, performed super-resolution STED microscopy to precisely assess the cluster volumes of SNARE-associated proteins in the CA1 region of hippocampal sections. We observed a significant increase in the volume of Rab5<sup>+</sup> and Rab3a<sup>+</sup> clusters at presynaptic sites ([Figures 6A and 6C](#)). An earlier study reported an increase in Rab5-GFP puncta in cultured neurons, as

### Figure 6. Decrease in the Volume of SNARE-Regulating Proteins in cKO Mice

- (A) Representative images of Rab5 cluster intensity-coded by volume size (Imaris). The boxed region is shown at higher magnification in the bottom panels.
- (B) Rab3a clusters (left, coded by volume size) and two-color images (intensity-coded by volume) depicting the close proximity of Rab3a clusters to synaptotagmin (right).
- (C) Bar graphs representing the analysis of Rab5 and Rab3a cluster volumes in Ctl and cKO mice. ANOVA with Fisher's LSD post hoc test revealed a significant increase in the cluster volume sizes for Rab5 ( $F_{(2, 47322)} = 60.43$ ,  $p < 0.0001$ , Ctl versus *Syn*  $p = 0.0018$ , Ctl versus *Emx*  $p < 0.0001$ ) and Rab3a ( $F_{(2, 25397)} = 50.55$ ,  $p < 0.0001$ , Ctl versus *Syn* and Ctl versus *Emx*  $p < 0.0001$ ).
- (D) Stacked area plot of the difference in Rab5 cluster-size frequency distribution between Ctl and cKO. The difference in distribution was calculated as  $[(\text{number of Rab5 clusters/volume})^{\text{Ctl}} - (\text{number of Rab5 clusters/volume})^{\text{cKO}}]$ , with a bin size of  $0.001 \mu\text{m}^3$ . For clarity, only the clusters ranging from  $0.001$  to  $0.05 \mu\text{m}^3$  are shown in the plot. The inset shows an enlarged area of the plot.
- (E and F) Representative images of presynaptic (E, blue) and postsynaptic (F, red) protein clusters generated with Imaris and intensity-coded by volume (from  $0.001$  to  $0.05 \mu\text{m}^3$ ).
- (G) Heatmap representation of the mean variation of presynaptic and postsynaptic protein cluster volumes.
- See also [Figures S7 and S8](#) and [Table S3](#).





(legend on next page)



observed by conventional microscopy, following the knockdown of BIN1 expression (Calafate et al., 2016). Histogram analysis was performed to reveal better how the loss of BIN1 in cKO brains alters the size distribution of Rab5<sup>+</sup> clusters, as observed by super-resolution microscopy. This analysis revealed an overall increase in the volume of Rab5<sup>+</sup> clusters in *Syn* and *Emx* mice (Figure 6D). STED microscopy analysis also disclosed a significant decrease in the cluster size of synaptotagmin-1 ( $p < 0.0001$ ), a protein involved in presynaptic endocytosis and known to be associated with BIN1 (Yao et al., 2010). Furthermore, we observed a significant reduction in the volume of Munc13-1 clusters ( $p < 0.0001$ ), a SNARE-associated protein, whose cluster density was also reduced in *Emx* mice. However, we observed a significant increase in the cluster volume of synaptophysin ( $p = 0.0003$ ), an abundant synaptic vesicle membrane protein (Figure 6G; Table S3). Moreover, the cluster volume of synaptojanin ( $p < 0.0001$ ), a protein involved in ultra-fast endocytosis at the presynaptic site (Watanabe et al., 2018), was increased in the *Syn* model. Since immunoblot analysis of synaptosomes indicated no difference in the overall abundance of presynaptic proteins (Figure S7), the results of STED analyses would denote a difference in the local protein density at the synapse or indicate a change in the size of the vesicles where they are detected. The above results suggest a role for BIN1 in the organization of SNARE-regulating proteins and presynaptic vesicle dynamics. At the postsynaptic site, an increase in pCaMKII cluster volumes was recorded in cKO mice as compared with Ctl animals ( $p < 0.0001$ ). The volumes of PSD95 clusters remained unchanged, and the volumes for GluA1 clusters were significantly increased only in the *Emx* model ( $p < 0.0001$ ) (Figures 6F and 6G; Table S3). These results correlate well with the electrophysiology observations described above and suggest a role for BIN1 in the clustered organization of presynaptic machinery, and likely, compensatory changes occurring at the postsynaptic sites. Altered organizations of the presynaptic site, in particular, crucial active zone proteins in *Bin1* cKO mice, could account for the robust differences in release probability and vesicular depletion that we observed in CA1 synapses.

To examine the possibility that the loss of BIN1 in cKO mice alters neurotransmitter vesicle properties and/or distribution, we used serial section conventional EM. An automatic tape-collecting ultramicrotome was used to generate serial ultra-thin sections, which were then imaged at nanoscale resolution to reconstruct the presynaptic and postsynaptic compartments of CA3-CA1 axospinous synapses from Ctl, *Syn*, and *Emx* mice in their entirety (Kasthuri et al., 2015). This approach allowed us to measure the size of the PSD as well as count the number of docked neurotrans-

mitter vesicles and the reserve pool in each presynaptic terminal of Ctl and *Bin1* cKO mice. We found that the number of docked vesicles was significantly elevated in the cKO mice, as compared to their Ctl counterpart (Figure 7). Furthermore, the reserve pool within the synaptic bouton was significantly elevated in the cKO mice, as compared with Ctl animals (Figure 7). Although there was no measurable difference in Nq calculated from electrophysiological measurements, the slower time constant for the depletion of synaptic events observed in CA1 pyramidal neurons from cKO mice during sustained presynaptic stimulation is consistent—or at least not contradictory—with the EM data. Finally, the PSD areas of the different mouse lines did not differ from one another (Figure 7), a finding that is in agreement with the enrichment of BIN1 in the presynaptic compartment and the absence of apparent postsynaptic changes in cKO neurons. In aggregate, the electrophysiology and 3D EM analysis of hippocampal synapses show that the lack of BIN1 expression affects synaptic transmission and suggest that BIN1 may be an important component of presynaptic vesicular dynamics.

## DISCUSSION

Despite the identification of *BIN1* as the second most common susceptibility gene for LOAD, the underlying cellular mechanism by which BIN1 elevates the risk for AD remains poorly understood because the normal role of BIN1 in neurons has not been fully elucidated. Using behavioral and electrophysiological approaches, we characterized *Bin1* cKO models to reveal that the loss of BIN1 expression in neurons leads to impaired spatial memory consolidation and deficits in synaptic transmission, particularly at the level of presynaptic vesicle release. Using super-resolution microscopy and immuno-EM analyses, we show that BIN1 localizes to presynaptic sites in excitatory synapses and document reduced synaptic density, altered presynaptic protein clustering, and neurotransmitter vesicle distribution in *Bin1* cKO mice. These findings are highly relevant to synaptic dysfunction and cognitive decline in human LOAD because BIN1 has been implicated in memory impairments. For example, recent studies have uncovered a correlation between LOAD-associated *BIN1* rs744373 single nucleotide polymorphism and memory impairment (Franzmeier et al., 2019) or poor performance in high-load working memory task and lower functional connectivity in healthy carriers (Zhang et al., 2015) and memory impairment in patients with temporal lobe epilepsy (Bungenberg et al., 2016).

BIN1 is an essential component of the endocytic machinery as evidenced by (1) the recruitment dynamics of BIN1 to clathrin-coated pits before scission (Taylor et al., 2011); (2) the critical

### Figure 7. *Bin1* Deletion Increases Docked Neurotransmitter Vesicles and Reserve Pool but Does Not Affect Synapse Size

(A) Schematic representation of the tape-based pipeline for electron microscopic reconstruction of hippocampal synapses (Kasthuri et al., 2015). (B) Electron micrographs of serial sections through representative synapses are shown along with a 3D reconstruction of the dendritic spine head (yellow), the postsynaptic density (PSD; blue), and the presynaptic terminal (pink) containing docked neurotransmitter vesicles (dark pink) and the reserve pool of vesicles (aqua). (C) The PSD area was not significantly different among the three mouse genotypes, but the number of docked vesicles per synapse and the number of vesicles in the reserve pool was significantly higher among synapses in both cKO mouse strains as compared with Ctl by multivariate analysis of covariance. Multivariate analysis of variance (MANOVA) on PSD area:  $F_{(2, 436)} = 0.001$ ,  $p = 0.42$ ; multivariate analysis of covariance (MANCOVA) on docked vesicles:  $F_{(2, 436)} = 8.81$ ,  $p = 0.0002$ ; MANCOVA on reserve pool:  $F_{(2, 436)} = 9.53$ ,  $p = 0.00009$ . See also Figure S8.

function of the BIN1 SH3 domain in recruiting dynamin and regulating its self-assembly at the clathrin-coated pits (Owen et al., 1998); (3) BIN1 interaction with clathrin-coated pits-associated proteins Amph1, AP-2 complex, clathrin, synaptojanin 1, endophilin, and multiple other proteins involved in regulating endocytosis (Prokic et al., 2014). Moreover, the activation of the calcium-dependent phosphatase calcineurin coordinately dephosphorylates dynamin, synaptophysin, Amph1, and BIN1 during synaptic vesicle recycling (Marks and McMahon, 1998). However, the involvement of BIN1 in hippocampal synaptic transmission has not been experimentally assessed *in vivo*. By characterizing *Bin1* cKO mice, we demonstrate that the ablation of BIN1 expression in neurons results in (1) reduced vesicular release probability concomitant with an increase in the number of docked vesicles; and (2) an increase in the number of reserve pools of synaptic vesicles, which is consistent with a slower depletion time course of synaptic responses during sustained activity in excitatory synapses in the CA1 region of the hippocampus. These results indicate an altered synaptic vesicle turnover in *Bin1* cKO mice that could reflect a critical role of BIN1 in the regulation of both exocytic and endocytic mechanisms associated with synaptic transmission. The presynaptic changes in hippocampal synapses of *Bin1* cKO mice are schematically represented in Figure S8.

In the pan-neuronal *Syn* cKO model, the frequency of Cre-mediated recombination varied in different brain regions, with the hippocampus showing the striking loss of BIN1 expression in all subregions. Consistent with lineage tracing and fluorescence reporter analyses using the *Emx1*-IRES-Cre line (Gorski et al., 2002), BIN1 expression was ablated in forebrain glutamatergic neurons and glia in the *Emx1* cKO model. Both *Syn* and *Emx1* cKO models displayed impaired spatial learning, whereas the *Emx1* cKO showed an additional impairment in recognition memory. The latter finding likely represents the extent to which BIN1 expression is lost in the *Emx1* cKO cortex as compared with *Syn* cKO mice (Figure 1). Of note, unlike the concomitant loss of BIN1 expression in Amph1 KO mice reported previously (Di Paolo et al., 2002), neither the *Syn* nor the *Emx1* cKO model showed a diminution of Amph1 expression. Thus, the phenotypes we observe are entirely attributable to the primary loss of BIN1 expression. The selectivity of *Emx1*-IRES-Cre to target forebrain glutamatergic neurons suggests an essential role for BIN1 in excitatory synaptic transmission. This notion is experimentally supported by the enrichment of BIN1 at presynaptic sites observed by sub-diffraction-limited dSTORM microscopy and immuno-EM analyses. BIN1 localization is not exclusive at the presynaptic site, as we observed BIN1 immunofluorescence staining along axons and found occasional BIN1<sup>+</sup> immunogold particles in the periphery of the postsynaptic density in hippocampal CA3-CA1 synapses.

Genetic deletion of endocytic proteins results in vesicular recycling deficits (Lou et al., 2012; Rizzoli and Betz, 2005). However, in contrast to our findings in *Bin1* cKO mice, loss of endocytic proteins typically results in faster depletion of synaptic responses during sustained activity, which is a shared phenotype in mouse lines lacking synaptojanin1 (Cremona et al., 1999), dynamin1 (Ferguson et al., 2007), endophilins (Milosevic et al., 2011), and most important, Amph1 (Di Paolo et al.,

2002). It is important to note that loss of Amph1 expression also results in a drastic reduction in BIN1, which may indicate that phenotypes observed in *Amph1* KO neurons represent the effect of the loss of both Amph1 and BIN1 function (Di Paolo et al., 2002). Double KO mice lacking neuronal dynamin isoforms, dynamin1, and dynamin3, demonstrate slower vesicle depletion and reduced release probability similar to what we have observed in *Bin1* cKO mice (Lou et al., 2012). However, unlike in dynamin double KO mice (Lou et al., 2012), there was no detectable alteration in the size of the functional RRP in *Bin1* cKO mice (Figure 4). Therefore, it seems that the primary role of BIN1 in the presynapse is to regulate release probability. Based on the 3D EM data, the defect in vesicular release in *Bin1* cKO mice appears to be in a step that follows initial vesicle docking. The intriguing finding that BIN1 regulates exocytosis of the synaptic vesicles does not rule out an additional role for BIN1 in presynaptic endocytic processes. Following the knockdown of BIN1 expression in cultured neurons, Rab5 is overactivated, leading to increased endocytic flux and suggesting that the BIN1 function limits the kinetics of endocytosis (Calafate et al., 2016). In agreement, our STED analyses revealed an increase in the size of Rab5<sup>+</sup> clusters in the hippocampus of *Bin1* cKO mice.

Paradoxically, a recently published study reported that BIN1 regulates postsynaptic trafficking and glutamatergic signaling in cultured cortical neurons (Schürmann et al., 2019). Despite the focus on postsynaptic BIN1, the results of the published study included presynaptic BIN1 localization documented by the immuno-EM analysis of cortical tissue and images showing the colocalization between synapsin1 and BIN1 in cultured cortical neurons, as observed by immunofluorescence labeling. Moreover, although the authors focused on changes in mEPSC amplitude that were found to be statistically significant in their electrophysiological characterization, acute BIN1 knockdown increased the average AMPA mEPSC inter-event interval by 2-fold, which was found to be non-significant due to a large spread in the data (Schürmann et al., 2019). In contrast to the findings reported by Schürmann et al. (2019), our investigation of *Bin1* cKO mice reveals reduced synapse density and a significant role for BIN1 in synaptic vesicle dynamics, as well as no effect on quantal size based on a comprehensive characterization of multiple presynaptic parameters by hippocampal slice physiology. The presynaptic organization changes revealed by our investigation were confirmed by sophisticated 3D-EM reconstruction of hippocampal slices from two cKO models.

Our characterization of hippocampal synapses in *Bin1* cKO mice using super-resolution STED microscopy uncovered a significant reduction in the cluster volume of two presynaptic proteins that are essential for presynaptic vesicle docking and neurotransmitter release, namely synaptotagmin-1 and Munc13-1 (Li et al., 2017; Sakamoto et al., 2018; Südhof and Rizo, 1996). Moreover, there was a significant increase in pCaMKII cluster volume, but not that of GluA1 or PSD95. The increase of pCaMKII cluster volume could be the result of a compensatory response to presynaptic changes or represent a specific role for BIN1 in postsynaptic sites in the brain. Thus, the findings reported here reveal that the loss of BIN1 expression *in vivo* has a more robust impact in the presynaptic terminal,

leading to synaptic defect and memory impairments. We surmise that the reduction in the cluster volume of SNARE-related presynaptic proteins may contribute to the accumulation of docked vesicles and the lower release probability observed in CA1 synapses in *Bin1* cKO mice. A range of synaptic dysfunction, including presynaptic vesicular defects, have been described in mouse models of AD. For example, presynaptic functions have been detailed for APP (Priller et al., 2006), A $\beta$  (Fogel et al., 2014), PS1 (Barthet et al., 2018; Zhang et al., 2009), and APP/tau/PS1 (Chakroborty et al., 2019). Finally, a recent in-depth proteomic analysis found unbiased evidence to support presynaptic alterations that are more important than postsynaptic changes in the early stages of the disease in human AD ((Haytural et al., 2020).

Using cKO models, our study demonstrates the presynaptic BIN1 localization and a role for BIN1 in synaptic physiology that correlates with effects on learning and memory. BIN1 is known to interact with several proteins found in synaptic boutons, including Amph1, dynamin, clathrin, AP-2, endophilin, synaptojanin, and phospholipase D (Prokic et al., 2014). How BIN1 mechanistically participates in exocytosis at the presynaptic site and whether the dysregulation of the presynaptic vesicle dynamics is indirectly responsible for the altered cluster organization of SNARE-associated proteins in hippocampal neurons in *Bin1* cKO are key questions to address in future studies. In understanding the involvement of BIN1 in LOAD, only correlative data are available thus far to link BIN1 expression or the presence of LOAD-associated *BIN1* single nucleotide polymorphisms with Tau pathology and memory impairment (Franzmeier et al., 2019; Holler et al., 2014; Zhang et al., 2015). Large-scale analysis of transcriptomes and proteomes have identified an AD-associated decrease in BIN1 expression as part of the oligodendrocyte gene coexpression network dysfunction (McKenzie et al., 2017) and alternative splicing of *BIN1* in AD (Johnson et al., 2018). With the revelation that BIN1 may be an essential component of presynaptic vesicular dynamics, our report opens new paths for future investigation on the precise role of BIN1 as a risk factor in AD pathophysiology.

## STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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- Post-embedding immunogold EM
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- DATA AND CODE AVAILABILITY

## SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.celrep.2020.02.026>.

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## AUTHOR CONTRIBUTIONS

Conceptualization, P.D.R., T.N., N.K., D.A.N., A.C., and G.T.; Investigation, P.D.R., T.N., R.J.A., V.S., T.F.M., T.L.M., M.T.H., H.-N.S., S.V.K., V.P.B., N.K., and D.A.N.; Formal Analysis, P.D.R., T.N., N.Y.M., T.F.M., A.S., D.J.F., D.A.N., A.C., and G.T.; Writing – Original Draft, P.D.R., T.N., N.K., D.A.N., and G.T.; Writing – Review & Editing, P.D.R., D.A.N., A.C., and G.T.; Visualization, P.D.R., T.N., T.F.M., A.J.R., and G.T.; Supervision, N.K., D.A.N., A.C., and G.T.; Project Administration, G.T.; Funding Acquisition, P.D.R., R.J.A., D.A.N., A.C., and G.T.

## DECLARATION OF INTERESTS

The authors declare no competing interests.

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## STAR★METHODS

### KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Antibodies</b>		
Rabbit monoclonal anti-BIN1 (clone EPR13463)	AbCam	Cat# ab182562
Rabbit polyclonal anti-BIN1	Protein Tech	Cat# 14647-1-AP; RRID:AB_2243396
Rabbit polyclonal anti-BIN1 BSH3	Thinakaran Lab	<a href="#">De Rossi et al., 2016a</a>
Rabbit polyclonal anti-BIN1 B1415	Thinakaran Lab	This paper
Mouse monoclonal anti-Actin (clone 2D4H5)	Protein Tech	Cat# 66009-I-Ig; RRID:AB_2687938
Mouse monoclonal anti-Amph1 (clone 8)	Santa Cruz	Cat# SC21710; RRID:AB_673386
Rabbit polyclonal anti-APLP1 CT11	Thinakaran Lab	<a href="#">Thinakaran et al., 1998</a>
Rabbit polyclonal anti-APLP2 CT12	Thinakaran Lab	<a href="#">Thinakaran et al., 1995</a>
Guinea Pig polyclonal anti-Bassoon	Synaptic Systems	Cat# 141-004; RRID:AB_2290619
Mouse monoclonal anti-CaMKII (clone 6G9)	Santa Cruz	Cat# SC32288; RRID:AB_626787
Mouse monoclonal anti-CASK (clone K56A/50)	NeuroMab	Cat# O14936; RRID:AB_2750669
Mouse monoclonal anti-CNPase (clone 11-5B)	Millipore	Cat# MAB326; RRID:AB_2082608
Rabbit polyclonal anti-Flotillin-2	Thinakaran Lab	<a href="#">Meckler et al., 2010</a>
Mouse monoclonal anti-GluA1(clone N355/1)	NeuroMab	Cat# P19490; RRID:AB_2750785
Mouse monoclonal anti-MAP2 (clone HM-2)	Sigma	Cat# M4403; RRID:AB_477193
Mouse monoclonal anti-MBP (clone SMI-94)	Covance	Cat# SMI-94R; RRID:AB_510039
Mouse monoclonal anti-Mint2	BD Biosciences	Cat# M76120
Rabbit polyclonal anti-Munc13-1	Synaptic Systems	Cat# 126-102; RRID:AB_887734
Mouse monoclonal anti-NeuN (clone A60)	Chemicon	Cat# MAB377; RRID:AB_2298772
Rabbit monoclonal anti-pCaMKII (clone D21E4)	Cell Signaling	Cat# 12716S; RRID:AB_2713889
Rabbit polyclonal anti-pGluA1-831	Millipore	Cat# AB5847; RRID:AB_92077
Rabbit polyclonal anti-pGluA1-845	Millipore	Cat# AB5849; RRID:AB_92079
Mouse monoclonal anti-PSD95 (clone K28/43)	UC Davis	Cat# K28/43; RRID:AB_2750659
Mouse monoclonal anti-Rab3a (Clone9)	Santa Cruz	Cat# Sc-136050; RRID:AB_2300792
Rabbit monoclonal anti-Rab5 (clone C8B1)	Cell Signaling	Cat# 3547S; RRID:AB_2300649
Rabbit polyclonal anti-RIM1 $\alpha$	Synaptic Systems	Cat# 140-003; RRID:AB_887774
Mouse monoclonal anti-Synapsin1 (clone 46.1)	Synaptic Systems	Cat# 106 011BT; RRID:AB_2271476
Rabbit polyclonal anti-Synaptojanin1	Synaptic Systems	Cat# 145-103; RRID:AB_887820
Mouse monoclonal anti-Synaptophysin1 (clone SVP38)	Sigma	Cat# S5768; RRID:AB_477523
Rabbit polyclonal anti-Synaptoporin	Synaptic Systems	Cat# 102-003; RRID:AB_2619748
Chicken polyclonal anti-Synaptotagmin1	Synaptic Systems	Cat# 105-106; RRID:AB_2661782
<b>Experimental Models: Organisms/Strains</b>		
Mouse: C57BL/6J <i>Bin1</i> <sup>fl/fl</sup>	Dr. George C. Prendergast	Lankenau Institute for Medical Research
Mouse: B6.Cg-Tg(Syn1-cre)671Jxm/J	The Jackson Laboratory	JAX stock #003966, RRID:IMSR_JAX:003966
Mouse: B6.129S2- <i>Emx1</i> <sup>tm1(cre)Krl/J</sup>	The Jackson Laboratory	JAX stock #005628, RRID:IMSR_JAX:005628
<b>Software and Algorithms</b>		
FreezeFrame 4 software	Coulbourn	<a href="https://www.coulbourn.com/product_p/act-100a.htm">https://www.coulbourn.com/product_p/act-100a.htm</a>
EthoVision XT, v. 8.5 and v.14	Noldus	<a href="https://www.noldus.com/ethovision-xt">https://www.noldus.com/ethovision-xt</a> , RRID:SCR_000441
GraphPad Prism 8	Graphpad software	<a href="https://www.graphpad.com/scientific-software/prism/">https://www.graphpad.com/scientific-software/prism/</a> , RRID:SCR_002798

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<b>Continued</b>		
REAGENT or RESOURCE	SOURCE	IDENTIFIER
SPSS® software platform v. 21	IBM	<a href="https://www.ibm.com/analytics/spss-statistics-software">https://www.ibm.com/analytics/spss-statistics-software</a> , RRID:SCR_002865
Origin Pro 9.0 software	Origin Lab	<a href="https://www.originlab.com/">https://www.originlab.com/</a> , RRID:SCR_014212
Odyssey Infrared Imaging System v. 3.0.30	LI-COR Biosciences	RRID:SCR_013430
MiniAnalysis 6.0.3	Synaptosoft, Inc.	<a href="http://www.synaptosoft.com/MiniAnalysis/">http://www.synaptosoft.com/MiniAnalysis/</a> , RRID:SCR_002184
pClamp 10	Molecular devices	<a href="https://www.moleculardevices.com/products/software/pclamp.html">https://www.moleculardevices.com/products/software/pclamp.html</a> , RRID:SCR_011323
Leica LAS-X suite software	Leica	<a href="https://www.leica-microsystems.com/products/microscope-software/p/leica-las-x-ls/">https://www.leica-microsystems.com/products/microscope-software/p/leica-las-x-ls/</a> , RRID:SCR_013673
Fiji ImageJ 1.52	<a href="#">Rueden et al., 2017</a>	<a href="https://imagej.net/Fiji">https://imagej.net/Fiji</a> , RRID:SCR_002285
Spots colocalization (ComDet)	ImageJ plugin	<a href="https://imagej.net/Spots_colocalization_(ComDet)">https://imagej.net/Spots_colocalization_(ComDet)</a>
Thunderstorm	ImageJ plugin	<a href="https://github.com/zitmen/thunderstorm">https://github.com/zitmen/thunderstorm</a> , RRID:SCR_016897
Python v. 3.6.8	Python.org	<a href="https://www.python.org/">https://www.python.org/</a> , RRID:SCR_008394
SciPy v. 1.2.1	SciPy.org	<a href="https://www.scipy.org/">https://www.scipy.org/</a> , RRID:SCR_008058
Bitplane Imaris software v. 9.2	Oxford instrument	<a href="https://imaris.oxinst.com/packages">https://imaris.oxinst.com/packages</a> , RRID:SCR_007370
ATLAS 5 software	Fibics / Zeiss	<a href="https://www.zeiss.com/microscopy/int/products/microscope-software/atlas.html">https://www.zeiss.com/microscopy/int/products/microscope-software/atlas.html</a>
TrackEM2	ImageJ plugin	<a href="https://imagej.net/TrakEM2">https://imagej.net/TrakEM2</a>
dSTORM analysis code	This paper	<a href="https://github.com/nmasse/Storm-Microscopy-Analysis">https://github.com/nmasse/Storm-Microscopy-Analysis</a>

## LEAD CONTACT AND MATERIALS AVAILABILITY

Further information and requests for reagents should be directed to and will be fulfilled by the Lead Contact, Gopal Thinakaran ([thinakaran@usf.edu](mailto:thinakaran@usf.edu)).

This study did not generate new transgenic mouse lines. The antibody generated in this study is available from the Lead Contact without restriction.

## EXPERIMENTAL MODEL AND SUBJECT DETAILS

*Bin1<sup>fl/fl</sup>* animals were obtained from Dr. George C. Prendergast (Lankenau Institute for Medical Research) ([Chang et al., 2007](#)). *Syn1-Cre* (JAX stock #003966) and *Emx1-IRES-Cre* (JAX stock #005628) lines were obtained from The Jackson Laboratory (Bar Harbor, ME). We crossed *Bin1<sup>fl/fl</sup>* strain with *Syn1-Cre* or *Emx1-IRES-Cre* driver lines to generate *Bin1<sup>fl/fl</sup>;Syn1-Cre* (*Syn* mice), *Bin1<sup>fl/fl</sup>;Emx1-IRES-Cre* (*Emx* mice), and *Bin1<sup>fl/fl</sup>* or *Cre* littermates (referred to as Ctl mice). Female mice carrying the *Syn-Cre* transgene were used in breeding to prevent possible germline ablation of *Bin1*. Mice were maintained in a C57BL/6J background. All mice were housed under standard conditions with free access to food and water. All procedures related to animal care and treatment conformed to the policies of the Institutional Animal Care and Use Committee at the University of Chicago.

## METHOD DETAILS

### Behavioral tests

Behavioral assays were conducted using 6- to 10-week-old neuronal *Bin1* cKO mice and their littermate controls to test memory and motor functions. Male and female cohorts were used, but no sex differences were recorded, and the data presented in this study reports the effect of the genotype in both males and females. In addition, animals as young as P15 were subject to rotarod analysis. The number of animals used per genotype for individual experiments is indicated in the figure legends.

### Y-Maze spontaneous alternation

Testing occurred in a Y-shaped maze with three white, opaque plastic arms at a 120° angle from each other. Each mouse was placed in the middle of the symmetrical Y-maze and allowed to explore freely through the maze during an 8-min session. The movements of the mice were recorded using a 180° fisheye camera (Yi 4K Action and Sports Camera, 12MP) to avoid manipulator-stress induced interference, and analyzed *a posteriori*. An entry occurred when all four limbs were within the arm. The sequence and the total number of arm entries were recorded. An alternation is defined as entries into all three arms on consecutive occasions. The percentage of

alternations was calculated as: the number of triads containing entries into all three arms / maximum alternations) X 100. A minimum of 12 entries/mouse was required. The results were compared by ANOVA.

#### **Novel object recognition (NOR)**

NOR test was conducted in an open field arena (11.7" H X 15.7" W X 19.7" D) using objects that were similar in height and volume but were different in shape and appearance. During habituation, the animals were allowed to explore an empty arena. 24 h after habituation, mice were exposed to the familiar arena with two identical objects placed at an equal distance for a total period of 8 min. The next day, the mice were allowed to explore the open field in the presence of one familiar object (Old) and a novel object (Novel) to test long-term recognition memory. Mice that did not interact with the objects on day 1 were excluded from the study. The time spent exploring each object, and the discrimination index percentage were recorded and analyzed by ANOVA. The discrimination index was calculated as  $[(\text{time with Novel object} - \text{time with Old object}) / \text{total time interaction}]$ .

#### **Contextual fear conditioning**

The experiment was performed using an Actimetrics fear conditioning chamber, and the footshock was delivered by a Coulbourn Instrument shocker. The mouse movements were recorded by FreezeFrame 4 software. On day 1, each mouse was placed in a conditioning chamber and allowed to explore. After a 2 min habituation period, a brief, mild foot shock (1.0 mA, 2 s) was delivered through a grid floor at the bottom of the cage. After an interval of 2 min, the animal was presented again with the foot shock to strengthen the association. The mouse remained in the chamber for an additional 3 min before it was returned to its home cage. On the following day, the animal was brought to the same conditioning chamber, and its movements were recorded with a video camera for 5 min to test for contextual conditioning. The data were analyzed using FreezeFrame 4 software, which automatically monitors baseline motor activity and expression of fear. For contextual fear conditioning extinction, the animals were returned to the conditioning chamber (no foot shock) 3, 7, 15, 21, 28, 35, and 42 days after the initial fear conditioning and their movements were monitored for 5 min (as above) to determine extinction of the fear memory. Multiple unpaired t tests were used to compare cumulating freezing time between genotypes, with multiple comparison adjustments made using the Holm-Sidak method.

#### **Morris water maze spatial learning**

The water maze apparatus (a 122 cm tank) was filled with room temperature water (22–25°C), which was made opaque by the addition of non-toxic, white tempera paint. The water temperature was monitored using a thermometer before beginning the trials. Several high-contrast visual cues were placed around the apparatus in the room. Mice were placed on the platform for 10 s prior to the first training trial of each day to reduce stress. Over a period of 7 days, mice were trained to swim to a 14 cm diameter circular clear Plexiglas platform submerged 1.5 cm beneath the surface of the water and invisible to the mice while swimming. If they did not find the platform within the 120 s trial limit, they were guided to the hidden platform. The platform location (positioned in the middle of the Target (T) quadrant) was selected randomly and kept constant for each cohort throughout the training. On each trial, the mouse was placed into the tank at one of three designated start points (excluding the T quadrant) in a semi-random order. To ensure that memory differences are not due to lack of task learning, each mouse was undergoing a daily five-trial session (120 s for each with a 1 min interval). 24 h after the last training trial, the retention of the spatial memory was assessed in a probe trial, which consisted of a 60 s free swim in the pool without the platform. An overhead-mounted video tracking system (EthoVision XT, v.8.5 and v.14) was used to monitor and quantify different parameters during the probe trial. The corrected time spent in each quadrant (*total time per quadrant – the time of immobility in each quadrant*) was measured, and the results were compared by ANOVA. Other swimming parameters (speed, mobility, immobility, and velocity) were analyzed to control for each mouse's ability to swim. The following day, mice were submitted to a visible platform test to ensure no physical impairment influenced the performance during the test. The parameters were kept the same as for the training days, except that the location of the platform was indicated by a visible cue directly placed on the platform in the pool. The platform was placed at a different location than that during the probe trial to avoid any spatial memory-related success. The distance traveled to reach the platform was significantly longer in the *Emx* cohort ( $p < 0.01$ ). Hence, the swim distance was included as a covariate and the training data were analyzed by a linear mixed effects model (SPSS, v. 21). The probe trial data were analyzed using ANOVA.

#### **Rotarod**

Mice were examined for deficits in motor coordination and balance by the accelerating rotarod test and fixed rotarod test. Mice were placed on rotating drums (3-cm diameter) of an accelerating rotarod with 4 lanes (Harvard Apparatus). Adult mice were given 5 trials per day, with a 60 s inter-trial interval for 4 consecutive days. Younger mice were tested [5 trials per day, with a 60 s inter-trial interval] twice a week (P15 to 1 month) and then less frequently (i.e., once a week or once every two weeks) till approximately 4 months of age. The animal's latency to fall and rotation speed when the fall occurs were recorded using an automated system. The data were analyzed by a linear mixed effects model (SPSS). Mice were also tested on the rotarod using fixed speeds of 5, 10, 15, or 20 rpm for a maximum of 60 s per trial. The latency to fall was recorded as above. Each mouse was given 3 trials per speed, with a 60 s inter-trial interval and at least 5 min rest between the different speeds (Carter et al., 2001).

#### **DigiGait**

The DigiGait trials were performed using the DigiGait Imaging System (Mouse Specifics, Inc.). Each animal was placed on a stationary treadmill, which was then accelerated to the test speeds of 20, 30, and 40 cm/s. At their first trials, mice were allowed to run on the treadmill by gradually increasing the speed of the treadmill. Ten trials were attempted at each speed, and the animals were rested at least 10 s between trials. The successful trials (in which the animal is able to stay on the treadmill for > 5 s) were counted and recorded. If a mouse failed to gain speed and run, the trial was repeated after 5 min of rest with the treadmill held at the next lower speed.



### Immunoblot analysis

Hemibrain and sub-regions of the central nervous system were dissected and homogenized (20% weight/volume) in ice-cold lysis buffer [150 mM NaCl, 50 mM Tris-HCl pH 7.4, 0.5% NP-40, 0.5% sodium deoxycholate, 5 mM EDTA, 0.25 mM PMSF and protease inhibitor cocktail (Sigma)]. An equal volume of lysis buffer containing 2% SDS was then added to each sample and briefly sonicated. Protein samples were resolved on 4%–20% SDS-PAGE gels and analyzed by immunoblots using the antibodies listed in [Table S1](#). Rabbit polyclonal antiserum B1415 was generated against a fusion protein containing the BIN1 CLAP domain (residues encoded by exons 14 and 15). The blots were developed with IR680 anti-rabbit and IR800 anti-mouse secondary antibodies and imaged with the Odyssey Infrared Imaging System (Li-COR Biosciences). Quantification was performed using Fiji software ([Rueden et al., 2017](#)).

### Synaptosome fractionation

Synaptosome fractions were prepared following the protocol adapted from [De Rossi et al. \(2016b\)](#). Briefly, hippocampi were dissected from the brains of 4-month-old animals and resuspended in cold buffer containing 0.32 M sucrose and 10 mM HEPES at pH 7.4 and centrifuged twice (at 770 x g) to remove nuclei and large debris, followed by centrifugation at 12000 x g to obtain the synaptosome fraction. The synaptosomes were washed and pelleted in EDTA buffer to chelate calcium (4 mM HEPES, 1 mM EDTA, pH 7.4, 20 min at 12000 x g). The synaptosomes were then incubated in a low-Triton buffer for 30 minutes on ice (20 mM HEPES, 100 mM NaCl, 0.5% Triton X-100, pH 7.2) and centrifuged at 12000 x g to separate the postsynaptic density fraction (PSD; pellet) from the non-postsynaptic density fraction (non-PSD; supernatant). Protein samples were resolved on 4%–20% SDS-PAGE gels and analyzed by immunoblots using the antibodies listed in [Table S1](#). The blots were developed with IR680 anti-rabbit and IR800 anti-mouse secondary antibodies and imaged with the Odyssey Infrared Imaging System (Li-COR Biosciences). Quantification was performed using Fiji software ([Rueden et al., 2017](#)).

### Immunofluorescence staining

Mice were anesthetized by isoflurane inhalation before perfusion with PBS containing 4% paraformaldehyde and 4% sucrose. Brains were harvested and post-fixed overnight in the same fixative and then stored at 4°C in PBS containing 30% sucrose. Fifty  $\mu\text{m}$ -thick coronal sections were cut on a cryostat and processed for free-floating immunofluorescence staining. Brain sections were incubated with the indicated primary antibodies ([Table S1](#)) for 48 h at 4°C, followed by secondary antibodies for 3–4 h at room temperature. The antibodies were diluted in a 1X Tris-buffered saline solution containing 10% donkey serum, 3% BSA, and 0.25% Triton X-100. Nuclei were labeled using Hoechst stain (Molecular Probes) before mounting the sections on slides with Vectashield mounting medium (VectorLabs) for confocal microscopy or Prolong Diamond (Life Technologies) for STED microscopy.

### Electrophysiology

Acute hippocampal slices from 2–3-month-old Ctl and *Syn* mice were prepared as described previously ([Nomura et al., 2016](#)). Briefly, animals were deeply anesthetized with an intraperitoneal injection of xylazine (10 mg/kg) and ketamine (100 mg/kg). Mice were transcardially perfused with an ice-cold sucrose artificial cerebrospinal fluid (ACSF) solution containing (in mM): 85 NaCl, 2.5 KCl, 1.25  $\text{NaH}_2\text{PO}_4$ , 25  $\text{NaHCO}_3$ , 25 glucose, 75 sucrose, 0.5  $\text{CaCl}_2$ , and 4  $\text{MgCl}_2$ , including 10  $\mu\text{M}$  DL-APV and 100  $\mu\text{M}$  kynurenate, equilibrated with 95%  $\text{O}_2$  and 5%  $\text{CO}_2$ . Horizontal sections (350  $\mu\text{m}$ -thick) were made in the same ice-cold sucrose ACSF on a Leica Vibratome (Leica Microsystems, Inc). Slices were transferred to, and incubated in, a recovery chamber containing the same sucrose ACSF at 30–32°C for  $\sim$ 30 min, then the solution was gradually exchanged for a recovery ACSF containing (in mM): 125 NaCl, 2.4 KCl, 1.2  $\text{NaH}_2\text{PO}_4$ , 25  $\text{NaHCO}_3$ , 25 glucose, 1  $\text{CaCl}_2$ , and 2  $\text{MgCl}_2$ , including 10  $\mu\text{M}$  DL-APV and 100  $\mu\text{M}$  kynurenate at room temperature. After at least 1.5 h of recovery, slices were transferred to a recording chamber and visualized using Dodt contrast optics. Slices were continuously perfused with normal ACSF containing (in mM): 125 NaCl, 2.4 KCl, 1.2  $\text{NaH}_2\text{PO}_4$ , 25  $\text{NaHCO}_3$ , 25 glucose, 2  $\text{CaCl}_2$ , and 1  $\text{MgCl}_2$ , equilibrated with 95%  $\text{O}_2$  and 5%  $\text{CO}_2$ . For extracellular recordings, electrodes were manufactured from borosilicate glass pipettes and had tip resistances of 3–5  $\text{M}\Omega$  when filled with regular ACSF. Extracellular field excitatory postsynaptic potentials (fEPSPs) were elicited by an extracellular tungsten bipolar electrode placed in the *Stratum Radiatum*. For whole-cell recordings, recording electrodes were filled with an internal solution containing (in mM) 75 Cs-methanesulfonate, 60 CsCl, 1  $\text{MgCl}_2$ , 0.2 EGTA, 10 HEPES, 2 Mg-ATP, 0.3 GTP- $\text{Na}_2$ , 10  $\text{Na}_2$ -phosphocreatine, 10 TEA, 5 QX-314. Cells were voltage clamped at  $-70$  mV for EPSC recordings. EPSCs were pharmacologically isolated by the treatment of the slices with the GABA<sub>A</sub> receptor antagonist picrotoxin (PTX) (50  $\mu\text{M}$ ). The coefficient of variation (CV) of EPSCs was defined as the ratio of the standard deviation to the mean of the amplitude of EPSCs at given stimulation intensity in each cell. Miniature EPSCs (mEPSCs) were recorded in the presence of the  $\text{Na}^+$  channel blocker tetrodotoxin (TTX) (1  $\mu\text{M}$ ) in the extracellular solution. Data were collected and analyzed using pClamp 10 (Molecular Devices, Sunnyvale, CA) or MiniAnalysis (Synaptosoft Inc.) software. The recordings were discarded when the series resistance changed by  $> 20\%$  during experiments.

Depletion plots were constructed by normalizing the responses to the amplitude of the peak response during each train (60 s at 20 Hz) ([Vargas et al., 2014](#)). Depletion time constants for the fEPSP amplitude of 90–10% (10% increment) were measured in each train. Data were collected and compared at each point between genotypes. Recovery plots were generated by normalizing the responses to the 1<sup>st</sup> responses of the depletion trains and were fit logarithmically to the equation  $y = \ln(mx) + b$  where  $m$  and  $b$  represent the slope and the y axis intersect obtained in each recording (5 s at 2 Hz) ([Vargas et al., 2014](#)).  $m$  was measured in each recording, and collected data were compared between genotypes. The readily releasable pool (RRP) was estimated by

analyses of the cumulative amplitudes of the fEPSPs during trains of stimuli used for the depletion paradigm. The RRP was quantified as  $Nq$ , where  $N$  and  $q$  represent the total number of releasable vesicles and the quantal size of synaptic responses, respectively.  $Nq$  can be estimated from the zero-time y axis intersect of the linear regression fit to the cumulative fEPSP amplitude plot (Fernandes et al., 2015; Fioravante and Regehr, 2011).  $Nq$  was analyzed in each recording, and grouped data were compared between genotypes. Statistical analyses were conducted with GraphPad Prism 8, SPSS, and Origin Pro 9.0 software. Two sample comparisons were made using the Kolmogorov–Smirnov (K-S) test or Mann-Whitney test. For multiple comparisons, repeated two-way analysis of variance (ANOVA) followed by post hoc Bonferroni correction or a linear mixed effects model was employed. Differences were considered significant when  $p < 0.05$ . Data are shown as mean  $\pm$  SEM.

### Direct Stochastic Optical Reconstruction Microscopy (dSTORM)

Super-resolution images were acquired on a Leica SR Ground State Depletion 3D / 4 color TIRFM microscope with an Andor iXon Ultra EM-CCD camera (Andor Technology PLC). DIV18-21 rat primary neurons were fixed for 20 min in 4% PFA and 4% sucrose. Primary antibodies were incubated overnight at 4% in PBS containing 10% donkey serum, 3% BSA, and 0.25% Triton X-100. Secondary antibodies were incubated at RT for 3 h in the same buffer. After 3 washes in PBS, the cells were re-fixed with 4% PFA for 5 min. The coverslips were then washed over a period of 2 days at 4°C in PBS to remove the non-specific binding of the secondary antibodies. Coverslips were mounted extemporarily in a monoethanolamine (MEA) solution to limit oxidation of the fluorophores during image acquisition. The areas of capture were blindly selected by direct observation in DIC. Images were acquired using a 160X (NA 1.43) objective in the TIRF mode North direction with a penetration of 200 nm. Far-red channels (Alexa 647 or 660) were acquired using a 642 nm laser and 664LP filter. Red channels (Alexa 568 or 555) were acquired using a 532 nm laser and a 582–636 nm band-pass filter. Green channel (Alexa 488) was acquired using a 488 nm laser and a 503–547 nm bandpass filter. Images were acquired using GSD 3D astigmatism mode in the Leica LAS-X suite software. The irradiation intensity was adjusted until the single-molecule detection reached a frame correlation  $< 0.25$ . Detection particle threshold was defined between 20–60 photons depending on the marker and adjusted to obtain a number of events per frame between 0 and 25. The exposure was maintained at 7 ms, and the EM gain was set at 300. The power of depletion and acquisition was defined for each marker and kept constant during acquisition. The number of localizations collected was maintained constant per markers and between experiments. At least 4 independent cultures or coverslips were imaged per marker.

### GSD Super-resolution image processing and analysis

Raw GSD images were processed using a custom-made macro in Fiji to remove background by subtraction of a running median of frames (300 renewed every 300 frames) and subtracting the previously processed image once the background was removed (Hoo-gendoorn et al., 2014). A blur (0.7-pixel radius) per slice prior to median subtraction was applied to reduce the noise. Finally, a 3D Gaussian blur (0.7-pixel radius) was applied to decrease noise further and improve fitting certainty. These images were then processed using Thunderstorm plugin in ImageJ (Ovesný et al., 2014). Blink detection was performed using a Wavelet filter (B-spline, order 3/scale2.0). The molecules were localized using the centroid of connected components, and the peak intensity threshold was determined per marker/dye to maintain an XY uncertainty  $< 50$ . Sub-pixel localization of molecules was performed using PSF elliptical Gaussian (3D astigmatism) and least squared fitting methods with a fitting radius of 5 pixels and an initial sigma of 1.6 pixels. 3D location was mapped from  $-500$  nm to  $+500$  nm from the focus point.

To analyze the localization of BIN1 and synaptic marker clusters, we used the Density-Based Spatial Clustering of Applications with Noise (DBSCAN) data clustering algorithm (Ester et al., 1996; Yue et al., 2004). In this method, a coordinate belongs to a cluster when a minimum number of neighbors required to form a dense region (minPts) exists within a defined distance  $\epsilon$ . We set minPts to 3 and  $\epsilon$  to 50 nm, although values of minPts between 2 and 4, and values of  $\epsilon$  between 50 and 100 nm produced qualitatively similar results. We discarded clusters that contained fewer than 50 coordinates, or if the diameter (as measured between the two furthest apart coordinates) was less than 25 nm. Furthermore, to ensure we only analyzed clusters that were confined to a single synaptic bouton, we discarded clusters with diameters greater than 1000 nm. For BIN1 and synaptic marker clusters, we calculated the centroid of each cluster and then calculated the distance between each BIN1 cluster and each synaptic marker cluster. We considered two clusters to be in close proximity if the distance between their centroids was less than 30 nm, and calculated “experimental colocalization” as the proportion of BIN1 clusters found with a synaptic marker cluster within 30 nm distance, averaged across all the samples; since it is a proportion, the values range from 0–1. We also estimated the cluster proximity by setting the distance as 40 nm or 50 nm. We next measured how the proportion of estimated synaptic and BIN1 clusters differed from chance. To do so, we shifted all coordinates belonging to a single synaptic cluster by a common vector, in which the x, y, and z components of this common vector were randomly sampled from a uniform distribution from  $-250$  to  $250$  nm. We randomly shifted all synaptic clusters and then recalculated the proportion of synaptic clusters that were co-located with a BIN1 cluster. We repeated this process 100 times to obtain a null distribution of colocalization proportions. We then calculated the z-score between the experimental colocalization proportion and the null distribution:

$$z = (\text{experimental colocalization} - \text{mean}[\text{null distribution}] / (\epsilon + \text{std}[\text{null distribution}]),$$

where epsilon =  $1e-4$  (results were qualitatively similar for different values of  $\epsilon$ ). Negative z-scores were set to zero.

To confirm these results, images were also analyzed using Bitplane Imaris software v.9.2 (Andor Technology PLC). Volumes for each marker were generated using smooth surfaces with details set up at 0.005. The diameter of the largest sphere was set up at 0.75  $\mu\text{m}$ . A threshold background subtraction method was used to create the surface, and the threshold was calculated and applied to all the images of the same experiment. Surfaces were then filtered by setting up the number of voxels > 10 and area between 0.01–1  $\mu\text{m}^2$ . The nearest neighbor distance was processed using the integrated distance transformation tool in Imaris. Distances were then organized and statistically analyzed using median comparison and ANOVA with Fisher's LSD post hoc test. The nearest neighbor distances greater than 1000 nm were excluded to facilitate calculating the frequency distribution of clusters located within 500 nm from each side of the synaptic cleft. A mask of the spines was generated based on CaMKII staining using smooth surfaces with details set up at 0.1. Imaris filament tracing was performed for MAP2 and BIN1.

### Confocal image acquisition and quantification

Confocal images were acquired on Leica SP5 or Leica SP8 microscopes using 10X (NA 0.4), 20X (NA 0.75), 40X (NA 1.25), 60X (NA 1.4), or 100X (NA 1.4) objectives. For Figure 1B, four tiled images each were acquired using a 10X objective and stitched using LAS-AF software (Leica). Images acquired on a Leica SP5 using a 100X (NA 1.4) objective were used to quantify the density of synaptic markers. Images were acquired in the CA1 region  $\sim$ 50  $\mu\text{m}$  from the soma at the bifurcation of the apical dendrite of pyramidal cells, using the same parameters for Ctl and cKO animals. Z-stacks of fifty 2048 X 2048-pixel images were acquired at a scan zoom of 2.5, yielding 30.28 nm pixels.

Confocal microscopy data were quantified from at least 2 image stacks acquired from each of two immunostained sections per animal in at least 3 animals per genotype. Confocal Z stacks were deconvolved using Huygens Professional software (Scientific Volume Imaging) and analyzed using the built-in particle analysis function in Fiji (Rueden et al., 2017). The size of the particles was defined according to previously published studies (MacGillavry et al., 2013; Nair et al., 2013). To assess the number of clusters, images were thresholded (same threshold per marker and experiment), and a binary mask was generated. As image pixel size was 0.000917  $\mu\text{m}^2$ , a low size threshold of 0.01  $\mu\text{m}$  diameter was applied to eliminate individual pixels from the quantification. A high pass threshold of 0.4  $\mu\text{m}$  diameter was applied to avoid the quantification of non-clustered structures. For the analysis, the number of clusters per stack was summed. The density was normalized by the control of each experiment to eliminate variability between experiments. The densities were compared by one-way ANOVA with Fisher's LSD post hoc test. The Spots Colocalization plugin ComDet v.0.3.7 was used to analyze the % colocalization between BIN1 and Bassoon clusters, utilizing the counting mask generated by the cluster quantification. The detection of particles was set up independently for each channel (channel 1: Bassoon; channel 2: BIN1). The default parameters were applied to quantify the colocalization: particle size 3, intensity threshold 3, the maximum distance between colocalized particles 5 pixels (= 150 nm). The results were expressed as the % of Bassoon clusters colocalizing with BIN1 clusters.

### Super-resolution STED image acquisition and quantification

Super-resolution STED images of BIN1 and synaptic markers were acquired on a Leica SP8 3D, 3-color gated STED laser scanning confocal microscope. Images were acquired in the CA1 area, at the bifurcation of the apical dendrite of pyramidal cells. A 775 nm depletion laser was used to deplete both 647 and 594 dyes, and a 592 nm laser was used for the depletion of 488 dyes. The powers used for depletion lasers, the excitation laser parameters, and the gating parameters necessary to obtain STED resolution were assessed on control sections, and the same parameters were applied for sections from Ctl and cKO mice. 3  $\mu\text{m}$ -thick Z stacks of 1248 X 1248-pixel images at 50 nm step size were acquired at 8 kHz bidirectional scan rate with a line averaging of 32 and 2–4 frame accumulation, using a 100X (1.45) objective with a digital zoom factor of 5, yielding 18.64 nm pixels.

STED microscopy data were quantified from at least 2 image stacks acquired from each of two immunostained sections per animal in at least 3 animals per genotype. The STED images were deconvolved using Huygens Professional software using the gated 3D-STED module (Scientific Volume Imaging). The images were analyzed using Imaris software. Each image was cropped 3D X = 50 to 1200 and Y = 50 to 1200 and Z = 1 to 60 to obtain a 21.5  $\mu\text{m}$  X 21.5  $\mu\text{m}$  X 3  $\mu\text{m}$  block to avoid edge artifacts. Volumes for each marker were generated using smooth surfaces with details set up at 0.01  $\mu\text{m}$ . The diameter of the largest sphere was set up at 0.75  $\mu\text{m}$ . Threshold background subtraction methods were used to create the surface, and the threshold was calculated with control images and applied to all the images of the same experiment. Surfaces were then filtered by setting up the number of voxels > 10 and area between 0.01–1  $\mu\text{m}^2$ . The volume of each surface was then exported and analyzed by ANOVA with Fisher's LSD post hoc test. Each surface was then color-coded for the volume in Imaris using the same scale in both control and cKO images for the purpose of illustration. The extent of colocalization between two proteins was determined by calculation of the nearest neighbor distance using the integrated distance transformation tool in Imaris. Distances were then organized and statistically analyzed using ANOVA with Fisher's LSD post hoc test.

### Post-embedding immunogold EM

Reagents used for EM were from Electron Microscopy Sciences (Hatfield, PA) unless otherwise noted. Tissue was processed as described previously (Neuman et al., 2015). Briefly, mice were perfused transcardially with 4% paraformaldehyde/0.5% glutaraldehyde, after which the hippocampus was dissected and sectioned into 300  $\mu\text{m}$ -thick slices. Slivers containing CA1 stratum radiatum were isolated from these sections and then plunge-frozen in liquid propane ( $-184^\circ\text{C}$ ), and then processed using freeze-substitution,

first stained *en bloc* in 1.5% uranyl acetate at  $-90^{\circ}\text{C}$ , then infiltrated with Lowicryl HM20 (Electron Microscopy Sciences) at progressively escalated temperatures, and finally cured with ultraviolet light at  $0^{\circ}\text{C}$ . Ribbons of serial sections (20–55) were then mounted on formvar-coated, carbon-coated nickel slotted grids, and immunolabeled using pAb BSH3 (10  $\mu\text{g}/\text{ml}$ ) overnight at  $4^{\circ}\text{C}$ , followed by secondary immunogold staining with 1:20 dilution of 10-nm immunogold particles (Ted Pella, Inc.). Images were acquired with a Zeiss Sigma SE-EM in STEM mode with an accelerating voltage of 29 kV at 1.8 nm/pixel.

### Conventional electron microscopy

Large-volume electron microscopy datasets from Ctl and cKO mouse brains were performed as previously described (Hua et al., 2015; Kasthuri et al., 2015). Briefly, mice were anesthetized and transcardially perfused with buffered aldehydes (2% paraformaldehyde and 2.5% glutaraldehyde in 0.1 M cacodylate buffer). The brain was dissected out and post-fixed overnight at  $4^{\circ}\text{C}$  in a similar fixative solution as described above. The brain was then coronally sectioned at 500  $\mu\text{m}$  thickness using a vibratome, and the CA1 region of the hippocampus was removed. Tissues were stained with heavy metals (Hua et al., 2015), dehydrated, infiltrated with epon resin, and cured in an oven at  $60^{\circ}\text{C}$  for 48 h. The cured block was trimmed, and then 200–300 ultra-thin sections (55 nm) from CA1 stratum radiatum were collected on Kapton tape using the automatic tape-collecting ultramicrotome (ATUM). The serial sections were imaged using the backscattered electron detector on Gemini SEM 300 (Carl Zeiss) equipped with ATLAS 5 software (Fibics) (Kasthuri et al., 2015). The high-resolution (6 nm pixel size) images were aligned using trackEM2 and analyzed using ImageJ. A total of 444 asymmetric axospinous synapses were analyzed, with measured parameters being PSD area, number of docked vesicles, and the number of vesicles in the reserve pool (both of the latter in the presynaptic terminal). PSD area was estimated as the product of section thickness (55 nm) and the sum, total length of the PSD. Docked and reserve pool vesicles were defined using the parameters as defined in Schikorski and Stevens (1997). Briefly, docked vesicles were those neurotransmitter vesicles making contact with the presynaptic active zone, immediately apposed to the PSD. Reserve pool vesicles were all other vesicles present in the presynaptic terminal in sections harboring a PSD profile on the dendritic spine head. Quantifications were performed blinded with respect to genotype, and the results were statistically analyzed by multivariate analysis of covariance.

### QUANTIFICATION AND STATISTICAL ANALYSIS

The software used for behavior analysis, electrophysiology, image quantification are listed under each section above. Statistical analyses were conducted with GraphPad Prism 8, SPSS v.21, and Origin Pro 9.0 software. Two sample comparisons were made using the Kolmogorov–Smirnov (K-S) test or Mann-Whitney test. Analysis of variance (ANOVA), two-way ANOVA, or repeated two-way ANOVA were used for multiple comparisons. In some experiments, the data was analyzed using MANOVA, MANCOVA, or a linear mixed effects model, as stated under Results or figure legends. Individual tests and post hoc analyses can be found in the Results and figure legends. ANOVA F statistic, degrees of freedom between groups (DFn) and within groups (DFd), and the  $p$  value are listed in results, figure legends, or supplementary tables. The mean  $\pm$  SEM are plotted in each graph. Differences were considered significant when  $p < 0.05$ . The bar graphs depict mean  $\pm$  SEM and the violin plots depict the median and quartiles. \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ , \*\*\*\* $p < 0.0001$ .

### DATA AND CODE AVAILABILITY

The code generated during this study are publically available at <https://github.com/nmasse/Storm-Microscopy-Analysis>



**Cell Reports, Volume 30**

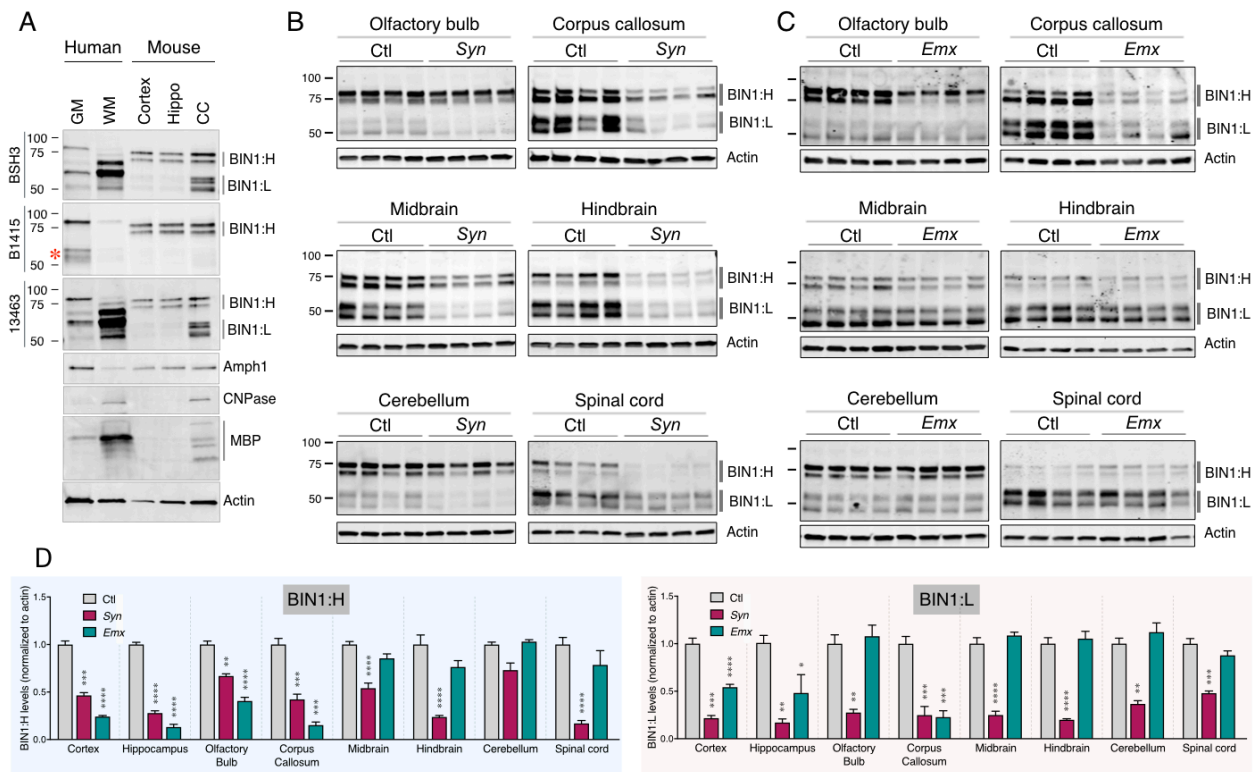
## **Supplemental Information**

### **Neuronal BIN1 Regulates**

### **Presynaptic Neurotransmitter Release**

### **and Memory Consolidation**

**Pierre De Rossi, Toshihiro Nomura, Robert J. Andrew, Nicolas Y. Masse, Vandana Sampathkumar, Timothy F. Musial, Ari Sudwarts, Aleksandra J. Recupero, Thomas Le Metayer, Mitchell T. Hansen, Ha-Na Shim, Sofia V. Krause, David J. Freedman, Vytas P. Bindokas, Narayanan Kasthuri, Daniel A. Nicholson, Anis Contractor, and Gopal Thinakaran**



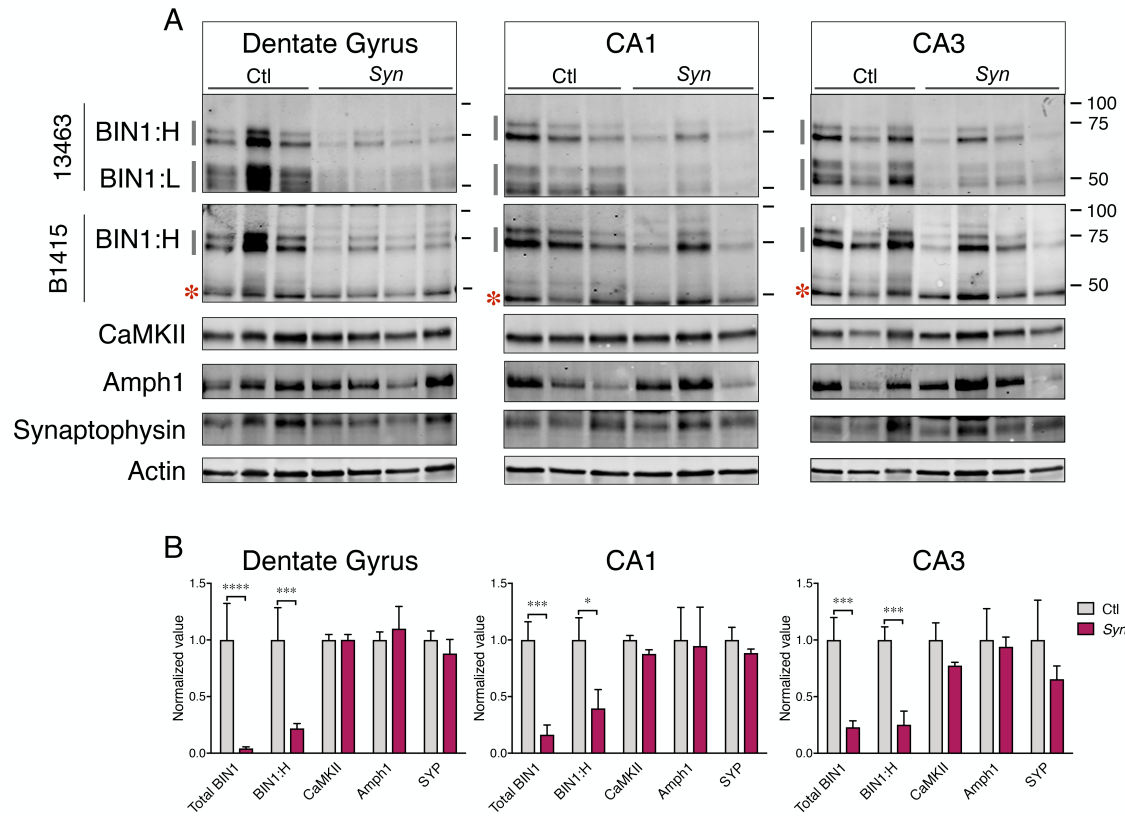
**Figure S1.** Related to Figure 1: **Analysis of BIN1 expression in brain.**

**(A)** A comparison of BIN1 expression in human and mouse brain. Antibodies BSH3 and 13463 are capable of detecting all BIN1 isoforms, whereas pAb B1415, raised against the CLAP domain, can only recognize the BIN1 isoforms containing the CLAP domain (BIN1:H). Only low levels of BIN1:H isoforms were detected in the grey matter in the human brain, whereas they are relatively more abundant in the mouse brain. The isoforms lacking the CLAP domain (BIN1:L) are abundant in the human brain white matter and mouse brain corpus callosum. Amph1 (neurons) and CNPase and MBP (oligodendrocytes) were used as markers. An *asterisk* indicates a non-specific signal observed in human brain grey matter.

**(B)** Microdissection analysis of BIN1 levels in the central nervous system regions of *Syn* mice and littermates. BIN1 was detected using pAb 14647 (detects all BIN1 isoforms), and actin was used as the loading control.

**(C)** Microdissection analysis of *Emx* mice and littermates.

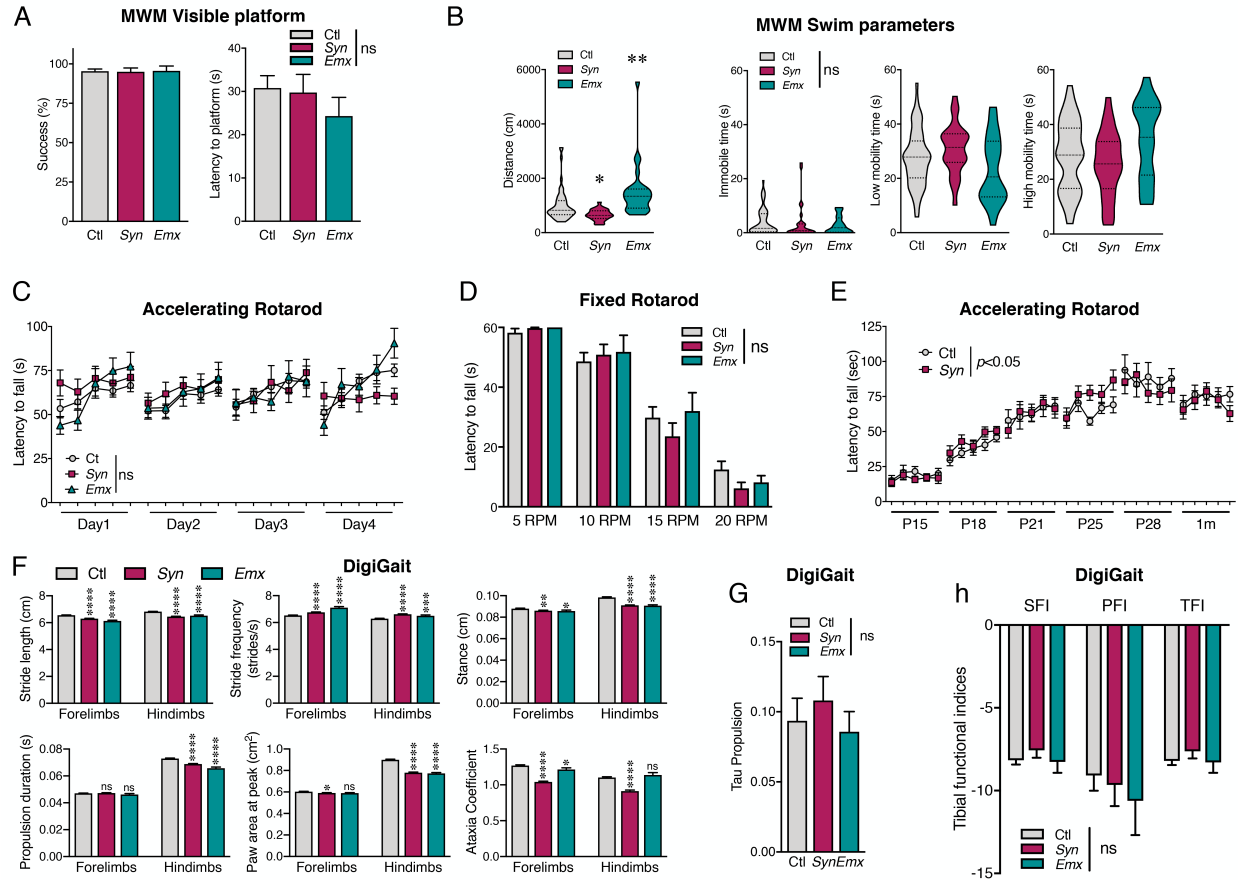
**(D)** Quantification of relative BIN1:H and BIN1:L isoforms normalized to actin. The mean  $\pm$  SEM are plotted in each graph.  $*p < 0.05$ ,  $**p < 0.01$ ,  $***p < 0.001$ ,  $****p < 0.0001$ .



**Figure S2.** Related to Figure 1: **Microdissection analysis of BIN1 levels in the hippocampus of Ctl and *Syn* mice.**

(A) Western blot analysis of BIN1 and neuronal marker levels in the dentate gyrus (left), CA1 (middle), and CA3 (right) regions. BIN1 was analyzed using antibodies 13463 and B1415. Red *asterisks* indicate a non-specific protein detected by antibody B1415. The levels of CaMKII, Amph1, and synaptophysin were also quantified.

(B) Quantification of relative protein levels normalized to actin. The levels of BIN1 were significantly reduced in *Syn* mice, but the levels of CaMKII, synaptophysin, and Amph1 were not altered in cKO mice.



**Figure S3.** Related to Figure 2: **Neuronal knock-out of *Bin1* does not induce physical impairments.**

(A) Graphs representing performances (success and time to escape) recorded during Morris water maze visible platform test (see fig1e). The mean  $\pm$  SEM are plotted. No significant difference was observed between the groups by ANOVA. Success:  $F_{(2,104)}=0.01446$ ,  $p=0.9856$ ; latency:  $F_{(2,104)}=0.8371$ ,  $p=0.4358$ ;  $n=52$  Ctl, 28 *Syn* and 27 *Emx*.

(B) Violin plots depict the distribution of swimming distance and mobility in the probe trial. Swimming parameters during the probe trial, analyzed by ANOVA, revealed significant differences between the Ctl and cKO groups in the swim distance ( $F_{(2,106)}=11.99$ ,  $p<0.0001$ ). There were also significant differences in the duration of low-mobility ( $F_{(2,106)}=3.738$ ,  $p=0.027$ ) and duration of high-mobility ( $F_{(2,106)}=30127$ ,  $p=0.0479$ ), but Tukey's post-hoc analysis revealed that the significance existed only between the two cKO groups and not the Ctl group.

(C) The line graph represents the performance (latency to fall) of 6 weeks-old mice on the accelerating rotarod (mean  $\pm$  SEM). No significant difference between the groups was observed ( $F_{(2,68)}=0.166$ ,  $p=0.847$ ,  $n=39$  Ctl, 19 *Syn*, and 13 *Emx*).

(D) The graph represents performance (latency to fall) of 6 weeks-old mice on the fixed rotarod (mean  $\pm$  SEM). No significant difference between the groups was observed ( $F_{(2,68)}=0.303$ ,  $p=0.74$ ,  $n=39$  Ctl, 19 *Syn*, and 13 *Emx*).

(E) The line graph represents the performance of a cohort of *Syn* mice on the accelerating rotarod (mean  $\pm$  SEM) during postnatal development. A small but significant difference between the groups was observed ( $F_{(1,993)}=5.135$ ,  $p=0.024$ ;  $n=18$  Ctl and 19 *Syn*).

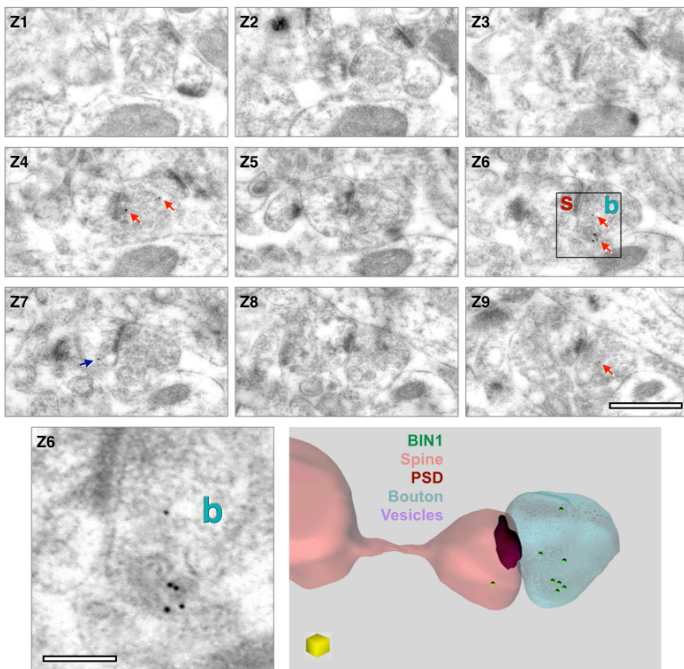


**(F)** Significant differences between the groups were observed on multiple parameters recorded by DigiGait (stride length and frequency, stance, propulsion duration, paw area at peak, and ataxia coefficient) in 1.5-month-old mice are plotted (mean  $\pm$  SEM). However, these differences were too small to impact on the animals' general mobility.

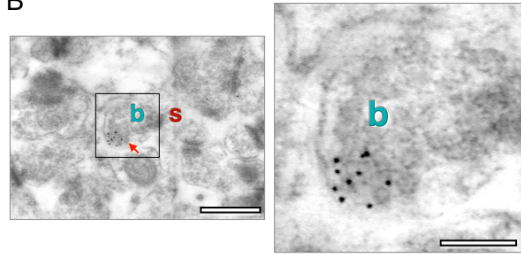
**(G)** The graph represents the Tau propulsion (mean  $\pm$  SEM). No difference between the groups was recorded (ANOVA,  $F_{(2, 94)}=0.3009$ ,  $p=0.7409$ ).

**(H)** The graph represents sciatic, peroneal, and posterior tibial nerve parameters (Bain et al., 1989) recorded using DigiGait analysis (mean  $\pm$  SEM). No difference between the groups was recorded (ANOVA,  $F_{(2, 282)}=0.4422$ ,  $p=0.6430$ ).

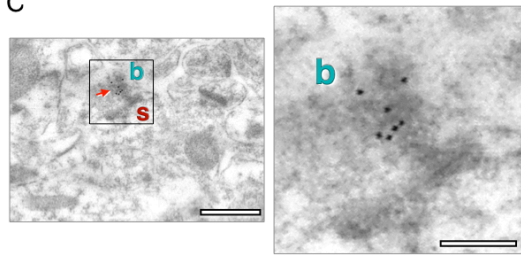
A



B



C

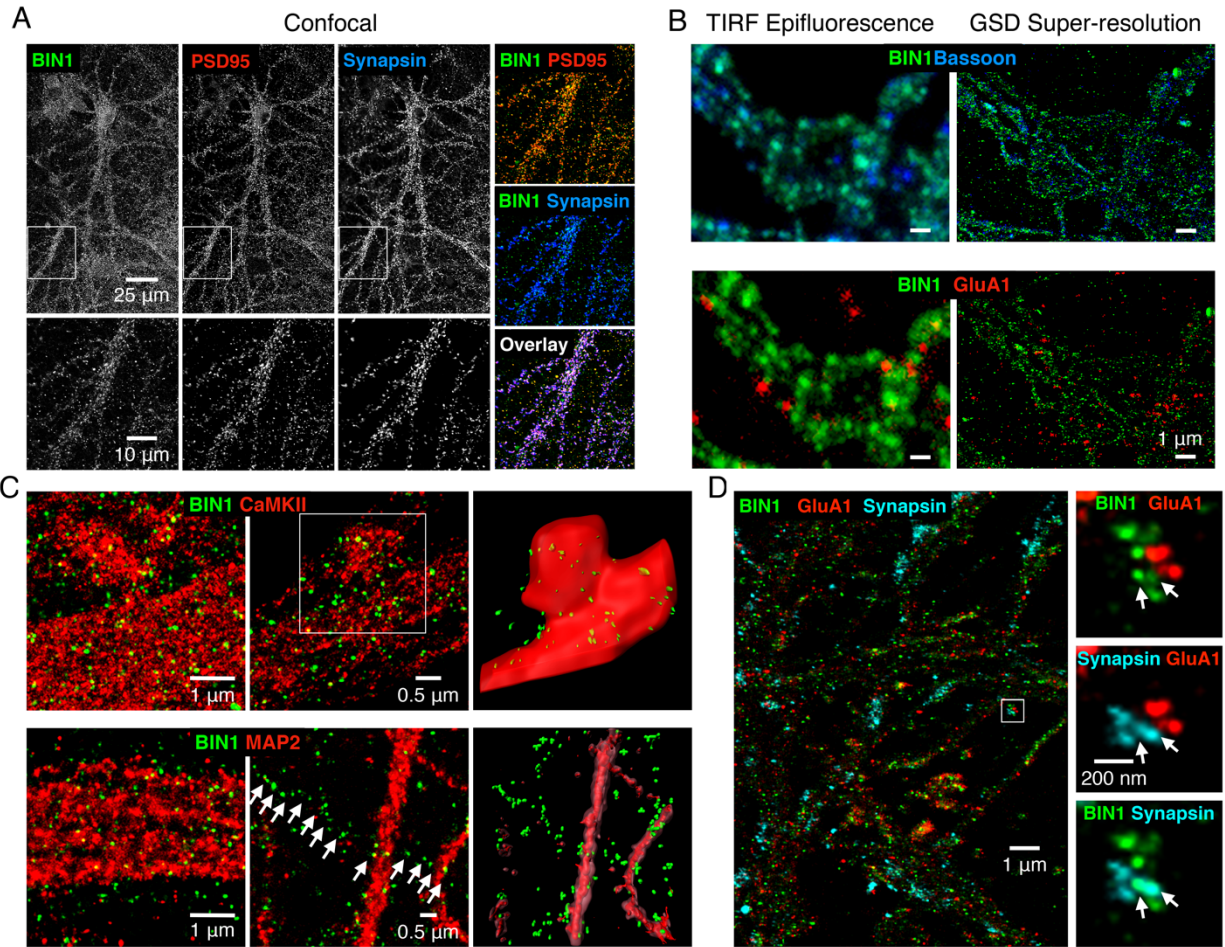


**Figure S4. Related to Figure 5: Presynaptic enrichment of BIN1.**

**(A)** Micrographs of nine serial sections through an axospinous synapse in CA1 *stratum radiatum* (Z1-Z9) illustrating that BIN1 can be detected at both pre- and postsynaptic sites. Note that BIN1 immunogold localization in the presynaptic axonal bouton (*b*) is considerably higher than that in the postsynaptic dendritic spine (*s*). Eight immunogold particles for BIN1 can be seen in the presynaptic terminal (red arrows), as compared to one in the postsynaptic dendritic spine (blue arrow). The boxed region of the presynaptic terminal in Z6 is shown at a higher magnification at the bottom (left). At the bottom right is a 3-dimensional reconstruction of the synapse and BIN1 immunogold particles as visualized in Z1-Z9.

**(B and C)** Two more examples of presynaptic localization of BIN1, captured in micrographs of single sections through a small non-perforated synapse (top) and a larger perforated synapse (bottom). Panels on the right are higher magnification images of the regions marked by the boxes.

Scale bars main images: 500 nm; higher magnification images: 150 nm.



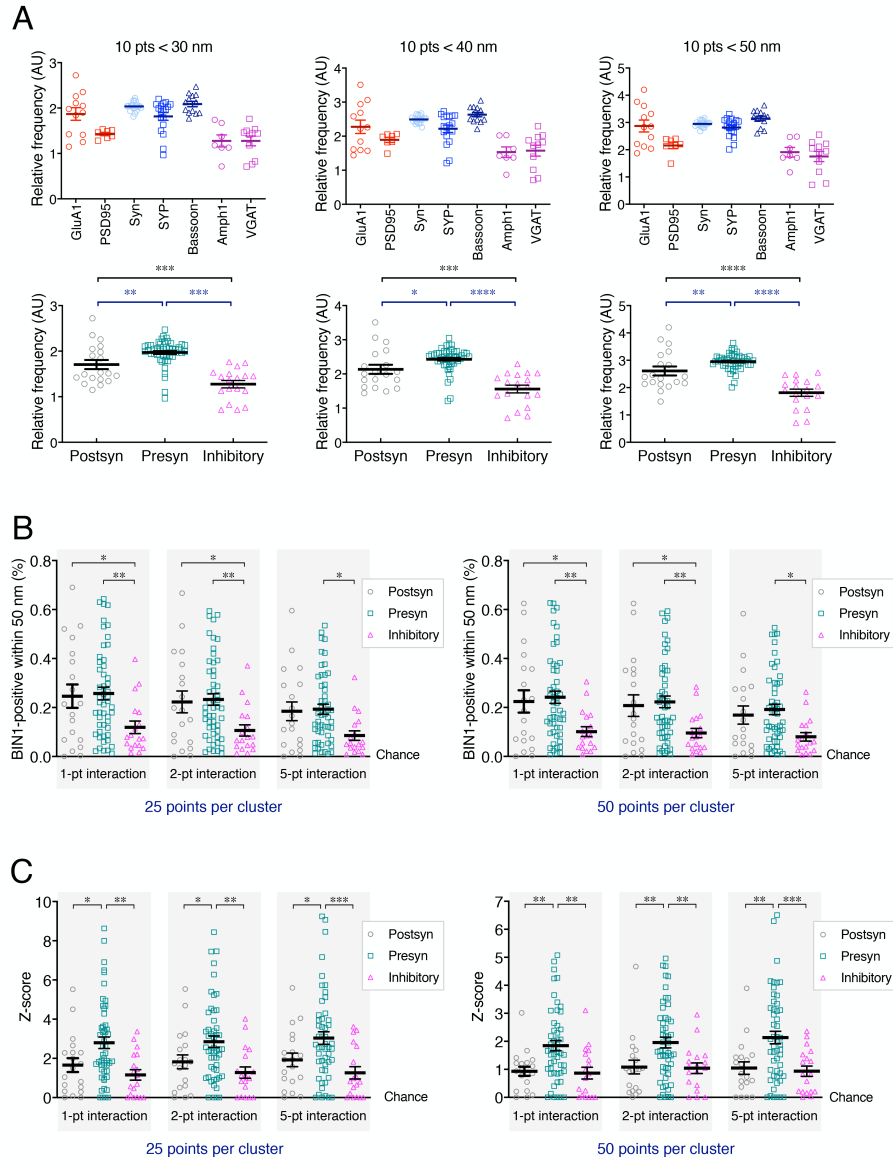
**Figure S5. Related to Figure 5: Super-resolution analysis of BIN1 presynaptic localization.**

(A) Representative confocal images for BIN1 along with synapsin (presynaptic marker), and PSD95 (postsynaptic marker). The overlays show a high level of colocalization between BIN1 and synaptic markers.

(B) The gain of resolution obtained using dSTORM microscopy for BIN1 (green) as well as synaptic proteins Bassoon (blue) and GluA1 (red).

(C) Top: localization of BIN1 (green) within CaMKII-positive structures (red). Imaris reconstruction (right panel) shows a low frequency of BIN1 signal within the spines. Bottom: localization of BIN1 (green) in MAP2-positive dendrites (red). A lower incidence of BIN1 staining was observed in the dendritic shaft. BIN1 puncta were found in MAP2-negative structures (white arrows), highlighted by filament tracing in Imaris (right panel).

(D) dSTORM analysis of BIN1 colocalization with pre- and postsynaptic markers. The boxed region is shown at a higher magnification on the right as two-color overlays. *White arrows* indicate the overlap between BIN1 and synapsin.



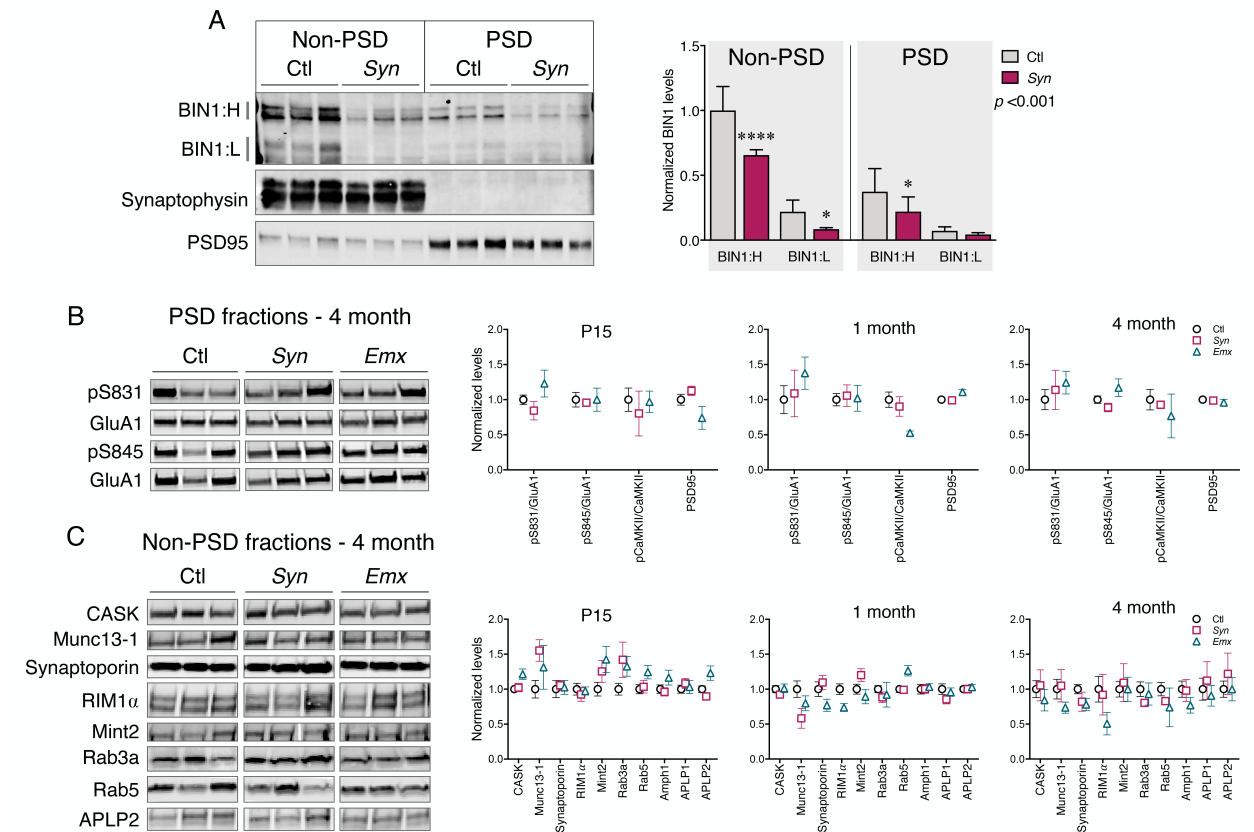
**Figure S6. Related to Figure 5: Quantification of the distance between BIN1 and different synaptic markers.**

**(A)** DBSCAN analysis of cluster proximity showing proportion of BIN1 within 30 nm, 40 nm, and 50 nm of each synaptic marker (top graphs) or grouped postsynaptic, presynaptic, and inhibitory presynaptic markers (bottom graphs). Note that the top graphs in panel A are color coded differently from the rest of the graphs.

**(B)** Graph representing the proportion of BIN1-positive structures within 50 nm of synaptic markers. BIN1 was found associated more with general presynaptic sites (25-pt clustering  $F_{(8, 255)}=3.819$ ,  $p=0.0003$ ; 50-pt clustering  $F_{(8, 255)}=3.784$ ,  $p=0.0003$ ).

**(C)** Graph representing randomization analysis of BIN1 proximity to synaptic markers. When the Z score is equal to 0, the association between BIN1 and the markers is not greater than chance. The results of the Z score analysis showed specific association of BIN1 with the different markers analyzed (25-pt clustering  $F_{(8, 255)}=4.706$ ,  $p<0.0001$ ; 50-pt clustering  $F_{(8, 255)}=5.446$ ,  $p<0.0001$ ).



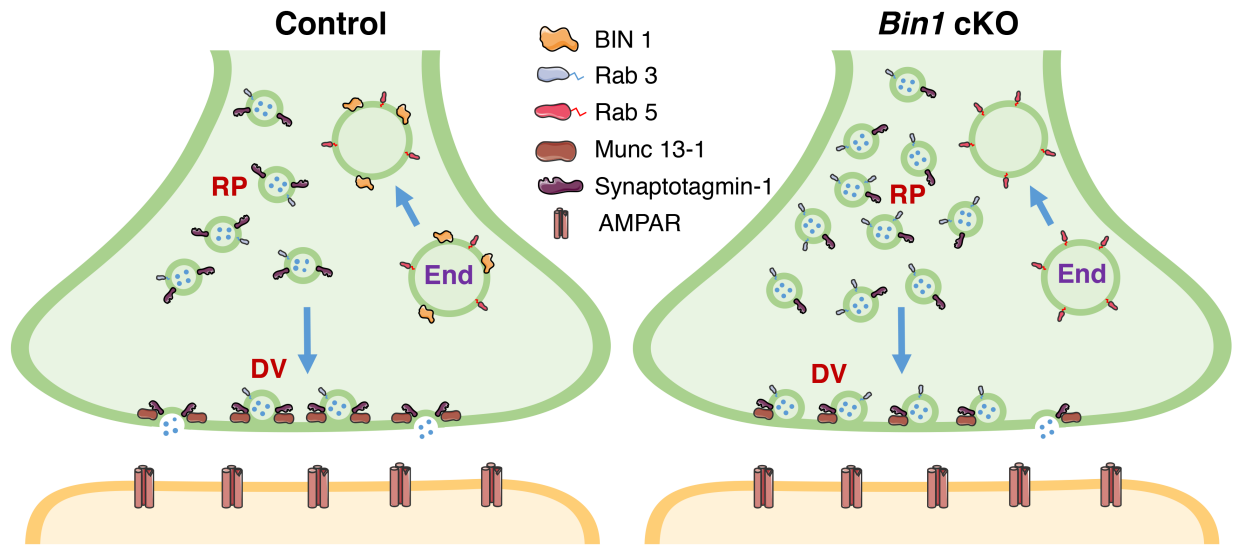


**Figure S7.** Related to Figures 5 and 7: **Synaptosome fractionation analysis of synaptic protein levels in *Bin1* cKO mice.**

**(A)** Western blot analysis of non-PSD and PSD fractions of hippocampus tissue from Ctl and *Syn* mice. The blots were probed with antibodies against BIN1 (mAb 13463), synaptophysin and PSD95. The graph (right) represents quantification of BIN1 levels in each fraction normalized to BIN1 in Ctl non-PSD fractions.

**(B)** The levels of GluA1 phosphorylation in PSD fractions of Ctl, *Syn*, and *Emx* mice analyzed at 4 months of age. The graphs on the right represent the ratio of p831/GluA1, p845/GluA1, and pCaMKII/CaMKII, and PSD95 levels [normalized to Ctl mice] in mice analyzed at P15 ( $F_{(6, 101)}=0.8136, p=0.5618$ ), 1 month ( $F_{(6, 55)}=1.428, p=0.2208$ ) and 4 months of age ( $F_{(6, 76)}=0.7103, p=0.6423$ ).

**(C)** The levels of CASK, Munc13-1, Synaptoporin, RIM1, Mint2, Rab3a, Rab5 and APLP2 in the non-PSD fraction in Ctl, *Syn*, and *Emx* mice at 4 months of age. On the right, graph representing level of presynaptic proteins at P15 (left) ( $F_{(18, 230)}=1.397, p=0.1338$ ), 1 month (middle) ( $F_{(18, 140)}=2.109, p=0.0116$ ) and 4 months of age (right) ( $F_{(18, 150)}=0.4135, p=0.9835$ ).



**Figure S8.** Related to Figures 7 and 8: **Schematic representation of presynaptic changes *Bin1* cKO mice.** Our results suggest that BIN1 localizes to the synapse and participates in synaptic vesicle dynamics in glutamatergic neurons. In the absence of BIN1, excitatory synapses onto hippocampal CA1 pyramidal neurons have increased number of docked synaptic vesicles (*DV*) concomitant with a significant reduction in presynaptic vesicular release probability. The decrease in the cluster organization of SNARE-related presynaptic proteins may underlie this defect. The loss of BIN1 also leads to an increase in reserve pool of synaptic vesicles (*RP*), consistent with a slower depletion time course of synaptic responses. In addition, the loss of BIN1 leads to an increase in the size of Rab5 clusters in *Bin1* cKO mice, in agreement with the data from cultured neurons (Calafate et al., 2016). We propose that the altered synaptic vesicle dynamics in *Bin1* cKO mice reflects a critical role of BIN1 in the regulation of exocytosis associated with synaptic transmission in addition to its previously described role in endocytosis. *End*, endosome.

**Table S1. List of antibodies used in this study, Related to STAR Methods**

<b>Target</b>	<b>Species / clone</b>	<b>Catalog no.</b>	<b>Source</b>	<b>WB</b>	<b>IF</b>
BIN1	mAb EPR13463	ab182562	AbCam	1:500	1:200
BIN1	Rabbit pAb	14647-1-AP	Protein Tech	1:500	
BIN1	Rabbit pAb BSH3		Thinakaran Lab	1:1000	1:2500
BIN1	Rabbit pAb B1415		Thinakaran Lab	1:2000	
Actin	mAb 2D4H5	66009-I-Ig	Protein Tech	1:20000	
AmphI	mAb 8	SC21710	Santa Cruz	1:2000	1:1000
APLP1	Rabbit pAb CT11		Thinakaran Lab	1:1000	
APLP2	Rabbit pAb CT12		Thinakaran Lab	1:1000	
Bassoon	Guinea Pig pAb	141-004	Synaptic Systems		1:500
CaMKII	mAb 6G9	SC32288	Santa Cruz	1:500	
CASK	mAb K56A/50	O14936	NeuroMab	1:1000	
CNPase	mAb 11-5B	MAB326	Millipore	1:2000	
Flotillin-2	Rabbit pAb		Thinakaran Lab	1:1000	
GluA1	mAb N355/1	P19490	NeuroMab	1:100	1:50
MAP2	mAb HM-2	M4403	Sigma		1:10000
MBP	mAb SMI-94	SMI-94R	Covance	1:20000	
Mint-2	mAb	M76120	BD Biosciences	1:500	
Munc13-1	Rabbit pAb	126-102	Synaptic Systems	1:1000	1:500
NeuN	mAb A60	MAB377	Chemicon		1:2000
pCaMKII	Rb mAb D21E4	12716S	Cell Signaling	1:2000	1:500
pGluA1-831	Rabbit pAb	AB5847	Millipore	1:500	
pGluA1-845	Rabbit pAb	AB5849	Millipore	1:500	
PSD95	mAb K28/43	K28/43	UC Davis	1:5000	
Rab3a	mAb (Clone9)	Sc-136050	Santa Cruz	1:500	1:100
Rab5	Rb mAb C8B1	3547S	Cell Signaling	1:1000	1:500
RIM1 $\alpha$	Rabbit pAb	140-003	Synaptic Systems	1:1000	1:500
Synapsin 1	mAb 46.1	106 011BT	Synaptic Systems		1:5000
Synaptojanin 1	Rabbit pAb	145-103	Synaptic Systems		1:200
Synaptophysin 1	mAb SVP38	S5768	Sigma	1:20000	1:500
Synaptoporin	Rabbit pAb	102-003	Synaptic Systems	1:1000	1:500
Synaptotagmin 1	Chicken IgY	105-106	Synaptic Systems		1:300

**Table S2. Statistical analysis of cluster density quantified by confocal imaging,  
Related to Figure 1**

Protein	ANOVA F (DFn, DFd)	Ctl vs <i>Syn</i>	Ctl vs <i>Emx</i>
<b>BIN1</b>	F (2, 107) = 50.70, $p < 0.0001$	<0.0001	<0.0001
<b>RIM1</b>	F (2, 28) = 0.2967, $p = 0.7456$	0.6426	0.4507
<b>Munc13-1</b>	F (2, 26) = 2.935, $p = 0.0709$	0.5138	0.0247
<b>Bassoon</b>	F (2, 152) = 6.403, $p = 0.0021$	0.0014	0.0058
<b>Synapsin1</b>	F (2, 95) = 0.5160, $p = 0.5986$	0.4142	0.8883
<b>Synaptojanin</b>	F (2, 29) = 0.8146, $p = 0.4527$	0.3184	0.2612
<b>Synaptoporin</b>	F (2, 31) = 4.054, $p = 0.0273$	0.0162	0.0258
<b>Synaptotagmin</b>	F (2, 66) = 1.101, $p = 0.3385$	0.1884	0.2269
<b>GluA1</b>	F (2, 69) = 2.330, $p = 0.1049$	0.1644	0.0398
<b>pCaMKII</b>	F (2, 29) = 3.010, $p = 0.0649$	0.0288	0.7094
<b>PSD95</b>	F (2, 93) = 0.4385, $p = 0.6463$	0.4927	0.3887
<b>Rab5</b>	F (2, 128) = 7.376, $p = 0.0009$	0.0004	0.0060
<b>Rab3a</b>	F (2, 61) = 0.3307, $p = 0.7197$	0.5214	0.4664
<b>Clathrin</b>	F (2, 30) = 1.209, $p = 0.3127$	0.1589	0.2464
<b>Dynamin</b>	F (2, 30) = 30632, $p = 0.0387$	0.0168	0.0547

The table reports statistical analysis of data from a minimum of 7 images from at least 3 animals per genotype. ANOVA F statistic, degrees of freedom between groups (DFn) and within groups (DFd), and the  $p$  value are listed. The right two columns list the  $p$  value of post-hoc Fisher's LSD after ANOVA for Ctl *versus* each *Syn* or *Emx* mice.



**Table S3. Statistical analysis of cluster volume quantified by STED imaging,  
Related to Figure 7**

Protein	ANOVA F (DFn, DFd)	Ctl vs <i>Syn</i>	Ctl vs <i>Emx</i>
<b>BIN1</b>	F (2, 8298) = 31.61, $p < 0.0001$	<0.0001	0.0706
<b>Rab5</b>	F (2, 47322) = 60.44, $p < 0.0001$	0.0018	<0.0001
<b>Rab3a</b>	F (2, 25397) = 50.55, $p < 0.0001$	<0.0001	<0.0001
<b>Synaptotagmin-1</b>	F (2, 84385) = 145.8, $p < 0.0001$	<0.0001	<0.0001
<b>Munc13-1</b>	F (2, 8375) = 14.60, $p < 0.0001$	<0.0001	0.0021
<b>Synaptophysin</b>	F (2, 3695) = 8.192, $p = 0.0003$	0.0011	0.0001
<b>Synaptoporin</b>	F (2, 7953) = 39.13, $p < 0.0001$	<0.0001	<0.0001
<b>Bassoon</b>	F (2, 19454) = 18.41, $p < 0.0001$	0.9496	<0.0001
<b>Synaptojanin</b>	F (2, 4914) = 52.24, $p < 0.0001$	<0.0001	0.2331
<b>RIM1</b>	F (2, 17957) = 1.475, $p = 0.2289$	0.0891	0.2457
<b>Synapsin1</b>	F (2, 39891) = 3.562, $p = 0.0284$	0.0638	0.3574
<b>GluA1</b>	F (2, 17662) = 249.5, $p < 0.0001$	0.1483	<0.0001
<b>pCaMKII</b>	F (2, 7224) = 64.07, $p < 0.0001$	<0.0001	<0.0001
<b>PSD95</b>	F (2, 6966) = 1.058, $p = 0.3471$	0.1811	0.2759

The table reports statistical analysis of data from a minimum of 6 images from at least 3 animals per genotype. ANOVA F statistic, degrees of freedom between groups (DFn) and within groups (DFd), and the  $p$  value are listed. The right two columns list the  $p$  value of post-hoc Fisher's LSD after ANOVA for Ctl *versus* each *Syn* or *Emx* mice.