Evidence for opposing roles of Celsr3 and Vangl2 in glutamatergic synapse formation

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The signaling mechanisms that choreograph the assembly of the highly asymmetric pre- and postsynaptic structures are still poorly defined. Using synaptosome fractionation, immunostaining, and coimmunoprecipitation, we found that Celsr3 and Vangl2, core components of the planar cell polarity (PCP) pathway, are localized at developing glutamatergic synapses and interact with key synaptic proteins. Pyramidal neurons from the hippocampus of Celsr3 knockout mice exhibit loss of ~50% of glutamatergic synapses, but not inhibitory synapses, in culture. Wnts are known regulators of synapse formation, and our data reveal that Wnt5a inhibits glutamatergic synapses formed via Celsr3. To avoid affecting earlier developmental processes, such as axon guidance, we conditionally knocked out Celsr3 in the hippocampus 1 week after birth. The CA1 neurons that lost Celsr3 in vivo also showed a loss of ~50% of glutamatergic synapses without affecting the inhibitory synapses assessed by miniature excitatory postsynaptic current (mEPSC) and electron microscopy. These animals displayed deficits in hippocampus-dependent behaviors in adulthood, including spatial learning and memory and fear conditioning. In contrast to Celsr3 conditional knockouts, we found that the conditional knockout of Vangl2 in the hippocampus 1 week after birth led to a large increase in synaptic density, as evaluated by mEPSC frequency and spine density. PCP signaling is mediated by multiple core components with antagonizing functions. Our results document the opposing roles of Celsr3 and Vangl2 in glutamatergic synapse formation.

Celsr3 | Vangl2 | glutamatergic | synapse formation

Glutamatergic synapses, the predominant excitatory synapses in the brain, are asymmetric cell–cell junctions formed from distinct pre- and postsynaptic components involving highly organized complexes of hundreds of proteins across the 20-nm synaptic cleft (1, 2). The signaling pathway that directly assembles these asymmetric protein complexes has not been well understood. Understanding mechanisms of glutamatergic synapse formation will provide important insights into the function and plasticity as well as dysfunction of glutamatergic synapses, which underlie numerous nervous system disorders.

Many epithelial tissues show planar cell polarity, the global asymmetry of cellular and tissue morphology and/or structure along the tissue plane (3, 4). The conserved core planar cell polarity (PCP) components, Frizzled, Dishevelled, Diego, Prickle, Vang(l), and Flamingo (Fmi)/Celsr, form asymmetric complexes at the cadherin-mediated adherens junctions that connect neighboring epithelial cells (3, 4). Recent studies suggest that mutations of some components of the PCP signaling pathway, Celsr3/Fmi and Vangl2, affect GABAergic circuit development in zebrafish retina, GABAergic motoneuron synapse development in Caenorhabditis elegans, and hippocampal/cortical glutamatergic and GABAergic synapse formation (5–11). PCP components are critical regulators of neuronal migration and axon guidance, which take place before synapse formation and their mutations may secondarily affect synapse formation (8–10, 12–19). Therefore, evaluating the specific functions of PCP components in synapse formation requires

conditionally knocking out these components in defined synapses after the development of axons and dendrites.

In this study, we directly address the role of Celsr3 and Vangl2 in glutamatergic synapse formation by deleting Celsr3 and Vangl2 in hippocampal pyramidal neurons after the first postnatal week (postnatal day 7; P7). We found that at the peak of synapse formation (P14), Celsr3 and Vangl2 are specifically localized in developing glutamatergic synapses and colocalized with pre- and postsynaptic proteins. In the absence of Celsr3, hippocampal neurons showed ~50% reduction in the number of glutamatergic synapses in vitro and in vivo. Inhibitory synapses were not affected. Noncanonical Wnt signaling inhibits glutamatergic synapse formation, whereas canonical Wnt signaling promotes glutamatergic synapse formation in the hippocampus (20). We found here that Celsr3, a key component of this noncanonical Wnt signaling pathway, the PCP pathway, mediates responses to Wnt5a, which negatively regulates synapse formation. Additionally, the postnatal deletion of Celsr3 induced at P7 lead to deficits in hippocampus-dependent learning and memory formation. Some PCP components are known to exert opposing biochemical functions as well as exclude each other in subcellular localization. Indeed, we found that the postnatal deletion of Vangl2 initiated at P7 resulted in an increase in glutamatergic synapses, measured at P14. Because mutations of some PCP components, such as Prickle1 and Prickle2, have been implicated in autism and epilepsy, precisely pinpointing the function of PCP signaling components will lead to better understanding of the synaptopathy underlying many neurological and neuropsychiatric disorders (21–24).

Results

Celsr3 and Vangl2 Are Localized in Excitatory Synapses. To characterize the role of Celsr3 and Vangl2 in developing synapses, we

Significance

The signaling mechanisms mediating glutamatergic synapse assembly are fundamental to our understanding of neural circuit function, plasticity, and disorders, but have remained elusive. We provide direct evidence that two components of the conserved planar cell polarity signaling pathway, which assembles asymmetric cell–cell junctions, have opposing functions in glutamatergic synapse formation. Celsr3 promotes assembly whereas Vangl2 inhibits assembly, suggesting that this signaling mechanism is accessible for both positive and negative regulation and is also a candidate pathway for mediating synaptic plasticity.

focused on the glutamatergic synapses formed between the Schaffer collaterals and the hippocampal CA1 pyramidal neuron apical dendrites spanning the mouse stratum radiatum (Fig. 1A). This region is well characterized and a commonly used model for studying synapse formation. The onset of synaptogenesis occurs during the first postnatal week and continues for approximately 2 more weeks (Fig. 1B).

We first analyzed PCP components in the synaptic membrane fraction (SMF) and the postsynaptic density (PSD) using subcellular fractionation of synaptosome preparations from P14 wild-type mouse forebrains (Fig. 1C) (25, 26). As internal controls, synaptophysin and SV2, two presynaptic proteins, were examined and found enriched in the SMF, whereas PSD-95 and NMDA receptor subunit NR2B, two postsynaptic proteins, were enriched

in the PSD fraction. We found that Celsr3 and Celsr2 were present in both fractions. Dvl1 was enriched in the SMF, whereas Dvl2 and Vangl2 were enriched in the PSD fraction. The hyperphosphorylated form of Fzd3 (top band) was more abundant in the SMF than in the PSD, whereas the unphosphorylated form of Fzd3 (bottom band) was enriched in the PSD fraction. This is consistent with our previous findings that Dvl1 increases hyperphosphorylation of Fzd3 to prevent its endocytosis and that Vangl2 and Dvl2 antagonize this by reducing Frizzled3 phosphorylation and promoting its internalization (14, 16). These results suggest that PCP components are asymmetrically localized at developing glutamatergic synapses in similar fashion to epithelial cell–cell junctions being planar polarized.

Fig. 1. PCP components are localized in developing excitatory synapses. (A) Schematic diagram of the CA3 to CA1 Schaffer collateral projections of the hippocampus. (B) Developmental progression of CA1 pyramidal neuron dendrites and CA3 axons. (C, Left) Diagram of the subcellular fractionation procedure. (C, Right) Western blot analysis of the distribution of PCP proteins by subcellular fractionation. P2, crude synaptosome; S1, supernatant. Tuj1 controls for loading. (D and E) Celsr3 (D) and Vangl2 (E) colocalization with postsynaptic marker PSD-95 and presynaptic marker bassoon. Arrows mark colocalized puncta in each image. (Scale bars, 1 μm.) (F) Immunoprecipitation (IP) assays using a P2 fraction of P14 wild-type mouse brains show interaction of endogenous Celsr3 and Frizzled3 proteins with endogenous synaptic proteins. (G) PCP components are distributed in glutamatergic synapses analogous to their organization in asymmetric epithelial cell junctions. SLM, stratum lacunosum moleculare; SO, stratum oriens; SP, stratum pyramidale; SR, stratum radiatum.

We then performed immunofluorescence staining of Celsr3 and Vangl2 in P14 hippocampal neurons in vivo. Celsr3 and Vangl2 were costained with either postsynaptic marker (PSD-95) or presynaptic marker (bassoon) on cryosections of hippocampus and visualized with confocal microscopy. We found at the peak of synaptogenesis that Celsr3 and Vangl2 are specifically colocalized in the developing synapses, as we observed colocalized puncta (denoted by arrows) of the Celsr3 and Vangl2 (red) puncta together with the synaptic marker (green) (Fig. 1 *D* and *E*). Although at the resolution of confocal microscopy it is not possible to separate the pre- and postsynaptic compartments, we observed highly specific colocalization (yellow) of these proteins together with the synaptic markers inside the synapses rather than distributed diffusely throughout the membranes of axons and dendrites. These results suggest that at this stage of postnatal development, Celsr3 and Vangl2 are likely specifically dedicated to regulating synapse formation. To visualize the dendrites and spines, we used a mouse line endogenously expressing yellow fluorescent protein (YFP) in the cytosol of a subset of CA1 pyramidal neurons (Fig. S1).

To test whether endogenous PCP components interact with specific synaptic proteins, we performed coimmunoprecipitation and Western blot using protein extracts from P14 mouse brain (Fig. 1*F*). After a series of centrifugation steps (26), the crude synaptosomal membrane pellet was solubilized and used for the coimmunoprecipitation. We found that endogenous Celsr3 interacts with SV2, PSD-95, and Frizzled3 (16), whereas endogenous Frizzled3 interacts with synaptophysin and SV2 (Fig. 1*F*). Recent studies also showed that other PCP proteins, Vangl2 and Prickle2, are found in the postsynaptic compartment and may interact with PSD-95 (5, 6). These data indicate that PCP components interact with a number of key synaptic proteins, suggesting a direct role in glutamatergic synapse assembly/function. The localization of PCP components and their interaction with key synaptic proteins are analogous to their organization in asymmetric epithelial cell junctions in PCP signaling (Fig. 1*G*).

Celsr3 Is Required for Glutamatergic Synapse Formation in Hippocampal Culture. To test the function of Celsr3 in synapse formation, we examined hippocampal neurons in 14-DIV (days in vitro) cultures prepared from embryonic day (E)18.5 *Celsr3* knockout mice (KOs) (27). Compared with wild-type (WT), *Celsr3*$^{-/-}$ cultures contained 24 ± 4% fewer presynaptic puncta as revealed by vGlut1 immunostaining, 36 ± 6.7% fewer postsynaptic puncta as revealed by PSD-95 staining, and 38 ± 5% fewer colocalized puncta characteristic of glutamatergic synapses (Fig. 2 *A* and *B*; **P < 0.01 for pre- and postsynaptic puncta; *P < 0.05 for colocalized puncta; Mann–Whitney U-statistic test). Knockout of *Celsr3* did not affect the mean area of the glutamatergic synapses marked by colocalized vGlut1+/PSD-95+ puncta (Fig. 2 *A* and *C*). No deficits were seen in the number or area of inhibitory synapses revealed by immunostaining *Celsr3*$^{-/-}$ cultures, indicating that inhibitory synapses are not regulated by Celsr3 (Fig. 2 *D–F*).

To assess the effects on early stages of synaptogenesis, we cultured E18.5 hippocampal neurons from *Celsr3*$^{-/-}$ embryos for only 6 days and examined synapse formation. We found that after 6 days of culture, when synapses are just beginning to form, the reduction of excitatory synapse puncta number was already significant (Fig. 2 *G* and *H*), suggesting that *Celsr3* is required at the onset of synapse formation. Although the areas of colocalized presynaptic vGlut1+ puncta were not different from the 14-DIV cultures (Fig. 2*C*), the areas of colocalized presynaptic vGlut1+ puncta were smaller in the 6-DIV cultures, further suggesting that Celsr3 is required for synapse assembly but not for maintenance. The areas of colocalized postsynaptic PSD-95+ puncta were unaffected in *Celsr3*$^{-/-}$ 6-DIV cultures (Fig. 2 *G* and *I*).

The reduction in vGlut1+/PSD-95+ glutamatergic synapses as early as 6 DIV and persisting at least until 14 DIV provides evidence supporting the possible role of Celsr3 in glutamatergic synapse formation.

Wnt5a inhibits glutamatergic synapses formed by Celsr3. Previous studies have shown that Wnt signaling regulates synapse development in various neuronal cell types (28). Wnt5a is robustly expressed in the mouse hippocampus during postnatal development and reduces the number of presynaptic puncta in neurons via a noncanonical Wnt signaling pathway (20). Therefore, we sought to determine whether Celsr3 mediated Wnt5a signaling. Hippocampi isolated from E18 *Celsr3*$^{+/+}$ and *Celsr3*$^{-/-}$ mice were treated with Wnt5a after 12 DIV (100 ng/mL for 36 h). *Celsr3*$^{+/+}$ neurons showed a 56% reduction in the number of colocalized puncta (Fig. 2 *J* and *K*; **P < 0.01, two-way ANOVA) at 14 DIV. *Celsr3*$^{-/-}$ neurons not treated with Wnt5a showed a 52% reduction compared with untreated *Celsr3*$^{+/+}$ neurons in colocalized puncta (Fig. 2 *J* and *K*; **P < 0.01, two-way ANOVA). Wnt5a addition to *Celsr3*$^{-/-}$ neurons did not produce a significant difference compared with untreated *Celsr3*$^{-/-}$ neurons (Fig. 2 *J* and *K*), suggesting that Wnt5a inhibits synapse formation via a PCP component, Celsr3. To determine whether the reduction of synaptic puncta for vGlut1 and PSD-95 was due to changes in transcription, translation, or stability of these proteins, cell lysates for *Celsr3*$^{+/+}$ and *Celsr3*$^{-/-}$ cultured hippocampal neurons 14 DIV were collected to probe for vGlut1, PSD-95, vGAT, and gephyrin. There was no reduction in their total protein levels in *Celsr3*$^{-/-}$ (Fig. 2*L*).

Celsr3 Is Required for Excitatory Synapse Formation in Vivo. Because *Celsr3* is essential for axon guidance (12–14, 17, 18), it is critical to delete *Celsr3* after axons have reached their proper target area using *Celsr3* conditional knockout mice (cKOs) to test its role in synapse formation in vivo. We crossed a *Celsr3* cKO with an inducible Cre line (SLICK) so that we could delete *Celsr3* postnatally (12, 29). The SLICK-A line expresses constitutively active YFP and a tamoxifen-inducible form of Cre recombinase, $CreER^{T2}$, from two separate *Thy-1* promoters. The Thy-1 promoter of SLICK-A restricts Cre and YFP expression to neurons. There is no YFP fluorescence in glial cells, including GFAP+ astrocytes and Olig2+ oligodendrocytes or interneurons (29, 30). As a result, we can specifically assess the function of *Celsr3* in pyramidal neurons.

We first characterized the expression of YFP and the efficiency of recombination over the course of the first 2 postnatal weeks. SLICK-A is expressed in 59% of CA1 neurons indicated by the expression of YFP, and expressed sparsely in the cortex and other regions (Fig. S1*A*). In the absence of tamoxifen injections, SLICK-A–positive mice show no recombination at P7, although modest recombination at P14 due to a low level of leaky Cre expression (Fig. S1 *B* and *C*). However, administering tamoxifen at P7 to P8 resulted in robust recombination by P14, as indicated by 99% coexpression of tdTomato and YFP (Fig. S1 *A* and *D–F*; n = 4 mice, n = 460 neurons). For all of the in vivo experiments described below, control mice (SLICK-A–negative or SLICK-A–positive;*Celsr3*$^{+/+}$) were littermates and tamoxifen-injected at P7 and P8 and compared with tamoxifen-injected at P7 and P8 *Celsr3* cKOs (SLICK-A–positive;*Celsr3*$^{flox/flox}$). No changes were observed in the gross hippocampal anatomy, dendritic complexity, hippocampal commissure, mossy fiber projections, or CA1/CA3 cell-layer thickness and density, confirming that *Celsr3* was deleted late enough to avoid defects in axon guidance, neuronal migration, and dendrite morphogenesis (Fig. S1 *E–L*; control and *Celsr3* cKO, n = 3 mice). To characterize the cellular morphology, we analyzed dendritic complexity in the *Celsr3* cKOs by injecting Alexa Fluor hydrazide 555 into CA1 pyramidal cell bodies in acute hippocampal slices to label the
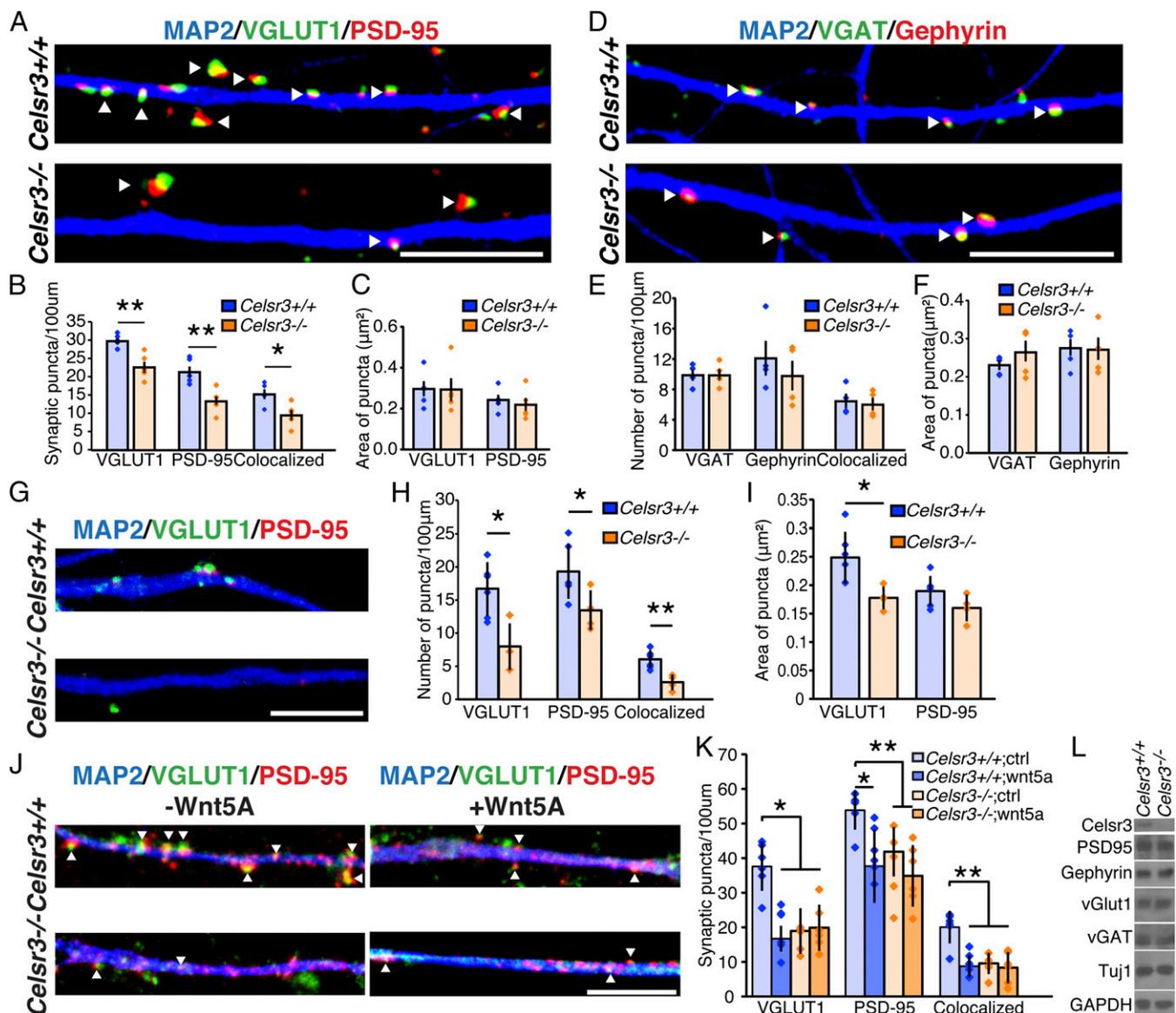
Fig. 2. *Celsr3* is required for glutamatergic synapse formation in hippocampal culture. (*A*) Immunostaining for pre- (green) and postsynaptic puncta (red) of glutamatergic synapses in 14-DIV hippocampal cultures from *Celsr3*$^{+/+}$ and *Celsr3*$^{-/-}$ E18.5 embryos. (*B* and *C*) Quantification of the density and area of glutamatergic synapses (*Celsr3*$^{+/+}$ $n = 5$ experiments, $n = 50$ neurons; *Celsr3*$^{-/-}$ $n = 5$ experiments, $n = 49$ neurons; *$P < 0.05$, **$P < 0.01$, Mann–Whitney *U*-statistic test). *Celsr3*$^{-/-}$ showed $24 \pm 4\%$, $36 \pm 6.7\%$, and $38 \pm 5\%$ reduction in presynaptic, postsynaptic, and colocalized puncta, respectively. The synaptic area was not significantly different. (*D*) Staining for pre- (green) and postsynaptic puncta (red) of inhibitory synapses in 14-DIV hippocampal cultures from *Celsr3*$^{+/+}$ or *Celsr3*$^{-/-}$ E18.5 embryos. (*E* and *F*) Quantification of the density and area of inhibitory synapses (*Celsr3*$^{+/+}$ $n = 4$ experiments, $n = 46$ neurons; *Celsr3*$^{-/-}$ $n = 4$ experiments, $n = 47$ neurons). Neither the density nor the area of inhibitory synapses was significantly different. (*G*) Staining for pre- (green) and postsynaptic puncta (red) of glutamatergic synapses in 6-DIV hippocampal cultures from *Celsr3*$^{+/+}$ and *Celsr3*$^{-/-}$ E18.5 hippocampal neurons. (*H* and *I*) Quantification of glutamatergic synaptic density showed a significant reduction in presynaptic, postsynaptic, and colocalized puncta. The area of post-synaptic puncta showed no change, whereas the area of presynaptic puncta showed a statistically significant change (*Celsr3*$^{+/+}$ $n = 127$ neurons, $n = 6$ experiments; *Celsr3*$^{-/-}$ $n = 85$ neurons, $n = 4$ experiments; *$P < 0.05$, **$P < 0.01$, Mann–Whitney *U*-statistic test). (*J*) Coimmunostaining for MAP2, PSD-95, and vGlut1 on *Celsr3*$^{+/+}$ or *Celsr3*$^{-/-}$ 14-DIV hippocampal neuronal dendrites after application of Wnt5a peptide. (*K*) Quantification of synaptic density after Wnt5a addition to *Celsr3*$^{+/+}$ or *Celsr3*$^{-/-}$ neurons. Colocalized vGlut1 and PSD-95 puncta density drops 56.4% after application of Wnt5a to *Celsr3*$^{+/+}$ neurons ($n = 6$ experiments; *$P < 0.05$, **$P < 0.01$, two-way ANOVA). *Celsr3*$^{-/-}$ neurons show a 52.2% decrease in the number of colocalized puncta compared with control neurons (**$P < 0.01$, two-way ANOVA). No significant change is observed after the addition of Wnt5a to *Celsr3*$^{-/-}$ neurons. (*L*) There is no change in synaptic protein expression in *Celsr3*$^{-/-}$ compared with *Celsr3*$^{+/+}$ hippocampal cultures at 14 DIV. Arrowheads denote colocalized puncta. All data are expressed as mean $\pm$ SD. (Scale bars, 10 μm.)

entire cell body and dendritic branching pattern. Completely labeled neurons without artificially broken branches 200 μm from the soma were analyzed with Sholl analysis (ImageJ plugin). We found no significant difference between control and *Celsr3* cKO basal dendrites or apical dendrites spanning the stratum radiatum (Fig. S1 *M* and *N*; Student's *t* test at each 10-μm interval;

control: $n = 11$ neurons, $n = 3$ mice; *Celsr3* cKO: $n = 10$ neurons, $n = 3$ mice).

We performed electron microscopy (EM) to examine synaptic density on distal CA1 apical dendrites, 150 to 200 μm from pyramidal cell bodies in P14 littermates (tamoxifen-injected controls, and *Celsr3* cKOs on P7 and P8). Synapses in this region are

formed by the CA1 dendrites and Schaffer collateral fibers from CA3 pyramidal neurons (Fig. 1A). Images were taken at equally spaced regions spanning the stratum radiatum. Because 59% of the CA1 neurons expressed SLICK-A and 99% of the SLICK-A–positive neurons had undergone Cre recombination, Celsr3 should be deleted in 58% of the CA1 neurons in tamoxifen-injected Celsr3 cKO mice. We observed a significant 36% decrease (Fig. 3 A and A′; *P < 0.05, Mann–Whitney U-statistic test) in the density of asymmetric (excitatory) axospinous synapses, whereas the density of asymmetric axodendritic synapses and symmetric (inhibitory) synapses remained unchanged (Fig. 3 A–B′). Due to the overlapping expression of Celsr2 with Celsr3 in hippocampal neurons (31, 32), we hypothesize that the remaining synapses in neurons lacking Celsr3 could be assembled by Celsr2 or a separate signaling pathway. These data show that Celsr3 is critical for the formation of excitatory synapses formed on spines and not the dendritic shaft or inhibitory synapses in vivo.

To assess the formation of functional synapses in vivo, we measured miniature excitatory postsynaptic current (mEPSC) and miniature inhibitory postsynaptic current (mIPSC) frequency and amplitude in acute hippocampal slices. Patch-clamp recordings of mEPSCs were obtained from CA1 pyramidal neurons of P13 to P15 littermates, tamoxifen-injected controls (n = 20 cells, n = 7 mice), and Celsr3 cKOs (n = 12 cells, n = 5 mice) on P7 and P8 and in the presence of TTX (1 mM) and gabazine (10 μM). We found that loss of Celsr3 caused a 45% reduction in the mean frequency of mEPSCs in CA1 neurons that express YFP (Fig. 3 C and D; **P = 0.0018, Mann–Whitney U-statistic test), with no change in the mean mEPSC amplitude (P = 0.77182, Mann–Whitney U-statistic test) or decay time constant (P = 0.42372, Mann–Whitney U-statistic test) of the remaining synapses (Fig. 3 C–E). There were no significant changes in mIPSC frequency (control n = 18 cells, n = 5 mice; Celsr3 cKO n = 14 cells, n = 3 mice; P = 0.89656, Mann–Whitney U-statistic test) or amplitude (P = 0.72786, Mann–Whitney U-statistic) or decay time constant (P = 0.95216, Mann–Whitney U-statistic test) (Fig. 3 F–H). These data indicate that Celsr3 plays a critical role in the development of excitatory, not inhibitory, synapses, contrary to previous studies that found embryonic loss of Celsr3 also affected the formation of inhibitory synapses (10). The decrease in mEPSC frequency observed in the Celsr3 cKO correlates with the observed decrease in excitatory synaptic density (asymmetric axospinous synapses) (Fig. 3 A–B′).

Celsr3 Conditional Knockout Mice Display Impaired Hippocampus-Dependent Behaviors. To test whether the reduction of excitatory synapse formation in the hippocampal CA1 pyramidal neurons resulted in behavioral deficits, we performed hippocampus-dependent behavioral tasks, including the Barnes maze and contextual fear conditioning (33, 34). Barnes maze tests were performed to evaluate hippocampus-dependent spatial memory in 3–4-month-old animals injected with tamoxifen at P7 and P8 (35). The Barnes maze consists of a circular field with 20 holes evenly spaced near its edge, one of which contains an escape route to a darkened escape chamber. Additional visual cues are placed on the wall for spatial orientation and a bright light is shined on the test area, giving the mouse an incentive to escape quickly. Because the maze does not involve water, it avoids complications due to the poorer swimming ability of mice (36). After initial habituation to the escape route and test area, the animals were subjected to four daily training sessions to form a stable memory of the location of the escape hole, followed by a probe test where the escape was removed.

We evaluated the latency to the first encounter of the escape hole by quantifying the latency to locate the hole with the escape chamber under it (37, 38). The controls had a significant learning curve (Fig. 4A; P = 0.0022, ANOVA, n = 18 male mice), whereas the Celsr3 cKOs only had a trend toward a learning curve (Fig. 4A; P = 0.0830, ANOVA, n = 17 male mice). In addition, the controls had a significantly shorter latency to locate the hole with the escape

chamber compared with Celsr3 cKOs (Fig. 4A; P = 0.0113, ANOVA). These data suggest that Celsr3 cKOs had impaired learning and memory in a hippocampus-dependent behavioral task.

In the probe test, control mice spent significantly more time in the quadrant originally housing the escape chamber than in the average of the other three quadrants, as expected (Fig. 4B; *P = 0.016, ANOVA), indicating that they were correctly remembering spatial cues in this test. On the other hand, the difference in percent time spent in the target vs. the average of the other quadrants was not statistically significant in Celsr3 cKOs (Fig. 4B; P = 0.54, ANOVA), indicating that Celsr3 cKOs had impaired spatial memory.

The mice were then subjected to a fear-conditioning paradigm that evaluates context- and cue-dependent learning. Because the fear-conditioning test sometimes has adverse effects on other behavioral tests, the fear-conditioning test was the final behavioral test performed on the mice, after the Barnes maze and the control behavioral tests. The strong aversive foot-shock stimulus provides a robust learning and memory motivation and behavioral response. Contextual-fear conditioning is a hippocampus- and amygdala-dependent behavior, whereas cued-fear conditioning is an amygdala-dependent behavior (39, 40). In this experimental paradigm, freezing behavior—which was defined as the absence of all except respiratory movement—is quantified after mice learn to associate context (the chamber environment) and a conditioned stimulus (a previously neutral tone stimulus) with an aversive foot shock. After initial habituation, mice are placed in the context of the chamber and exposed to the condition of a buzzer alarm sound/light in association with the aversive foot-shock stimulus. The next day, mice were submitted to the context test, in which mice were placed in the same chamber but in the absence of the tone, light, and aversive foot shock. On the final day, the cued stimulus test was performed, in which the mice were placed in a novel chamber exposed to the light and tone in the absence of the aversive foot shock. The Celsr3 cKOs displayed significantly weaker contextual conditioning compared with controls, which depends on hippocampal inputs (Fig. 4C; *P = 0.0491, ANOVA), whereas cued-fear conditioning was completely intact (Fig. 4C; P = 0.8825). These data suggest Celsr3 cKOs have a specific hippocampus-dependent behavioral defect.

Because SLICK-A is highly expressed in the hippocampus compared with other brain regions, including the cortex, at the time of tamoxifen injection (Fig. S1A), we expected behavioral defects would be restricted to those that are hippocampus-dependent. Indeed, there were no differences between control and Celsr3 cKOs in visual function as measured in the optomotor test, in anxiety-like behavior as measured using the light/dark transfer test, or in activity levels assessed in the locomotor activity test (Fig. 4 D–J).

Vangl2 Inhibits Excitatory Synapse Formation in Vivo. The localized expression of Vangl2 protein in the postsynaptic density suggests a role in synapse formation. However, Vangl2 is also an important signaling component in axon guidance. Germline Vangl2 mutations, either complete loss-of-function or gain-of-function (looptail), lead to massive axon projection defects (as well as other earlier developmental defects, such as an open neural tube), which will secondarily cause synapse formation defects (14, 41, 42). In two earlier studies, these germline Vangl2 mutants show a decrease in dendritic complexity and spine density (7, 11). To identify the biological function of Vangl2 in synapse formation, which occurs postnatally, we crossed a Vangl2 cKO allele (43) with the SLICK-A Cre line to delete Vangl2 postnatally. To evaluate the role of Vangl2 in functional synapses in vivo, we first measured mEPSC frequency and amplitude in acute hippocampal slices. Patch-clamp recordings of mEPSCs for P14 to P21 controls (n = 17 neurons, n = 7 mice) and Vangl2 cKOs (n = 18 neurons, n = 6 mice) were performed the same way as for Celsr3 cKO acute slices. In contrast to the germline Vangl2 mutations, we observed a large,

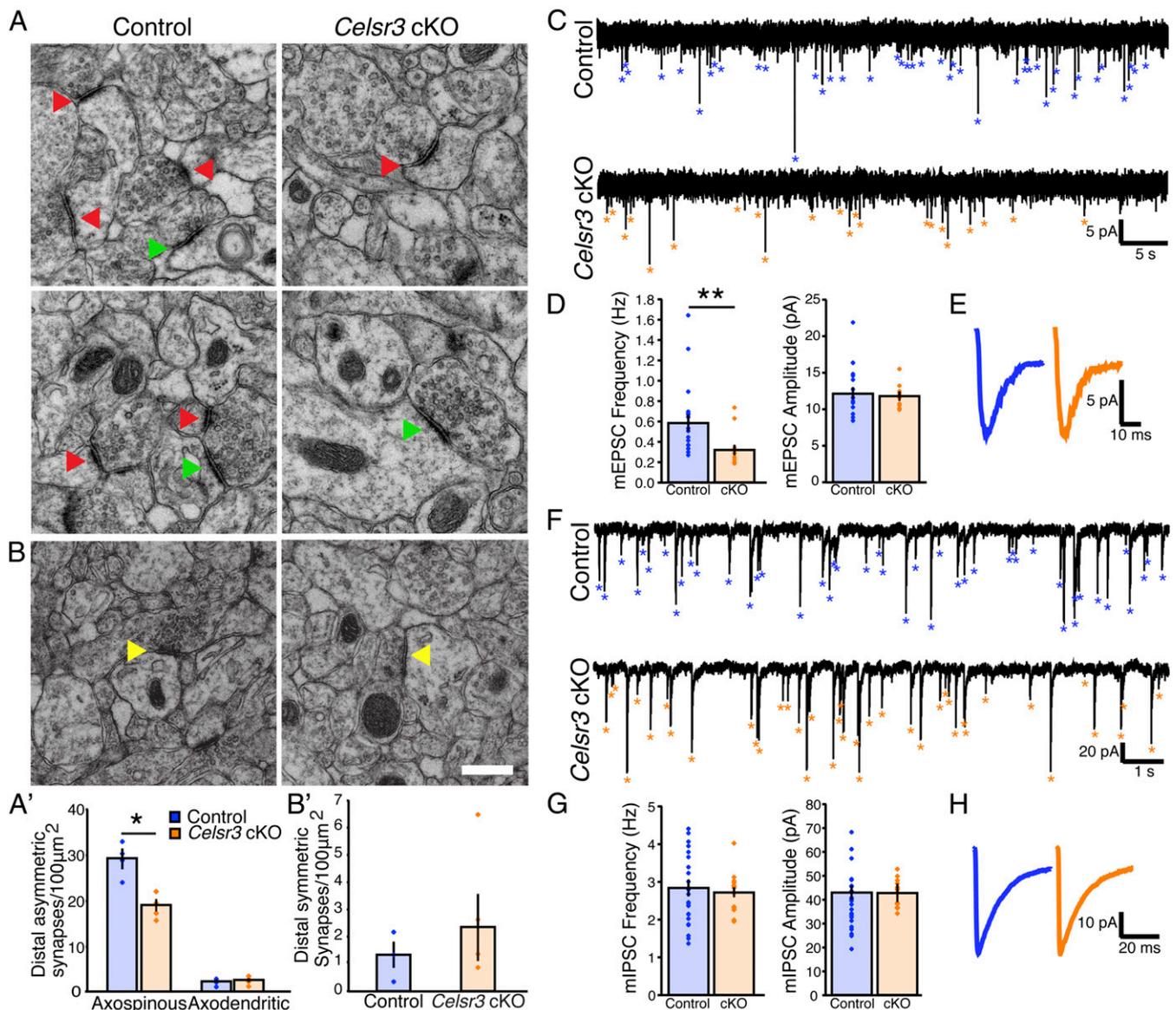**Fig. 3.** *Celsr3* is required for excitatory synapse formation in vivo. (*A–B'*) Representative EM images and quantification showing a 36% decrease in distal asymmetric axospinous synaptic density (indicated by red arrowheads) on the apical CA1 dendrites in the stratum radiatum of *Celsr3* cKOs. Controls and *Celsr3* cKOs (*n* = 4 mice per group; **P* < 0.05, Mann–Whitney *U*-statistic test). There was no significant change in axodendritic synaptic density (indicated by green arrowheads) or in distal symmetric synaptic density (indicated by yellow arrowheads). (Scale bar, 400 nm.) (*C*) Representative traces showing mEPSCs (control, blue asterisks; cKO, orange asterisks). (*D*) Quantification of mean mEPSC frequency (*Left*) and amplitude (*Right*) recorded in control (*n* = 20 cells, *n* = 7 mice) and *Celsr3* cKO littermates (*n* = 12 cells, *n* = 5 mice) (***P* < 0.01, Mann–Whitney *U*-statistic test). (*E*) Average event trace for control and *Celsr3* cKO shows no significant difference in decay time constant (*P* = 0.42372, Mann–Whitney *U*-statistic test) of the remaining synapses. (*F*) Representative traces showing mIPSCs (control, blue asterisks; cKO, orange asterisks) recorded in CA1 hippocampal pyramidal neurons from controls (*Upper*) or *Celsr3* cKOs (*Lower*). (*G*) Quantification of mean mIPSC frequency (*Left*) and amplitude (*Right*) recorded in control (*n* = 18 cells, *n* = 5 mice) and cKO littermates (*n* = 14 cells, *n* = 3 mice) shows no significant difference. (*H*) Average event trace for control and *Celsr3* cKO shows no significant difference in decay time constant (*P* = 0.95216, Mann–Whitney *U*-statistic test). All data are expressed as mean ± SEM.

58% increase in the mean mEPSC frequency (Fig. 5 *A* and *B*; **P = 0.00252, Mann–Whitney *U*-statistic test) with no change in the mean mEPSC amplitude (Fig. 5 *A* and *B*) or decay time constant (Fig. 5*C*; *P* = 0.50286, Mann–Whitney *U*-statistic test). Recording of the mIPSCs (control *n* = 13 neurons, *n* = 3 mice; *Vangl2* cKO *n* = 11 neurons, *n* = 3 mice) was performed and revealed no significant changes in mIPSC mean frequency (Fig. 5 *D* and *E*; *P* = 0.45326, Mann–Whitney *U*-statistic test), amplitude (Fig. 5 *D* and *E*; *P* = 0.48392, Mann–Whitney *U*-statistic test), or decay time constant (Fig. 5*F*; *P* = 0.47152, Mann–Whitney *U*-statistic test). These data suggest Vangl2 inhibits excitatory synapse formation.

We then analyzed spine density and morphology in P14 slices. Spines are dynamic structures that rapidly respond to environmental cues and molecular signaling. To preserve the spine density and morphology as close to the live mouse as possible, mice were anesthetized and quickly transcardially perfused with room temperature 4% paraformaldehyde (PFA). After processing the tissue (44), CA1 neurons and their dendritic branches were injected with Alexa Fluor hydrazide 555 to visualize spines. Only YFP+ neurons were included in the analysis, to ensure Cre-expressing dendrites were evaluated in the SLICK-A+;*Vangl2* cKO. CA1 oblique apical dendrites located 100 to 200 μm from
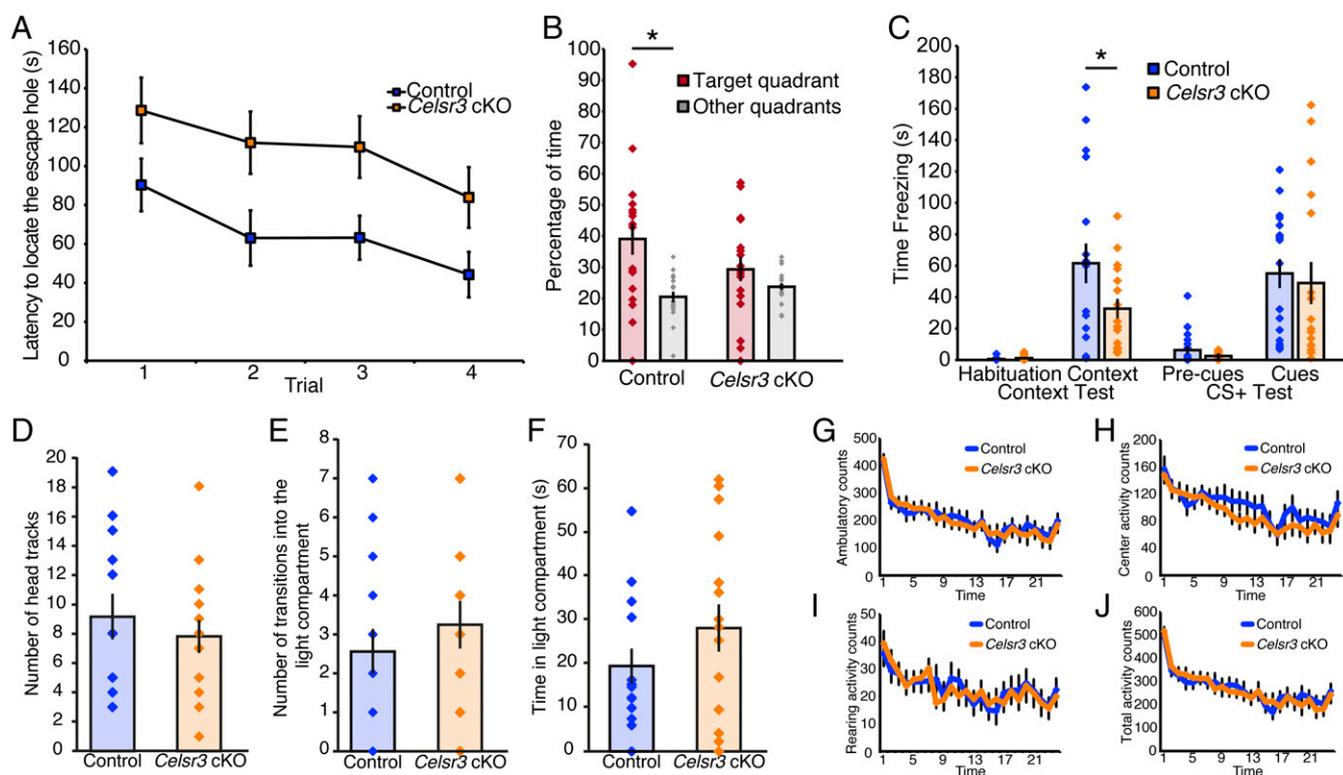
Fig. 4. *Celsr3* conditional knockout mice displayed impaired hippocampus-dependent behaviors. (*A*) *Celsr3* cKO littermates showed a longer latency in locating the escape chamber than control littermates in the Barnes maze assay (control $n = 18$; cKO $n = 17$; $P = 0.0113$, ANOVA). The controls had a significant learning curve ($n = 18$ male mice; $P = 0.0022$, ANOVA), whereas the *Celsr3* cKOs only showed a trend toward a learning curve ($n = 17$ male mice; $P = 0.0830$, ANOVA). (*B*) In the probe test, control mice spent significantly more time in the quadrant originally housing the escape chamber than the average of the other three quadrants [$F_{(1,17)} = 7.14$, *$P = 0.016$, ANOVA]. *Celsr3* cKOs showed no significant difference in the percent time spent in the target vs. other quadrants [$F_{(1,15)} = 0.40$, $P = 0.54$, ANOVA]. (*C*) *Celsr3* cKOs had significant impairment of hippocampus-dependent contextual conditioning (*$P < 0.05$, unpaired Student's *t* test) as reflected by the 50% reduced time spent freezing in the fear conditioning test. Hippocampus-independent cued conditioning remained the same. (*D–J*) Behavioral tests for non–hippocampus-dependent tasks showed no differences between control and *Celsr3* cKO mice as observed in (*D*) visual ability in the optomotor test, (*E* and *F*) anxiety-like behavior as measured using the light/dark transfer test, or (*G–J*) activity levels assessed in the locomotor activity test. All data are expressed as mean ± SEM.

the CA1 pyramidal neuronal cell bodies spanning the stratum radiatum were imaged using confocal microscopy, and spine density and morphology were analyzed. *Vangl2* cKO lead to 29% increase of spine density (Fig. 5*G*; 15.6 ± 0.06 spines per 10-μm dendritic segment; $n = 24$ neurons, $n = 4$ mice; ***$P < 0.001$, Student's *t* test) compared with controls (Fig. 5*G*; 12.1 ± 0.04 spines per 10-μm dendritic segment; $n = 31$ neurons, $n = 3$ mice). The average percentage of stubby-shaped spines in *Vangl2* cKOs decreased by 36% (Fig. 5*G*; ***$P < 0.001$, Student's *t* test) and the average percentage of thin-shaped spines increased by 18% (Fig. 5*G*; **$P = 0.00104$, Student's *t* test). The increase in mEPSC frequency observed in the *Vangl2* cKO correlates with the increase in excitatory synaptic density (as observed with quantifying spine density) in the *Vangl2* cKO. Together, these data suggest Vangl2 normally inhibits initial spine formation and potentially indirectly or directly promotes spine elimination, contrary to previous reports (7, 11).

Discussion

Opposing Roles of Celsr3 and Vangl2 in Glutamatergic Synapse Formation.
We found that PCP components are specifically localized at excitatory synapses as they form during early postnatal development and are associated either directly or indirectly with a number of key synaptic proteins. Hippocampal neurons lacking *Celsr3* develop approximately half as many excitatory synapses, as revealed by immunostaining, electron microscopy, and patch-clamp recording of mEPSCs both in cultured hippocampal

neurons and in acute hippocampal slices. Both the *Celsr3* KO and cKO showed no change in the number of inhibitory synapses, and the *Celsr3* cKO displayed no change in the activity of inhibitory synapses. The remaining excitatory synapses after the loss of *Celsr3* may be formed by other *Celsr* isoforms, such as Celsr2 expressed in an overlapping pattern in the hippocampus (31, 32), or a separate signaling pathway. These synaptic defects manifested in hippocampus-dependent behavioral deficits in adult *Celsr3* cKOs. Therefore, this study precisely identifies the essential role of a core PCP protein, Celsr3, in glutamatergic synapse formation. Because PCP components often have opposing functions and mutually exclusive subcellular localization, we anticipated that if PCP-like signaling is responsible for synapse formation, there should be some components that will inhibit synapse formation. Indeed, we found that Vangl2 inhibits glutamatergic synapse formation in the same hippocampal pyramidal neurons. Our paper provides evidence that components of the PCP signaling pathway have opposing functions in glutamatergic synapse formation, suggesting that PCP signaling mediates both positive and negative regulation of synapse formation and may be a candidate for a novel regulatory mechanism of synaptic plasticity.

Direct Role of Celsr3 and Vangl2 in Synapse Formation.
Because PCP signaling components play important roles in multiple steps of neural development, including neuronal migration and axon guidance, the proper experimental design is required to precisely dissect the function of each component and assign their role in
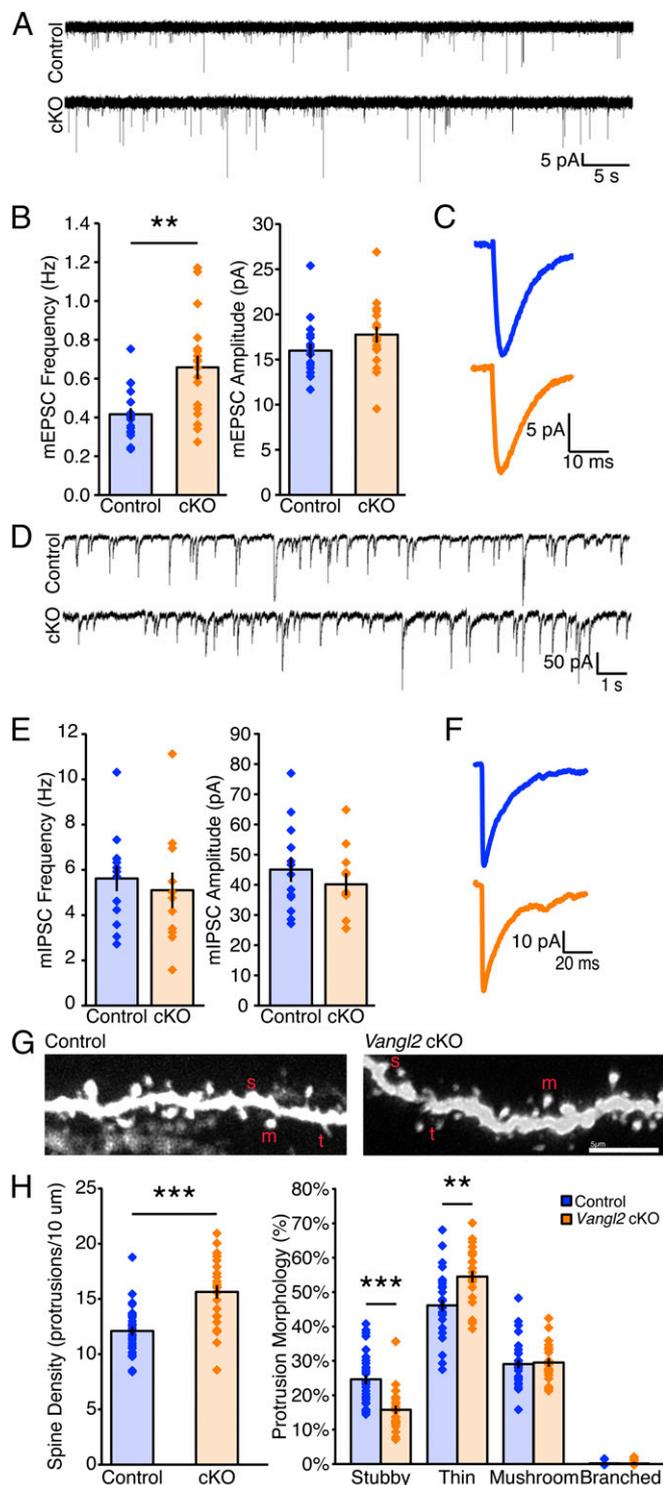
Fig. 5. *Vangl2* inhibits excitatory synapse formation in vivo. (*A*) Representative traces showing mEPSCs. (*B*) Quantification of mean mEPSC frequency (*Left*) and amplitude (*Right*) recorded in control ($n = 17$ cells, $n = 7$ mice) and *Vangl2* cKO littermates ($n = 18$ cells, $n = 6$ mice; **$P < 0.01$, Mann–Whitney U-statistic test). (*C*) Average event trace for control and *Vangl2* cKO shows no significant difference in decay time constant ($P = 0.50286$, Mann–Whitney U-statistic test). (*D*) Representative traces showing mIPSCs recorded in CA1 hippocampal pyramidal neurons from controls (*Upper*) or *Vangl2* cKOs (*Lower*). (*E*) Quantification of mean mIPSC frequency (*Left*) and amplitude (*Right*) recorded in control ($n = 13$ cells, $n = 3$ mice) and *Vangl2* cKO littermates ($n = 11$ cells, $n = 3$ mice) shows no significant difference. (*F*) Average event trace for control and cKO shows no significant difference in decay time constant ($P = 0.47152$, Mann–Whitney U-statistic test). (*G*) Represen-

synapse formation (12–18). Previous studies deleted the *Celsr3* gene either from early embryonic development or early embryonic brain development (8–10). When *Celsr3* was deleted from the embryonic forebrain using *Emx1-Cre*, a number of defects were found in neuronal cell numbers, migration, and axon guidance. These severe early defects will have an impact on later phases of development, such as synapse formation, which rely on correct axon guidance and dendrite formation. In addition to the reduction of glutamatergic synapses, inhibitory synapse numbers were found to be increased, probably due to the cell migration and axon guidance defects (10). In our study, *Celsr3* was deleted postnatally and we observed no change in tissue/cellular arrangement or axon guidance, therefore avoiding the early developmental confounds and allowing us to analyze the precise role of *Celsr3* in synapse formation. We did not observe any changes in inhibitory/symmetric synapse formation, consistent with our observation that Celsr3 and Vangl2 were not present in symmetric synapses. Similarly, another recent study implicated Vangl2 in glutamatergic synapse formation using the *looptail* mouse line containing a germline point mutation in the *Vangl2* gene (7). The authors found that Vangl2 binds to N-cadherin and promotes the endocytosis of N-cadherin, suggesting that Vangl2 normally disassembles synapses because N-cadherin promotes synapse formation. However, glutamatergic synapse formation was found to be greatly reduced using the *looptail* heterozygous mutant. The *looptail* mouse line has a point mutation in the *Vangl2* gene, which is embryonic-lethal in homozygotes, and heterozygotes show a number of severe neural developmental defects, including open neural tube, neurogenesis, neuronal migration, and axon guidance defects (14, 41). Therefore, the reduction of glutamatergic synapse formation is probably secondary to these earlier defects, misrepresenting the true function of Vangl2. Furthermore, the *looptail* mutation has been proposed to be a gain-of-function mutation, which could lead to other unexpected artifacts (42). Therefore, our study correctly assigns the function of Celsr3 and Vangl2 in glutamatergic synapse formation, laying down the foundation for future studies in this important pathway for synapse formation and potentially plasticity.

Wnt signaling has long been implicated in synapse formation (28). However, due to the complexity of Wnt signaling mechanisms, some published papers may report apparently conflicting results. In our study, we show that the Celsr3-mediated signaling pathway responds to the inhibitory function of Wnt5a in glutamatergic synapse formation. We also revealed the opposing roles of Wnt-regulated PCP components, Celsr3 and Vangl2, in glutamatergic synapse formation. Therefore, the precise function of Celsr3 and Vangl2 in synapse formation that we report here will provide important clues to fully understand the mechanisms of the assembly and plasticity of glutamatergic synapses in health and disease.

Materials and Methods

Celsr3 KO and cKO mice were provided by Andre Goffinet, Université Catholique de Louvain, Brussels (12, 13). *Vangl2* cKO mice were provided by Yingzi Yang, Harvard Medical School (43). The SLICK-A (JAX; 007606) line (29) constitutively expresses YFP and expresses a tamoxifen-inducible form of Cre recombinase, CreER[T2]. P7 and P8 pups were administered tamoxifen via intraperitoneal injection once daily. All experiments were performed with littermate, tamoxifen-injected controls. All animal work in this research was approved by the University of California, San Diego (UCSD) Institutional Animal Care and Use Committee.

tative images of control and *Vangl2* cKO dendritic spines from P16 CA1 neurons labeled by cell filling with Alexa Fluor 555 hydrazide. m, mushroom; s, stubby; t, thin. (Scale bar, 5 μm.) (*H*) *Vangl2* cKO has a significant 29% increase in average spine density (***$P < 0.001$, Student's t test). cKO has a significant 36% decrease in stubby-shaped spines (***$P < 0.001$, Student's t test) and an 18% increase in thin-shaped spines (**$P < 0.01$, Student's t test). All data are expressed as mean $\pm$ SEM.

Subcellular fractionation and coimmunoprecipitation data were collected from P14 wild-type mouse brains. All hippocampal culture data are from E18.5 embryos collected from littermate *Celsr3* KO mice (*Celsr3*$^{+/+}$ and *Celsr3*$^{-/-}$). Celsr3 and Vangl2 immunofluorescence staining was performed at P14 in SLICK-A–positive; *Celsr3*$^{+/+}$ mice. Celsr3 cKO EM was performed in P14 mice imaging the CA1 apical dendrites 150 to 200 μm from the cell body spanning the stratum radiatum. *Celsr3* cKO and *Vangl2* cKO mEPSC and mIPSC recordings were from acute hippocampal slices taken from 2- to 3-week-old mice. *Vangl2* cKOs were from mice that were killed and perfused at room temperature and then neurons individually labeled using Alexa Fluor hydrazide 555 (Invitrogen). All behavioral experiments were performed in mice 3 to 4 months old. Additional details regarding methods for all experiments described here are available in *SI Materials and Methods*.

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Supporting Information

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SI Materials and Methods

Subcellular Fractionation. Subcellular fractionation was performed as previously described with modifications (25). Forebrains from P14 wild-type mice were homogenized to 10% (wt/vol) in ice-cold 0.32 M sucrose buffer containing 1 mM $MgCl_2$, 0.5 mM $CaCl_2$, 1 mM $NaHCO_3$, and protease inhibitors using 16 strokes with a glass dounce. The homogenates were spun at $710 \times g$ for 30 min at 4 °C to pellet out nuclei and large debris. The supernatant was further centrifuged at $13,800 \times g$ for 10 min at 4 °C to get pellets (P2). P2 pellets were resuspended in the sucrose buffer and layered on top of a discontinuous sucrose gradient containing 1.0 and 1.4 M sucrose in 4 mM Hepes buffer (pH 7.4). The gradient was centrifuged at $82,500 \times g$ for 1 h at 4 °C. Synaptosomes were recovered from the cloudy band between 1.0 and 1.4 M sucrose, resuspended in 1 mM $NaHCO_3$ (1:9, vol/vol), and lysed by hypoosmotic shock using 3 strokes with a glass dounce. The lysates were then incubated with an equal volume of 0.32 M sucrose buffer containing 1% Triton-X shaking at 4 °C for 15 min and spun at $82,500 \times g$ for 1 h. The PSD fraction was obtained from the resulting pellet. The synaptic membrane fraction (SMF) was precipitated from the supernatant using the methanol/chloroform/H_2O method (45). Both the PSD fraction and SMF were solubilized in 3% SDS for Western blot analysis.

Celsr3 and Vangl2 Colocalization with Synaptic Protein Immunostaining. SLICK-A mice were perfused with PBS and 4% PFA at P14. Brains were removed and postfixed in 4% PFA overnight at 4 °C. Brains were then cryoprotected in 30% sucrose for 2 days before being embedded in OCT and frozen on dry ice. Coronal free-floating sections were prepared at 30 μm in a cryostat and used for immunofluorescence staining for Celsr3, Vangl2 (Santa Cruz; sc-46561), PSD-95 (Thermo Fisher; MA1-045), Bassoon (SYSY; 141004), and YFP (Millipore; AB16901). Before staining, tissues were treated with 1% SDS for 5 min at room temperature for antigen retrieval. Fluorescent z-stack images were acquired with an LSM 510 Zeiss confocal microscope using a 63× oil-immersion lens with 2× zoom-in.

Coimmunoprecipitation. A crude synaptosomal membrane fraction was obtained from mouse P14 brain (26). Briefly, the snap-frozen brain was homogenized in ice-cold Hepes buffer containing 0.32 M sucrose. After centrifuging the homogenate for 10 min at $800 \times g$ and 4 °C, the supernatant was centrifuged for 15 min at $9,200 \times g$ and 4 °C. The pellet (P2) contains crude synaptosomal membrane. The P2 pellet was solubilized and the supernatant was used for coimmunoprecipitation assay. Control IgG (Sigma), anti-Frizzled3 antibodies (R&D), or anti-Celsr3 antibodies (17) were combined with precleared crude synaptosomal membrane fraction and protein A/G agaroses. The proteins retained on the beads were then analyzed by Western blotting.

Hippocampal Neuron Culture. Hippocampal neuron culture was performed as described previously (27). Hippocampi were dissected from E18 mice. Cells were pelleted and resuspended in Neurobasal medium supplemented with 2% B27 (Invitrogen), 5% FBS (Invitrogen), penicillin/streptomycin (CellGro), and GlutaMAX (Invitrogen) and plated on poly-D-lysine– (Millipore) and laminin- (Invitrogen) coated glass coverslips in a 24-well plate at a density of 2×10^4 cells per square centimeter for immunostaining or 6×10^4 cells per square centimeter for electrophysiology. Medium was changed every 3 days. Arabinofuranosyl cytidine (4 μM) was added at day 6 in vitro to prevent glial cell proliferation. Cultures were grown for 6 or 14 to 15 DIV at 37 °C with 5% carbon dioxide atmosphere.

Hippocampal Culture Immunofluorescence and Image Analysis. For puncta density analysis, the cultured neurons (6 and 14 DIV) from littermate $Celsr3^{+/+}$ and $Celsr3^{-/-}$ hippocampi were immunostained with the following primary antibodies: chicken anti-MAP2 (Abcam) to visualize dendrites, mouse anti–PSD-95 (Affinity BioReagents), and rabbit anti-VGLUT1 (Synaptic Systems). $Celsr3^{+/+}$ and $Celsr3^{-/-}$ hippocampal cultures were treated with the same staining process. Z-stacked images were obtained with a Carl Zeiss microscope using a 63× oil-immersion lens. Three or more neurons with pyramidal morphology and at least two diameters' distance from the neighboring neurons were selected per coverslip. Three coverslips were used for each group per experiment ($n = 5$ experiments for excitatory synapses and $n = 4$ experiments for inhibitory synapses). Secondary dendrites were chosen for puncta analysis. Number and area of puncta were analyzed using the ImageJ Puncta Analyzer program (46, 47), and the length of the dendrite was analyzed by ImageJ (NIH).

Mouse Lines. Celsr3 KO and cKO mice were provided by Andre Goffinet (12, 13). Vangl2 cKOs were provided by Yingzi Yang (43). The strain backgrounds between the Celsr3 cKOs and Vangl2 cKOs were different, with Celsr3 cKOs being 100% C57BL/6 whereas the Vangl2 cKOs were mixed 129;C57BL/6. The SLICK-A (JAX; 007606) line (29) has a dual Thy-1 promoter system, with one promoter constitutively expressing YFP and the other Thy-1 promoter expressing a tamoxifen-inducible form of Cre recombinase, $CreER^{T2}$. For postnatal injections, tamoxifen (Sigma; T5648) was dissolved in corn oil at 10 mg/mL by rocking overnight at room temperature and used immediately the following day. P7 and P8 pups were administered ~50 to 60 μL tamoxifen via i.p. injection once daily for 2 days. All experiments were performed with littermate tamoxifen-injected controls. Controls for Celsr3 cKO experiments: SLICK-A–positive; $Celsr3^{+/+}$, or SLICK-A–negative $Celsr3^{+/+}$, $Celsr3^{fl/+}$, or $Celsr3^{fl/fl}$ mice; for Vangl2 cKO experiments: SLICK-A–positive;$Vangl2^{+/+}$, or SLICK-A–negative $Vangl2^{+/+}$, $Vangl2^{fl/+}$, or $Vangl2^{fl/fl}$ mice. For the cKO group, tamoxifen-injected SLICK-A–positive;$Celsr3^{fl/fl}$ and SLICK-A–positive;$Vangl2^{fl/fl}$ mice were used.

Electron Microscopy. P14 Celsr3 cKOs and their littermate control mice were anesthetized i.p. with a ketamine/xylazine mixture and perfused with modified Karnovsky's fixative (2.5% glutaraldehyde and 2% PFA in 0.15 M sodium cacodylate buffer, pH 7.4). After postfixation in the same solution overnight at 4 °C, whole brains were cut coronally (200-μm) with a vibratome. Sections were postfixed with 1% osmium tetroxide for 1 h on ice. Following en bloc staining in 2% uranyl acetate in distilled water for 1 to 2 h, sections were dehydrated in a graded ethanol series and then treated twice in acetone for 10 min each. Sections were embedded by being infiltrated in Durcupan resin (Sigma-Aldrich) and then placed in an oven at 60 °C for 48 h. Ultrathin sections from the CA1 region were cut and stained with uranyl acetate and Sato's lead. Images were captured using an FEI Tecnai G2 Spirit BioTWIN transmission electron microscope equipped with an Eagle 4k HS digital camera (FEI). Twelve to 20 fields from the distal stratum radiatum (150 to 200 μm from CA1 pyramidal cell bodies) were selected under 6,800× magnification, and synapses were counted in each field.

Electrophysiology in Acute Slices. For *Celsr3* cKOs at P13 to P15, mice were anesthetized by isoflurane and decapitated, and their brains were quickly removed and placed in ice-cold dissection buffer containing 110 mM choline chloride, 2.5 mM KCl, 1.2 mM NaH$_2$PO$_4$, 25 mM NaHCO$_3$, 20 mM glucose, 1.3 mM sodium ascorbate, 2.4 mM sodium pyruvate, 0.5 mM CaCl$_2$, and 7 mM MgCl$_2$ (pH 7.3). For *Vangl2* cKO acute brain slices ranging over 2 to 3 weeks old, bath solution and internal solutions were prepared similar to *Celsr3* cKO.

Sagittal acute slices were cut in 300-µm-thick sections on a vibratome (Leica; VT1200). Slices were allowed to recover at 34 °C for 15 min and then at room temperature for 45 min to 6 h in carbogenated artificial cerebrospinal fluid (ACSF) containing 119 mM NaCl, 2.5 mM KCl, 26.2 mM NaHCO$_3$, 1 mM NaH$_2$PO$_4$, 2.5 mM CaCl$_2$, 1.3 mM MgCl$_2$, and 11 mM glucose (pH 7.3). Individual slices were transferred to a recording chamber and then continuously perfused at a rate of 2 to 3 mL/min with ACSF containing 1 µM TTX (Tocris). To isolate mEPSCs we added 10 µM gabazine, and for mIPSCs we added 20 µM CNQX (6-cyano-7-nitroquinoxaline-2,3-dione), to the ACSF. CA1 pyramidal cells were visualized by infrared differential interference microscopy and voltage-clamped at −70 mV.

Whole-cell recordings for mEPSCs were made using 3- to 5-MΩ pipettes filled with an internal solution that contained 115 mM CsMeSO$_4$, 20 mM CsCl, 10 mM Na$_2$ phosphocreatine, 2.5 mM MgCl$_2$, 4 mM Na$_2$ATP, 0.3 mM Na$_3$GTP, 0.6 mM EGTA, 10 mM Hepes, and 5 mM QX-314 (pH 7.3 adjusted with CsOH, and osmolarity was maintained at 285 to 290 mOsm). Whole-cell recordings for mIPSCs were made using 3- to 5-MΩ pipettes filled with an internal solution that contained 130 mM CsCl, 4 mM NaCl, 5 mM Na$_2$ phosphocreatine, 2 mM MgCl$_2$, 2 mM Na$_2$ATP, 0.6 mM Na$_3$GTP, 1.1 mM EGTA, and 5 mM Hepes (pH 7.3 adjusted with CsOH, and osmolarity was maintained at 285 to 290 mOsm).

Analysis of the recording started after 5 min to allow for stabilization of the established whole-cell configuration. Signals were recorded with a 10× gain, low-pass Bessel-filtered at 2 kHz (Axopatch 200B; Molecular Devices) with Clampex 10 software (pCLAMP; Molecular Devices), and analysis for *Celsr3* cKOs was performed with Clampfit software (pCLAMP; Molecular Devices). *Vangl2* cKOs were analyzed using the Mini Analysis Program (Synaptosoft). Automatic detection was verified by visual inspection. Series and input resistances were monitored during the recording, and experiments were removed if total series resistance changed more than 20% or if the current required to hold the cell at −70 mV varied by more than 50 pA. Values are presented as mean ± SEM. Mann–Whitney *U*-statistic test was used to compare changes with the control.

Mouse Behavioral Assessments. All behavioral tests were performed on mice 3 to 4 months of age, and tamoxifen was injected as described above using littermate controls.

Barnes maze test. The Barnes maze is a spatial learning and memory test (35, 48, 49) sensitive to impaired hippocampal function (50). This task has the benefit of minimizing pain and distress to the animal. Traditional spatial tests involve swimming in opaque water to locate a hidden platform (the Morris water maze). The Barnes maze is less physically taxing than the Morris water maze, making it more suitable for mice, which are not as strong as rats, and also suitable for studies in all ages and abilities of mice (48). In addition, the strategies used by the animals to perform the task are readily revealed. The Barnes maze used was an opaque Plexiglas disk 75 cm in diameter elevated 58 cm above the floor by a tripod. Twenty holes, 5 cm in diameter, were located 5 cm from the perimeter, and a black Plexiglas escape box (19 × 8 × 7 cm) was placed under one of the holes. Distinct spatial cues were located all around the maze and kept constant throughout the study.

On the first day of testing a training session was performed, which consisted of placing the mouse in the escape box and leaving it there for 1 min. One minute later, the first session was started. At the beginning of each session, the mouse was placed in the middle of the maze in a 10-cm-high cone-shaped silver start chamber. After 10 s the start chamber was removed, a buzzer (80-dB) and a light (400-lx) were turned on, and the mouse was set free to explore the maze. The session ended when the mouse entered the escape tunnel or after 3 min elapsed. When the mouse entered the escape tunnel, the buzzer was turned off and the mouse was allowed to remain in the dark for 1 min. If the mouse did not enter the tunnel by itself it was gently put in the escape box for 1 min. The tunnel was always located underneath the same hole (stable within the spatial environment), which was randomly determined for each mouse. Mice were tested once a day for 4 days for the acquisition portion of the study.

For the probe test (day 5), the escape tunnel was removed and the mouse was allowed to freely explore the maze for 3 min. The time spent in each quadrant was determined and the percent time spent in the target quadrant (the one originally containing the escape box) was compared with the average percent time in the other three quadrants. This is a direct test of spatial memory, as there is no potential for local cues to be used in the mouse's behavioral decision.

Each session was videotaped and scored by an experimenter blind to the genotype of the mouse. Measures recorded included the latency to escape the maze and the number of errors made per session. Errors were defined as nose pokes and head deflections over any hole that did not have the tunnel beneath it.

Conditioned fear test. In this procedure, mice learn to associate a novel environment (context) and a previously neutral stimulus (conditioned stimulus, a tone) with an aversive foot-shock stimulus (51). It allows for the assessment of both hippocampus-dependent and amygdala-dependent learning processes in the same mouse (52, 53). Testing then occurs in the absence of the aversive stimulus. Conditioned animals, when exposed to the conditioned stimuli, tend to refrain from all but respiratory movements by freezing. Freezing responses can be triggered by exposure to either the context in which the shock was received (context test) or the conditioned stimulus (CS+ test). Conditioning took place in Freeze Monitor chambers (Med Associates) housed in sound-proofed boxes. The conditioning chambers (26 × 26 × 17 cm) were made of Plexiglas, with speakers and lights mounted on two opposite walls and shockable grid floors.

On day 1, mice were placed in the conditioning chamber for 5 min to habituate them to the apparatus. On day 2, the mice were exposed to the context and conditioned stimulus (30-s, 3,000-Hz, 80-dB sound + white light) in association with foot shock (0.60-mA, 2-s, scrambled current). Specifically, the mice received three shock exposures in their 6-min test, each in the last 2 s of a 30-s tone/light exposure. On day 3, contextual conditioning (as determined by freezing behavior) was measured in a 5-min test in the chamber where the mice were trained (context test). On the following day, the mice were tested for cued conditioning (CS+ test). The mice were placed in a novel context for 3 min, after which they were exposed to the conditioned stimuli (light + tone) for 3 min. For this test, the chamber was disguised with new walls (white opaque plastic creating a circular compartment in contrast to a clear plastic square compartment) and a new floor (white opaque plastic in contrast to a metal grid). Freezing behavior (i.e., the absence of all voluntary movements) was measured in all of the sessions by real-time digital video recordings calibrated to distinguish between subtle movements, such as whisker twitches and tail flicks, and freezing behavior. Freezing behavior in the context and cued tests (relative to the same context before shock and an altered context before tone, respectively) is indicative of the formation of an association

between the particular stimulus (either the environment or the tone) and the shock, namely that learning has occurred.

Optomotor test of visual ability. The optomotor is a stationary elevated platform surrounded by a drum with black and white striped walls. The mouse was placed on the platform to habituate for 1 min and then the drum rotated at 2 rpm in one direction for 1 min, stopped for 30 s, and then rotated in the other direction for 1 min. The total number of head tracks (15° movements at the speed of the drum) was recorded. In our hands, mice that have intact vision track 5 to 15 times, whereas blind mice do not track at all.

Light/dark transfer test. The light/dark transfer procedure has been used to assess anxiety-like behavior in mice by capitalizing on the conflict between exploration of a novel environment and the avoidance of a brightly lit open field (49). The apparatus is a rectangular box made of Plexiglas divided by a partition into two environments. One compartment (14.5 × 27 × 26.5 cm) is dark (8 to 16 lx) and the other compartment (28.5 × 27 × 26.5 cm) is highly illuminated (400 to 600 lx) by a 60-W light source located above it. The compartments are connected by an opening (7.5 × 7.5 cm) located at floor level in the center of the partition. Mice were placed in the dark compartment to start the 5-min test. The time spent in the lit compartment was used as a predictor of anxiety-like behavior, namely a greater amount of time in the lit compartment would be indicative of decreased anxiety-like behavior.

Locomotor activity test. Locomotor activity is measured in polycarbonate cages (42 × 22 × 20 cm) placed into frames (25.5 × 47 cm) mounted with two levels of photocell beams at 2 and 7 cm above the bottom of the cage (San Diego Instruments). These two sets of beams allowed for the recording of both horizontal and vertical (rearing) behavior. Horizontal locomotor activity was further separated into ambulation (consecutive beam breaks, indicative of walking or running), center activity (all beam breaks in the center of the apparatus), and total horizontal activity (all lower-level beam breaks). A thin layer of bedding material was applied to the bottom of the cage. Mice were tested for 120 min.

Spine Density and Morphology. Mice were perfused and sectioned, and individual neurons and their dendrites were labeled with Alexa Fluor hydrazide 555 (Invitrogen) as previously described (44). SLICK-A:*Vangl2* cKO pups were i.p. injected with tamoxifen at P7 and P8. At P16, pups were anesthetized with a ketamine/xylazine mixture followed by transcardial perfusion with 20 mL of room temperature 4% PFA, postfixed for 30 to 45 min in 4% PFA, vibratome-sectioned at 100 μm, and then postfixed again for 10 min. Alexa Fluor hydrazide 555 (10 mM in 200 mM KCl) was injected into dendritic segments ionophoretically by filling the cell with Alexa Fluor hydrazide dye. Sections were briefly fixed for

15 min to preserve the fluorescent label. To confirm the labeled dendrite was from a SLICK-A–positive Cre-expressing neuron, colabeling with YFP was achieved by immunolabeling with GFP antibody (Invitrogen). Thirty- to 40-μm segments of CA1 oblique apical dendrites, which are located 100 to 200 μm from the CA1 pyramidal neuronal cell bodies in the stratum radiatum, were analyzed. These spines represented the postsynaptic structures of the Schaffer collateral–CA1 synapses.

The maximal spine length and head width were measured manually with ImageJ (NIH) to characterize spine shape as previously described (54–59). Spines with heads equal to or less than the head width were categorized as "thin," and those with heads greater than the neck width were categorized as "mushroom." Spines without a neck that had a width longer than their lengths were termed "stubby." Spines with one neck and a branch point resulting in two heads were termed "branched." Protrusions not clearly seen or with lengths >5 μm were excluded from analysis. Both image acquisition and morphometric analyses were blinded. Results were compared between littermates and then pooled by genotype to assess the influence of prenatal care. The results are presented as mean ± SEM. Statistical differences were determined using Student's *t* test for two-group comparisons. Analyses of spine density and morphology were done by experimenters blinded to the genotypes of the animals.

Sholl Analysis. Acute brain slices 300 μm thick were prepared as described in the electrophysiology experiments. Slices were transferred from a room temperature recovery chamber to the electrophysiology rig, where slices were maintained at 32 °C to aid the neuron to take in the dye. CA1 pyramidal neurons about 50 μm from the slice surface were individually injected with Alexa Fluor hydrazide 555 diluted in internal solution for 10 to 15 min. After filling two or three cells per slice, the slice was fixed in 4% PFA for 30 min followed by a brief PBS wash and mounting onto slides.

Neurons that were completely filled by the dye and not artificially broken within 200 μm of the soma were used for Sholl analysis. Neurons were imaged using the 40× objective on a Carl Zeiss confocal microscope. The cell body, basal dendrites, and apical dendrites were reconstructed using Adobe Illustrator CS2. The Sholl analysis plugin available from ImageJ software was used to quantify the number of dendritic crossings made with concentric spheres around the soma center with radii increasing in increments of 10 μm. CA1 pyramidal cells were analyzed independently, rather than averaging all cells within an animal, because it has been shown that subsets of cells within an anatomical nucleus, such as CA1 hippocampus, can participate independently in behaviors and that different subsets have different functions (60).
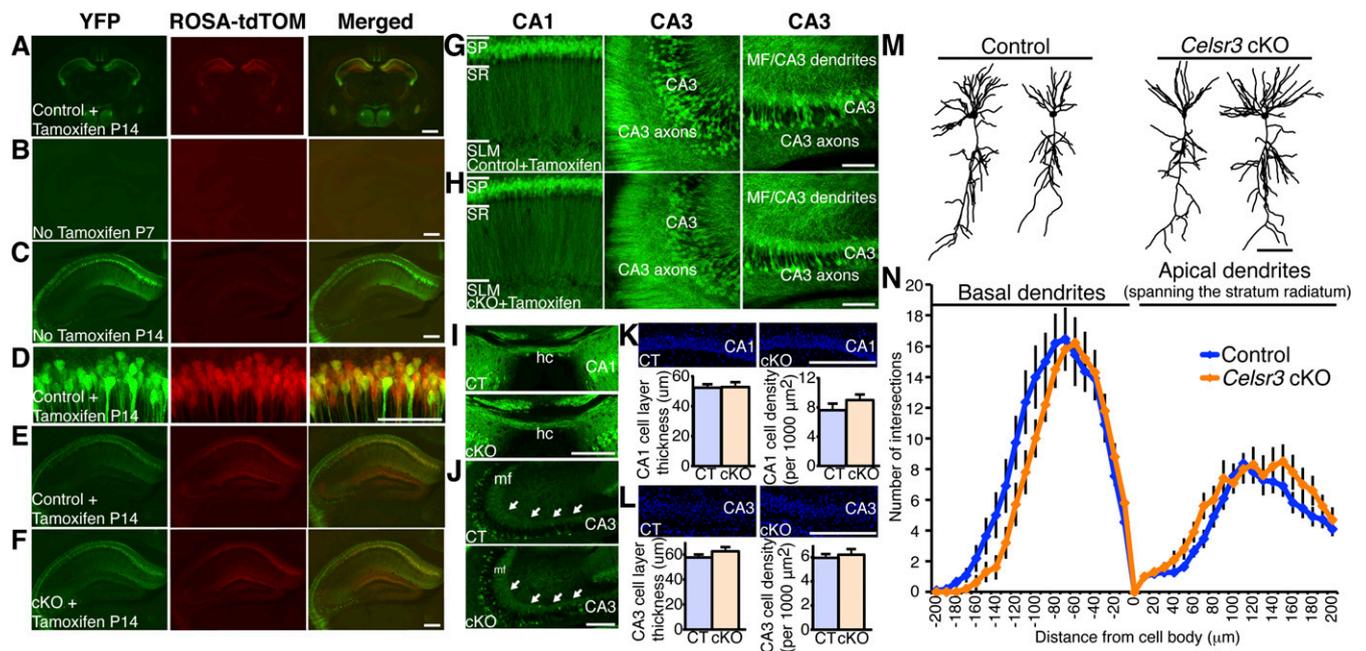
Fig. S1. Characterization of recombination and hippocampal development in SLICK-A–positive;*Celsr3* cKO. (*A*) SLICK-A–positive;*ROSA26-tdTomato*+ mice tamoxifen-injected at P7 and P8 show coexpression of YFP and tdTomato in a subset of neurons at P14 in the brain. (Scale bar, 1 mm.) (*B* and *C*) SLICK-A–positive mice not injected with tamoxifen show no recombination at P7 and modest recombination at P14. (Scale bars, 100 μm.) (*D*) SLICK-A–positive; *ROSA26-tdTomato*+ mice tamoxifen-injected at P7 and P8 show coexpression of YFP and tdTomato in a subset of neurons at P14 in CA1 pyramidal neurons. The recombination efficiency is 99% in SLICK-A–positive CA1 hippocampal neurons ($n = 460$ cells, $n = 4$ animals). (Scale bar, 100 μm.) (*E* and *F*) Tamoxifen-injected P14 SLICK-A–positive;*Celsr3*^{+/+};*ROSA26-tdTomato* and SLICK-A–positive;*Celsr3*[flox/flox];*ROSA26-tdTomato* mice exhibit constitutively active YFP reporter expression as well as tdTomato reporter expression to visualize gross hippocampal anatomy showing no defects in the *Celsr3* cKO. (Scale bar, 100 μm.) (*G* and *H*) Tamoxifen-injected P14 SLICK-A–positive;*Celsr3*^{+/+} and SLICK-A–positive;*Celsr3*[flox/flox] YFP expression showed no abnormal cell-body migration/localization or dendritic branching defects. (Scale bars, 100 μm.) (*I*) YFP expression shows no defects in *Celsr3* cKO hippocampal commissure. (Scale bar, 200 μm.) (*J*) YFP expression shows no abnormal mossy fiber projections in tamoxifen-injected *Celsr3* cKO. (Scale bar, 200 μm.) Arrows denote mossy fiber projections. (*K* and *L*) DAPI staining in tamoxifen-injected SLICK-A–positive;*Celsr3*^{+/+} and SLICK-A–positive;*Celsr3*[flox/flox] shows no difference in CA1 or CA3 cell-layer thickness or cell density. (Scale bars, 200 μm.) (*M*) Representative reconstructed traces of control and *Celsr3* cKO CA1 pyramidal neurons. (*N*) Sholl analysis of basal dendrites and apical dendrites spanning the stratum radiatum shows no significant difference between tamoxifen-injected control and *Celsr3* cKOs (Student's *t* test; control $n = 11$ neurons, $n = 3$ mice; cKO $n = 10$ neurons, $n = 3$ mice). All data are expressed as mean ± SEM. CT, control; hc, hippocampal commissure; MF, mossy fiber; SLM, stratum lacunosum moleculare; SP, stratum pyramidale; SR, stratum radiatum.