Endophilin A1 promotes Actin Polymerization in response to Ca^{2+} /calmodulin to Initiate Structural Plasticity of Dendritic Spines

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Abstract

Dendritic spines of excitatory neurons undergo activity-dependent structural and functional plasticity, which are cellular correlates of learning and memory. However, mechanisms underlying the rapid morphological changes immediately after NMDAR-mediated Ca^{2+} influx into spines remain poorly understood. Here we report that endophilin A1, a neuronal N-BAR protein, orchestrates membrane dynamics with actin polymerization to initiate spine enlargement in the induction phase of long-term potentiation (LTP). Upon LTP induction, Ca^{2+} /calmodulin enhances its binding to both membrane and p140Cap, a cytoskeleton regulator. As a result, endophilin A1 rapidly associates with the relaxed plasma membrane and promotes actin polymerization, leading to acute expansion of spine head. Moreover, not only the p140Cap-binding, but also calmodulin- and membrane-binding capacities of endophilin A1 are required for LTP and long-term memory. Thus, endophilin A1 functions as calmodulin effector to drive spine enlargement in response to Ca^{2+} influx in the initial phase of structural plasticity.

Keywords: endophilin A1; calcium; calmodulin; dendritic spines; synaptic plasticity; actin polymerization; membrane expansion

1 **Introduction**

2 Long-term potentiation (LTP) of synaptic strength contributes to neural mechanisms
3 underlying learning and memory (Nabavi et al., 2014). In the mammalian brain, most
4 glutamatergic synapses are located on dendritic spines, micron-sized protrusions from
5 dendrites. In response to input activity, spines undergo changes in both morphology
6 (structural plasticity) and function (functional plasticity), which are tightly correlated during
7 LTP (Harvey and Svoboda, 2007; Matsuzaki et al., 2001; Matsuzaki et al., 2004). Imaging
8 studies have revealed that induction of LTP triggers a large transient increase in spine volume
9 (1-5 min after stimulation, early or transient phase) that decays to a long-lasting spine size
10 expansion (> 40 min, late or sustained phase) (Harvey and Svoboda, 2007; Matsuzaki et al.,
11 2004), a process termed structural LTP (sLTP), which would allow physical enlargement of
12 glutamatergic synapses to accommodate more
13 α -amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid receptors (AMPA) for synaptic
14 potentiation (Herring and Nicoll, 2016). Moreover, recent studies have established a direct
15 link between spine morphological changes and memory trace *in vivo* by demonstrating
16 disruption of acquired motor learning by optical shrinkage of potentiated spines in the motor
17 cortex (Hayashi-Takagi et al., 2015). Although sLTP has been studied intensively, with
18 calcium signaling-regulated actin remodeling being the central process that governs the
19 stabilization and consolidation of spine enlargement (Nakahata and Yasuda, 2018),
20 mechanism(s) initiating rapid spine head expansion remains largely unexplored due to limited
21 spatiotemporal resolution of the molecular events in the acute phase (0-1 min) of LTP
22 induction.

23 Endophilin A1 is a member of the endophilin A protein family characterized by an
 24 N-terminal BIN/amphiphysin/Rvs (BAR) domain and a C-terminal Src homology 3 (SH3)
 25 domain. The gene encoding endophilin A1 (*EEN1*, a.k.a *sh3gl2*) is almost exclusively
 26 expressed in brain (Ringstad et al., 1997) and has been implicated in epilepsy, Alzheimer's
 27 disease and schizophrenia (Corponi et al., 2019; Ren et al., 2008; Yu et al., 2018a; Yu et al.,
 28 2018b). Originally identified as a component of the endocytic machinery, endophilin As
 29 function in synaptic vesicle recycling at the presynaptic site in two distinct processes: ultrafast
 30 endocytosis from the plasma membrane following synaptic vesicle fusion and clathrin
 31 uncoating of regenerated synaptic vesicles (Milosevic et al., 2011; Ringstad et al., 1997;
 32 Schuske et al., 2003; Verstreken et al., 2003; Watanabe et al., 2018). Recently endophilin A2
 33 has also been found to mediate fast clathrin-independent endocytosis in mammalian epithelial
 34 cells (Boucrot et al., 2015; Renard et al., 2015). Other studies have implicated endophilin As
 35 in autophagosome formation and protein homeostasis at presynaptic terminals (Murdoch et al.,
 36 2016; Soukup et al., 2016). In dendrites, both endophilin A2 and A3 interact with Arc/Arg3.1
 37 to accelerate endocytosis of AMPARs at the postsynaptic membrane during late-phase
 38 synaptic plasticity (Chowdhury et al., 2006).

39 Previously we found that during synaptic development, endophilin A1 contributes to
 40 dendritic spine morphogenesis and stabilization through interaction with p140Cap, an actin
 41 cytoskeleton regulator (Yang et al., 2015). We also found that *EEN1* gene knockout (KO) in
 42 the hippocampal CA1 region of mouse brain causes impairment of LTP of the Schaffer
 43 collateral-CA1 pathway and long-term memory (Yang et al., 2018). At the cellular level,
 44 endophilin A1, not A2 or A3, is required for N-methyl-D-aspartate receptor

(NMDAR)-mediated synaptic potentiation of dendritic spines in mature CA1 pyramidal cells (Yang et al., 2018). Intriguingly, overexpression of p140Cap, its downstream effector, fails to rescue the structural and functional plasticity of spines in *EENI* KO (*EENI*^{-/-}) neurons (Yang et al., 2018), suggesting the necessity of spatiotemporal coordination of membrane expansion and actin polymerization during synaptic potentiation. In this study, we investigated the mechanistic role(s) of endophilin A1 in sLTP with a combination of cell biological, biochemical, electrophysiological and genetic approaches. We present evidence that endophilin A1 serves as an immediate effector of Ca²⁺/calmodulin to promote actin polymerization-dependent membrane expansion in the initial phase of spine structural plasticity.

Results

Endophilin A1 is Required for the Acute Structural Plasticity of Dendritic Spines

Knockout of the *EENI* gene causes inhibition of structural and functional plasticity of dendritic spines in hippocampal neurons (Yang et al., 2018). To investigate the mechanistic role(s) of endophilin A1 in synaptic plasticity, first we determined at which temporal stage(s) of LTP it functions by rescuing the morphological phenotype of *EENI* KO neurons with overexpressed endophilin A1 in the induction phase of chemically-induced LTP (cLTP) (Figure 1A). Quantification of spine size indicated that endophilin A1 is required for spine enlargement as early as 1 minute after application of the NMDAR co-agonist glycine (Figure 1B and 1C).

Actin polymerization in dendritic spines is crucial for not only spine morphogenesis

67 during synaptic development but also sLTP of mature neurons. Imaging studies on
68 hippocampal neurons have shown that actin polymerization in spines starts as early as 20 s
69 after LTP induction (Okamoto et al., 2004). As endophilin A1 is required for acute
70 LTP-induced spine enlargement, we reasoned that it might function to promote actin
71 polymerization in the initial phase of structural plasticity. To monitor morphological changes
72 and actin dynamics of spines simultaneously, we performed super-resolution live imaging of
73 *EENI* wild-type (WT) and KO neurons expressing membrane-anchored GFP (mGFP) and the
74 F-actin probe LifeAct-mCherry by Grazing Incidence Structured Illumination Microscopy
75 (GI-SIM) (Guo et al., 2018). In WT neurons, consistent with our previous study (Guo et al.,
76 2018), we observed rapid increase in both spine size and F-actin signal intensity in dendritic
77 spines within 1 minute upon glycine application (Video 1 and Figure 1D-1F). In contrast, no
78 significant changes in spine size were detected in *EENI* KO neurons even though the shape of
79 spines changed constantly (Video 2, Figure 1D and 1E). These live imaging data indicate that
80 endophilin A1 is required for spine head enlargement during the initial phase of sLTP. Notably,
81 although the spine heads of *EENI* KO neurons were as motile as those of WT cells, the
82 LTP-induced net increase in F-actin content was also abolished (Figure 1D and 1F), indicating
83 that endophilin A1 is also required for spine actin polymerization in the initial phase of sLTP.

84 Consistent with previous findings (Guo et al., 2018; Honkura et al., 2008), we observed
85 membrane expansion of spine head accompanied by local increase in F-actin content in the
86 initial phase of sLTP (Figure 1G and Video 1). Quantitative analysis of the images clearly
87 revealed that actin polymerization and plasma membrane protrusion of spine head are tightly
88 coupled spatially and temporally (Figure 1H-K and Video 3), suggesting that local actin

polymerization provides propulsive force for spine growth. Notably, compared with WT, the membrane protrusion of spine head and increase in F-actin content was much less coupled in *EENI* KO neurons (Figure 1H-1K and Video 4), implicating endophilin A1 in actin polymerization-dependent membrane expansion in the initial phase of sLTP.

93

Membrane Unfolding and Branched Actin Polymerization are Required for Acute Structural Plasticity

The observation that endophilin A1 is required for rapid spine membrane expansion upon glycine application prompted us to investigate mechanism(s) underlying initiation of sLTP.

Although it was postulated that the membrane source for LTP-induced spine enlargement comes from transport of intracellular recycling endosomes to the neuronal plasma membrane

(Park et al., 2004), imaging studies revealed that spine head expansion precedes most of the

AMPA exocytotic events during the early phase of cLTP (Kopeck et al., 2006), and that the

light chain of botulinum toxin type B (BoTox), a neurotoxin that binds to the SNARE

complex and inhibits exocytosis, had no effect on the initial spine expansion after the theta

burst pairing protocol of LTP induction (Yang et al., 2008). To determine whether spine

membrane expansion requires fusion of exocytosed vesicles with the plasma membrane in

cLTP, we tested the effect of Tetanus toxin (TeTx), another inhibitor of SNARE-mediated

membrane fusion, on glycine-induced acute increase in spine size. Consistent with previous

studies (Hiester et al., 2018), treatment of hippocampal neurons with TeTx did not affect spine

enlargement in the first 3 minutes of glycine treatment (Figure 2A and 2B), indicating that

vesicle fusion is not the direct source of membrane supply for the rapid structural expansion

111 of spines in the initial phase of sLTP.

112 Both electron microscopy and super-resolution imaging reveal that the surface of mature
113 spines is not smooth but rather convoluted (Arellano et al., 2007; Harris and Stevens, 1989;
114 Smith et al., 2014). To determine whether the membrane folds or invaginations in spine head
115 contribute to structural expansion, we increased membrane tension by exposing neurons to
116 hypo-osmotic buffer and found that cLTP-induced spine enlargement was abolished (Figure
117 2C and 2D). As membrane tension increases fusion efficiency (Kliesch et al., 2017), these
118 findings corroborate that membrane fusion does not contribute to rapid spine enlargement.
119 Conversely, incubation of neurons with hyper-osmotic buffer, which shrunk the spines and
120 generated membrane folds, also antagonized cLTP-induced spine enlargement (Figure 2C and
121 2D). These data together indicate that, similar to the formation of membrane expansion in
122 migrating primordial germ cells (Goudarzi et al., 2017), the membrane supply for rapid spine
123 enlargement in the initial phase of sLTP comes from plasma membrane invaginations, which
124 are local unfolding of spine surface convolutions.

125 Previous studies on activity-dependent structural remodeling of dendritic spines indicate
126 that glutamate uncaging-induced rapid spine enlargement requires NMDAR, calmodulin and
127 actin polymerization, whereas long-lasting spine enlargement also requires the activity of
128 Ca^{2+} /calmodulin-dependent protein kinase II (CaMKII) (Matsuzaki et al., 2004). Indeed,
129 inhibition of calmodulin but not CaMKII abolished glycine-induced spine enlargement of
130 hippocampal neurons during the initial phase of sLTP (Figure 2E and 2F). The
131 Ca^{2+} /calmodulin-CaMKII pathway triggers several signaling cascades to promote actin
132 polymerization and AMPAR trafficking to the plasma membrane during synaptic potentiation

133 (Murakoshi and Yasuda, 2012). In line with previous studies (Honkura et al., 2008; Matsuzaki
134 et al., 2004), inhibition of actin polymerization with latrunculin A (LatA) abolished
135 enlargement of spine head during the initial phase of sLTP (Figure 2G and 2H). Moreover,
136 inhibition of Arp2/3 had a similar effect to that of LatA, whereas inhibition of Formin had no
137 effect (Figure 2G and 2H), indicating that branched rather than linear actin polymerization is
138 required for acute spine expansion.

139 The Rho family members of small GTPases are known downstream effectors of
140 Ca^{2+} /calmodulin and regulators of actin reorganization and structural plasticity (Hedrick and
141 Yasuda, 2017; Spiering and Hodgson, 2011). Next we determined whether they are required
142 for cLTP-induced acute spine enlargement by treating hippocampal neurons with inhibitor for
143 RhoA, Rac1 or Cdc42. In agreement with previous findings that rapid structural remodeling
144 requires actin severing and nucleation by ADF/cofilin, inhibition of RhoA, the upstream
145 activator of ADF/cofilin (Hedrick et al., 2016; Murakoshi et al., 2011), abolished
146 glycine-induced spine enlargement (Figure 2G and 2H). Notably, although both Rac1 and
147 Cdc42 can activate actin polymerization (Hedrick and Yasuda, 2017), only Rac1 is required
148 for the initial spine growth of sLTP (Hedrick et al., 2016; Murakoshi et al., 2011) (Figure 2G
149 and 2H, this study). Moreover, inhibition of Rac1 only partially inhibited acute spine
150 expansion (Figure 2G and 2H), suggesting the presence of other factor(s) that promotes
151 branched actin polymerization in response to Ca^{2+} influx immediately upon sLTP induction.
152 Collectively, these data indicate that Ca^{2+} /calmodulin-regulated branched actin polymerization
153 plays an essential role in acute spine membrane expansion in the initial phase of sLTP.

154

Ca²⁺/calmodulin Enhances Endophilin A1-p140Cap Interaction to Promote Actin

Polymerization in Spines

Having established that endophilin A1 is required for both spine head enlargement and actin polymerization in the initial phase of sLTP, we speculated that endophilin A1 promotes actin polymerization in spines to provide pushing force for membrane expansion. Since CaMKII is not required for acute spine enlargement, we reasoned that endophilin A1 might function earlier than CaMKII in molecular events triggered by NMDAR-mediated Ca²⁺ influx upon LTP induction. Previously studies reported that Ca²⁺ binding changes the conformation of endophilin A2 and regulates its interaction with dynamin and voltage-gated Ca²⁺ channels (VGCC) (Chen et al., 2003). Nevertheless, no direct binding between endophilin A1 and Ca²⁺ was detected by isothermal titration calorimetry (ITC) and fluorescence spectrometry (Figure 3-figure supplement 1).

Recent studies reported that calmodulin binds to mammalian N-BAR proteins including endophilin A1 and A2 (Myers et al., 2016). Indeed, GST-pull down assay showed that endophilin A1 binds to calmodulin via its N-terminal BAR domain and the interaction is strengthened by Ca²⁺ (Figure 3A-3F). We reasoned that Ca²⁺/calmodulin might function as upstream regulator of endophilin A1 function(s). As expected, co-immunoprecipitation (co-IP) from both transiently transfected HEK293 cells and mouse hippocampal neurons showed that the interaction between endophilin A1 and p140Cap is Ca²⁺-dependent (Figure 3G-3J). Moreover, co-IP from cultured hippocampal neurons revealed that LTP induction enhanced the interaction between endophilin A1 and p140Cap in a Ca²⁺- and NMDAR-dependent manner (Figure 3K and 3L). Further, the association of endophilin A1 with not only

177 calmodulin but also p140Cap was enhanced acutely upon LTP induction, which was abolished
178 with the calmodulin inhibitor W-7 (Figure 3M and 3N), indicating that Ca^{2+} /calmodulin
179 enhances endophilin A1–p140Cap interaction during the initial phase of LTP.

180 As regulator of actin remodeling, p140Cap recruits cortactin to drive Arp2/3-mediated
181 branched actin polymerization (Jaworski et al., 2009; Schnoor et al., 2018; Yang et al., 2015).
182 Indeed, co-IP of not only p140Cap but also cortactin by antibodies against endophilin A1 was
183 enhanced by Ca^{2+} (Figure 3I and 3J). To test the idea that endophilin A1 functions via
184 p140Cap and cortactin during the initial phase of sLTP, first we determined whether they are
185 recruited to dendritic spines in an LTP- and endophilin A1-dependent manner. Quantitative
186 analysis of immunofluorescence confocal images indicated that enrichment of p140Cap and
187 cortactin in spines upon LTP induction requires not only endophilin A1 but also activities of
188 calmodulin and NMDAR (Figure 3O-3R).

189 Next, to determine whether Ca^{2+} /calmodulin promotes actin polymerization via the
190 endophilin A1–p140Cap pathway, we generated a calmodulin-binding deficient mutant of
191 endophilin A1 (I154A/L158A, DM) that has much lower affinity for calmodulin than the WT
192 protein (Figure 4A and 4B; and Figure 4-figure supplement 1), and tested whether it can
193 rescue the sLTP phenotype of *EEN1* KO neurons. Indeed, WT but not the DM mutant of
194 endophilin A1 restored the rapid increase in spine size and F-actin content in spines upon LTP
195 induction (Figure 4C-4F). Moreover, Y343A, the p140Cap-binding deficient mutant of
196 endophilin A1 (Yang et al., 2015), also failed to rescue the sLTP phenotype of *EEN1* KO
197 neurons (Figure 4G-4I). Together, these data indicate that Ca^{2+} /calmodulin enhances the

198 recruitment of p140Cap by endophilin A1 to promote actin polymerization during the initial
199 phase of sLTP.

200

201 **Ca²⁺/calmodulin-Dependent Increase in Plasma Membrane-Associated Endophilin A1**

202 **Nanodomains Correlates with Spine Size During the Initial Phase of sLTP**

203 Single protein tracking and super-resolution imaging revealed dynamic changes in the
204 nanoscale organization of branched F-actin regulators in spines during synaptic plasticity
205 (Chazeau et al., 2014). Intriguingly, although the interaction between endophilin A1 and
206 p140Cap is required for sLTP, overexpression of p140Cap could not rescue plasticity
207 phenotypes of *EENI* KO neurons (Yang et al., 2018), suggesting the existence of
208 spatiotemporal regulation of their interaction during LTP induction. To this end, we analyzed
209 the sub-spine localization of endophilin A1 by immunofluorescence staining and
210 3D-structured illumination microscopy (3D-SIM). Interestingly, in spines endophilin A1 was
211 organized into nanoscale objects (mean area 0.014 μm^2 , referred to as nanodomains) which
212 did not overlap with PSD95, marker for the postsynaptic density (PSD) structure (Figure
213 5-figure supplement 1A). Quantitative analysis revealed an NMDAR-dependent increase in
214 the number of endophilin A1 nanodomains in spines undergoing sLTP (Figure 5-figure
215 supplement 1A-1C). Moreover, the size of spine head correlated with the number but not the
216 area of endophilin A1 nanodomains (F Figure 5-figure supplement 1E and 1F). In contrast, the
217 number of endophilin A1 nanodomains did not correlate with the size of the PSD95-labeled
218 PSD structures (Figure 5-figure supplement 1D and 1G). Further, treatment of hippocampal
219 neurons with inhibitor of calmodulin but not CaMKII abolished the increase in not only spine

size but also the number of endophilin A1 puncta in spines (Figure 5-figure supplement 1H-1J). These data together suggest that the calmodulin-regulated subsynaptic localization of endophilin A1 is required for spine enlargement.

As endophilin A1 contains the positive membrane curvature-sensing and binding N-BAR domain (Gallop et al., 2006) that enables it associate with invaginated plasma membrane, next we investigated whether its association with the spine plasma membrane is also regulated in the initial phase of sLTP. To distinguish plasma membrane-localized endophilin A1 from those localized to intracellular structures, we permeabilized cell membrane with the mild detergent saponin to limit access of antibodies to the cytosolic leaflet of the plasma membrane. Indeed, most endophilin A1 nanodomains were plasma membrane-localized (Figure 5A). Moreover, there was an increase in the number of plasma membrane-localized endophilin A1 nanodomains as early as 1 minute after LTP induction (Figure 5B). Further, the number but not the area of the nanodomains correlated with the size of spine head (Figure 5B-5D). As endophilin A1 functions via p140Cap and its downstream effector cortactin (Yang et al., 2015) to activate Arp2/3 which in turn induces branched actin polymerization (Urano et al., 2001; Weaver et al., 2001), these data suggest that upon LTP induction, endophilin A1 localizes to the periphery of spine head and promotes local actin polymerization underneath the plasma membrane.

Given that sub-spine accumulation of endophilin A1 requires activation of NMDAR and calmodulin (Figure 5-figure supplement 1), we reasoned that upon NMDAR-mediated Ca^{2+} influx, binding of Ca^{2+} -activated calmodulin to endophilin A1 not only enhances its interaction with p140Cap, but also facilitates its association with the invaginated plasma

242 membrane. Indeed, inhibition of NMDAR or calmodulin, not CaMKII, abolished
243 glycine-induced increase in the number of plasma membrane-localized endophilin A1
244 nanodomains (Figure 5A, 5C and 5E). Notably, although latrunculin A treatment inhibited
245 spine enlargement, it had no effect on LTP-induced increase in plasma membrane-localized
246 endophilin A1 (Figure 5F-5I), indicating that Ca^{2+} /calmodulin-enhanced association of
247 endophilin A1 with the plasma membrane precedes actin polymerization during sLTP
248 initiation.

249 To corroborate that Ca^{2+} /calmodulin regulates the association of endophilin A1 with the
250 plasma membrane, we performed *in vitro* liposome sedimentation assays and found that,
251 whilst calmodulin alone did not change the membrane-binding capacity of WT endophilin A1,
252 Ca^{2+} /calmodulin enhanced it significantly (Figure 5J and 5K). In contrast, Ca^{2+} /calmodulin
253 had no effect on the membrane association ability of the calmodulin-binding deficient DM
254 mutant of endophilin A1 (Figure 5J and 5K). Collectively, these data indicate that
255 Ca^{2+} /calmodulin enhances association of endophilin A1 with the plasma membrane of
256 dendritic spines in the initial phase of sLTP.

257

258 **Endophilin A1 Promotes Branched Actin Polymerization Underneath the Plasma** 259 **Membrane**

260 Based on the findings that endophilin A1 accumulates into spine plasma membrane-associated
261 nanodomains and recruits p140Cap and cortactin in the initial phase of sLTP, we further
262 reasoned that it might transduce the Ca^{2+} signals instantaneously to enable plasma membrane
263 expansion of spines by promoting branched actin polymerization. If it is true, we should be

264 able to detect more endophilin A1 associated with both membranes and the actin cytoskeleton
 265 upon LTP induction. To test this possibility, first we analyzed subcellular fractions of mouse
 266 hippocampi from animals subjected to fear conditioning, a physiological learning paradigm
 267 associated with synaptic plasticity. Indeed, compared with naïve mice, we detected significant
 268 increase in the amount of endophilin A1 and p140Cap in both membrane and cytoskeletal
 269 fractions from trained animals (Figure 6A and 6B). Consistently, although the levels of either
 270 protein remained unchanged (Figure 6C and 6D), their association with membrane and
 271 cytoskeleton fractions also increased in dissociated cultured hippocampal neurons in the
 272 initial phase of cLTP (Figure 6E and 6F). Moreover, the enhanced association of endophilin
 273 A1 and p140Cap with both membrane and cytoskeleton was inhibited by W-7 but not KN-62
 274 (Figure 6E and 6F), indicating that calmodulin is the immediate upstream regulator of their
 275 subcellular redistribution. Notably, subcellular distribution of endophilin A2, another member
 276 of the endophilin A family, was not affected by either neural activity or Ca^{2+} /calmodulin
 277 (Figure 6A-6F).

278 We then determined whether the membrane-binding capacity of endophilin A1 is
 279 required for the rapid spine enlargement of neurons undergoing sLTP. Compared with WT, the
 280 membrane-binding deficient mutant of endophilin A1 (KKK-EEE) (Gallop et al., 2006) was
 281 unable to rescue the deficit in acute spine enlargement of glycine-treated *EEENI* KO neurons
 282 (Figure 6G-6I). All together, these data indicate that rapid enlargement of spines in the initial
 283 phase of sLTP requires Ca^{2+} /calmodulin-dependent enhancement of not only endophilin
 284 A1-p140Cap interaction but also the association of endophilin A1 with membrane.

285 Since the small size of dendritic spines and the wide distribution of F-actin and actin

polymerization regulators in spines prevent us to better visualize the spatiotemporal relationship between plasma membrane association of endophilin A1 and its effectors, we tested whether there are changes in the subcellular distribution of endophilin A1 and Arp2/3 in response to Ca^{2+} /calmodulin using HeLa cell as a heterologous model system. In HeLa cells ectopically co-expressing endophilin A1 and p140Cap, upon Ca^{2+} influx induced by the calcium ionophore ionomycin, we detected enrichment of endophilin A1 signals underneath the plasma membrane by confocal microscopy (Figure 6-figure supplement 1A). Moreover, preincubation with W-7 but not latrunculin or CK-666 abolished ionomycin-induced endophilin A1 recruitment to the cell periphery (Figure 6-figure supplement 1A), indicating that Ca^{2+} -activated calmodulin directly regulates accumulation of endophilin A1 at the plasma membrane. Further, although the strong intrinsic signals for cortical actin did not allow us to quantify changes in F-actin content underneath the plasma membrane, the Arp2/3 complex (labeled with fluorescently tagged Arp1b) was also enriched in the cell periphery upon ionomycin application (Figure 6-figure supplement 1B). In contrast, ionomycin treatment failed to cause recruitment of either the membrane-binding (KKK-EEE) or the calmodulin-binding deficient mutant of endophilin A1 to the cell periphery (Figure 6-figure supplement 1B). Intriguingly, although ionomycin treatment increased plasma membrane association of the p140Cap-binding mutant Y343A, no enrichment of Arp1b signals at the cell periphery was observed (Figure 6-figure supplement 1B), indicating that recruitment of the Arp2/3 complex underneath the plasma membrane requires the interaction between endophilin A1 and p140Cap. Collectively, these results support that Ca^{2+} /calmodulin enhances association of endophilin A1 with the plasma membrane and promotes branched actin

308 polymerization in spines.

309

310 **Ca²⁺/calmodulin-Regulated Functions of Endophilin A1 is Required for LTP and** 311 **Long-term Memory**

312 It was proposed that spine enlargement enables formation of a stable F-actin:cofilin complex
313 that serves as a synaptic tag to capture postsynaptic constituent proteins for maintenance and
314 consolidation of the potentiated state (Bosch et al., 2014). To determine the functional
315 significance of endophilin A1-mediated structural plasticity, next we tested whether the
316 molecular functions of endophilin A1 in structural plasticity are also required for LTP by
317 electrophysiological analyses. To this end, we performed molecular replacement in a small
318 subset of neurons by injection of the hippocampal CA1 region of *EENI*^{fl/fl} mice with
319 adeno-associated viral vectors encoding the Cre recombinase (AAV-mCherry-Cre) together
320 with those encoding either WT or mutant endophilin A1 (AAV-EGFP-2A-EEN1 WT, Y343A,
321 KKK-EEE or DM) at postnatal day 0 (P0), and induced LTP in Schaffer-collateral synapses
322 by double patch whole-cell recording of non-infected (control, *EENI*^{fl/fl}) and virus-infected
323 CA1 pyramidal cells in acute slices from virus-injected animals at P14-21. In line with our
324 previous findings (Yang et al., 2018), Cre-mediated knockout of *EENI* caused significant
325 impairment of LTP, which was fully rescued by re-expression of WT endophilin A1 (Figure
326 7A, 7B and 7F). In contrast, none of the mutants could rescue the magnitude or the
327 maintenance of LTP in *EENI*^{-/-} neurons (Figure 7C-7F), indicating that indeed, molecular
328 functions of endophilin A1 to initiate sLTP is required for expression and stabilization of
329 synaptic potentiation.

330 Finally, we determined the physiological significance of the mechanistic roles of
 331 endophilin A1 in LTP by testing whether endophilin A1 mutants can rescue the learning and
 332 memory deficits in *EENI* KO mice. Indeed, whilst AAV-mediated expression of WT
 333 endophilin A1 in the CA1 region restored the long-term memory in KO mice in both Morris
 334 water maze and fear conditioning tests, neither of the three mutants ameliorated the
 335 phenotypes (Figure 7G-O). Collectively these data indicate that in CA1 pyramidal cells, not
 336 only the p140Cap-binding, but also the calmodulin-interaction and membrane association
 337 capacities of endophilin A1 are required for long-term synaptic potentiation and memory.

338

339 Discussion

340 The temporal phases of sLTP include initiation of spine expansion (≤ 1 min after LTP
 341 stimulation, initial phase), transient (early phase) and sustained spine enlargement (late phase)
 342 (Harvey and Svoboda, 2007; Matsuzaki et al., 2004). Although actin polymerization is
 343 essential for sLTP (Matsuzaki et al., 2004; Obashi et al., 2019), little is known about the
 344 relationship between actin remodeling and membrane dynamics during the initial phase due to
 345 limited spatiotemporal resolution of light microscopy. Moreover, although imaging studies
 346 have revealed subspine organization and dynamics of F-actin pools as well as nanoscale
 347 segregation of branched F-actin regulators in dendritic spines and suggested that their
 348 localization might be spatially and temporally controlled during activity-dependent
 349 morphological changes (Chazeau et al., 2014; Honkura et al., 2008), the precise sequence of
 350 molecular events leading to rapid structural expansion of spines remains to be defined. In this
 351 work, we uncover a novel mechanism for initiation of sLTP. We show that in direct response

352 to Ca^{2+} /calmodulin, endophilin A1 drives acute spine enlargement in NMDAR-mediated sLTP
353 by localizing to spine plasma membrane and recruiting p140Cap to promote branched actin
354 polymerization (Figure 8).

355 Endophilin A2 and A3, two other members of the endophilin A family, have been found
356 to interact with the immediate early protein Arc/Arg3.1 to accelerate AMPAR endocytosis and
357 might contribute to late-phase synaptic plasticity (Chowdhury et al., 2006). The impairment of
358 structural and functional plasticity of potentiated dendritic spines caused by ablation of
359 endophilin A1 cannot be rescued by overexpression of endophilin A2 or A3 (Yang et al.,
360 2018). Although both endophilin A1 and A2 interact with calmodulin (Myers et al., 2016)
361 (and Figure 3, this study), only endophilin A1, but not A2 or A3, binds to and recruits
362 p140Cap to dendritic spines to promote actin polymerization (Yang et al., 2015). Moreover, *in*
363 *vivo* and *in vitro* LTP stimuli induce increased association of endophilin A1, but not
364 endophilin A2, with both membrane and cytoskeleton (Figure 6, this study). Together these
365 findings indicate that different interaction partners for the endophilin A family members
366 confer them distinct mechanistic roles in the induction and expression of synaptic plasticity in
367 dendritic spines.

368 Consistent with previous reports (Yang et al., 2008), we found that SNARE-mediated
369 membrane fusion is not required for rapid spine expansion in the initial phase of sLTP (Figure
370 2). Upon LTP induction, Ca^{2+} influx through the NMDAR ion channel activates calmodulin,
371 which in turn activates CaMKII and its downstream signaling cascades to trigger various
372 subcellular events including actin remodeling (Chazeau et al., 2014). Previous live imaging
373 studies have revealed calmodulin-dependent formation of an F-actin pool that associates with

374 spine enlargement (hence referred to as “enlargement pool”) (Honkura et al., 2008). Although
 375 the observation that the membrane ruffling of the spine head synchronizes with the
 376 enlargement pool of F-actin has prompted the authors to conclude that spine enlargement is
 377 induced by the propulsive force generated by calmodulin-regulated actin polymerization
 378 (Honkura et al., 2008), the mechanistic link between Ca^{2+} /calmodulin and actin
 379 polymerization was still missing. In this study, we demonstrate that membrane-associated
 380 endophilin A1 is the direct molecular target of Ca^{2+} /calmodulin, which enhances both its
 381 membrane association and its interaction with the downstream effector p140Cap (Figure 3
 382 and 5). Our data further indicate that coordination of the membrane-association and
 383 p140Cap-binding capacities of endophilin A1 provides the protrusive force for rapid structural
 384 remodeling of dendritic spines by promoting actin polymerization underneath the plasma
 385 membrane (Figure 4, 6, 7 and 8), which is also in good agreement with most recent studies
 386 that the interplay between membrane tension and branched actin polymerization could
 387 produce membrane deformations (Simon et al., 2019).

388 Most recent studies revealed that although Rac1 activity is not required for the initial
 389 spine expansion induced by glutamate uncaging, formation of a “reciprocally activating
 390 kinase-effector complex” (RAKEC) between CaMKII, and Tiam1, a guanine exchange factor
 391 for Rac (RacGEF), converts the transient Ca^{2+} signal triggered by LTP induction into a
 392 persistent kinase signal required for the maintenance of sLTP (Saneyoshi et al., 2019). Given
 393 the spinous dynamic reorganization of nanoscale distribution of various F-actin regulators that
 394 are downstream of Ca^{2+} /calmodulin and CaMKII (Chazeau and Giannone, 2016; Chazeau et
 395 al., 2014), including the F-actin severing protein cofilin (Noguchi et al., 2016), there might be

396 crosstalk between Ca^{2+} /calmodulin-dependent, endophilin A1-mediated actin polymerization
397 and other Ca^{2+} /calmodulin and/or CaMKII effector-regulated pathways (e.g., Rac1 and
398 ADF/cofilin), which enables spatiotemporally controlled actin reorganization during the
399 initial phase and/or the transition to the early phase of sLTP.

400 Copine-6, a Ca^{2+} sensor, relocates from the cytosol of dendrites to lipid raft-enriched
401 postsynaptic plasma membrane in response to NMDAR-mediated Ca^{2+} influx, and activates
402 Rac1 to spines during synaptic potentiation (Reinhard et al., 2016). It is also required for
403 spine structural plasticity and LTP, probably contributing to stabilization of the actin
404 cytoskeleton by inhibiting ADF/cofilin via the Rac1-PAK-LIMK1 pathway (Reinhard et al.,
405 2016). In addition, by keeping cortactin active to prevent branched actin filaments from
406 severing by ADF/cofilin, the fast Ca^{2+} sensor caldendrin stabilizes an F-actin pool at the spine
407 base required for the structural remodeling of spines in the transition from early to late phase
408 LTP (Mikhaylova et al., 2018). In our study, while repetitive expansion and shrinkage of spine
409 heads were also observed in *EENI* KO neurons, the F-actin content and spine size remained
410 constant during the initial phase of sLTP (Figure 1). Rather than stabilizing F-actin,
411 endophilin A1 responds to Ca^{2+} /calmodulin and promotes branched actin polymerization
412 beneath the plasma membrane to achieve rapid spine enlargement. While they all function via
413 CaMKII-independent mechanisms to regulate actin dynamics, how Copine-6, caldendrin and
414 endophilin A1 coordinate with each other to ensure spine potentiation and stabilization awaits
415 further investigation.

416

417

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426

427 **Author Contributions**

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430 Y.Y., J.C., X.C., J.H., S.Z., S.D., X.L. and S.D.; Writing – Original Draft, Y.Y., J.C., S.Z., X.Y.,
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432 Supervision, Dong.L., C.M., X.L., Y.S.S., and J-J.L.; Project Administration, J-J.L.

433

434 **Declaration of Interests**

435 The authors declare no competing interests.

436

437 **Materials and Methods**

438 **Ethics statement**

439 All animal experiments were approved by and performed in accordance with the guidelines of

the Animal Care and Use Committee of Institute of Genetics and Developmental Biology, Chinese Academy of Sciences (Approval code: AP2013003 and AP2015002), and the Animal Care and Use Committee of the Model Animal Research Center, the Host for the National Resource Center for Mutant Mice in China, Nanjing University (Approval code: AP#SY06). All animals were housed in standard mouse cages at 22-24 °C on a 12 h light/dark cycle with access to food and water freely.

446

Animals

Generation of *EEN1*^{fl/fl} and *EEN1*^{-/-} mice on the C57BL/6J background was as previously described (Yang et al., 2018). Briefly, the targeting vector for *EEN1* was obtained from European Mouse Mutant Cell Repository (EuMMCR, PRPGS00060_A_A02). The endophilin A1 KO first and *EEN1*^{fl/fl} C57BL/6J mice were generated at Nanjing Model Animal Research Center of Nanjing University. Genotyping of mouse lines was performed by genomic PCR of tail prep DNA from offspring with the following primer pairs: loxPF/loxPR: 5'-CAAGGACTCCCAGAGACCTAGCATC-3' and 5'-GAGATGGCGCAACGCAATTAAT-3' (A PCR product of 375 base pairs in *EEN1* KO first mice but none in wild-type mice).

zptF/zptR: 5'-GTAAGCGGCTCTAGCGCATGTTCT-3' and 5'-GCAGGGGCATGTAGGTGGCTCAAC-3' (A PCR product of 466 base pairs in WT mice, none in *EEN1* KO first mice, and of 627 base pairs in *EEN1*^{fl/fl} mice).

460

Constructs

462 The putative amino acid residues of EEN1 involved in calmodulin binding were predicated
463 using the Binding Site Search and Analysis tool provided at the Calmodulin Target Database:
464 <http://calcium.uhnres.utoronto.ca/ctdb/ctdb/home.html>. pCMV-Tag2B-EEN1 DM and
465 pET28a(+)-EEN1 single mutants and DM (mutation I154 and L158 to A) were created by
466 site-directed mutagenesis using pCMV-Tag2B-EEN1 and pET28a(+)-EEN1 as template,
467 respectively. pET28a(+)-EEN1 Δ BAR (Δ aa 6-242) and pET28a(+)-EEN1 Δ SH3 (Δ aa
468 295-346) were subcloned from pGEX4T-1-EEN1 Δ BAR and pGEX4T-1-EEN1 Δ SH3. The
469 pAAV-CaMKII α -EGFP-2A-MCS-3FLAG-EEN1 or mutant of EEN1 AAV viral constructs
470 (KKK-EEE, Y343A and DM) were generated by cloning EEN1 cDNA amplified from
471 pCMV-Tag2B-EEN1 or mutant constructs into the pAAV-CaMKII α -EGFP-2A-MCS-3FLAG.
472 The bacterial expression construct for His-tagged p140Cap fragment (aa 351-1051) was
473 generated by PCR amplification of the cDNA encoding p140Cap (aa 351-1051) and insertion
474 into pET-28a(+). Bacterial expression constructs for calmodulin were generated by PCR
475 amplification of cDNA for mouse calmodulin from mouse brain by RT-PCR and insertion into
476 pGEX4T-1 and pET28a(+). Arp1b-mCherry construct was a generous gift from Drs. Na Mi
477 and Li Yu (Tsinghua University, China). All other constructs used in this study
478 (pCMV-Tag2B-EEN1 WT, pCMV-Tag2B-EEN1 Y343A, pCMV-Tag2B-EEN1 KKK-EEE,
479 mGFP and LifeAct-mCherry) were described previously (Guo et al., 2018; Yang et al., 2018).
480 Viral particles of adeno-associated virus (AAV) carrying pAOV-CaMKII α -EGFP-2A-Cre,
481 pAAV-CaMKII α -EGFP-2A-MCS-3FLAG-EEN1 or mutants of EEN1 and the control
482 construct pAAV-CaMKII α -EGFP-2A-MCS-3FLAG were purchased from OBiO Technology
483 (Shanghai) Corp. Ltd., (Shanghai, China).

484

485 **Antibodies**

486 The following antibodies were obtained from commercial sources: goat anti-endophilin A1
487 (S-20), endophilin A2 (E-15), mouse anti-SYP (D-4), and mouse anti-cortactin (E-4) (Santa
488 Cruz Biotechnology, Santa Cruz, CA, USA) for WB and anti-cortactin (EMD Millipore,
489 Temecula, CA, USA) for staining; rabbit anti-endophilin A1 (Synaptic Systems GmbH,
490 Germany); rabbit anti-Myc, mouse anti-GST, rabbit and mouse anti-GFP, rabbit and mouse
491 anti-RFP which recognizes DsRed and mCherry (Medical & Biological Laboratories,
492 Naka-ku Nagoya, Japan); mouse anti-M5 DYKDDDDK-Tag (Mei5 Biotechnology, Beijing,
493 China), mouse anti- α -tubulin and mouse anti- β -actin (Sigma-Aldrich, St. Louis, MO, USA);
494 mouse anti-His (CoWin Biosciences, Jiangsu, China), rabbit anti-calmodulin (Boster
495 Biological Technology, Pleasanton, CA, USA) for WB and mouse anti-calmodulin (Invitrogen,
496 Carlsbad, CA, USA) for IP; Rabbit anti-p140Cap was described previously (Yang et al., 2015).
497 Secondary antibodies for immunofluorescence staining were from Molecular Probes
498 (Invitrogen, Carlsbad, CA, USA).

499

500 **Cell culture, transfection and drug treatment**

501 Primary hippocampal neurons were cultured as previously described (Yang et al., 2018).
502 Briefly, mouse hippocampi were dissected from P0 C57BL/6J mice, dissociated with 0.125%
503 trypsin in Hank's balanced salt solution without Ca^{2+} and Mg^{2+} at 37°C for 15 min, triturated
504 in DMEM, 10% F12, and 10% FBS (Gibco, Carlsbad, CA, USA). Hippocampal neurons were
505 plated on poly-D-lysine-coated coverslips in 24-well plates or 30-mm dishes at a density of

506 $2.5\text{-}3.0 \times 10^4$ cells/well in 24-well plate or $1.0\text{-}1.2 \times 10^5$ cells/35 mm dish. The medium was
507 replaced with the serum-free Neurobasal A (NB-A) media supplemented with 2% B27
508 supplement, GlutaMAX (Gibco, Carlsbad, CA, USA) and 0.3% glucose 4 h after plating. Half
509 of the media were replaced every 3 days until use.

510 For neuronal morphology and immunofluorescence staining, neuronal transfections were
511 performed using Lipofectamine LTX according to the manufacturer's instructions (Invitrogen,
512 Carlsbad, CA, USA) on 12-14 days *in vitro* (DIV) after plating. Briefly, DNA (0.5 μg /well)
513 was mixed with 0.5 μl PLUS reagent in 50 μl Neurobasal A medium, then mixed with 1.0 μl
514 Lipofectamine LTX in 50 μl NB-A medium, incubated for 20 min and then added to the
515 neurons in NB-A at 37 °C in 5% CO₂ for 1 h. Neurons were then rinsed with NB-A and
516 incubated in the original medium at 37 °C in 5% CO₂ for 4-5 days. For co-transfection,
517 neurons were transfected with 1.0 μg of DNA consisting of two plasmids (0.50 μg each).

518 For HeLa cell culture or HEK293T cell culture, DMEM supplemented with 10% FBS
519 were used. Cell transfections were performed using Lipofectamine2000 according to the
520 manufacturer's instructions (Invitrogen) after plating.

521 For inhibitor treatment, HeLa cells or primary neurons cultured on coverslips were
522 pre-incubated with MK801 (10 μM , Sigma-Aldrich), Latrunculin A (100 nM, Sigma-Aldrich),
523 NSC 23766 trihydrochloride (100 μM , Abcam), CK-666 (100 μM , Sigma-Aldrich), W-7 (20
524 μM , TOCRIS), KN-62 (4 μM , TOCRIS), BAPTA-AM (10 μM , Sigma-Aldrich) for 30 min.
525 Neurons were pre-incubated with ML141 (15 μM , Sigma-Aldrich) or SMIFH2 (30 μM ,
526 Millipore) for 2 h or with CT04 (2 $\mu\text{g}/\text{ml}$, Cytoskeleton) for 3 h. For tetanus toxin treatment
527 (10 nM, Sigma-Aldrich), neurons were pre-incubated for 10 min. These drugs were

528 maintained during glycine or ionomycin application.

529 For ionomycin treatment of HeLa cells, cells were pre-incubated with modified
530 Krebs-Ringer Hepes buffer (containing 120 mM NaCl, 4.8 mM KCl, 1.2 mM KH₂PO₄, 1.2
531 mM MgSO₄, 1.3 mM CaCl₂, 5.5 mM glucose, 25 mM HEPES, pH 7.4, at 37 °C) for 30 min
532 then treated with ionomycin (2 μM, Beyotime Biotechnology) in KRH buffer for 20 min.

533

534 **Chemically-induced LTP (cLTP)**

535 Chemical induction of LTP was performed as previously described (Fortin et al., 2010; Park
536 et al., 2006). Briefly, neurons were treated with glycine (200 μM) in Mg²⁺-free extracellular
537 iso-osmotic solution (mM: 125 NaCl, 2.5 KCl, 2 CaCl₂, 5 HEPES, 33 glucose, 0.2 glycine,
538 0.02 bicuculline, and 0.003 strychnine, pH 7.4) (Yang et al., 2018). For experiments
539 performed in the absence of extracellular Ca²⁺, 10 μM BAPTA-AM was substituted for 2 mM
540 CaCl₂ in the Mg²⁺-free extracellular solution.

541 For experiment to determine the role of membrane tension in spine expansion, neurons
542 were pretreated with the hypo-osmotic solution (mM: 80 NaCl, 2.5 KCl, 2 CaCl₂, 5 HEPES,
543 33 glucose, pH 7.4, OSM 210) or hyper-osmotic solution (mM: 125 NaCl, 2.5 KCl, 2 CaCl₂,
544 5 HEPES, 33 glucose, 250 mM sucrose, pH 7.4, OSM 600) for 20 min and chemically
545 induced LTP in the same solution.

546

547 **Immunostaining, image acquisition and analysis**

548 Neurons were fixed in 4% PFA/4% sucrose in PBS at RT for 15 min. After blocking with 1%
549 BSA in PBS containing 0.4% Triton X-100 for 40 min at RT, neurons were incubated with

550 primary antibodies for 1 h at RT or overnight at 4°C, and appropriate secondary antibodies
551 conjugated with Alexa Fluor 488, Alexa Fluor 555, or Alexa Fluor 647 were applied for
552 detection.

553 Confocal images were collected using the Spectral Imaging Confocal Microscope Digital
554 Eclipse C1Si (Nikon, Tokyo, Japan) with a 100× Plan Apochromat VC (NA 1.40) oil
555 objective. Images were z projections of images taken at 0.2 μm step intervals. The number of
556 planes, typically 4-6, was chosen to encompass the entire dendrite from top to bottom.

557 The procedure for morphometric analysis of dendritic spines was described previously
558 (Yang et al., 2015) (Yang et al., 2018). DsRed was used as a cell-fill. The final reconstructed
559 spines were obtained using a maximum-intensity projection strategy provided by
560 NIS-Elements AR software (Nikon). DsRed-labeled spines were outlined manually. Dendritic
561 segments 40-120 μm from the neuronal cell body were selected for analysis. All
562 morphological experiments were repeated at least three times with an $n \geq 11$ for individual
563 experiments.

564

565 **Super-resolution live cell imaging and data analysis**

566 The GI-SIM live imaging experiments were performed as described by Guo et al (Guo et al.,
567 2018). Briefly, mouse hippocampal neurons cultured on 25-mm coverslips were transfected
568 with constructs expressing membrane-bound GFP (mGFP) and LifeAct-mCherry on DIV12
569 and imaged on DIV16 in Mg^{2+} -free extracellular solution (Yang et al., 2018). Time-lapse
570 images were obtained with acquisition time of 110 ms for each channel at 5 s intervals. To
571 quantify the area of each spine head and enrichment of F-actin in spines, we measured the

fluorescence mean intensity of LifeAct-mCherry within the spines and normalized each measurement by the fluorescence signal along the adjacent dendritic shaft with the NIH ImageJ software. The mGFP-labeled dendrites or spines were outlined manually.

For analysis of the spatiotemporal relationship between actin polymerization and membrane expansion in dendritic spines, spines were segmented from raw GI-SIM images using Otsu's method (Otsu, 1979). The area of each spine and the mean fluorescence signal of F-actin inside each spine were then quantified using NIH ImageJ. To facilitate the visualization of instantaneous spine growth and localized actin polymerization, differential images of both membrane (mGFP) and actin (LifeAct-mCherry) channels were calculated by subtracting the image at time point t from that at $t+1$ through the entire time lapse movie (1 min duration before and 3 min duration post glycine application with 5 s intervals) using Matlab (R2018a, Mathworks). To highlight the regions with spine growth or increased F-actin signals, only pixels with a positive difference were displayed in the final differential images. We then measured the overlap between differential images in the membrane channel and those in the actin channel at individual time points, and determined the extent of overlap by calculating the ratio of overlapped to total changes in the membrane channel using the JACoP plug-in of the ImageJ software. Then we obtained fluctuations in the extent of overlap at different time points before or post glycine treatment to evaluate the randomness of overlap between membrane expansion and actin polymerization.

Immunohistochemical analyses

Mice were anesthetized with 1% sodium pentobarbital and transcardially perfused with

594 normal saline followed by 4% paraformaldehyde (PFA) in 0.01 M phosphate-buffered saline
595 (PBS). Mouse brain was dissected out and post-fixed with 4% PFA/PBS for 4 h at 4 °C. Fixed
596 brain was incubated with 20% sucrose overnight and then 30% sucrose overnight. The brain
597 was embedded in OCT and stored at -80°C until usage. Thirty-micron cyrosections were
598 made using cryostat and collected.

599 For immunostaining of brain sections, floating 30 µm-thick slices were rinsed with PBS
600 and permeabilized in 0.4% Triton X-100 in 0.01M PBS for 30 min. Cyrosections were
601 blocked with 1% BSA in PBS containing 0.4% Triton X-100 for 1 h at RT, then incubated
602 with primary antibodies overnight at 4 °C. Secondary antibodies conjugated with Alexa Fluor
603 555 were used for detection. Sections were then incubated with DAPI (Roche,
604 Grenzach-Wyhlen, Germany) for nuclear staining for 5 min at RT. Following rinsing,
605 cyrosections were mounted on gelatin-coated slides and covered with coverslip with
606 mounting medium. Confocal images were collected using the Spectral Imaging Confocal
607 Microscope Digital Eclipse C1Si (Nikon, Tokyo, Japan) with a 10×Plan Apochromat DIC N1
608 0.45 objective or 40×Plan Fluo (NA 1.30) oil objective (Yang et al., 2018).

609

610 Protein expression and purification

611 His-EEN1, His-calmodulin, His-p140Cap fragment (aa 351-1051) or GST-Calmodulin was
612 expressed in *E. coli* BL21 (DE3). Cells were grown at 37 °C in LB (g/L: tryptone 10, yeast
613 extract 5, NaCl 10) supplemented with ampicillin or kanamycin. Cells were induced at OD₆₀₀
614 of ~0.6 with 0.4 mM isopropyl β-D-1-thiogalactopyranoside (IPTG) for 4 h at 30 °C or 16 h
615 at 16 °C. Cells were harvested and stored at -80 °C until purification.

616 For His-tagged proteins, cells were resuspended in lysis buffer (50 mM NaH₂PO₄, 300
617 mM NaCl, 15 mM imidazole, pH 8.0) supplemented with 1% Triton X-100 and 0.1 mM
618 PMSF. Protein were purified with Ni-NTA resin according to the manufacturer's instructions
619 (R90115, Invitrogen). For GST-tagged proteins, cells were resuspended in PBS supplemented
620 with 0.2% Triton X-100 and 0.1 mM PMSF. Protein were purified with Glutathione
621 Sepharose 4B (GE17-0756-01, Sigma-Aldrich, St. Louis, MO, USA) according to the
622 manufacturer's instructions.

623 For liposome sedimentation assay, purified His-calmodulin was dialyzed in a
624 Spectra/Por™ 4 RC Dialysis Membrane Tubing (08-667D, Thermo Fisher Scientific,
625 Waltham, MA, USA) against 1,000 volumes of PBS supplemented with 0.5 mM EGTA at
626 4 °C for 4 h and replaced with fresh 1,000 volumes of PBS twice.

627

628 Isothermal titration calorimetry (ITC)

629 ITC was performed on a MicroCal PEAQ-ITC ((Malvern Panalytical, U.S.A) calorimeter.
630 Calmodulin and EEN1 were purified on a Superdex-200 16/600 column (GE Healthcare,
631 U.S.A) in solution buffer containing 20 mM Tris-Cl at pH 7.0, 150 mM NaCl. Solution buffer
632 containing 1 mM calcium was injected into the calorimeter cell fullfilled with
633 protein solution, cell temperature set to 25°C. Each analysis involved 20 injections of 4 s
634 duration (2 µL per one injection), 120 s spacing, stir with 750 rpm, 5 µcal/s reference power
635 and high gain feedback mode. Data were processed by Origin software to obtain
636 thermodynamic profiles.

637

638 **GST-pull down, co-immunoprecipitation (IP) and immunoisolation**

639 For GST-pull down assays, 5 μ g of GST-tagged protein conjugated with
640 glutathione-Sepharose beads was incubated with 1 μ g of His-tagged protein in 0.01 M PBS
641 supplemented with 1% NP-40 at 4°C for 1 h. Beads then was washed five times with PBS
642 supplemented with 0.3% Triton-X 100 and was boiled in SDS sample buffer.

643 For co-IP experiments, HEK293T cells, cultured neurons or mouse brain were lysed with
644 lysis buffer 1 (0.05% [vol/vol] NP-40, 15 mM Tris-HCl, pH 7.4, 50 mM NaCl) supplemented
645 with protease inhibitors for IP of endogenous proteins (endo-IP), or with lysis buffer 2 (0.1%
646 [vol/vol] NP-40, 50 mM Tris-HCl, pH 7.4, 150 mM NaCl) supplemented with protease
647 inhibitors for Flag-IP. Lysates were then centrifuged at $16,000 \times g$ for 15 min at 4°C. For Flag
648 IP, cell lysates were incubated with anti-Flag Affinity Gel (Sigma-Aldrich) at 4°C for 4 h on a
649 roller mixer. For endo-IP, antibody (1 μ g) was added to the cell lysates and incubated at 4°C
650 for 2 h on a roller mixer, followed by incubation with Protein G agarose (Santa Cruz)
651 pre-equilibrated in lysis buffer overnight at 4°C. Immunoprecipitates were washed four times
652 in lysis buffer and boiled in SDS sample buffer, then subjected to SDS-PAGE s and
653 immunoblotting.

654 For immunoisolation of membrane proteins, cultured neurons were homogenized with
655 lysis buffer (mM: Tris-HCl 20, HEPES pH 7.4 10, NaCl 150, sucrose 250) supplemented with
656 protease inhibitors and centrifuged at $800 \times g$ for 15 min. The supernatants were collected and
657 subjected to high-speed centrifugation at $100,000 \times g$ for 1h (TLS-55 rotor, OptimaTMMAX
658 Ultracentrifuge; Beckman Coulter, Germany). The supernatants (S100) and pellets (p100, the
659 membrane fraction) resuspended in lysis buffer were subjected to immunoisolation with

660 Dynabeads Protein G (Invitrogen, Carlsbad, CA, USA) coupled with 2 μ g of mouse
661 anti-calmodulin antibody. Bound proteins were eluted by boiling in 2 $\times$ SDS gel loading buffer
662 and subjected to SDS-PAGE and immunoblotting.

663

664 **Subcellular fractionation**

665 Cultured neurons or mouse hippocampi were homogenized to isolate the membrane and
666 cytoskeleton fractions with Subcellular Protein Fractionation Kit for Cultured Cells (Thermo,
667 77840) or for tissues (Thermo, 87790) according to the manufacturer's instructions. Proteins
668 in different fractions were subjected to SDS-PAGE and immunoblotting.

669

670 **Liposome co-sedimentation assay**

671 Brain extract from bovine brain (Sigma-Aldrich, B1502) were dissolved in chloroform and
672 dried under vacuum for 30 min. The solvent-free lipid films were rehydrated with liposome
673 buffer (150 mM NaCl, 20 mM Tris-HCl pH 7.4, 1 mM DTT) and subjected to 7 cycles of
674 flash freezing in liquid nitrogen and thawing in a 37°C bath. Liposomes were then extruded
675 21 times through a polycarbonate membrane with a 50 nm pore size (Mini-Extruder, Avanti
676 Polar Lipids). Extruded liposomes were centrifuged at 18,000 g for 5 min to remove insoluble
677 material and stored at 4°C. Liposomes (0.5 mg/ml) were then incubated with freshly purified
678 recombinant proteins (1 μ M His-EEN1 or 1 μ M His-EEN1 and 2 μ M His-calmodulin), in
679 100 μ l liposome buffer for 10 min at 30°C before sedimentation at 140,000g for 30 min at
680 4°C. The supernatant (unbound) and pellet (bound) were subjected to SDS-PAGE. Ratios of
681 binding to liposomes were determined using NIH ImageJ software.

682

683 **Three-dimensional Structured Illumination Microscopy (3D-SIM) imaging and image**
 684 **analysis**

685 3D-SIM images were acquired as previously described (Niu et al., 2013) on the DeltaVision
 686 OMX V4 imaging system (Applied Precision Inc, USA) with a 100 × 1.4 oil objective
 687 (Olympus UPlanSApo), solid state multimode lasers (488, 593 and 642 nm) and
 688 electron-multiplying CCD (charge-coupled device) cameras (Evolve 512×512, Photometrics,
 689 USA). Serial Z-stack sectioning was done at 125 nm intervals. The microscope is routinely
 690 calibrated with 100 nm fluorescent spheres to calculate both the lateral and axial limits of
 691 image resolution. SIM image stacks were reconstructed using softWoRx 5.0 (Applied
 692 Precision) with the following settings: pixel size 39.5 nm; channel-specific optical transfer
 693 functions; Wiener filter 0.001000; discard Negative Intensities background; drift correction
 694 with respect to first angle; custom K0 guess angles for camera positions. Pixel registration
 695 was corrected to be less than 1 pixel for all channels using 100 nm Tetraspeck beads. For
 696 clarity of display, small linear changes to brightness and contrast were performed on
 697 three-dimensional reconstructions.

698

699 **Electrophysiology**

700 *EEN1^{fl/fl}* mice within 24 h after birth were co-injected with high-titer AAV stock carrying
 701 pAOV-CAMKII-mCherry-2A-Cre (AAV-mCherry-2A-Cre) and
 702 pAAV-CaMKIIa-EGFP-2A-MCS-3FLAG-EEN1 or mutants of EEN1 (about 1 ~ 5 × 10¹³
 703 IU/ml). Newborns were anesthetized on ice for 5 minutes and then mounted in a custom

ceramic mold to make the head level in the X - and Y- axes. Lambda was set as (X, Y) = (0, 0).
Zero point of Z - axis was the position at which the injecting needle penetrated the skin.
About 10 nl viral solution was injected at each of the seven sites ((X, Y, Z) = (1.2, 1.2, 1.4/
1.0/ 0.6) and (1.5, 1.0, 1.7/ 1.3/ 0.9/ 0.5)) targeting the hippocampus at each cerebral
hemisphere with microsyringe (Sutter Instrument) and a beveled glass injection pipette.
Injected pups were returned to home cage and used for recording two to three weeks
afterward. Transverse 350 μ m hippocampal slices were cut from viral injected *EENI^{fl/fl}* mice
on a Leica vibratome (VT1000 S) in high sucrose cutting solution containing (in mM): KCl
2.5, NaH₂PO₄ 1.25, NaHCO₃ 25, CaCl₂ 0.50, MgSO₄ 7, sucrose 210, glucose 10, Na-ascorbic
acid 1.3. Freshly cut slices were placed in an incubating chamber containing ACSF, and
recovered at 32°C for about 20 min followed by 60 min at room temperature before recording.
The slices were perfused with ACSF containing GABAA receptor antagonists PTX (100 μ M)
/ Bic (10 μ M) and saturated with 95% O₂/5% CO₂ in whole-cell LTP experiments. CA1
pyramidal cells were voltage-clamped at -70 mV and AMPAR EPSCs were evoked by
stimulation at SC with concentric electrode (FHC CBBRC75). LTP was induced by
stimulating SC axons at 2 Hz for 90 s while clamping the cell at 0 mV, after recording a stable
3- to 5-min baseline, but no more than 6 min after breaking into the cell (Diaz-Alonso et al.,
2017; Granger et al., 2013). To minimize run-up of baseline responses during LTP, cells were
held cell-attached for about 1 to 2 min before breaking into the cell.

723

724 **Stereotaxic injection and Behavioral tests**

725 Nine-week-old sexually naive male and female mice were anesthetized and stereotactically

726 injected with viral particles in the hippocampal CA1 region as described (Yang et al., 2018).
 727 The virus-injected mice were tested for behavior two weeks later. Morris water maze and fear
 728 conditioning tests were performed as previously described (Yang et al., 2018). We observed
 729 no sex-related difference in behaviors and the results were pooled together.

730

731 **Statistical analysis**

732 All data were presented as the mean $\pm$ SEM. GraphPad Prism 5 (GraphPad Software, LaJolla,
 733 CA) was used for statistical analysis. For two-sample comparisons vs. controls, unpaired
 734 Student's t-test was used except where noted. One-way analysis of variance with a Dunnett's
 735 multiple-comparison or Newman-Keuls multiple comparison *hoc* test was used to evaluate
 736 statistical significance of three or more groups of samples. A *p* value of less than 0.05 was
 737 considered statistically significant.

738

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908 **Additional files**

909 Supplementary files:

910 (A) Figure supplements

911 **Figure 3 - figure supplement 1.** Endophilin A1 does not bind Ca^{2+} .

912 **Figure 4 - figure supplement 1.** I154 and L158 are required for endophilin A1 binding to
913 calmodulin.

914 **Figure 5 - figure supplement 1.** Calmodulin-dependent increase in the number of endophilin
915 A1 nanodomains correlates with spine size during the initial Phase of sLTP.

916 **Figure 6 - figure supplement 1.** In response to Ca^{2+} /calmodulin, endophilin A1 localizes to
917 the plasma membrane and recruits Arp2/3 via p140Cap.

918 (B) Supplementary Videos

919 **Video 1.** Morphological changes and actin dynamics in *EENI* WT dendrite before and during
920 glycine-induced sLTP.

921 Dendrites of DIV16 *EENI* WT mouse hippocampal neurons co-expressing mGFP (green) and
922 the F-actin probe LifeAct-mCherry (red) were imaged every 5 s by GI-SIM for 12 frames
923 before and 48 frames after glycine treatment. Video plays at 5 frames/s. Scale bars, 5 μm (left)
924 and 1 μm (right).

925 **Video 2.** Morphological changes and actin dynamics in *EENI* KO dendrite before and during
926 glycine-induced sLTP.

927 Dendrites of DIV16 *EENI* KO mouse hippocampal neurons co-expressing mGFP (green) and
928 the F-actin probe LifeAct-mCherry (red) were imaged every 5 s by GI-SIM for 12 frames
929 before and 48 frames after glycine treatment. Video plays at 5 frames/s. Scale bars, 5 μm (left)

930 and 1 μm (right).

931 **Video 3.** Spatiotemporal relationship of spine growth and actin polymerization in *EENI* WT

932 spines before and during glycine-induced LTP.

933 Increases in spine head area and F-actin signal intensity are color coded green and red,

934 respectively. Video plays at 5 frames/s. Scale bars, 5 μm (left) and 1 μm (right).

935 **Video 4.** Spatiotemporal relationship of spine growth and actin polymerization in *EENI* KO

936 spines before and during glycine-induced LTP.

937 Increases in spine head area and F-actin signal intensity are color coded green and red,

938 respectively. Video plays at 5 frames/s. Scale bars, 5 μm (left) and 1 μm (right).

939

940 **Figures**

941

Figure 1. Endophilin A1 is required for spine expansion and actin polymerization in the initial Phase of sLTP.

(A) Mouse hippocampal neurons were co-transfected with pLL3.7-DsRed (volume marker) and pCMV-Tag2B (Flag vector) or pCMV-Tag2B-endophilin A1 (Flag-EEN1) on DIV12, pre-treated with DMSO (vehicle control) or MK801 (NMDAR antagonist) and chemically induced LTP with glycine (Gly) on DIV16. Neurons were fixed 1 min or 5 min after glycine application, stained with antibodies to Flag and imaged by confocal microscopy. Bar = 5 μ m.

(B) Quantification of spine size in (A). $N \geq 13$ cells, $n \geq 563$ spines per group. (C) Changes of spine size in (A). (D) GI-SIM imaging of *EEN1* WT and KO hippocampal neurons expressing mGFP and LifeAct-mCherry. Glycine was applied after imaging for 1 min and imaging was continued for 5 more minutes. Shown are representative still images right before and 1 min after glycine application. Lower panels are magnification of boxed areas in upper panels. Bars: 5 μ m (upper) and 1 μ m (lower). (E, F) Quantitative analysis of spine size increase (E) or F-actin enrichment (F) in spines imaged by GI-SIM. (G) Spine membrane expansion overlapping with actin polymerization in D. Spine growth and increase in F-actin signals are color coded green and red, respectively. Bars: 5 μ m (upper) and 1 μ m (lower). (H) Quantification of fractions of expanded membrane overlapping with actin polymerization in individual spines of *EEN1* WT and KO neurons before and after glycine application at 5 s intervals. (I) Quantification of fractions of membrane expansion overlapping with actin polymerization at 1 min after glycine application in individual spines of *EEN1* WT and KO neurons. (J) Changes in the extent of overlap between membrane expansion and actin polymerization 1 min after glycine application in individual spines of *EEN1* WT and KO

964 neurons. Data are normalized to the time point right before glycine application. **(K)** Mean
 965 fluctuation of the overlap between membrane expansion and actin polymerization in
 966 individual spines within 1 min before and after glycine application for *EENI* WT and KO
 967 neurons. Data represent mean $\pm$ SEM in **(B)**, **(C)** and **(I)**. WT: N = 5 neurons and n = 24
 968 spines; KO: N = 6 neurons and n = 27 spines in **(E)**, **(F)** and **(H-K)**. * $p < 0.05$, ** $p < 0.01$,
 969 *** $p < 0.001$.
 970

Figure 2. Membrane unfolding and branched actin polymerization are required for initiation

of sLTP.

(A) Mouse hippocampal neurons transfected with pLL3.7-DsRed on DIV12 were pre-treated

with DMSO or tetanus toxin (TeTx) for 10 min and chemically induced LTP with glycine on

DIV16. Neurons were fixed at different time points after glycine application and imaged by

confocal microscopy. Bar = 5 μm . (B) Quantification of spine size in (A). $N \geq 12$ neurons, n

≥ 530 spines per group, *** $p < 0.001$. (C) Effect of osmotic shock on spine enlargement 1

min after glycine application. Neurons were pre-treated with iso-osmotic, hypo-osmotic or

hyper-osmotic solution respectively for 10 min and chemically induced LTP in the same

solution on DIV16. Neurons were fixed 1 min after glycine application and imaged by

982 3D-SIM microscopy. Bar = 4 μ m. **(D)** Quantification of spine size in **(C)**. $N \geq 12$ neurons, $n \geq$
983 382 spines per group. **(E)** Effects of W-7 (calmodulin inhibitor) and KN-62 (CaMKII
984 inhibitor) on spine enlargement 1 min after glycine application. Bar = 5 μ m. **(F)**
985 Quantification of spine size in **(E)**. $N \geq 12$ neurons, $n \geq 504$ spines per group. **(G)** Effects of
986 inhibitors for actin remodeling regulators on spine enlargement 1 min after glycine
987 application. Shown are representative confocal images. Bar = 5 μ m. CK-666: Arp2/3 inhibitor.
988 SMIFH2: Formin inhibitor. CT04: RhoA inhibitor. NSC 23766: Rac1 inhibitor. ML 141:
989 Cdc42 inhibitor. **(H, I)** Quantification of spine size (left) and changes in spine size (right) in
990 **(G)**. $N \geq 12$ neurons, $n \geq 516$ spines per group. Data represent mean $\pm$ SEM in **(B)**, **(D)**, **(F)**
991 and **(H)**. *** $p < 0.001$.

992

Figure 3. Ca²⁺/calmodulin enhances the interaction between endophilin A1 and p140Cap.

(A) Diagram showing the domain structure and fragments of endophilin A1 used in this study.

(B) Binding of His-tagged endophilin A1 full length, ΔSH3 and ΔBAR fragments to

997 GST-tagged calmodulin (CaM) in GST-pull down assay. **(C)** Effect of Ca^{2+} on binding of
 998 endophilin A1 to calmodulin in GST-pull down assay. **(D)** Quantification of Ca^{2+} -regulated
 999 endophilin A1 binding to calmodulin in **(C)**. N = 3 independent experiments. **(E)** Effect of
 1000 EGTA on binding of endophilin A1 to calmodulin in GST-pull down assay. **(F)** Quantification
 1001 of endophilin A1 binding to calmodulin in **(E)**. N = 5. **(G)** Effect of EGTA or Ca^{2+} on binding of
 1002 endophilin A1 to p140Cap in transiently transfected HEK293 cells in Flag IP assay. **(H)**
 1003 Quantification of endophilin A1 binding to p140Cap in **(G)**. N = 4. **(I)** Effect of EGTA or Ca^{2+}
 1004 on binding of endophilin A1 to p140Cap in cultured neurons. Endogenous IP assay was
 1005 performed from lysates of mouse neurons with antibodies to endophilin A1. **(J)** Quantification
 1006 of endophilin A1 binding to p140Cap in **(I)**. N = 3. **(K)** Effect of BAPTA or MK801 on the
 1007 binding of endophilin A1 to p140Cap in neurons upon cLTP induction. **(L)** Quantification of
 1008 endophilin A1 binding to p140Cap and total protein levels of endophilin A1 or p140Cap in
 1009 **(K)**. N = 5. **(M)** Effect of W-7 on interactions between calmodulin and endophilin
 1010 A1/p140Cap upon cLTP induction. DIV16 neurons were collected and the cytosolic (S100)
 1011 and membrane (P100) fractions were used for immunoisolation with antibodies to calmodulin.
 1012 **(N)** Quantification of endophilin A1 and p140Cap precipitated by antibodies to calmodulin in
 1013 **(N)**. N = 3. **(O)** Effects of W-7 and MK801 on the amount of p140Cap in spines upon LTP
 1014 induction. Neurons transfected with pLL3.7-DsRed on DIV12 were pre-treated with DMSO,
 1015 W-7 or MK801 and chemically induced LTP on DIV16. Neurons were fixed 1 min after
 1016 glycine application, immunostained with antibodies to p140Cap (green) and imaged by
 1017 confocal microscopy. Bar = 5 μm . **(P)** Quantification of changes in p140Cap mean intensity
 1018 in spines as compared with the control (- Gly) group in **(O)**. N $\geq$ 12 cells, n $\geq$ 464 spines per

1019 group. **(Q)** Same as **(O)** except that neurons were immunostained with antibodies to cortactin.
1020 Bar = 5 μ m. **(R)** Quantification of changes in cortactin mean intensity in spines as compared
1021 with the control (- Gly) group in **(Q)**. $N \geq 11$ cells, $n \geq 431$ spines per group. Data represent
1022 mean $\pm$ SEM in **(D)**, **(F)**, **(H)**, **(J)**, **(L)**, **(N)**, **(P)** and **(R)**. * $p < 0.05$, ** $p < 0.01$, *** $p <$
1023 0.001, when compared with Ctrl, NT or - Gly. ### $p < 0.001$ in **(L)**, when compared with +
1024 Gly. # $p < 0.05$, ## $p < 0.01$, ### $p < 0.001$ in **(N)**, when compared to + Gly 1'.
1025 **Figure 3 - figure supplement 1.** Endophilin A1 does not bind Ca^{2+} .
1026

Figure 4. Ca²⁺/calmodulin promotes actin polymerization in spines via the endophilin

A1-p140Cap pathway.

(A) Effect of Ca²⁺ on binding of EEN1 double mutant (DM) to calmodulin in GST-pull down

assay. (B) Quantification of EEN1 DM binding to GST-CaM as compared with WT in (A). Y

axis shows as two segments. N = 4, *** *p* < 0.001. (C) Cultured EEN1^{+/+} and EEN1^{-/-}

hippocampal neurons co-transfected with LifeAct-mCherry and pCMV-Tag2B, and EEN1^{-/-}

hippocampal neurons co-transfected with LifeAct-mCherry and pCMV-Tag2B-EEN1 WT or

pCMV-Tag2B-EEN1 DM on DIV12 were pre-treated with DMSO or W-7 and chemically

induced LTP (cLTP) on DIV16. Neurons were fixed 1 min after glycine application and

1037 imaged by confocal microscopy. Bar = 5 μ m. **(D, E)** Quantification of spine size and changes
 1038 in spine size in **(C)**. $N \geq 12$ neurons, $n \geq 450$ spines per group. **(F)** Quantification of F-actin
 1039 enrichment in spines in **(C)**. $N \geq 12$ neurons, $n \geq 450$ spines per group. **(G)** Cultured *EENI*^{+/+}
 1040 and *EENI*^{-/-} hippocampal neurons co-transfected with pLL3.7-DsRed and pCMV-Tag2B, and
 1041 *EENI*^{-/-} hippocampal neurons co-transfected with pLL3.7-DsRed and pCMV-Tag2B-EEN1
 1042 WT or pCMV-Tag2B-EEN1 Y343A on DIV12 were pre-treated with DMSO or MK801 and
 1043 chemically induced LTP (cLTP) on DIV16. Neurons were fixed 1 min after glycine
 1044 application and imaged by confocal microscopy. Bar = 5 μ m. **(H, I)** Quantification of spine
 1045 size and changes in spine size in **(G)**. $N \geq 12$ neurons, $n \geq 502$ spines per group. Data
 1046 represent mean $\pm$ SEM in **(B)**, **(D)**, **(E)**, **(F)**, **(H)**, and **(I)**. ** $p < 0.01$, *** $p < 0.001$ in **(B)**
 1047 when compared with 0 μ M Ca^{2+} .

1048 **Figure 4 - figure supplement 1.** I154 and L158 are required for endophilin A1 binding to
 1049 calmodulin.

1050

Figure 5. Ca^{2+} /calmodulin-enhanced plasma membrane association of endophilin A1

correlates with spine expansion.

(A) DIV16 neurons were pre-treated with DMSO, W-7 or KN-62 and chemically induced LTP.

Neurons were fixed 1 min after glycine application. Plasma membrane (PM)-localized

endophilin A1 was immunostained and imaged by 3D-SIM. Spines were outlined manually.

Bar = 4 μ m in left and center panels and 500 nm in magnified images in right panels. **(B)**

Quantification of spine size in **(A)**. **(C)** Quantification of the area and number of

PM-localized endophilin A1 nanodomains in spines in **(A)**. **(D)** Scatterplot of the number or

area of PM-localized endophilin A1 nanodomains versus the size of spine head for control (-

Gly, n = 278 spines) and cLTP (+ Gly, n = 254 spines) groups with linear fits. **(E)**

Quantification of changes in the mean intensity of PM-localized endophilin A1 in spines. **(F)**

DIV16 neurons were pre-treated with DMSO or LatA and chemically induced LTP. Neurons

were fixed at 30 s, 45 s, 60 s or 120 s after glycine application. PM-localized endophilin A1

was stained and imaged by 3D-SIM. Bar = 500 nm. **(G)** Quantification of spine size in **(F)**.

(H) Quantification of the mean intensity of PM-localized endophilin A1 in spines in **(F)**. **(I)**

Quantification of changes in the mean intensity of PM-localized endophilin A1 in spines in

(F). **(J)** Effect of Ca^{2+} /calmodulin on binding of EEN1 WT or DM protein to membrane in

liposome sedimentation assay. **(K)** Quantification of endophilin A1 binding to liposome in **(J)**.

N = 7. Data represent mean $\pm$ SEM in B-E, G-I, and K. *** $p < 0.001$, when compared with -

Gly in **(G)** and **(H)**. ### $p < 0.001$, when compared with - Gly + LatA treatment in **(H)**. N $\geq$

10 neurons, n $\geq$ 254 spines per group in **(B-E)**, and N $\geq$ 10 neurons, n $\geq$ 241 spines per group

in **(G-I)**.

Figure 5 - figure supplement 1. Calmodulin-dependent increase in the number of endophilin

A1 nanodomains correlates with spine size during the initial phase of sLTP.

Figure 6. Ca^{2+} /calmodulin regulates the association of endophilin A1 and p140Cap with both membrane and cytoskeleton upon LTP induction.

(A) Immunoblotting of subcellular fractionations of mouse hippocampi from animals subjected to fear training. (B) Quantification of protein levels of p140Cap, endophilin A1 or endophilin A2 in cytoplasmic, membrane and cytoskeleton fractions in (A). N = 6 animals. (C) Cultured neurons were pre-treated with DMSO, 10 μM MK801, W-7 or KN-62 and chemically induced LTP on DIV16. Cell lysates were prepared 1 min after glycine application and subjected to SDS-PAGE and immunoblotting. (D) Quantification of total protein levels in (C). N = 5. (E) Effects of MK801, W-7 and KN-62 on subcellular distribution of protein levels. Neurons were pre-treated with DMSO, MK801, W-7 or KN-62 and chemically induced LTP on DIV16. Cells were collected and subjected to subcellular fractionation 1 min after glycine application. (F) Quantification of protein levels in subcellular fractions in (E). N = 5. (G) *EENI*^{+/+} and *EENI*^{-/-} hippocampal neurons co-transfected with pLL3.7-DsRed and pCMV-Tag2B, and *EENI*^{-/-} hippocampal neurons co-transfected with pLL3.7-DsRed and pCMV-Tag2B-EEN1 WT or pCMV-Tag2B-EEN1 KKK-EEE on DIV12 were pre-treated with DMSO or MK801 and chemically induced LTP on DIV16. Neurons were fixed 1 min after glycine application and imaged by confocal microscopy. Bar = 5 μm . (H, I) Quantification of spine size or changes in spine size in (G). N $\geq$ 12 neurons, n $\geq$ 482 spines per group. Data represented are mean $\pm$ SEM in (B), (D), (F), (H) and (I). * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ when compared to Ctrl or - Gly in (B) and (F).

Figure 6 - figure supplement 1. In response to Ca^{2+} /calmodulin, endophilin A1 localizes to the plasma membrane and recruits Arp2/3 via p140Cap.

Figure 7. The calmodulin-, membrane- and p140Cap-binding capacities of endophilin A1 are required for LTP and long-term memory.

(A-F) Rescue of the LTP impairment phenotype in *EEN1* KO neurons by EEN1 WT and mutants. AAV viruses expressing the Cre recombinase (AAV-mCherry-2A-Cre) and GFP (AAV-EGFP) or Cre, GFP and EEN1 WT or mutant (AAV-EGFP-2A-EEN1) were stereotactically injected into the CA1 regions of *EEN1^{fl/fl}* mice at P0. Acute slices of hippocampi were prepared on P14-P21 for dual recording analysis of LTP. Shown are pairwise comparisons of LTP in non-infected (Ctrl) and infected neurons of the same slice. For Cre-expressing neurons (Cre) versus Ctrl in (A), 6 recording pairs from three mice, marked as N = 6/3, were analyzed. N = 10/5 in (B), N = 5/4 in (C), N = 5/3 in (D) and N = 4/3 in (E). Bar graphs shown percentage of baseline at 45 min after pairing. A summary of rescue effects of EEN1 WT and mutants is shown in (F). Data represent mean $\pm$ SEM. * $p < 0.05$, ** $p < 0.01$. (G) AAV virus was stereotactically injected into the CA1 regions of *EEN1^{+/+}* to express GFP alone, or *EEN1^{-/-}* mice to express GFP alone, EEN1 WT, KKK-EEE, Y343A or DM and GFP. Shown are images of GFP fluorescent signals and DAPI labeling of nuclei captured by fluorescence microscopy. Bar = 1 mm. (H) Immunofluorescence staining of endophilin A1 in CA1 neurons of brain slices from mice in (G). Bar = 50 μ m. (I-N) Morris water maze test of AAV-injected mice. Shown are escape latency or distance travelled before escaping to the platform in the visible platform training (I), escape latency and distance travelled before escaping to the platform in the invisible platform training (J), the swim trace in probe trial 3 and recall following training once again one month after training (K, L), number of crossings within the 1.5 $\times$ platform area in 5-day probe trial and recall test (M), and number of crossings within the 1.5 $\times$ platform area before and after recall (N). (O) Freezing behavior in AAV-injected *EEN1^{+/+}* or *EEN1^{-/-}* mice subjected to contextual fear conditioning.

1125 Data represent mean $\pm$ SEM in H-O (15 *EENI*^{+/+} + GFP, 10 *EENI*^{-/-} + GFP, 6 *EENI*^{-/-} +
1126 *EEN1*, 6 *EENI*^{-/-} + Y343A, 7 *EENI*^{-/-} + KKK-EEE, 7 *EENI*^{-/-} + DM), * $p < 0.05$, ** $p < 0.01$,
1127 *** $p < 0.001$ when compared with *EENI*^{+/+} + GFP, # $p < 0.05$, ## $p < 0.01$, ### $p < 0.001$
1128 when compared with *EENI*^{-/-} + GFP; & $p < 0.05$, && $p < 0.01$ when compared with before
1129 recall.
1130

1131

Figure 8. Model for endophilin A1-mediated initial expansion of spine head in sLTP.

LTP stimuli induce NMDAR-mediated Ca^{2+} influx into dendritic spines and activation of calmodulin. Ca^{2+} /calmodulin interacts directly with endophilin A1 and enhance its binding to both the plasma membrane and p140Cap. As a result, the plasma membrane-associated endophilin A1 recruits p140Cap, which in turn recruits cortactin to promote branched actin polymerization underneath the plasma membrane, generating propulsive force for rapid structural expansion of spine head during the initial phase of sLTP.