The APP-interacting protein FE65 is required for hippocampus-dependent learning and long-term potentiation

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FE65 is expressed predominantly in the brain and interacts with the C-terminal domain of β-amyloid precursor protein (APP). We examined hippocampus-dependent memory and in vivo long-term potentiation (LTP) at the CA1 synapses with isoform-specific FE65 knockout (p77FE65<sup>−/−</sup>) mice. When examined using the Morris water maze, p77FE65<sup>−/−</sup> mice were impaired for the hidden platform task but showed normal performance in the probe test. To further discriminate the role of FE65 in acquisition and memory consolidation, we examined p77FE65<sup>−/−</sup> mice with temporal dissociative passive avoidance (TPDA) and contextual fear conditioning (CFC). p77FE65<sup>−/−</sup> mice showed impaired short-term memory for both TPDA and CFC when tested 10 min after training. After multiple TPDA training sessions, the crossover latency of some p77FE65<sup>−/−</sup> mice reached the cutoff value, but it significantly decayed in 8 d. At the Schaffer collateral-CA1 synapses, p77FE65<sup>−/−</sup> mice showed defective early-phase LTP (E-LTP). These results demonstrate novel roles of FE65 in synaptic plasticity, acquisition, and retention for certain forms of memory formation.

[Supplemental material is available online at http://www.learnmem.org.]
FE65 and hippocampus-dependent plasticity

Results

p97FE65\(^{-/-}\) mice show normal hippocampus formation

We first examined the overall morphology of the hippocampus of isoform-specific p97FE65\(^{-/-}\) mice. Neutral red staining (Supplemental Fig. 1A) and calbindin staining (Supplemental Fig. 1B) revealed no significant difference between wild-type (WT) and p97FE65\(^{-/-}\) mice in gross hippocampal morphology and mossy fiber formation, respectively. Immunostaining for the neuronal and dendritic marker MAP2 also revealed no significant difference in cortex and different areas of the hippocampus including CA1, CA3, and dentate gyrus (Supplemental Fig. 1C). The expression levels of synaptic marker proteins Synapsin I (Supplemental Fig. 1D) and VGAT (Supplemental Fig. 1E) were comparable between WT and p97FE65\(^{-/-}\) mice. These data, together with a previous report (Wang et al. 2004a), implicate that there is no severe abnormality in hippocampal morphology in p97FE65\(^{-/-}\) mice.

p97FE65\(^{-/-}\) mice exhibit impaired acquisition, but normal memory retrieval, in the MWM task

Despite major efforts to probe the function of FE65 in AD pathogenesis, little has been investigated to study plasticity under normal and nonpathological conditions. A previous study documented very interesting memory phenotypes in spatial and associative memory with isoform-specific p97FE65\(^{-/-}\) mice at 14 and 27 mo (Wang et al. 2004a). Due to aging effects on plasticity, the phenotypes observed with aged p97FE65\(^{-/-}\) mice may be complicated by other factors. In addition, it is not clear whether FE65 regulates learning/acquisition or memory consolidation and retention. Here, we examined several forms of learning and memory with young adult mice (2–4 mo old).

WT and p97FE65\(^{-/-}\) mice were first trained by the visible platform version of the Morris water maze (MWM), during which a visible flag was attached to the platform. Both groups showed significant improvement in escape latency over the training sessions (Fig. 1A), and there was no significant difference between WT and p97FE65\(^{-/-}\) mice (F\(_{1,20}\) = 0.12, P = 0.5326). Furthermore, WT and p97FE65\(^{-/-}\) mice displayed similar swimming speed during visible platform training (Fig. 1B). These results indicate that p97FE65\(^{-/-}\) mice are normal in nonspatial learning, motivation, vision, and motor functions.

We then further trained the same animals by a hidden platform task, during which the platform was placed underneath opaque water, and animals learned to use extramaze spatial cues to navigate the circular pool and find the hidden platform. Although both WT and p97FE65\(^{-/-}\) showed significant improvement in escape latency during training (F\(_{8,20}\) = 13.88, P < 0.0001), repeated-measures of two-way ANOVA showed a significant difference between genotypes (F\(_{1,20}\) = 23.45, P < 0.0001; Fig. 1C). There was no significant difference in swimming speed between WT and p97FE65\(^{-/-}\) mice during the hidden platform training.

WT and p97FE65\(^{-/-}\) mice show defective hidden platform learning. (A) Both p97FE65\(^{-/-}\) (KO) and wild-type (WT) mice show comparable performance during the visible platform training. (B) p97FE65\(^{-/-}\) mice show comparable swim speed to that of WT mice during the visible platform training. (C) p97FE65\(^{-/-}\) mice show impaired improvement in escape latency during the hidden platform training. (D) WT and p97FE65\(^{-/-}\) mice show comparable swimming speed during the hidden platform training. (E) WT and p97FE65\(^{-/-}\) mice show comparable time spent in the target quadrant during the probe test. (F) p97FE65\(^{-/-}\) mice show normal acquisition during the reversal training. (G) WT and p97FE65\(^{-/-}\) mice show comparable swimming speed during the reversal training. (H) p97FE65\(^{-/-}\) mice show normal performance in the probe test after reversal training. The probe test was performed 1 d after the last reversal training session.
(P = 0.2034, two-way ANOVA; Fig. 1D). The travel distance for p97FE65−/− mice to find the platform was also significantly longer (F(1,16) = 11.27, P = 0.0031; Supplemental Fig. 2A).

Next, we examined the animals with the probe test, during which the hidden platform was removed. Although the time spent to first reach the hidden platform location was significantly longer for the p97FE65−/− mice during the probe test (P < 0.01, data not shown), the two groups showed no significant difference in the percentage of time spent in the target quadrant (P = 0.2594; Fig. 1E). There was also no significant difference between WT and p97FE65−/− mice in the crossing number for the hidden platform location (data not shown).

We then trained the animals with the reversal paradigm, during which the hidden platform was moved to the opposite quadrant. This task requires the animals to relearn the new platform location and suppress the pre-established memory for the old platform location. As shown in Figure 1F, WT and p97FE65−/− mice showed a similar decrease in escape latency during the reversal training. WT and p97FE65−/− mice also showed comparable swimming speed during reversal training (P = 0.1274, two-way ANOVA; Fig. 1G), and there was no significant difference in the travel distance either (P = 0.8794, two-way ANOVA; Supplemental Fig. 2B). When tested by the probe test, both groups showed significant preference for the new platform location (P < 0.05 for both genotypes; Fig. 1H).

p97FE65−/− mice show impaired learning and memory retention in TDPA

Because p97FE65−/− mice showed a slower rate of acquisition but normal performance in the probe test, it is possible that FE65 is required for learning rather than consolidation. To further confirm the possibility, we next trained animals with TDPA, during which multiple training sessions were involved and the learning/acquisition could be better examined. Compared with standard passive avoidance (PA), the animals received a mild electric footshock 10 min after they crossed over to the dark chamber in the TDPA paradigm. Therefore, the association between the dark chamber (the conditioned stimulus) and the aversive footshock (the unconditioned stimulus) was temporally separated by 10 min. Our previous results have demonstrated that animals gradually improve their crossover latency through the multiple trainings, and the task is dependent on the hippocampus (Zhang et al. 2008). As shown in Figure 2A, p97FE65−/− mice displayed significantly slower improvement in crossover latency than WT mice during TDPA training (P < 0.05). Furthermore, the percentage of animals reaching the cutoff value (300 sec) of crossover latency was higher for WT mice (Fig. 2B). After 5 d of TDPA training, all WT mice reached the cutoff value when tested on day 6. Only 60% of p97FE65−/− mice (six out of 10 animals) reached the cutoff value on day 6. To test long-term memory retention, we selected animals that reached the cutoff value and retested them 8 d later. As shown in Figure 2C, p97FE65−/− mice showed significant shorter crossover latency, indicating that p97FE65 is required for long-term retention of TDPA memory.

To further examine the ability of learning or acquisition, another cohort of WT and p97FE65−/− mice was trained by a single TDPA trial and tested shortly (10 min) after the training. As shown in Figure 2D, p97FE65−/− mice showed significantly shorter crossover latency (F(1,26) = 8.12, P = 0.0215). These results demonstrate that p97FE65−/− mice are defective in learning/acquisition, as well as retention, which may involve both consolidation and reconsolidation. Because p97FE65−/− mice showed comparable crossover latency to that of WT mice during the first TDPA training session (Fig. 2A,D), it is unlikely that their deficits during testing are attributed to an abnormal preference for darkness. We further performed the dark/light test, during which their preference for the lit and dark chambers was examined (Krishnan et al. 2008). The animals were first placed in the dark chamber; p97FE65−/− and WT mice displayed a similar latency to cross over to the lit chamber (Fig. 2E). During the 10-min dark/light test, animals were allowed to travel freely between the dark and lit chambers, and the two groups stayed for similar length of time in the lit chamber (Fig. 2F).

Figure 2. p97FE65−/− mice show deficits in TDPA. (A) p97FE65−/− (KO) mice show poor improvement in crossover latency during TDPA training. Mice were trained on day 1 and tested daily on days 2–6. From day 2 to day 5, mice were first scored for crossover latency. Then they received a mild footshock 10 min after entering the dark chamber. When a particular mouse reached the 300-sec arbitrary cutoff value, it was not further trained. The value of 300 sec was registered as crossover latency for the test and for all the tests thereafter. (B) The percentage of animals reaching the cutoff value of crossover latency is lower for p97FE65−/− mice during whole training sessions. (C) p97FE65−/− mice show impaired long-term memory retention. After 5 d of TDPA training, all WT mice reached the 300-sec cutoff value. Six out of 10 p97FE65−/− mice also showed crossover latency of 300 sec. These mice were tested 8 d later to measure their long-term memory retention. For example, if a particular mouse reached the 300 sec value on day 4, it was further tested on day 12. (D) p97FE65−/− mice show deficits in short-term memory for TDPA. Animals were trained by a single session of TDAP and tested 10 min after the training. (E) During the dark/light test, WT and p97FE65−/− mice showed comparable latency to cross over from the dark chamber to the lit chamber. (F) During the dark/light test, WT and p97FE65−/− mice spent comparable time in the lit chamber. The values for the first 5 min of the entire dark/light test are shown.
p97FE65−/− mice. However, p97FE65−/− mice displayed significantly less immediate freezing, indicating impaired acquisition/learning (P < 0.01, analysis with both two-way ANOVA and Bonferroni post-test; Fig. 3A). When tested 24 h after training, p97FE65−/− mice showed less freezing 24 h after the training. (B) p97FE65−/− mice and WT mice show comparable reaction during the footshock. Both genotypes show similar low velocity before the mild electric footshock. The velocity increased significantly during the footshock for both WT and p97FE65−/− mice. (C,D) The animals were introduced to the CFC training chamber for 10 min, and the time for ambulatory movement (C) or total movement (D) was measured by the TruScan system. Data for the first 5 min or last 5 min are shown. (E) p97FE65−/− mice showed normal freezing when they were tested 24 h after the hippocampus-independent CFC training.

Figure 3. p97FE65−/− mice show defective acquisition in CFC. (A) p97FE65−/− (KO) and WT mice show comparable freezing before the mild footshock (during training). p97FE65−/− mice showed less freezing (immediate freezing) after the footshock. p97FE65−/− mice also showed less freezing 24 h after the training. (B) p97FE65−/− mice and WT mice show comparable reaction during the footshock. Both genotypes show similar low velocity before the mild electric footshock. The velocity increased significantly during the footshock for both WT and p97FE65−/− mice. (C,D) The animals were introduced to the CFC training chamber for 10 min, and the time for ambulatory movement (C) or total movement (D) was measured by the TruScan system. Data for the first 5 min or last 5 min are shown. (E) p97FE65−/− mice showed normal freezing when they were tested 24 h after the hippocampus-independent CFC training.

Acute knockdown of FE65 by siRNA leads to defective learning in TDPA

To specifically reduce the expression of FE65 in the hippocampus, we infused FE65 siRNA to the dorsal hippocampus. We first demonstrated that the siRNA could effectively reduce both p97 and p60 by >70% 48 h after administration (Fig. 5A). We next trained the animals by TDPA 48 h after hippocampal infusion with control or FE65 siRNA. When tested 10 min after training, the FE65 siRNA-infused mice showed significantly shorter crossover latency (Fig. 5B).

Inhibition of phosphodiesterase activity rescues the learning deficits in the isoform-specific p97FE65−/− mice

As described earlier, we showed that the hippocampal morphology, as well as the expression levels of neuronal and synaptic marker proteins, is normal in p97FE65−/− mice. To further test whether some of the plasticity-related signaling pathways are intact in p97FE65−/− mice, we administered rolipram to elevate cAMP signaling. Rolipram is a specific inhibitor of the type 4 phosphodiesterase (PDE4), and it enhances the activity of cAMP/PKA/CREB and LTP (Vitolo et al. 2002; Gong et al. 2004). Compared with the vehicle-injected control, rolipram-injected p97FE65−/− mice showed significant improvement in TDPA. The crossover latency (tested 10 min after training) in the rolipram-injected p97FE65−/− mice was comparable to that of the WT mice (Fig. 6A). These data demonstrate that the function of an FE65-independent signal transduction pathway is intact in p97FE65−/− mice. Interestingly, rolipram treatment rescues memory deficits in a mouse model for AD (Gong et al. 2004). We also found that rolipram caused potentiation of fEPSP (Fig. 6B,C). However, administration of rolipram did not rescue the HFS-induced E-LTP

Figure 4. p97FE65−/− mice show defective in vivo E-LTP induced by a single tetanus. The field EPSP at the Schaffer collateral-CA1 synapses was recorded from anesthetized WT and p97FE65−/− (KO) mice. (A) p97FE65−/− and WT mice show comparable input–output curves. (Right) Representative traces of fEPSP from WT and p97FE65−/− (KO) mice. (B) p97FE65−/− and WT mice show comparable PPF. Representative traces for PPF (with 80-msec interval) are shown. (C) Early-phase LTP (E-LTP) was induced by a single train of HFS (100 Hz, 1-sec duration), p97FE65−/− mice showed lower potentiation. Representative traces before HFS (trace 1) and 60 min after HFS (trace 2) are shown.
Figure 5. siRNA-mediated knockdown of FE65 results in short-term memory deficits for TDPA. (A) Effective knockdown of FE65 by an siRNA approach. To determine the specific effects of siRNA on FE65 expression, neurons were transfected with control or FE65 siRNA. Samples were collected 48 h after transfection and analyzed by Western blot. The expression of both p97 and p60 was significantly reduced in neurons transfected with FE65 siRNA. (B) infographic shows the hippocampus results in defective short-term memory of TDPA. Animals infused with control or FE65 siRNA were trained by a single TDPA session and tested 10 min after training.

Discussion

The function of FE65 in learning and memory was addressed with isoform-specific p97FE65−/− mice, which express p60 but not the full-length p97. In a previous study, 14- and 27-mo-old p97FE65−/− mice were compared with their WT littermates. At 14 mo, p97FE65−/− mice showed defective performance in TDPA. At 27 mo, p97FE65−/− and WT mice showed similar performance that is significantly worse than 14-mo-old WT mice. Interestingly, the level of p97 but not p60 declined significantly in 27-mo-old WT mouse. Because p60 lacks part of the WW domain, its transcriptional activity is compromised (B Cool, G Zitnik, G Martin, and Q Hu, unpubl.). Therefore, it is suggested that the learning and memory deficits in the isoform-specific p97FE65−/− mice may be due to the lack of activity-dependent transcription mediated by p97. However, the previous study did not test whether p97FE65−/− mice were impaired for learning/ acquisition or memory consolidation/ retention. To avoid the complication of aging, we examined young adult p97FE65−/− mice (2–4 mo old) with a battery of hippocampus-dependent paradigms. In the TDPA paradigm, the p97FE65−/− mice can eventually learn and reach the cutoff value of crossover latency after multiple training sessions (Fig. 2A). However, they showed significantly less memory retention (Fig. 2B), indicating that p97 may regulate the transcription of memory-related genes. Because both consolidation and reconsolidation are triggered during the multiple training sessions, it is difficult to determine the function of FE65 in these distinct transcription-dependent processes. Our preliminary experiments detected that membrane depolarization by KCl, which induces calcium influx and gene transcription, stimulated the nuclear translocation of FE65 in primary hippocampal neurons (C. Moon and H Wang, unpubl.). To demonstrate the role of FE65-mediated transcription, future experiments need to identify the target genes of FE65 in stimulated neurons and during memory consolidation/reconsolidation.

Surprisingly, the major deficits with p97FE65−/− mice are related to short-term memory or acquisition, which is a transcription-independent process. Consistently, the isoform-specific p97FE65−/− mice showed severe impairment in E-LTP at the Schaffer collateral-C1 synapses. These data suggest that FE65 may be a multifunctional protein that is involved in both transcription-dependent and -independent phases of memory formation. Although cellular studies implicate APP function in transcription, which depends on FE65 (Cao and Sudhof 2001), the existing data suggest that APP may regulate neuronal function in transcription-independent manners. APP KO mice show similar phenotypes to that of p97FE65−/− mice. They display impaired acquisition in the conditioned avoidance test and the hidden platform paradigm (Dawson et al. 1999). APP KO mice also show defective potentiation shortly after HFS. When hippocampal slices from APP KO mice were stimulated by four 10-burst trains (at 100 Hz with a 20-sec interval), the potentiation started to decline immediately after HFS and was significantly lower than WT controls within 30 min (Dawson et al. 1999; Seabrook et al. 1999).

Analysis of data obtained from p97FE65−/− and APP KO mice implicates aging/developmental effects. The learning impairments are observed with 10-mo-old, but not 4-mo-old, APP KO mice (Dawson et al. 1999). The LTP impairments are observed with 8- and 20-mo-old APP KO mice (Dawson et al. 1999; Seabrook et al. 1999). LTP in younger APP KO mice has not been examined. Compared with 2- to 4-mo-old mice, p97FE65−/− mice show more severe impairments in spatial memory at 14 mo. The 14-mo-old p97FE65−/− mice show defective acquisition in both hidden platform and reversal training. Although p97FE65−/− mice (14-mo-old) do not show significant defects in the probe test after hidden platform training, they are severely impaired in the probe test after reversal training (Wang et al. 2004a). During reversal training, the animals need to actively suppress the previously established memory, and in the meantime acquire new information to relearn the new platform location. In this study, the young adult p97FE65−/− mice showed normal time spent in the target quadrant, but we noticed that they took a longer time to reach the platform location during the first probe test (after hidden platform training but before HFS) (Fig. 2B). HFS in the mutant mice may not be as precise as the WT animals. Therefore, memory suppression may be easier for the young adult p97FE65−/−.

Figure 6. PDE4 inhibitor rescues TDPA deficits, but not LTP, in p97FE65−/− (KO) mice. (A) Twenty minutes before TDPA training, mice were given rolipram (0.1 mg/kg in 1% DMSO/PBS) or vehicle (1% DMSO/PBS) through i.p. injection. Mice were trained by a single TDPA session. The animals were tested 10 min after training. (B) After baseline iEPSP was established, p97FE65−/− mice were given rolipram (0.1 mg/kg through i.p. injection). HFS (100 Hz, 1 sec duration) was delivered 20 min after rolipram administration. iEPSP was recorded for 60 min after HFS. Representative iEPSP traces before rolipram treatment but before HFS (trace 1), after rolipram treatment (trace 2), and 60 min after HFS (trace 3) are shown. (C) The relative values of iEPSP corresponding to baseline (values for trace 1), rolipram-treated (values for trace 2), and HFS-induced (values for trace 3) responses are shown.
mice, and their performance in the reversal test, as a balanced outcome of old memory suppression and new learning, appeared to be comparable to that of WT mice. It is intriguing that aged p97FE65 /− /− mice are severely impaired for the reversal test (Wang et al. 2004a). One possibility is that the expression of other members of the FE65 family also declines with age. Because the FE65 family members interact with similar sets of signaling molecules, including APP (McLoughlin and Miller 2008), the compensation from FE65L1 and FE65L2 in p97FE65 /− /− mice may dramatically diminish following aging.

How could APP and FE65 regulate the transcription-independent events involved in plasticity? One possibility is that the basal level of certain plasticity-related gene products is lower in APP KO and p97FE65 /− /− mice. Alternatively, it has been shown that both APP and FE65 are localized in the nerve terminals and are associated with the actin-rich lamellipodia in the growth cones (Sabo et al. 2003), suggesting their function in neurite growth and synapse modification. Consistently, APP-containing membranous cargos undergo rapid axonal transport (Lazarov et al. 2005). Moreover, knocking down presynaptic APP impairs synaptic activity (Herard et al. 2006). FE65 may also regulate synaptic plasticity by regulating Aβ production and secretion (Sabo et al. 1999; Wang et al. 2004a). Compared with Tg2576 neurons, which have a high level of Aβ40 and Aβ42, significantly reduced in p97FE65 /− /− /Tg2576 double-transgenic neurons. Although exogenous application of Aβ (at 200 nM) impairs LTP, a slight increase of Aβ (at 200 pM) enhances both LTP and spatial memory (Puzzo et al. 2008). Future experiments need to determine whether the endogenous Aβ is lower, and whether application of picomolar Aβ rescues the LTP and learning in p97FE65 /− /− mice.

FE65 may regulate neuronal function in an APP-independent manner, and vice versa. At least, the FE65 family members interact with all members of the APP family (Bressler et al. 1996; Guenette et al. 2006). Although MAP2 staining revealed no significant difference between WT and p97FE65 /− /− mice, APP KO mice showed abnormality in neuronal morphology and development (Seabrook et al. 1999). Other lines of evidence also imply that the interaction between FE65 and other cellular components may be functionally relevant. For example, the FE65/FE65L1 double-KO and Mena KO mice share some common phenotypes, such as hypertrophic neurons and defects in axon pathfinding (McLoughlin and Miller 2008). Many other FE65-interacting proteins have been shown to regulate neuroplasticity under nonpathological conditions. LRP may regulate LTP through its binding to tissue plasminogen activator (Zhuo et al. 2000). ApoE receptors are required for both LTP and CFC (Weeber et al. 2002). Interestingly, FE65 also interacts with the intracellular domain of Notch, and it has been suggested that Notch and APP may regulate a number of same target genes (Fischer et al. 2005). Mice with a reduced level of Notch display defects in both LTP and spatial learning (Costa et al. 2003; Wang et al. 2004b). Because certain regulatory functions may be shared by FE65s (FE65, FE65L1, and FE65L2), using neurons derived from double- or triple-mutants will facilitate the understanding of how the FE65 family participates in the complex signaling network.

Materials and Methods

Animals

Isoform-specific FE65 mutant mice (p97FE65 /− /− ) were generated by conventional gene-specific recombination by replacing exon 2 with a neo cassette and backcrossing into the C3HBL/6 background for at least 10 generations. p97FE65 /− /− mice were previously characterized, and the expression of p97, but not p60, was ablated (Wang et al. 2004a). Animals were housed in the University Laboratory Animal Research Facility, and all the manipulations were in compliance with the guidelines of Institutional Animal Care and Use Committee at Michigan State University. The mice had ad libitum access to water and food and were housed under 12-h dark/light cycles.

Analysis of hippocampal morphology

Fresh whole brains were obtained from age-matched WT and p97FE65 /− /− mice (2–4 mo old), and then cut into 0.5-mm-thick coronal slices by a vibratome. The slices were immediately fixed in PBS containing 6% paraformaldehyde, 50 mM HEPES pH 7.6 for 6 h, subsequently cryoprotected with 20% sucrose/PBS for 24 h, and further processed by a sliding microtome to obtain 40-μm-thick slices. Overall hippocampal histological analysis was performed by Neutral Red staining. Mossy fiber formation was stained by antibodies against Calbindin (1:1000). Antibodies against microtubule-associated protein 2 (1:500) and synapsin I (1:500) and vesicular GABA transporter (VGAT) (1:1000) were used to stain general presynaptic and inhibitory presynaptic markers, respectively. The incubation for primary and secondary (rhodamine red-conjugated) antibodies was 2 h in PBS with 0.1% triton X-100, 3% BSA, and 3% crude serum at room temperature. The samples were then examined by fluorescence detection by confocal microscopy.

Morris water maze (MWM)

The MWM was used for testing hippocampus-dependent spatial memory (Morris et al. 1982). Animal activity was measured by a video-based tracking system (WaterMaze; Coulbourn Instruments). The pool was filled with opaque water (by adding washable white paint) and surrounded by extramaze cues. The escape platform (10 cm in diameter) was placed in the center of a designated quadrant with its top positioned 1 cm below the water surface. During the visible platform training, the platform was marked by a flag, and mice were trained by four trials per day for 2 d. In the hidden-platform test, mice were first trained daily by two trials with a 4-h interval for 9 d. A probe test was given on day 10. At the beginning of each trial, mice were pseudorandomly dropped at one of the six start points. Mice were allowed to swim freely until they climbed onto the platform and stayed on the platform for 20 sec. If mice were unable to find the platform within 90 sec, they were guided to the platform and allowed to stay on the platform for 20 sec. During the probe test, the hidden platform was removed from the pool, and mice were allowed to swim for 60 sec. The time spent in each quadrant, as well as the number of platform crossings, was recorded. For the reversal task, the hidden platform was moved to the opposite quadrant. Mice were trained daily by two trials with a 4-h interval for 3 d. A probe test was performed on day 4.

Temporal dissociative passive avoidance (TDPA)

Training and testing for the TDPA paradigm was performed as described (Zhang et al. 2008). During training, the mouse was introduced to the lit half of the training chamber (Coulbourn Instruments) and allowed to explore for 1 min before the trap door was opened. After the animal entered the dark chamber, the trap door was closed and a mild footshock (0.7 mA for 2 sec) was delivered 10 min later. The animal was returned to its home cage 20 sec after the footshock. When tested, the trained mouse was reintroduced to the lit chamber. The latency before entering the dark chamber was scored as crossover latency and used as the index for memory formation. We chose 300 sec as the cutoff value for crossover latency. For the experiments in Figure 2A, animals were trained on day 1 and tested daily on days 2–6. On days 2–5, mice were first scored for crossover latency. They were then shocked 10 min after they entered the dark chamber. Therefore, the animals were also trained following the scoring. For the experiments in Figure 2D, the animals were trained once, tested 10 min after the training, and crossover latency was recorded.
Dark/light test

The same PA chamber and lighting conditions were used for the dark/light test as described previously (Krishnan et al. 2008). A mouse was introduced to the dark chamber and habituated for 2 min before the trap door was opened. The animals were allowed to move between the dark and light chamber for 10 min. The latency of animals’ initial crossover to the light chamber was recorded. The total time spent in the light and dark chamber was also scored.

Contextual fear conditioning (CFC)

During training, mice were placed in the operant chamber with a metal grid floor for 30 sec (TruScan chamber with photobeam sensors). A loudspeaker, after which a mild electric footshock was delivered (0.7 mA with 2-sec duration). Mice stayed in the training chamber for another 30 sec, during which “immediate freezing” was scored. Twenty-four hours after training, mice were tested and scored for freezing. The freezing behavior, defined by immobility except for breathing, was scored once every 5 sec for 3 min. The movement velocity before and during the mild electric footshock was monitored by the TruScan system. To measure the hippocampus-independent CFC memory (Wiltgen et al. 2006), a mouse was placed into the training chamber for 2 min before receiving three mild electric footshocks (1.5 mA, 2-sec duration) delivered once every 2 min. The animals remained in the training chamber for 30 sec after the last shock and were returned to their home cages. The animals were tested 24 hr after training, and the percentage of freezing was scored.

In vivo extracellular recording

Naïve male mice (2–4 mo of age) were anesthetized with sodium pentobarbital (100 mg/kg, i.p.), and placed in a stereotaxic frame (David Kopf). Recording and stimulating electrodes were made of a pair of Teflon-coated Pt/IR wires (50 μm inner diameter, 100 μm outer diameter, World Precision Instruments, Inc; cat. no. PT964-5). The coordinates for the recording electrode in the CA1 region were AP = −1.7–1.9 mm, ML = 1.2 mm, and DV = 1.5–1.9 mm from skull surface. The stimulating electrode was implanted into the ipsilateral Schaffer collateral fibers (AP = −1.7–1.9 mm, ML = 1.7 mm, and DV = 1.6–2.0 mm from skull surface). Constant current pulses (0.10 msec duration) were delivered every 3 sec. After a stable baseline (50% of maximal fEPSP) was established, E-LTP was induced by a single train of high frequency stimulation (HFS, at 100 Hz for 1 sec). The potentiation was calculated by comparing the slope of fEPSP with that of the baseline values. After electrophysiological recordings, electrotic currents (4 mA, 2 sec) were delivered to produce a thermolytic lesion at the electrode tips, which was used for histological confirmation of the electrode placement.

Hippocampal infusion of FE65 siRNA

Stealth siRNA duplex was purchased from Invitrogen. The sequences for the double-strand siRNA used for mouse FE65 are: CCA GAUGUCUAGAGCCAGCGUUGUU (upper strand) and AAACA CUUGCAUGUGCCAGUUGG (lower strand). For the control duplex siRNA, sequences with the same GC content are scrambled. CGUGGCACUUGAGCAUCUGG (lower strand). For the control GAUGCUCAAGUGCCACGUGUUU (upper strand) and AAACA

transfection, the effects of siRNA on FE65 expression were analyzed by Western blot analysis. A polyclonal rabbit antibody (FE518) was used to detect both p97 and p60 isoforms of FE65 (Wang et al. 2004a).

Data analysis

Two-way repeated-measures ANOVA was used for analyzing MWM, CFC (followed by Bonferroni post-test), and TDPA data (genotype and time/trial as between-/within-subject factor). The Student’s t-test was used to assess significance for data between two groups. Data were expressed as the average ± SEM. Differences with p-values < 0.05 were considered statistically significant.

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References


FE65 and hippocampus-dependent plasticity


