CPG15 regulates synapse stability in the developing and adult brain

Citation

As Published
http://dx.doi.org/10.1101/gad.176172.111

Publisher
Cold Spring Harbor Laboratory Press

Version
Final published version

Citable link
http://hdl.handle.net/1721.1/91540

Terms of Use
Article is available under a Creative Commons license; see publisher's site for details.

Detailed Terms
http://creativecommons.org/
CPG15 regulates synapse stability in the developing and adult brain

Tadahiro Fujino,1 Jennifer H. Leslie,1,2 Ronen Eavri,1 Jerry L. Chen,1,2 Walter C. Lin,1,3 Genevieve H. Flanders,1 Erzsebet Borok,4,5 Tamas L. Horvath,4,5 and Elly Nedivi1,2,3,6

1The Picower Institute for Learning and Memory, 2Department of Biology, 3Department of Brain and Cognitive Sciences, Massachusetts Institute of Technology, Cambridge, Massachusetts 02139, USA; 4Department of Obstetrics, Gynecology, and Reproductive Sciences, 5Department of Neurobiology, Division of Comparative Medicine, Yale University School of Medicine, New Haven, Connecticut 06520, USA

Use-dependent selection of optimal connections is a key feature of neural circuit development and, in the mature brain, underlies functional adaptation, such as is required for learning and memory. Activity patterns guide circuit refinement through selective stabilization or elimination of specific neuronal branches and synapses. The molecular signals that mediate activity-dependent synapse and arbor stabilization and maintenance remain elusive. We report that knockout of the activity-regulated gene cpg15 in mice delays developmental maturation of axonal and dendritic arbors visualized by anterograde tracing and diolistic labeling, respectively. Electrophysiology shows that synaptic maturation is also delayed, and electron microscopy confirms that many dendritic spines initially lack functional synaptic contacts. While circuits eventually develop, in vivo imaging reveals that spine maintenance is compromised in the adult, leading to a gradual attrition in spine numbers. Loss of cpg15 also results in poor learning. cpg15 knockout mice require more trails to learn, but once they learn, memories are retained. Our findings suggest that CPG15 acts to stabilize active synapses on dendritic spines, resulting in selective spine and arbor stabilization and synaptic maturation, and that synapse stabilization mediated by CPG15 is critical for efficient learning.

[Keywords: mouse; brain; synapse; dendrite; two-photon microscopy; learning]

Supplemental material is available for this article.

Received August 4, 2011; revised version accepted November 1, 2011.

During development, neuronal processes extend and retract as they explore their environment to identify appropriate synaptic partners. The establishment of pre- and post-synaptic contact is an early event in an ordered progression that can lead to formation of stable mature synapses. It has been proposed that synapse formation consequently acts as a stabilizing force on growing axonal and dendritic processes [Meyer and Smith 2006; Ruthazer et al. 2006]. While studies have delineated some aspects of synaptogenesis and synaptic maturation [McAllister 2007], the signals at the contact point of axon and dendrite that determine whether a synapse will stabilize and persist have not been fully elucidated.

Although excitatory synaptogenesis can occur in the absence of neural activity [Gomperts et al. 2000; Verhage et al. 2000], strong evidence suggests that experience plays a critical role in biasing the formation and stabilization of synapses that transmit appropriately patterned activity [Hua and Smith 2004] and that NMDA-type glutamate receptors mediate this activity-dependent synapse selection [Gomperts et al. 2000]. Activation of NMDA receptors allows Ca\(^{2+}\) influx into the post-synaptic cell, turning on kinase signaling cascades, which in turn initiate transcription factor activation and new gene expression [for reviews, see Flavell and Greenberg 2008; Loebrich and Nedivi 2009]. Yet, how this set of events leads to local implementation of synaptic stabilization is not clear. In fact, little is known about the molecular mechanisms regulating selective synapse and dendrite stabilization in response to activity.

cpg15 (also termed neuritin) was first identified in a screen for activity-regulated genes in rats [Nedivi et al. 1993; Hevroni et al. 1998] and is a downstream target of the classic synaptic plasticity signaling cascade, involving the NMDA receptor, MAPK, CaMK, and CREB [Fujino et al. 2003]. During development, cpg15 expression is induced in presynaptic neurons upon contact with their target [Diaz et al. 2002] and is spatially and temporally correlated with synapse formation and activity-dependent plasticity [Nedivi et al. 1996; Corriveau et al. 1999; Lee and Nedivi 2002]. cpg15 encodes a small extracellular protein anchored to the cell surface via a glycosyl-phosphoinositide
Cpg15 overexpression in the developing Xenopus enhances dendritic and axonal elaboration in a non-cell-autonomous manner, as well as synapse formation and maturation (Nedivi et al. 1998; Cantallops et al. 2000; Javaherian and Cline 2005).

To investigate the requirement for CPG15 in a mammalian system, we generated a knockout mouse for cpg15 (cpg15 KO). We found that in the cpg15 KO, there is a developmental delay in axonal and dendritic arborization and maturation of excitatory synapses. In the dentate gyrus [DG] of the hippocampus, as many as 30% of spines initially lack synapses. Chronic in vivo imaging of cortical pyramidal neurons through a cranial window shows that while dendritic spine dynamics in cpg15 KO mice are comparable with controls, fewer events in these mice are stabilized and thus favor persistent spine loss. These results suggest that the in vivo developmental deficits in the cpg15 KO mouse derive from lack of a synaptic stabilization signal, perhaps supplied in an activity-dependent manner. While adult circuits appear normal, they are functionally suboptimal, leading to poor performance in learning tasks. These findings establish a role for CPG15 in efficient circuit formation and function and provide a potential molecular mechanism for selective synapse stabilization.

Results

Generation of a cpg15 KO mouse

To investigate the in vivo role of cpg15, we generated a mouse lacking cpg15 using a conditional knockout approach based on the Cre-loxP system. We first made a “floxed” cpg15 mouse with loxP sites flanking cpg15 exons two and three (Fig. 1A). Exons two and three contain the entire coding sequence for the mature form of CPG15, and excision of the sequence between the loxP sites by Cre recombinase would generate a null mutation. Floxed cpg15 mice were crossed to a global Cre deleter line that expresses Cre recombinase in all tissues, including the cpg15 Cre recombinase would generate a null mutation. Floxed mice were crossed to a global Cre deleter line to generate the floxed-neo allele without neo, and the null allele. Indicated are three cpg15 exons (closed boxes), the neo and the diphertheria toxin A gene (DT-A) serving as positive and negative selection markers, respectively [open boxes], loxP sites (closed triangles), FRT sites [open triangles], and XhoI [X] and EcoRV [RV] restriction sites. Homologous recombination between the wild-type allele and the targeting vector generated a floxed neo allele and eliminated the DT-A gene. The neo gene flanked by two FRT sites was deleted by injection of flippase RNA into eggs harvested from the floxed neo mice. Mice after flippase recombination were crossed with a Cre deleter line to generate the cpg15-null allele. Positions of the 3’ probe used for Southern blot analysis and expected band sizes are indicated. (B) Southern blot analysis of wild-type (WT) cpg15 allele, the targeting vector, the floxed allele with the neomycin-resistant gene (neo), the floxed allele without neo, and the null allele. Indicated are three cpg15 exons (closed boxes), the neo and the diphertheria toxin A gene (DT-A) serving as positive and negative selection markers, respectively. (C) Nissl-stained coronal sections from wild-type and cpg15 KO mice. Bar, 1 mm.

Delayed axonal and dendritic arbor development in cpg15 KO mice

In the developing Xenopus, CPG15 overexpression promotes arborization of retinal ganglion cell axonal projections in the tectum by reducing branch retractions (Cantallops et al. 2000). To test for deficits in axon arbor...
development in cpg15 KO mice, we bulk-labeled retinal ganglion cell projections as these axons were growing and elaborating their arbors in the lateral geniculate nucleus (LGN) of the thalamus by injecting each eye with an anterograde tracer conjugated to a different fluorophore. In wild-type mice, the total area of LGN covered by projections from both eyes increased dramatically between postnatal day 4 (P4) and P21 (Fig. 2A,B). Since neuronal proliferation in the LGN occurs during embryogenesis (Brückner et al. 1976), the postnatal increase in LGN volume is largely due to axon ingrowth and elaboration of both the axonal and dendritic neuropil. In the cpg15 KO mice, the retinogeniculate projection area is similar in size to that of wild-type mice at P4, but the projection area does not grow to the same extent as controls by P9 (Fig. 2A,B). LGN cell count and density in the cpg15 KO mice at P9 were comparable with controls (Supplemental Table S2), suggesting that the difference in projection areas at this age is not due to a decrease in LGN cell numbers, but rather delayed development of the neuropil, including retinal ganglion cell axons and LGN cell dendrites. By P90, the retinogeniculate projection area was indistinguishable between genotypes, with similar levels of segregation of ipsilateral and contralateral arbors, as measured by the degree of overlap (Fig. 2C). While the delay in retinogeniculate projection development in the LGN of cpg15 KO mice indicates that there may be deficits in axonal and perhaps dendritic elaboration, the normal appearance of the LGN at P90 suggests that these deficits are overcome with age.

CPG15 overexpression studies showing enhanced dendritic arborization [Nedivi et al. 1998] and the potential delay in LGN neuropil development seen in cpg15 KO mice led us to examine whether dendritic arbor development was also delayed in cpg15 KO neurons. Using diolistic labeling [Grutzendler et al. 2003], we visualized neurons in the granule cell layer of the DG of the hippocampus [Fig. 3A], a region that normally shows high cpg15 expression throughout development and has been extensively characterized in the context of synaptic plasticity and paradigms of learning and memory.

Granule cell dendritic arbors in the hippocampal DG predominantly develop in two stages [Rahimi and Claiborne 2007]. The bulk of neurogenesis occurs embryonically through P14, peaking during the second postnatal week, with the majority of cells born postnatally. During this period, granule cell dendritic trees are initially established. The second phase begins at the end of the second postnatal week and lasts until ~2 mo of age. It is during this time that the dendritic trees are sculpted and refined. At the onset of this second phase, granule cell dendritic trees have many segments, including short terminal branches, which are largely pruned by 2 mo. The remaining branches increase in length over this period, resulting in a conservation of total dendrite length but a reduction in segments and terminal ends. To assess the contribution of CPG15 at these two phases, we compared granule cell dendritic morphology in the cpg15 KO and wild-type littermates at P15 and P60. At P15, DG granule cells in the cpg15 KO showed significantly fewer branch tips per neuron as compared with wild type, suggesting an initial lack of complexity due to reduced numbers of short terminal branches. Between P15 and P60, we observed branch tip pruning that was more...
pronounced in the wild type as compared with knockout animals, such that the difference between them was diminished by 2 mo of age and dendritic arbors appeared normal [Fig. 3B]. This is similar to our findings regarding neuropil development in the LGN.

**Delayed synaptic development and maturation in cpg15 KO mice**

Next, we tested for deficits in the cpg15 KO that might derive from the effects of CPG15 on synapse formation and maturation. We examined formation of functional synapses by performing whole-cell patch clamp recordings in the granule cell layer of the DG in acute hippocampal slices and recorded spontaneous miniature post-synaptic currents (mEPSCs) [Fig. 3C]. We found that in the DG of cpg15 KO mice at P7, mEPSC amplitudes were reduced; however, this deficit was no longer observed by 2 mo of age [Fig. 3D]. These results suggest that during development, cpg15 KO mice have fewer and less mature functional synapses.

To examine the synaptic structural correlates of reduced mEPSC frequency and amplitude in the developing hippocampus, we performed electron microscopy (EM) in different subfields of the hippocampal formation. No apparent qualitative differences in synaptic structure were seen in any of these areas, as assessed by the presence of presynaptic terminal zones with vesicles and apposed post-synaptic densities [PSDs] [Fig. 4A]. However, when the density of asymmetric synapses on dendritic spines was measured by unbiased stereology, we found that cpg15 KO mice had spine synapse densities in the DG that were 26% lower than wild-type controls at 2 mo of age [Fig. 4B]. This change appeared specific to the DG and was not observed in CA1 at this age. The decrease in DG spine density correlates with our findings of decreased mEPSC frequency in the cpg15 KO DG [Fig. 3C] but not in

---

**Figure 3.** Delayed dendritic arbor and synapse development in DG of cpg15 KO mice. (A) Representative traces of reconstructed DG granule cells from the wild type (WT) (left) and cpg15 KO (right) at 2 mo of age. [B] Average number of dendritic branch tips per cell at P15 (left) \( n = 39 \) wild-type cells; \( n = 32 \) knockout cells and 2 mo (right) \( n = 32 \) cells for wild-type and knockout. \( * \) P-value < 0.05. (C) Cumulative probability plots of intervals of mEPSCs of DG granule cells at ages P7 (left) \( n = 9 \) wild-type cells; \( n = 7 \) knockout cells; P-value < 0.001 by K-S test) and 2 mo (right) \( n = 13 \) wild-type cells; \( n = 7 \) knockout cells, P-value = 0.003 by K-S test). (D) Average mEPSC amplitudes of DG granule cells at ages P7 (left) \( n = 9 \) wild-type cells; \( n = 7 \) knockout cells) and 2 mo (right) \( n = 13 \) wild-type cells, \( n = 7 \) knockout cells. [Right inset] Averaged mEPSCs are plotted for both ages. \( * \) P-value = 0.01.

**Figure 4.** Delayed spine synapse development and lack of age-dependent synaptic pruning in cpg15 KO mice. (A) Representative EM images from the molecular layer of the DG of wild-type (WT) and cpg15 KO mice. Arrowheads indicate spine synapses, and asterisks mark spine or spine-like protrusions. Spine synapse density [B], spine or spine-like protrusion density [C], and PSD length [D] of indicated regions \( n = 4 \) mice for wild-type and knockout. \( * * \) P-value < 0.05. (E) Age-dependent change in spine density of DG region. Spine density per mouse is represented by circles, and the average for each genotype is indicated by a horizontal line \( n = 4 \) mice each for wild type and knockout at 2 mo; \( n = 4 \) mice for wild-type and \( n = 5 \) mice for knockout at 9 mo]. P-value < 0.001 for 2 mo versus 9 mo for wild-type and knockout DG. [F] Age-dependent change in spine synapse density in the DG region. Spine synapse density per mouse is represented by circles, and the average for each genotype is indicated by a horizontal line \( n = 4 \) mice for 2 mo and \( n = 5 \) mice for 9 mo for wild type and knockout]. P-value < 0.001 for 2 mo versus 9 mo for wild type, n.s. for knockout. Bar, 1 mm.
CA1 (Supplemental Fig. S2) at 2 mo of age and may be due to late development of the DG as compared with other hippocampal regions. Interestingly, when we measured the density of spines and spine-like protrusions at 2 mo regardless of whether they contained a synaptic specialization, there were no significant differences between cpg15 KO mice and wild-type controls even in the DG [Fig. 4C]. These results suggest that in the developing DG of the cpg15 KO, a large fraction of spine-like protrusions lack a synaptic structure. EM examination of synapse size at 2 mo, as measured by PSD length, showed that PSDs of cpg15 KO mice were 33% larger than those of wild-type mice in the DG but not in CA1 [Fig. 4D]. Perhaps the increased synapse size is a compensatory response to the decrease in spine synapse numbers in the DG of cpg15 KO mice.

In the retinogeniculate pathway as well as the hippocampus, neuronal development in the cpg15 KO is initially delayed but eventually reaches wild-type levels. We examined whether the deficit in dendritic spine synapse number also recovers with age by comparing spine and spine synapse densities between 2- and 9-mo-old cpg15 KO mice and wild-type littermates. Between 2 and 9 mo, both cpg15 KO mice and wild-type controls prune spines in a similar manner [Fig. 4E]. Wild-type controls also prune spine synapses over this period, consistent with ongoing synapse remodeling and refinement [Markus and Petit 1987]. However, cpg15 KO synapse density, which starts out lower than wild type at 2 mo, is not further reduced by 9 mo of age and remains relatively constant during this period [Fig. 4E]. Thus, we see that neurons in the late developing DG of cpg15 KO mice initially form fewer spine synapses than wild-type neurons. Over time, these synapses are less likely to be pruned, so that by 9 mo, synapse numbers are similar in wild type and knockout.

Increased loss of persistent spines in cpg15 KO mice

In vivo imaging studies followed by EM have demonstrated that spine sprouting and retraction are associated with synapse formation and elimination (Trachtenberg et al. 2002; Knott et al. 2006) and that synaptogenesis is inversely correlated with spine motility (Konur and Yuste 2004). In light of the unusually large fraction of dendritic spines lacking synapses in the developing DG of cpg15 KO mice, we asked whether CPG15 depletion also affects dendritic spine dynamics. To this purpose, we performed in vivo imaging of neurons in the visual cortices of adult cpg15 KO mice. The cortex was chosen to assay spine dynamics because it is optically accessible via implantation of cranial windows, allowing chronic monitoring of spine dynamics. In addition, previous studies have shown that even in the adult cortex, a significant fraction of dendritic spines remain dynamic, with the majority of events being either transient spine additions or reversible eliminations [Holtmaat et al. 2005, 2006]. Persistent-dynamic events, including spines that emerge and persist (new-persistent spines) as well as spines that disappear and do not re-emerge (lost-persistent spines), likely best represent events that correspond to synapse formation and elimination [Holtmaat et al. 2005, 2006].

We generated cpg15 KO mice expressing GFP in a random subset of neurons sparsely distributed within the neocortical layers by crossing the thy1-GFP-S line that expresses GFP in a random subset of neocortical neurons [Feng et al. 2000; Lee et al. 2006] to the cpg15 KO [see the Materials and Methods]. Adult thy1-GFP/cpg15 KO mice [homozygous for thy1-GFP and cpg15<sup>−/−</sup>] and thy1-GFP littermate controls were surgically implanted with bilateral cranial windows over the visual cortices. Following 3 wk of recovery, layer 5 [L5] pyramidal neurons were identified, and a two-photon imaging volume encompassing their apical tufts in L1 was acquired at 4-d intervals [Fig. 5A]. We found that the total rate of dynamic events, including both spine gain and loss, was not significantly different between wild-type and cpg15 KO mice [Fig. 5B]. However, in the cpg15 KO, the percentage of dynamic events that persisted was significantly higher than in the wild type. Conversely, the percentage of transient events was significantly lower [persistent-dynamic: 69.23% for knockout, 55.06% for wild type; transient: 30.76% for knockout, 44.94% for wild type; P-value < 0.05] [Fig. 5C]. For both the wild type and cpg15 KO, dynamic-persistent events that favor spine loss [lost-persistent] were higher than dynamic-persistent events that favor spine gain [new-persistent]. However, this difference was more pronounced for the cpg15 KO [lost-persistent: 70.79% for knockout, 60.64% for wild type; new-persistent: 29.21% for knockout, 39.36% for wild type; P-value < 0.001 for knockout, P-value < 0.05 for wild type] [Fig. 5D]. Since the cpg15 KO has a greater percentage of dynamic-persistent events overall, and persistent events favor loss, we observed a significant reduction in spine densities for individual cpg15 KO neurons over the course of the imaging period, which was not seen in wild-type neurons [Fig. 5E]. Overall, these results suggest that loss of cpg15 leads to a decrease in spine stabilization, resulting in reduced maintenance of both newly formed and existing spines.

Inefficient learning in cpg15 KO mice

Given the contribution of activity-regulated genes to plasticity [Leslie and Nedivi 2011], we next examined whether the deficits in cellular development and spine stabilization seen in cpg15 KO mice impacted behavioral plasticity in the adult, such as learning and memory. cpg15 KO mice were subjected to a fear conditioning test, a form of classical conditioning. Context-dependent fear conditioning is a hippocampal-dependent paradigm. We rationalized that due to cellular defects observed in the hippocampus, cpg15 KO mice may not perform as well as wild-type counterparts in this task. Mice were given a paired shock and tone in a conditioning chamber. Afterward, they were returned to the same chamber to test for contextual memory or were presented with the same tone in a different context to test for tone-dependent memory. Surprisingly, cpg15 KO mice exhibited less freezing than wild-type controls in response to both context and tone [Fig. 6A], suggesting deficits in both contextual and tone-dependent memory. Interestingly, cpg15 KO mice showed little freezing in response to context even 1 h after training,
suggested impaired short-term memory. In this test, it was impossible to discriminate whether there was a true deficit in long-term memory or whether it was secondary to the short-term memory deficit. To address this question, mice were trained using repeated conditioning sessions on days 1, 3, 5, and 7 and tested 1 d after each session on days 2, 4, 6, and 8. For tone-dependent memory, wild-type mice showed a robust freezing response after the first conditioning trial, with an additional small increase after the second trial (Fig. 6B). cpg15 KO mice showed little freezing after the first conditioning trial despite a slightly lower threshold for pain and higher anxiety than wild-type controls (Supplemental Fig. S1C,D), but showed a large increase after the second trial before reaching a plateau similar to wild-type controls by the third trial. To see whether stronger conditioning would improve learning, in the fourth conditioning session, three tone–shock pairs were given instead of one. This fourth conditioning session did not elicit increased freezing in either genotype, indicating that they were able to learn the task (Fig. 6D). However, cpg15 KO mice showed slower acquisition when compared with wild-type controls. The difference was most obvious on the second day of the training, when wild-type controls were close to their peak performance, but many of the cpg15 KO mice showed no improvement. Despite these deficits, improvement in performance and the directed swimming toward the platform after training suggests that cpg15 KO mice were able to recognize the visual cue. Thus, in the visible platform test, cpg15 KO mice showed slower acquisition of the task but improved with repeated training, similar to what we observed in the fear conditioning test.

Discussion

Here we show that a knockout mouse lacking cpg15 exhibits delayed axonal, dendritic, and synaptic development. Adult cpg15 KO mice display reduced spine maintenance, leading to gradual spine loss. Loss of cpg15 also has behavioral consequences, manifested as poor performance in learning tasks.

The finding that adult cpg15 KO mice have an apparently normal neocortex with regards to size and cell number was initially surprising, since acute RNAi-mediated knockdown in embryonic rat brains showed a requirement for CPG15 in cortical progenitor cell survival in a pool using only visual or spatial cues. In the visible platform version of the Morris water maze, a visual cue is placed on the submerged platform so that mice can locate the platform based on this cue. This task does not require spatial memory, but does require mice to make certain associations, such as those between the visual cue and the platform, in addition to basic visual and motor abilities to perform the task.

During the 7 d of training, swim distance to reach the platform decreased for both wild-type and cpg15 KO mice, indicating that they were able to learn the task (Fig. 6D). However, cpg15 KO mice showed slower acquisition when compared with wild-type controls. The difference was most obvious on the second day of the training, when wild-type controls were close to their peak performance, but many of the cpg15 KO mice showed no improvement. Despite these deficits, improvement in performance and the directed swimming toward the platform after training suggests that cpg15 KO mice were able to recognize the visual cue. Thus, in the visible platform test, cpg15 KO mice showed slower acquisition of the task but improved with repeated training, similar to what we observed in the fear conditioning test.
One-trial fear conditioning

Repeated-trial fear conditioning

Memory retention (2 w)

Visible platform test

Figure 6. cpg15 KO mice show slow learning in fear conditioning and visible platform tests. [A] Mice were given a paired tone and shock and then tested for contextual memory 1 h and 24 h later and for tone-dependent memory at 25 h after the shock. Fear memory is measured as percentage of time spent freezing (n = 23 for wild type; n = 27 for knockout). [*] P-value < 0.05; [**] P-value < 0.01. [B] Freezing response to tone after repeated training. Mice were given tone–shock pairings every other day and were tested 1 d after each training session (n = 10 for wild type and knockout). [**] P-value < 0.01. [C] Fear conditioning is stable for at least 2 wk. Mice were tested at 1 d and 2 wk after the repeated training shown in B. Percentage of retention was calculated as the ratio in freezing time at 2 wk to 1 d after training and was averaged for each genotype. Some mice showed longer freezing times at 2 wk, resulting in some retention scores >100% (n = 10 for wild type and knockout). [*] P-value < 0.05; [**] P-value < 0.001. (D) Swim distance with repeated training in the Morris water maze visible platform test (n = 12 for wild type and knockout). [*] P-value < 0.05; [**] P-value < 0.001.

A biphasic role for CPG15 in arbor and synapse development

During development, an increased rate of growth is often equated with maturation, but this interpretation may be too simplistic. There are many examples throughout the developing brain where maturation proceeds first through the initial establishment of exuberant and/or promiscuous dendritic and axonal branches and synapses, followed by a period of dynamic refinement and sculpting during which inappropriate contacts and branches are eliminated and appropriate ones are stabilized and elaborated (Kano and Hashimoto 2009). The cpg15 KO mouse exhibits a lack of initial exuberance in dendritic branch elaboration and synapse formation in areas of the developing brain investigated in this study. This results, in some cases, in the superficial appearance of “mature”-looking branches from an early age; however, these arbors have not matured in the sense that they have not undergone the same extensive remodeling as their wild-type counterparts. We would not interpret this as a precocious maturation, since a true precocious maturation would go through the same sequence of events as a normally developing animal, except earlier. Consistently, our data show that this is not the case for the cpg15 KO mouse. Instead, we show that the cpg15 KO never reaches the phase in development where processes and synapses become exuberant enough for large-scale sculpting and refinement to be observed.

This interpretation is also consistent with the developmental regulation of cpg15 expression. In the visual system early in development, cpg15 expression is activity-independent (Corriveau et al. 1999, Lee and Nedivi 2002). Promiscuous early expression of CPG15 may result in the establishment of large arbors and many synapses, effective substrates upon which activity-dependent mechanisms can later work to sculpt circuits by choosing which

Putz et al. 2005]. We were also surprised that despite the robust effect of CPG15 overexpression on dendritic and axonal arbor growth in the developing Xenopus retinotectal system [Nedivi et al. 1998; Cantallops et al. 2000], the effects of CPG15 deletion on both axonal and dendritic arborization are more subtle and are fully compensated for in the cpg15 KO mice by 2 mo of age. There are precedents for in vivo RNAi-mediated interventions and overexpression studies during development resulting in deficits not observed in knockout animals. For example, in the case of the doublecortin (DCX) gene, acute knockdown early in development results in severe cortical laminination deficits, but in the knockout mouse, cortical lamination is normal [Bai et al. 2003]. Even in the case of neurotrophins and their receptors, mutant mouse studies repeatedly show less overt phenotypes than might be expected for molecules considered critical for both neuronal survival and differentiation [Conover and Yancopoulos 1997]. Molecules such as neurtogins that exhibit robust synaptogenic activity in coculture assays [Scheiffele et al. 2000] when completely eliminated in knockout mice fail to affect synapse numbers in vivo [Varoqueaux et al. 2006]. Rather, neurtogin knockout mice show subtle synaptic deficits that are likely associated with selection of specific synapse types. When a gene product is missing from the outset, compensatory or redundant molecules and mechanisms may be brought into play before developmental programs are significantly compromised. This is likely to be particularly true for molecules involved in synapse formation and maturation due to the multiplicity of players involved [Brose 2009]. This does not mean that the function of these molecules is completely interchangeable, rather, combinations of these molecules likely instruct specificity and diversity of synaptogenesis. In the cpg15 KO, other molecules can apparently replace the trophic function of CPG15, resulting in normal cell number and structure. cpg15 KO neurons can form functional synapses with normal ultrastructure, indicating that CPG15 is also dispensable for synaptogenesis and synaptic growth. However, the delay in arbor growth and synapse development and the reduced stability of dendritic spines in the adult suggest that CPG15 plays a critical regulatory role in determining which synaptogenic events are stabilized and maintained.
synapses and branches to keep and which to prune. During these later periods, cpg15 expression becomes activity-dependent, perhaps allowing CPG15 to function in the stabilization of appropriate activity-selected synaptic partners and branches.

**CPG15 as a synapse stabilization factor**

Despite lower spine synapse density in the hippocampal DG of 2-mo-old cpg15 KO mice as compared with wild-type controls, spine counts in this region were normal. This suggests that cpg15 KO neurons have a larger percentage of spines without well-defined synaptic structures. Previous serial EM reconstruction studies have found little evidence of spines without synaptic specializations in adult animals [Harris et al. 1989; Harris and Kater 1994], and developmental studies in dissociated and organotypic slice cultures suggest that synaptic molecules can cluster at pre- and post-synaptic sites shortly after contact [Vardinon Friedman et al. 2000; Okabe et al. 2001]. Imaging studies combined with retrospective EM show that the latency to formation of synaptic structures on nascent spines varies from several hours up to 4 d [Knott et al. 2006; Nägerl et al. 2007; Zito et al. 2009]. Thus, spine formation can lead to rapid synapse recruitment and stabilization, but does not necessarily do so. The choice between synapse stabilization and elimination is likely guided by specific signaling molecules that ensure the selection of optimal synaptic partners and efficient circuit wiring. Reduction in spine synapse density in the cpg15 KO mice without a change in spine number may reflect an increase in spine-like protrusions that fail to stabilize nascent synapses. Indeed, when spine and spine synapse densities are analyzed at a later age, cpg15 KO mice prune spines, but not spine synapses, suggesting selective pruning of unstabilized, synapseless spines. Furthermore, in vivo time-lapse imaging of dendritic spines on cortical L5 neurons of wild-type and cpg15 KO mice shows that while overall spine dynamics on cpg15 KO neurons are normal, there is a bias toward increased spine loss, suggesting that spines are less well maintained. The fact that this can be observed even in the adult on an individual cell basis further suggests an acute requirement for CPG15 in spine and perhaps synapse stabilization.

During development, synapses have been shown to act as anchoring points for growing dendritic and axonal branches, facilitating branch extension and growth into pruning to branch retractions, so that arbor dynamics are strongly influenced by synapse stabilization [Meyer and Smith 2006; Ruthazer et al. 2006]. It is therefore likely that delayed axon and dendrite development in the cpg15 KO is directly linked to the delay in functional synapse formation and maturation. This interpretation is consistent with findings from previous overexpression studies in the developing Xenopus that showed that CPG15 coordinately promotes synaptic maturation and arbor growth [Cantallops et al. 2000, Javaherian and Cline 2005] and that increased arbor growth is due to fewer branch retractions [Cantallops et al. 2000].

In rodents, exposure to an enriched environment improves performance in learning and memory tasks and increases dendritic spine density [van Praag et al. 2000], suggesting that changes in connectivity may be a cellular correlate of improved performance. In vivo imaging studies show that dendritic spines retain dynamic qualities even in the adult brain [Grutzendler et al. 2002; Trachtenberg et al. 2002] and that activity appears to play an instructive role in this remodeling. Spines appear and disappear in hippocampal slices after electrical stimulation as well as in vivo in response to experience [for review, see Bhatt et al. 2009]. Thus, it seems that the dynamic nature of dendritic spines provides the capacity for local circuit restructuring in response to normal day-to-day levels of activity. In the same way that synapse formation on growing axons and dendrites provides an anchor for arbor stabilization during development, synapse formation on dynamic spines could serve to consolidate nascent connections in a mature circuit. This suggests that CPG15 could continue to play a role as a synaptic stabilizing factor, extending past developmental circuit wiring and into optimization of adult circuitry. cpg15 KO neurons may be unable to stabilize connections in the adult and thus may have to rely on an initial set of synapses stochastically formed during development.

Based on our analysis of the cpg15 KO mouse, we propose that CPG15 acts to stabilize nascent synapses on dendritic spines, resulting in spine and arbor stabilization and synaptic maturation. Neuroligins were recently proposed to act as a signal for the validation of excitatory versus inhibitory synapses. Rather than mediating synapse formation per se, neuroligins signal whether a transiently initiated connection stays or goes [Chubykin et al. 2007]. It is intriguing to consider that since CPG15 is expressed and can be externalized in response to activity [Cantallops and Cline 2008], it could provide a saliency signal for selective stabilization of active synapses. While synaptic connections can form without CPG15, they are slower to form and are not optimized by activity patterns that are required for efficient learning. cpg15 KO mice are capable of learning with increased trials, and their sensory and motor functions appear to be within the normal range, suggesting that many behaviors required for survival are hardwired through highly overlapping and redundant mechanisms. Only when peak performance is desired does the role of activity-dependent tuning mediated by molecules such as CPG15 become starkly evident.

The subtle deficit we observed in relation to synapse stabilization allowed us to probe the consequences of CPG15 functional deletion on circuit properties and behavior without the confounding aspects of complex phenotypes commonly found in general knockout mice. Our results lead us to speculate that the inefficient learning seen in cpg15 KO mice derives from a diminished capacity for selective synapse stabilization.

**Materials and methods**

**Generation of the cpg15 KO mice**

All animal work was approved by the Massachusetts Institute of Technology Committee on Animal Care and conforms to NIH
guidelines for the use and care of vertebrate animals. To allow conditional deletion of the cpg15 gene using the Cre-loxP system, we generated floxed cpg15 mice. The targeting vector was constructed starting with a 13.7-kb genomic fragment spanning a region of the cpg15 gene from the EcoRV site 0.2 kb upstream of exon 1 to the AflII site 4.7 kb downstream from exon 3, cloned from a C57BL/6 mouse BAC library (Genomic Systems). Exons 2 and 3 were floxed by replacing a 19-base-pair (bp) sequence between the SacI and XpnI sites 0.5 kb upstream of exon 2 with a 40-bp fragment containing loxP and Xhol sites and inserting a 2.9-kb LFNT-TK cassette (gift of Kazu Nakawaza), consisting of a neomycin resistance (neo) gene flanked by two loxP sites and two FRT sites, into the SacI site 0.9 kb downstream from exon 3. The diptheria toxin A gene from pMCI DT-A (Yagi et al. 1990) were used to amplify a 230-bp wild-type allele and neo gene from cpg15loxneo/+ mice, thus generating floxed cpg15 mice without the neo gene in pure C57BL/6 background (cpg15lox−/−). To generate general cpg15−/− null mice, cpg15lox+/+ mice were crossed to an adenovirus Ella promoter-driven Cre transgenic line in C57BL/6 background [Jackson Laboratory, B6.FVB-TgN(EIIa-cre)C5379Lmgd] (Lakso et al. 1996). Progeny of this cross that were heterozygous for the cpg15−/− null allele (cpg15lox−/− ) were intercrossed to obtain homozygous cpg15−/− null mice (cpg15−/−). The absence of floxed cpg15 genomic sequence, cpg15 mRNA, and CPG15 protein was confirmed by Southern blots of tail genomic DNA (Sanbrock et al. 1989), Northern blots on brain RNA [Fujino et al. 2003], and Western blots, respectively. For Western blots, membrane protein–enriched fractions were prepared from the cerebral cortex and hippocampus of adult mice using the Mem-PER kit (Pierce) and PAGEprep advance kit (Pierce). Forty micrograms of protein was resolved by 15% SDS-PAGE, transferred to a nitrocellulose membrane, incubated with rabbit anti-CPG15 (1:100) and then HRP-conjugated goat anti-rabbit IgG (1:50,000; Jackson ImmunoResearch), and visualized by chemiluminescence (Pierce). Antibodies against CPG15 were generated by immunizing rabbits with peptides corresponding to amino acids 28-40 and 99–116 and were affinity-purified with the immunizing peptides (Open Biosystems).

(cpg15lox−/−) × cpg15lox−/− crosses. Genotyping was done by tail DNA PCR. Wild-type forward primer (5′-CGCAGGCCCATTGCTTTTC-3′; 0.13 pmol/μL), cpg15-null forward primer (5′-GGTTGGAAGGCACAAAC-3′; 0.5 pmol/μL), and common reverse primer (5′-GGGCGAGGGCGATTCCCTT-3′; 0.5 pmol/μL) were used to amplify a 230-bp wild-type band and a 350-bp cpg15-null band. Mice were housed under a 12-h light/dark cycle with ad libitum access to food and water.

Data collection and analysis

All quantifications comparing wild-type and cpg15 KO mice were done blind to genotype. Statistical analysis was done using StatView software (SAS Institute). Unless otherwise stated, Student’s unpaired t-test was used for two-group comparisons, and analysis of variance (ANOVA) and Student-Newman-Keuls post hoc analysis were used for comparisons involving more than two groups. Error bars are standard error of the mean unless otherwise stated.

Brain measurements

Age-matched males were perfused with phosphate-buffered saline (PBS) and then with 4% paraformaldehyde in PBS. Brains were dissected out, weighed, and measured with vernier calipers. After overnight post-fixation at 4°C, brains were cryoprotected in 30% sucrose in PBS, frozen in powdered dry ice, and sectioned coronally at 40 μm with a cryostat (Leica). Every sixth section was stained with cresyl violet. For volume measurements, the area on each section was measured using the point-counting method, and the total volume was estimated based on Cavalieri’s rule (Williams et al. 2003). Cell density was measured by the three-dimensional counting method (Williams and Rakic 1988).

Labeling retinal ganglion projections in the LGN

Mice younger than P9 were anesthetized by hypothermia or with 2.5% Avertin (250 mg/kg intraperitoneally [i.p.]) at later ages. Animals received an intravitreal injection of wheat germ agglutinin (WGA) conjugated to Alexa Fluor 555 (1 mg/mL, Molecular Probes) in the left eye and WGA-Alexa Fluor 488 in the right eye. For injections prior to natural eye opening, fused eyelids were separated or cut to expose the temporal region of the eye. After 24 h, animals were perfused with 4% paraformaldehyde in PBS. Brains were removed and post-fixed overnight and then coronally sectioned at 75 μm using a vibratome.

LGN images were acquired with an epifluorescence microscope [Nikon] using a 10×/N.A. 0.3 objective lens (Nikon). Four successive sections, representing the middle third of the LGN, were selected. Background fluorescence was subtracted, and grayscale images were normalized (0–255) using ImageJ [http://rsb.info.nih.gov/ij]. Grayscale images were converted into binary high-contrast black-and-white images by employing a threshold procedure that distinguishes signal from residual background fluorescence [Muir-Robinson et al. 2002; Torborg and Feller 2004] Contralateral [pseudo-colored green] and ipsilateral signals [pseudo-colored magenta] were superimposed (Fig. 2). An outline of the entire LGN was drawn, and the area was measured. The extent of overlapping projections was determined by counting pixels that contained both green and magenta signal, represented as white.

Diolistic labeling in the hippocampus

Brain sections from P7, P15, and 2-mo-old mice were processed for diolistic labeling as described (Grutzendler et al. 2003) with the following modifications. After rapidly perfusing P15 and 2-mo-old mice with 4% paraformaldehyde in PBS (40 mL in 2 min), brains were dissected out and post-fixed for 10 min at room temperature. P7 brains were removed without perfusion and then fixed in 4% PFA for 10 min at room temperature. All brains were coronally sectioned at 100 μm with a vibratome and stored in 30% sucrose in PBS. Bullets were prepared by coating tungsten particles (1.7 μm, Bio-Rad) with Dil [Molecular Probes], then loaded into Tefzel tubing (Bio-Rad) pretreated with polycrystalline-pyrolidone [1 mg/mL in isopropanol, Sigma], and dried with N2. Brain sections were covered with a tissue culture insert with a 3-μm pore size (Greiner) and then shot with Dil-coated particles using a Helios gene gun system (Bio-Rad) set at 160 psi. Sections were post-fixed in 4% paraformaldehyde/30% sucrose in PBS overnight and mounted on glass slides with Fluoromount-G (Southern Biotech). Neurons were imaged with a Nikon PCM 2000 confocal microscopy system controlled by Simple PCI software [Compix, Inc., Image System] or an Olympus Fluoview.
Electrophysiology

Hippocampi of P7 or 2-mo-old mice were isolated, and 300-μm slices were prepared with a vibratome (World Precision Instruments) in ice-cold solution containing 238 mM sucrose, 26 mM NaHCO₃, 10 mM glucose, 2.5 mM KCl, 1 mM NaH₂PO₄, 3 mM MgSO₄, and 1 mM CaCl₂ constantly bubbled with 95% O₂/5% CO₂. The slices were transferred to a holding chamber filled with artificial cerebrospinal fluid (ACSF) containing 119 mM NaCl, 26 mM NaHCO₃, 10 mM glucose, 2.5 mM KCl, 1 mM NaH₂PO₄, 1.3 mM MgSO₄, and 2.5 mM CaCl₂ constantly bubbled with 95% O₂/5% CO₂ and were recovered for half an hour at 32°C and then for half an hour at room temperature. Slices were placed in a recording chamber continuously perfused with 32°C ACSF bubbled with 95% O₂/5% CO₂ plus 100 μM picrotoxin and 1 μM tetrodotoxin to isolate mEPSCs. Whole-cell recordings were performed with patch pipettes (5–7 MΩ) containing 130 mM K-glucuronate, 4 mM KCl, 2 mM NaCl, 10 mM HEPES, 0.2 mM EGTA, 4 mM ATP-Mg, 0.3 mM GTP-Tris, 7 mM phosphocreatine-Tris, and 10 mM sucrose (pH 7.25, 290 mosm). Neurons in CA1 or dentate were patched under visual guidance with a Nikon microscope equipped with IR/DIC optics using a black-and-white CCD camera (CCD-300F, Dage-MTI). Current traces were collected using an AxoPatch 2B amplifier (Axon), digitized at 5 kHz by a Digidata 1322A (Axon), and analyzed offline using Clampfit software (Axon). At least 150 mEPSCs, thresholded at 6 pA, were recorded at ~70 mV from each cell. Twenty micromolar 6,7-Dinitroquinoxaline-2,3-dione was applied during some recordings to verify AMPAR mEPSCs.

EM

Two-month-old and 9-mo-old male littermates were sacrificed, and brains were fixed as described [Diano et al. 2006]. Ultrathin sections were prepared as described [Diano et al. 2006] from stratum radiatum of CA1 and CA3 and the molecular layer of the DG. Spine and spine synapse density was calculated by unbiased stereological methods based on the dissector technique [Diano et al. 2006]. PSD length was measured from 100–150 spine synapses in 10 nonoverlapping sections and averaged for each animal.

Two-photon imaging

Cranial window implantation and multiphoton imaging were done essentially as described [Lee et al. 2008]. Mice (6–8 wk of age) were implanted with cranial windows and imaged 2–3 wk after surgery at 4-d intervals. For imaging, mice were anesthetized with 1.25% avertin (250 mg per kilogram of body weight, i.p.). Anesthesia was monitored by breathing rate and foot-pinch reflex, and additional doses of anesthetic were administered during the imaging session as needed. Two-photon imaging was performed using a custom-built microscope modified for in vivo imaging by inclusion of a custom-made stereotactic restraint affixed to a stage insert and custom acquisition software. The light source for two-photon excitation was a commercial Mai Tai HP titanium:sapphire laser (Spectra-Physics) pumped by a 14-W solid-state laser delivering 100-fsec pulses at a rate of 80 MHz, with the power delivered to the objective ranging from ~37 to 50 mW, depending on imaging depth. Z resolution was obtained with a piezo actuator positioning system (Piezosystem Jena) mounted to the objective. The excitation wavelength was set to 950 nm, with the excitation signal passing through a 20×, 1.0 numerical aperture water immersion objective (Plan-Apochromat, Zeiss) and collected after a barrier filter by a photomultiplier tube.

Given the sparse density of GFP expression in the thy1-GFP-S line, typically, a dendritic segment from one L5 pyramidal neuron (and a maximum of two) was imaged per animal. Cells mapped to the visual cortex were then identified as L5 pyramidal neurons based on their depth from the pial surface and morphology. Blood vessel pattern maps were used to locate the cells and dendritic branches throughout the imaging sessions. Low-resolution image stacks (512 × 512 pixels, 1 μm/pixel X–Y resolution, 5-μm Z-step size) were used to identify cell type and depth. High-resolution image stacks (768 × 768 pixels, 0.2 μm/pixel X–Y resolution, 0.7-μm Z-step size) were used to capture spine dynamics on L5 dendrites in L1.

Dendritic spine analysis

Image stacks were aligned such that the presence or absence of an individual spine at a given location on the dendrite could be determined in each session. Spines were scored only if they protruded >0.4 μm from the dendritic shaft and were projecting into the XY plane. Long thin spines without bulbous heads were excluded from analysis, since they were always transient and were not significantly different in number or dynamics between genotypes. The rate of spine gain and spine loss was defined as the percentage of spines appearing (spine gain) or disappearing (spine loss) in a session compared with the total number of spines in the previous session. Dynamic spines were sorted into the following categories: Transient spines were those that appeared for no more than one or two imaging sessions then disappeared, or that disappeared and reappeared through the imaging period; new-persistent spines were defined as those that were not present in the first session and subsequently gained for at least the last two sessions; and lost-persistent spines were defined as those that were present in at least the first two sessions and then were lost for the remaining sessions. Analysis was performed using V3D software [Peng et al. 2010] by an observer blind to genotype.

Behavioral tests

Male littermates between 3 and 6 mo of age were used for behavioral experiments, unless otherwise noted.

Grip strength test Mice were allowed to hold on to a wire attached to a spring scale (Pesola) and were pulled horizontally by their tail. Maximal pulling force before releasing the wire was recorded. Five trials were done for each mouse, and the average was calculated for best three.

Rotarod test Mice were placed on an accelerating rotarod, and latency to falling from the rod was recorded. Each trial lasted for a maximum of 8 min, during which the rotation speed increased linearly from 2.5 rpm to 40 rpm. Three trials were done per day for two consecutive days.

Open field test Mice were placed in the center of an open field (40 × 40 cm) and allowed to move freely for 30 min. Their activity was monitored by a Digiscan system in 1-min bins and was analyzed by Versa Max software [AccuScan Instruments]. Data were averaged for 30 min.
Hot plate test  Mice were placed on a hot plate preheated to 55°C. Latency to paw lifting or sudden movement was recorded. All mice responded within 15 sec.

Fear conditioning  One-trial fear conditioning was done as described [Zeng et al. 2001] with modifications described in the Supplemental Material. Memory was tested after 1 h and 24 h of the conditioning trial for contextual memory and after 25 h for tone-dependent memory. For repeated training, conditioning sessions with one tone–shock pairing were given on days 1, 2, and 3 with three tone–shock pairings (each pair starting at 1, 2, and 3 min after placement in chamber) was given on day 7. Context and tone tests were done 24 h and 25 h, respectively, after each conditioning session. For one-trial fear conditioning, 4- to 9-mo-old male mice were used.

Morris water maze visible platform test  The visible platform test was done essentially as described [Zeng et al. 2001]. The pool was 150 cm in diameter, with water at room temperature (21°C–22°C). The platform was 10 cm in diameter. Each mouse was trained three trials per day, with intertrial intervals of ~30 sec before being picked up. The platform was 150 cm in diameter, with water at room temperature (21°C–22°C). The platform position was changed for each trial. In each trial, the mouse was allowed to swim until it found the platform or 22 sec had elapsed, at which point the platform was guided to the platform. The mouse was allowed to sit on the platform for 30 sec before being picked up.

Acknowledgments

We thank Matthew Wilson, Richard Morris, and many members of the Tonegawa, Bear, and Nedivi laboratories for helpful discussions, Kazu Nakazawa and Heathier Hinds from the Tonegawa laboratory for providing the LFNT cassette and C57BL/6 embryonic stem cell lines and suggestions on knockout mouse generation, Susan Su for help with BrdU labeling experiments, Marla Feller for advice on LGN analysis, Albert Hung for teaching diolistic labeling, Weifeng Xu, Gordon Smith, Arnold Heynen, and Kensiou Futai for advice on electrophysiology experiments, Thomas McGugh, Guiquan Chen, and Tsuoyoshi Miyakawa for suggestions on behavioral experiments; Michael Ragion, David Nguyen, and Matthew Wilson for sharing unpublished results on in vivo recordings; and members of the Nedivi laboratory for critical reading of the manuscript. This research was supported by funding to E.N. from the National Eye Institute [RO1 EY018949] and to T.L.H. from the National Institute of Health Director's Pioneer Award (DP1 OD006850) and the National Institute of Diabetes and Digestive and Kidney Diseases (DK080000).

References


CPEG15 is a synapse stabilizer