AMPA Receptor Auxiliary Subunit GSG1L Suppresses Short-Term Facilitation in Corticothalamic Synapses and Determines Seizure Susceptibility

Graphical Abstract

Highlights
- Input-specific synaptic phenotypes in the anterior thalamic neurons of GSG1L KO mice
- Short-term plasticity in corticothalamic synapses is regulated by GSG1L
- Stargazin is functionally absent from corticothalamic synapses
- GSG1L KO mice have increased hyperexcitability and seizure susceptibility

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In Brief
Kamalova et al. report the phenotypes of GSG1L KO mice. The synaptic function of AMPAR auxiliary subunit GSG1L in the anterior thalamus is input specific. GSG1L suppresses short-term facilitation and decreases AMPAR activity specifically in corticothalamic synapses, where stargazin is functionally absent. GSG1L KO mice exhibit hyperexcitability and seizure susceptibility.

Kamalova et al., 2020, Cell Reports 32, 107921
July 21, 2020 © 2020 The Author(s).
https://doi.org/10.1016/j.celrep.2020.107921
AMPA Receptor Auxiliary Subunit GSG1L Suppresses Short-Term Facilitation in Corticothalamic Synapses and Determines Seizure Susceptibility

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SUMMARY

The anterior thalamus (AT) is critical for memory formation, processing navigational information, and seizure initiation. However, the molecular mechanisms that regulate synaptic function of AT neurons remain largely unexplored. We report that AMPA receptor auxiliary subunit GSG1L controls short-term plasticity in AT synapses that receive inputs from the cortex, but not in those receiving inputs from other pathways. A canonical auxiliary subunit stargazin co-exists in these neurons but is functionally absent from corticothalamic synapses. In GSG1L knockout mice, AT neurons exhibit hyperexcitability and the animals have increased susceptibility to seizures, consistent with a negative regulatory role of GSG1L. We hypothesize that negative regulation of synaptic function by GSG1L plays a critical role in maintaining optimal excitation in the AT.

INTRODUCTION

Regulation of excitatory synaptic transmission is essential for synaptic plasticity, learning, and memory. AMPA-type ionotropic glutamate receptors (AMPARs), which are ligand-gated ion channels activated by the neurotransmitter glutamate, play a key role in this process by mediating most fast excitatory neurotransmission in the brain (Bowie, 2008; Traynelis et al., 2010; Huganir and Nicoll, 2013). GluA1–GluA4 are the pore-forming subunits of AMPARs that assemble into functional ligand-gated ion channels consisting of homo- and heterotetramers (Greger et al., 2017). The canonical structural units of native AMPARs are complexes composed of the core tetramers of GluA subunits and their auxiliary subunits (Nakagawa et al., 2005; Schwenk et al., 2012; Zhao et al., 2019).

AMPA auxiliary subunits are membrane proteins that regulate ion channel gating and trafficking of AMPARs (Jackson and Nicoll, 2011). The most extensively studied among these are the stargazin/TARPs (transmembrane AMPAR regulatory proteins) (Tomita et al., 2003). Other auxiliary subunits include cornichon homologs 2 and 3 (CNHI2/3) (Schwenk et al., 2009), CKAMP44 (also known as Shisa9) (von Engelhardt et al., 2010), Shisa6 (Klaassen et al., 2016), SOL-1 (Zheng et al., 2004), and GSG1L (Schwenk et al., 2012; Shanks et al., 2012). Each class of auxiliary subunits is structurally unrelated to the others except for GSG1L and TARPs, which are both claudin homologs. Modulation of AMPARs by auxiliary subunits is predicted to substantially affect brain function (Jackson and Nicoll, 2011). Mutations in one or more AMPAR auxiliary subunits lead to neurological and cognitive deficits in both mice and humans (Everett et al., 2007; Hamdan et al., 2011; Floor et al., 2012).

Most AMPAR auxiliary subunits positively modulate AMPAR function by promoting synaptic trafficking and/or modifying gating toward increasing net charge transfer (Jackson and Nicoll, 2011). A subset of auxiliary subunits has mixed effects on gating. For example, CKAMP44 slows AMPAR deactivation, increasing net charge transfer during synaptic transmission, but also delays recovery from desensitization so that the channel is not immediately re-usable (von Engelhardt et al., 2010). Similarly, TARP γ-8 slows AMPAR desensitization and delays recovery from AMPAR desensitization of GluA2 and GluA3 specifically (Cais et al., 2014). Among all auxiliary subunits, GSG1L stands out for having a strong negative modulatory function (McGee et al., 2015; Gu et al., 2016; Mao et al., 2017) (summarized in Figure 1A). Although GSG1L also slows desensitization, it stabilizes the desensitized state (Twomey et al., 2017) and dramatically delays recovery from desensitization over a magnitude slower than other auxiliary subunits (Schwenk et al., 2012; Shanks et al., 2012). In addition, GSG1L reduces single-channel conductance and calcium permeability of calcium-permeable AMPARs (McGee et al., 2015). Consistently, overexpression of GSG1L decreases the amplitude of evoked excitatory postsynaptic potentials.
GSG1L Is Sufficient to Express Short-Term Plasticity In Vitro

The most significant action of GSG1L is the slowing of AMPAR recovery from desensitization (Figure 1A) (Schwenk et al., 2014; Shanks et al., 2012). Therefore, in synapses in which AMPARs desensitize, GSG1L should prevent immediate re-use of AMPARs during repetitive activation and as a result promote short-term depression (STD) or attenuate short-term facilitation. Conversely, elimination of GSG1L should relieve STD or, in an extreme case, convert to short-term facilitation. Consistent with this hypothesis, GSG1L was sufficient to induce STD in a reconstituted system (Figures 1B–1E). Indeed, we found that outside-out patches pulled from HEK293 cells that co-express GSG1L and GluA2 (flip splice variant with unedited Q pore) show significantly depressing currents in response to 1 mM glutamate pulses of 1 ms at 20 Hz, whereas in the absence of GSG1L, the amplitudes remained constant (Figures 1B–1E).

High and Persistent Expression of GSG1L in the AT

We determined the spatiotemporal expression of GSG1L using a transgenic rat that expresses a lacZ reporter under the control of the GSG1L promoter (Figures S1A and S1B). High expression of GSG1L was found in the AT throughout development (Figures 1F and S1C). GSG1L is restricted to AD and AV but absent from anteromedial nucleus (AM) or lateral dorsal nucleus (LD), in agreement with in situ hybridization data from the Allen Brain Atlas (Figures 1G, S1D, and S1E) (Lein et al., 2007). AD and AV share progenitor origin (Shi et al., 2017), expression profile of neurotransmitter receptors (Phillips et al., 2019), and afferent and efferent projections (Jankowski et al., 2013). Therefore, we treated the two as a single nucleus, AD/AV, in our investigation. AD/AV receives dense projections from the retrosplenial cortex, the subiculum, and the MB via the mammillothalamic tract (MTT) (Wright et al., 2010) (Figures 1H and 1I). By directly stimulating each afferent, we could selectively record evoked postsynaptic responses from individual inputs (see STAR Methods and Figure 1I) (Petrof and Sherman, 2009; Oh et al., 2014). Analogous to the sensory thalamic relay circuits (Sherman and Guillery, 2004), the inputs to AD/AV neurons are classified into driver and modulator inputs. Specifically, MB inputs drive the firing of AD/AV neurons, whereas cortical and subiculum inputs modulate the action of the MB inputs (Petrof and Sherman, 2009).

Input-Specific Modulation of Short-Term Plasticity by GSG1L

Inputs to AD/AV neurons undergo varying degrees of short-term plasticity upon repetitive stimulation (Petrof and Sherman, 2009).

Figure 1. GSG1L Input Specifically Regulates Short-Term Plasticity at AD/AV

(A) Ribbon diagram of GluA2 (red) with or without GSG1L (blue) (PDB: 5WEK) (Twomey et al., 2017a). Schematic summarizing the effects of GSG1L, emphasized with black arrows (right). AMPAR+GSG1L (blue lines) exhibits decreased amplitude and slower recovery from desensitization compared with AMPAR alone (red lines).

(B–E) Recordings from outside-out patches. Representative averaged traces for (B) GluA2iQ (i.e., flip/Q pore) alone and (C) GluA2iQ+GSG1L. OTR, open tip response. (D) Ratio of each pulse over the first pulse (Ix/I1, where x = 2, 3, and 4). A2iQ (n = 5), A2iQ+GSG1L (n = 4) (p < 0.001 for pulses 2–4, two-way ANOVA).

(F) GSG1L KO rat brain coronal sections at P21 (scale bar, 1,000 μm). The lacZ expression is represented by dark blue stain (arrowheads show AD/AV).

(G) In situ hybridization data from the Allen Brain Atlas. The lower signal intensity in AV reflects lower cell density in AV compared with AD (Morel et al., 1997); see also Figure S1E.

(H) Inputs received by AD/AV.

(i) Input-dependent stimulation paradigm of AD/AV (scale bar, 420 μm). Brain images were derived from Allen brain connectivity (scale bar, 1,000 μm) (Oh et al., 2014).

(J, M, and P) Schematic of mammillothalamic, M-T pathway (J), subiculum-thalamic, S-T pathway (M), and corticothalamic, C-T pathway (P).

(K) Pulse ratios of electrically evoked EPSCs from AD/AV neurons at a holding potential of −70 mV of GSG1L KO (P21–P33, n = 15 cells, N = 6 mice). Het (n = 13, N = 5), and WT (n = 20, N = 8), S1–S5 are amplitudes of each stimulus in a train of five stimuli (p = 0.9956, two-way ANOVA). S(n)/S1, where n = 2, …, 5, is the paired-pulse ratio of the 2nd–5th pulse divided by the 1st pulse.

(L, O, and R) Superimposed sample traces of whole-cell recordings of currents in AD/AV in response to stimulating M-T (L), S-T (O), or C-T (R) pathway.

(N) No changes in pulse ratios of GSG1L KO (P21–P33, n = 18, N = 6), Het (n = 20, N = 6), and WT (n = 18, N = 7) (p = 0.7288, two-way ANOVA) in response to subiculum stimulation at 20 Hz.

(Q) Pulse ratios are significantly different for GSG1L KO (P21–P33, n = 13, N = 6), Het (n = 14, N = 6), and WT (n = 19, N = 7) at 20 Hz (p < 0.0001, two-way ANOVA) in response to cortical stimulation. Post hoc Sidak comparisons: **p < 0.01, ***p < 0.001, ****p < 0.0001.
(legend on next page)
Given the high expression of GSG1L in AD/AV neurons, we wondered whether GSG1L is responsible for short-term plasticity. To address this, the extent of short-term plasticity in AD/AV neurons was compared between GSG1L KO and wild-type (WT) mice (Figures S1F–S1I). Sagittal sections were used to record from mammillothalamic (M-T) and subiculum-thalamic (S-T) synapses, whereas coronal sections were used to obtain corticothalamic (C-T) synapses (Figure 1I). We electrically stimulated the MB inputs (i.e., MT) at 20 and 50 Hz while whole-cell recording from AD/AV neurons (Figures 1J–1L and S2A–S2C). The degree of synaptic depression in the M-T synapses was quantified by taking amplitude ratios of the first stimulus (S1) and following four stimuli (S2–S5) responses. There was no statistical difference between GSG1L KO and WT control littermates (Figures 1J–1L and S2A–S2C). Similar results were obtained when inputs from the subiculum, the S-T synapses, were examined at 20 and 50 Hz (Figures 1M–1O and S2D–S2F). In contrast, stimulation of the C-T pathway of GSG1L KO, at both 20 and 50 Hz, resulted in enhanced synaptic facilitation, relative to WT control and heterozygous littermates (Figures 1P–1R and S2G–S2I). Similar findings were obtained when only AD synapses were examined, consistent with the close cellular pedigree of neurons in AD and AV (Shi et al., 2017) (Figures S2J–S2M).

Changes in the presynaptic glutamate release would affect both AMPARs and NMDA receptors (NMDARs). If short-term plasticity were mediated by a presynaptic mechanism, one would expect to find the effect of the genotype in both AMPAR- and NMDAR-mediated EPSCs. Therefore, to exclude the possibility of a presynaptic effect, we examined short-term plasticity of NMDAR-mediated EPSCs in the C-T synapses in response to 20 and 50 Hz stimulations (Figures S3A–S3E). We found no differences between GSG1L KO and control littermates, indicating that the effect is postsynaptic and AMPAR dependent. Input-specific stimulation of the C-T pathway unveiled the postsynaptic role of GSG1L, where its deletion is associated with enhanced synaptic short-term facilitation. Collectively, these results suggest that GSG1L input specifically regulates synaptic plasticity at AD/AV.

**AMPAR Desensitization Occurs at C-T Synapses of AD/AV Neurons**

If GSG1L regulates C-T synapses by altering the rate of recovery from desensitization of postsynaptic AMPARs, desensitization of AMPAR must be occurring during synaptic transmission. A simple way to test this is to examine sensitivity of evoked EPSCs to cyclothiazide (CTZ), a blocker of AMPAR desensitization. In GSG1L WT mice, whole-cell recordings from AD/AV neurons detected a significant increase in the degree of synaptic facilitation upon addition of 100 μM CTZ while stimulating the cortical inputs at 20 and 50 Hz in the presence picROTOX, a blocker of γ-amino-butyric acid (GABA) A receptors (Figures S3F–S3J). Importantly, CTZ sensitivity in C-T synapses requires GSG1L, because it is absent from GSG1L KO (Figures S3F–S3J). This substantiates our hypothesis that GSG1L affects short-term plasticity in these synapses through a postsynaptic mechanism of slowing AMPAR recovery from desensitization.

**GSG1L Modulates Basal Synaptic Transmission at C-T Synapses**

Another predicted signature of synapses regulated by GSG1L is the reduction of amplitude of AMPAR-mediated responses (Figure 2A) (McGee et al., 2015). Thus, we next investigated whether GSG1L also regulates basal AMPAR-mediated synaptic transmission by recording AMPAR-mediated miniature EPSCs (mEPSCs) in AD/AV neurons from GSG1L KO and WT mice (Figures 2A–2D). We found no significant differences in the average amplitude or the frequency of AMPAR-mediated mEPSCs. However, upon careful inspection, we found that there was a subset of GSG1L KO neurons with enhanced amplitude of mEPSCs (Figure 2C). This is in agreement with GSG1L functionally regulating a subset of synapses, specifically those formed by C-T inputs (Figure 1).

To directly test whether GSG1L regulates basal synaptic transmission at C-T synapses, we recorded asynchronous quantal EPSCs (qEPSCs) evoked in the presence of strontium (4 mM SrCl2) (Figures 2E–2L). Synaptic responses produced by isolated asynchronous quantal events represent postsynaptic AMPAR activity in the evoked pathway (Goda and Stevens, 2004).
Figure 3. GSG1L Is a Dominant Auxiliary Subunit at C-T Synapses at AD/AV

(A) Mean amplitude of AMPAR-mediated mEPSCs in AD/AV of γ-2 KO and γ-2 WT (p = 0.0006, Mann-Whitney U test). γ-2 KO (P15–P22, n = 24, N = 3); γ-2 WT (n = 24, N = 3).

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(open access)
We found that the amplitude of asynchronous qEPSCs is enhanced in GSG1L KO animals relative to WT control (Figures 2E–2H). The frequency of qEPSCs was also higher in GSG1L KO C-T synapses, most likely because of an increase in apparent detection sensitivity caused by an increase in the amplitudes. In addition, consistent with GSG1L’s role in modulating AMPAR gating kinetics (Schwenk et al., 2012; Shanks et al., 2012), the kinetics of C-T qEPSCs are accelerated in GSG1L KO. Although the difference in the rise time of qEPSCs was not significant (p = 0.0900), the decay time was significantly reduced in GSG1L KO (Figures 2I–2L).

Consistent with GSG1L having an input-specific function at AD/AV, there were no significant differences in the amplitude or frequency of asynchronous qEPSCs between GSG1L KO and WT in M-T synapses (Figures 2M–2O) or S-T synapses (Figures 2P–2R). The kinetics of qEPSCs in these two pathways were unaltered between GSG1L KO and WT (Figures S3K–S3P).

**Functional Co-expression of GSG1L and Stargazin in AD/AV Neurons**

Using the results of the Allen Brain Atlas as a reference, we inferred that stargazin is the dominant type I TARP expressed in AD/AV neurons (Figures S4A–S4E). A previous study has shown that stargazin outcompetes GSG1L in AMPAR gating modulation in Xenopus oocytes (Schwenk et al., 2012). To test the hypothesis that stargazin and GSG1L both modulate AMPARs at synapses in AD/AV neurons, we investigated the effect of deleting stargazin using a γ-2 KO mouse generated in our lab (Figures S4F–S4J).

In stargazer mice (i.e., spontaneously occurred γ-2 KO mice), there is complete loss of functional synaptic AMPARs in cerebellar granule cells (Hashimoto et al., 1999; Chen et al., 2000). In contrast, we found that in AD/AV neurons, there is residual AMPAR activity in γ-2 KO mice. Consistent with stargazin being a positive AMPAR regulator, both the amplitude and the frequency of AMPAR-mediated mEPSCs are lower in γ-2 KO relative to WT control (Figures 3A–3D). The residual mEPSCs indicate the existence of a subset of synapses with functional AMPARs. We predicted that these are C-T synapses and did not have stargazin regulation instead are controlled by GSG1L.

We found no significant differences in the amplitude or the frequency of asynchronous qEPSCs between AD/AV neurons of GSG1L KO mice than in WT control, but with no current injection. We found that 60% of GSG1L KO AD/AV neurons spiked during a 3-min recording period, whereas only 15% spiked in WT neurons (Figures 4A–4C). There was no change in intrinsic excitability of neurons in GSG1L KO, because no differences were observed in the resting membrane potential, rheobase, and neuronal firing rate between GSG1L KO and WT control (Figures S5A–S5D). However, the input resistance is significantly reduced in GSG1L KO mice (Figure S5E), which is consistent with increased synaptic activity, although it may not be the sole cause of the effect. In addition, we found that spontaneous excitatory synaptic transmission is significantly enhanced in GSG1L KO mice (Figures 4D–4F), as well as in GSG1L KO mice (Figures S5F–S5I). Both the amplitude and the frequency of AMPAR-mediated spontaneous EPSCs (sEPSCs) are increased in GSG1L KO neurons. Altogether, neurons in AD/AV neurons of GSG1L KO mice are hyperactive overall.
Figure 4. Loss of GSG1L Results in Enhanced Excitability and Seizure Susceptibility

(A) Bar graph summarizing spontaneously firing AD/AV neurons in GSG1L KO (20/33 cells, N = 5) and WT control (5/33 cells, N = 5).

(B) Representative traces of two spiking GSG1L KO AD/AV neurons.

(C) Bar graph summarizing firing frequency distribution. Blue and red data points represent left and right traces in (B), respectively.

(D and E) Mean amplitude (p = 0.0037, Mann-Whitney U test) and frequency (p = 0.0084, Mann-Whitney U test) of AMPAR-mediated sEPSCs in AD/AV neurons of GSG1L KO (P15–P30, n = 17, N = 4) and WT (n = 18, N = 4).

(F) Representative sEPSC traces of GSG1L KO and WT.

(G–J) Time course of seizure severity following i.p. injection of 15 mg/kg (n = 13 each, 9–10 months) (G) or 25 mg/kg (n = 8 each) (I) of kainate. A higher score corresponds to a severer seizure status, with 7 denoting death. (H and J) Mortality rate during the 2-h recording period at 15 mg/kg (H) and 25 mg/kg (J) injection. Two-way ANOVA, p < 0.0001 for both doses. Post hoc Sidak comparisons: *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001.

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Next, we aimed to evaluate the effect of the loss of GSG1L at the organism level. Initial basic behavioral phenotyping of GSG1L KO mice revealed no differences in grip strength, rotarod performance, elevated zero maze, and gait parameters (Figures S5J–S5M). However, we found that GSG1L KO mice show deficits in a novel object recognition (NOR) task (Figure S5N), similar to the findings of previous reports using GSG1L KO rats (Gu et al., 2016). These results are consistent with the known role of AT in learning and memory (Parker and Gaffan, 1997), but a more drastic behavioral phenotype was revealed when the animals were challenged with seizure, a condition in which AT plays a critical role (Mirski and Ferrendelli, 1984; Bittencourt et al., 2010).

We found that GSG1L KO mice exhibit enhanced susceptibility to kainate-induced seizures (Figures 4G–4J). The seizure severity was determined on a scale of 0–7, a rating system with increasing severity in an ascending scale and with a score 7 denoting death (Morrison et al., 1996). At a lower-dose injection (15 mg/kg intraperitoneal [i.p.]) of kainate, GSG1L KO mice had a significantly higher seizure severity score than WT controls. Furthermore, 23% (8 of 13 mice) of GSG1L KO mice, and no GSG1L WT mice, died within 2 h after kainate administration. At a higher-dose injection (25 mg/kg i.p.) of kainate, 75% of GSG1L KO mice died within 2 h postinjection and only 25% of WT mice. These results highlight the in vivo importance of GSG1L in protecting against kainate-induced neurotoxicity and pathological hyperexcitability. Consistent with GSG1L being a negative regulator of AMPAR function, GSG1L KO mice have enhanced excitatory neurotransmission and susceptibility to seizures, along with increased spontaneous firing of AD/AV neurons.

**DISCUSSION**

AD/AV, where GSG1L is abundantly expressed, is at the core of the extended hippocampal-diencephalic network with crucial roles in memory (Aggleton and Brown, 1999); thus, AD/AV damage is associated with anterograde amnesia (Shikha-Schmid and Bogousslavsky, 2000; Gold and Squire, 2006). AD/AV contains neurons that sense head direction during spatial navigation (Clark and Taube, 2012) and have been implicated in seizure initiation and/or propagation (Mirski and Ferrendelli, 1984; Kerrigan et al., 2004). In contrast to the extensively studied lateral geniculate nucleus (LGN), molecular and functional characterization of the synapses of AD/AV is largely lacking.

We found that GSG1L and stargazin are functionally co-expressed in AD/AV. Among the synapses in AD/AV neurons formed by three distinct input projections, C-T synapses are controlled by GSG1L, but not by stargazin (Figures 1, 2, and 3). In C-T synapses, GSG1L reduces the amplitude and slows the kinetics of AMPAR-mediated qEPSCs and suppresses short-term facilitation by slowing recovery from desensitization.

GSG1L promotes surface expression of AMPARs in HEK cells and hippocampal neurons (Shanks et al., 2012; Gu et al., 2016), and such a mechanism may partially contribute to the properties of C-T synapses. However, we suggest that the effect of GSG1L on gating modulation plays a dominant role in regulating short-term plasticity given the CTZ sensitivity of these synapses in WT, but not in GSG1L KO (Figures S3F–S3J).

In contrast to GSG1L, stargazin plays a major role in M-T or S-T synapses (Figure 4K). The mechanism that restricts the functions of GSG1L and stargazin to specific afferent synapses remains to be determined. We speculate that presynaptic terminals may provide some trans-synaptic molecular cues, considering that GSG1L lacks a PDZ (PSD-95, dlg, ZO-1) domain binding motif in the cytoplasmic C terminus as it would normally facilitate synaptic anchoring.

In *Xenopus* oocytes, stargazin outcompetes GSG1L in AMPAR modulation (Schwenk et al., 2012), which agrees with GSG1L and stargazin sharing identical binding sites on AMPARs (Twomey et al., 2016, 2017b). In AD/AV, we also found that under basal conditions, stargazin outcompetes GSG1L, functionally, in most synapses (Figures 3H–3N). The effect of GSG1L is unmasked in GSG1L/γ-2 dKO mice. These observations also indicate that in AD/AV, the C-T synapses are the smaller population compared with the combined sets of M-T and S-T synapses.

Similar to GSG1L, CKAMP44 slows recovery from desensitization, modulates short-term plasticity, and is expressed in AD/AV (Figure S4E) (von Engelhardt et al., 2010; Chen et al., 2018). We found that M-T synapses in AD/AV undergo pronounced STD (Figures 1J–1L and S3A–S3C), which is highly sensitive to CTZ (data not shown). Given that GSG1L does not modulate AMPARs at M-T synapses, we speculate that CKAMP44 could be postsynaptically functional in these synapses, which has yet to be investigated. Again, input-specific functional expression of an AMPAR auxiliary subunit in the thalamus may extend to the case of CKAMP44.

Consistent with GSG1L being a negative regulator of AMPAR function, deletion of GSG1L results in hyperexcitability in the AT (Figure 4), with its impact observed in the downstream cortical neurons (data not shown). Moreover, GSG1L KO mice have enhanced susceptibility to kainate-induced seizures. We suggest that disturbed excitation/inhibition (E/I) balance in GSG1L KO is associated with seizure states (Bateup et al., 2013; Paz and Huguenard, 2015; Staley, 2015). Circumstantial evidence suggests GSG1L may serve a neuroprotective role against hyperexcitability and ischemia (Keum and Marchuk, 2009; Du et al., 2015). GSG1L was identified as one of the candidate genes at the locus that determines the extent of infarct volume in mouse models of focal cerebral ischemia (Keum and Marchuk, 2009). Furthermore, a suggestive single-nucleotide polymorphism in GSG1L was found to be associated with infarct volume upon ischemic stroke in mice and human patients (Du et al., 2015). However, these are correlative studies, and further investigation is warranted.
is needed to test the direct link between GSG1L and ischemia.

Previous investigations on the synaptic phenotypes of GSG1L KO rats in CA1 and DG (dentate gyrus) neurons (Gu et al., 2016; Mao et al., 2017) were both conducted at time points when the expression of the lacZ reporter for GSG1L promoter activity is low (Figures S1C and S1D). Specifically, GSG1L KO rat acute slices at postnatal day (P) 13–P19 were used. Although GSG1L protein is detectable in a subset of dendritic spines in older neurons (Schwenk et al., 2012; Shanko et al., 2012; Willems et al., 2020), the overall expression of GSG1L may be low in the hippocampus compared with the AT at young age (i.e., P13–P19). In agreement with such a hypothesis, a previous study on GSG1L KO rat reported a substantial contribution of another auxiliary subunit, CNII2, superimposed on the functional effect of GSG1L (Gu et al., 2016). However, because the lacZ reporter is an indirect indicator of GSG1L protein expression, future studies using more direct methods are necessary to determine whether its expression is low in young hippocampus.

AMPAR desensitization is critical for normal brain function (Christie et al., 2010). CKAMP44 regulates AMPAR desensitization and contributes to the underlying mechanism of STD in retinogeniculate synapses (Chen et al., 2018, 2000; Kielland and Heggelund, 2002; Budisantoso et al., 2012). The glomerular structure of the retinogeniculate synapse has large terminals and closely spaced release sites (Rafols and Valverde, 1973). The unique overall geometry prevents fast removal of glutamate and allows spillover, resulting in AMPAR desensitization (Budisantoso et al., 2012). However, the corticogeniculate synapses have single distant synaptic contacts (Erikir et al., 1997; Narushima et al., 2016) with low release probability (Granseth et al., 2002). In these synapses, AMPAR desensitization is predicted to not contribute to short-term plasticity, such as observed in the ventrobasal nucleus (Sun and Beierlein, 2011). CKAMP44 plays no role in short-term plasticity at corticogeniculate synapses in the LGN (Chen et al., 2018). In contrast, we found that AMPAR desensitization substantially contributes to short-term plasticity at C-T synapses in AD/AV. GSG1L is highly expressed in AD/AV (Figures 1 and S1), but not in the LGN or ventrobasal nuclei of the thalamus (Lein et al., 2007). Hence, it is uniquely positioned to regulate short-term plasticity in C-T synapses in AD/AV, despite its predicted ultrastructure and low release probability.

**STAR + METHODS**

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

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**SUPPLEMENTAL INFORMATION**

Supplemental information can be found online at https://doi.org/10.1016/j.celrep.2020.107921.

**ACKNOWLEDGMENTS**

Behavioral testing was performed at the Vanderbilt Mouse Neurobehavioral Laboratory, which is supported by the EKS NICHD of the NIH under award U54HD083211. We acknowledge the use of Vanderbilt Cell Imaging Shared Resource and Vanderbilt Transgenic Core (the current Vanderbilt Genome Editing Resource). We thank Elena Zaika for technical support. We also thank Kent Hamra for providing GSG1L KO rat. We thank Bob Matthews and John Allison for discussion and technical support. We also thank Jake Watson, Alexander Pinggera, Akihiro Harada, Brian Shonesy, Fiona Harrison, Roger Colbran, Sachin Patel, and Ege Kavalali for discussion. This work was funded by Vanderbilt University (to T.N.), NARSAD Independent Investigator Grant (to T.N.), and NIH grants R01HD061543 (to T.N.) and R01NS085215 (to K.F.).

**AUTHOR CONTRIBUTIONS**

T.N. conceived and supervised the project. A.K., E.D., and T.N. designed and conducted experiments. A.K. conducted slice physiology, histology, and behavioral experiments. T.N. conducted outside-out patch recordings. All electrophysiology experiments were conducted at the Nakagawa lab. E.D. provided expertise to generate GSG1L and -2 KO mouse lines. T.N. and E.D. generated the targeting construct for GSG1L KO mouse. K.F. provided intellectual and technical expertise in electrophysiology. All authors interpreted the results and analyzed data. T.N. and A.K. wrote the manuscript. All authors edited the manuscript. All authors read and commented on the final manuscript.

**DECLARATION OF INTERESTS**

The authors declare no competing interests.
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## STAR Methods

### Key Resources Table

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies** | | |
| Anti-GSG1L Ct-1 antibody | (Shanks et al., 2012) PMID: 22813734 |
| Anti-GSG1L Ct-2 antibody | (Shanks et al., 2012) PMID: 22813734 |
| Anti-GluA2CT antibody | (Nakagawa et al., 2005) PMID: 15690046 |
| Pan-TARP antibody | (Nakagawa et al., 2005) PMID: 15690046 |
| Anti-Stargazin antibody | Millipore AB9876; RRID: AB_877307 |
| **Chemicals, Peptides, and Recombinant Proteins** | | |
| PicROTOXIN | Tocris | 1128 |
| TETRODOTOXIN CITRATE | Abcam | Ab120055 |
| CYCLOTHIAZIDE | Tocris | 0713 |
| KAINIC ACID MONOHYDRATE | Sigma | K0250 |
| NBQX | Tocris | 1044 |
| **Experimental Models: Organisms/Strains, Cell lines** | | |
| Mouse: GSG1L / / C0 | This paper | N/A |
| Mouse: CACNG2 / / C0 | This paper | N/A |
| Mouse: GSG1L / / C0 / / CACNG2 / / C0 | This paper | N/A |
| Rat: GSG1L / / (Izsvák et al., 2010) PMID: 21193047 | | |
| TetON HEK cells | Clontech | 631182 |
| **Oligonucleotides** | | |
| Genotyping Primer GSG1L, mouse, left: aacagacaacatggcccctcagacct | Sigma | N/A |
| Genotyping Primer GSG1L, mouse right: gacctgtcccatcttagtatactccc | Sigma | N/A |
| Genotyping Primer GSG1L, rat, left: acgtgtgttagtgacctccaagc | Sigma | N/A |
| Genotyping Primer GSG1L, rat, right: tgcacgcatactaatga | Sigma | N/A |
| Genotyping Primer CACNG2, mouse, left: atggttgtgagaattcggctgtacc | Sigma | N/A |
| Genotyping Primer CACNG2, mouse, right: aaccctaaggactccaaagcatcc | Sigma | N/A |
| Recombinant DNA | | |
| DualpTREt-GluA2i(Q)+GSG1L | This paper | N/A |
| GluA2i(Q) | (Hawken et al., 2017) | N/A |
| **Software and Algorithms** | | |
| Prism 6.0 | Graphpad Software | [https://www.graphpad.com/scientific-software/prism/](https://www.graphpad.com/scientific-software/prism/) |
| PCLAMP10 | Molecular Devices | [https://www.moleculardevices.com/products/axon-patch-clamp-system/acquisition-and-analysis-software/pclamp-software-suite](https://www.moleculardevices.com/products/axon-patch-clamp-system/acquisition-and-analysis-software/pclamp-software-suite) |
| IGOR Pro | WaveMetrics | [https://www.wavemetrics.com/products/igorpro](https://www.wavemetrics.com/products/igorpro) |
| Neuromatics | WaveMetrics | [http://www.neuromatic.thinkrandom.com/index.html](http://www.neuromatic.thinkrandom.com/index.html) |
| LabView | National Instruments | [https://www.ni.com/en-us/shop/labview/labview-details.html](https://www.ni.com/en-us/shop/labview/labview-details.html) |
| Adobe Illustrator 2020 | Adobe | [https://www.adobe.com/products/illustrator.html](https://www.adobe.com/products/illustrator.html) |
RESOURCE AVAILABILITY

Lead contact
Further information and requests for resources and reagents should be directed to the Lead Contact, Terunaga Nakagawa (terunaga.nakagawa@vanderbilt.edu).

Materials availability
Requests that are related to mouse lines, should be directed to the lead contact.

Data and code availability
The data generated during this study are available from the lead contact upon reasonable request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

All animal procedures were approved by the Vanderbilt University Animal Care and Use Committee and were in agreement with the NIH and Vanderbilt University guidelines for the care and use of laboratory animals. Mature mice and rats were housed with same-sex litters and maintained on a 12 hr light/dark cycle under controlled temperature and humidity. Food and water were provided ad libitum.

A GSG1L knockout (KO) transgenic mouse line was generated by targeting exon 4 of GSG1L gene in 129/SvEvTac embryonic stem (ES) cells. Two independent clones were shown by Southern blot analysis and PCR to have undergone proper homologous recombination and germline transmission was confirmed from chimeras derived from clone 2H2. To remove the pGK-neo cassette, the 3-loxP mouse was crossed with FlpE line (Rodríguez et al., 2000) that ubiquitously expresses Flp recombinase, generating a 2-loxP animal. Finally, by crossing with Ella-cre mice (JAX: Stock 003724, which was backcrossed with C57BL6/J over 10 generations) that expresses cre-recombinase in the one cell zygote, we obtained the 1-loxP heterozygotes. All strains were backcrossed with C57BL6/J mice for 8 generations. All KO animals were viable with no overt issues with breeding and/or development.

The γ-2 KO mouse was generated as follows: A 20 bp sequence (TGAAACCAGCAAGAAGAACG, boxed in Figure S4) followed by AGG as proto-spacer adjacent motif (PAM), was selected within exon 1 of the CACNG2 (encoding γ-2) mouse gene to create a target-specific guide RNA molecule. The sequence flanked by BbsI sites was ligated in pX330, a vector expressing the guide RNA under a strong U6 promoter and cas9 under a hybrid chicken beta-actin (Cbh) promoter. The vector was injected alongside a 195 base repair single stranded oligonucleotide into 314 mouse embryos. The repair oligo contained 90 bp homology arms, a codon substituting lysine residue 53 to a stop codon, a unique AgeI restriction site, and a few additional third base mutations to prevent targeting of cas9 to the repaired DNA (Figure S4). Out of 314 embryos injected, 267 were transferred to 11 pseudo-pregnant females to generate 16 pups. At weaning, genotyping was done by amplifying a 926 bp fragment followed by sequencing. Out of 15 pups (one died prior to weaning), only one animal (female #12) carried the designed mutation. After several backcrossing to C57BL6/J mice to eliminate any possible off target events, heterozygote mice were crossed to generate homozygous KO animals.

A transgenic GSG1L KO rat line in the background of SD strain was among the mutants generated through transposon-based mutagenesis by Kent Hamra at UT Southwestern (Izsvák et al., 2010). The mutant rat line was purchased and maintained in house.

For each experiment, both male and female animals were used. Each cohort contained roughly equal number of each sex. For short-term plasticity experiments comparing the three pathways, GSG1L KO and WT mice P21-P33 were used. For short-term plasticity experiments of AD only, AD/AV combined, NMDAR-dependent, and CTZ recordings were done at P15-P30. For mEPSCs, sEPSCs, qEPSCs and current clamp recordings of GSG1L KO and WT mice at P15-P30 were used. For mEPSCs and sEPSCs of γ-2 KO and WT (and GSG1L/γ-2 dKO) mice at P15-P22 were used. For behavioral experiments, GSG1L KO and WT mice at 9-10 month old were used. LacZ staining of GSG1L KO rat and WT control littermates was done as indicated in the figures: P14, P21, P60, and P180.

Genotyping by PCR
PCR amplification was conducted with the following primers:

- GSG1L KO mice: GSG1L left: aaacagacaacatggccctcagactc; GSG1L right: gacttgtcccatctctaggatacctccc.
- GSG1L KO rats: Gsg1L left: acgttgtagtgaccccaagc; GSG1L right: tgcacgcatactacaatga; SFB2: tcatcaaggaaaccctggac
- γ-2 KO mice: CAC9: atgtgttgtgagaattcgggtgtgc; CAC10: aacccataaggactccaaagcatcc

METHOD DETAILS

Electrophysiology in acute slices
For all electrophysiology experiments that were done in the AT, mice of both sexes aged were used. Using the stria medullaris (sm) and the hippocampus as anatomical landmarks, we unambiguously identified AD/AV in 300 μm-thick coronal sections. For short-term plasticity experiments, the C-T synapse was isolated in a coronal slice configuration and the cortical inputs were stimulated...
outside of AD/AV (as illustrated in Figures 1H and 1I). M-T synapses were isolated in 300 μm-thick sagittal brain sections. The prominent MTT and the hippocampus were used as anatomical landmarks to identify the AD/AV. To stimulate this synapse, the stimulating electrode was placed directly on the MTT. The S-T synapse was isolated in 300 μm-thick sagittal brain sections. The MTT and the hippocampus were used as landmarks to isolate the AD/AV and the electric stimulator was placed on the subiculum inputs directly below the hippocampus.

Mice were decapitated and the brains were rapidly removed and placed in ice-cold N-methyl-D-glucamine (NMDG) cutting solution containing 92 mM NMDG, 2.5 mM KCl, 30 mM NaHCO₃, 1.25 mM NaH₂PO₄, 11 mM Glucose, 0.5 mM CaCl₂, 10 mM MgCl₂, 20 mM HEPES, 2 mM thiourea, 5 mM Na-ascorbate, 3 mM Na-pyruvate, with pH adjusted to 7.4 with HCl. Acute 300 μm-thick slices were prepared using a vibratome (Leica VT1200). Slices were then left to recover for 1 hr in artificial cerebro-spinal fluid (aCSF) solution containing 119 mM NaCl, 2.5 mM KCl, 26 mM NaHCO₃, 1 mM NaH₂PO₄, 11 mM Glucose, 1 mM MgCl₂, 2 mM CaCl₂, at pH 7.4 and 310 mOsM, saturated with 95%O₂/ 5% CO₂.

For AMPAR-mediated mEPSCs recordings, the aCSF was supplemented with tetrotodoxin (TTX, 0.5 μM) in order to suppress spontaneous excitation. To specifically isolate AMPAR-mediated mEPSCs at a holding potential of ~70 mV, GABA-A receptor activity was blocked with 100 μM picrotoxin (Tocris). For AMPAR-mediated sEPSCs recordings, the aCSF was supplemented with 100 μM picrotoxin to isolate the AMPAR component at ~70 mV. Pipettes were filled with the internal solution containing 115 mM Cs methanesulfonate, 20 mM CsCl, 10 mM HEPES, 2.5 mM MgCl₂, 4 mM Na₂ATP, 0.4 mM Na₃GTP, 10 mM Na phosphocreatine, and 0.6 mM EGTA. The osmolarity was adjusted to 295 mOsm and the pH was adjusted to 7.25 with CsOH. The patches were formed with borosilicate glass pipettes (O.D. 1.5mm, I.D. 0.86mm, Sutter) (3-5 MΩ tip resistance). Recordings were made using Multiclamp 700B Amplifier (Axon Instruments) operated by pCLAMP10 software, low-pass filtered at 10 kHz and digitized at 20 kHz using Digidata1440A (Axon Instruments). Obtained mEPSCs and sEPSCs events were detected and analyzed with NeuroMatic event detection software (IGOR-pro, WaveMetrics) and correct event detection was confirmed by manual visual inspection, while errors were corrected by manual inspection of the traces. All the recordings and analysis were done in a blind manner, with the experimenter blind to the genotype during data acquisition and analysis. Significance was determined with Mann-Whitney U-test or a two-way analysis of variance (Two-Way ANOVA) with Sidak’s multiple comparison post hoc test when applicable.

To evoke asynchronous quantal EPSCs (qEPSCs), Ca²⁺ was replaced with Sr²⁺ (4 mM SrCl₂). In order to avoid the multiquantal events, we only analyzed the qEPSCs that occurred >10 ms following the stimulation with the total sweep duration of 360 ms. Obtained qEPSC events were detected and analyzed with NeuroMatic event detection software (an extension to the IGOR-pro, WaveMetrics) and correct event detection was confirmed, while errors were corrected by manual inspection of the traces. The kinetic analysis of qEPSCs and mEPSCs were also done using NeuroMatic and IGOR-pro. From each corresponding cell, all isolated events with flat baseline were used for analysis. Typically, from each cell 15-30 representative events with monotonic rise and distinct decay were selected and averaged. The decay time of averaged qEPSCs was determined by fitting to double exponential function. The weighted decay time constant was further calculated as sum of slow and fast time constants and weighted by fractional amplitudes (Hawken et al., 2017).

For short-term plasticity experiments, whole-cell recordings AD/AV neurons were achieved in response to electrical stimulation of MTT, subiculum, or the cortical inputs at a holding potential of ~70 mV. For M-T recordings, 300 μm-thick acute sagittal slices with AD/AV and visible MTT were obtained. The MTT was electrically stimulated, and current changes were recorded in response to 5-pulse stimulation at 20 Hz and 50 Hz. For S-T recordings, 300 μm-thick acute sagittal slices with AD/AV were obtained. Similarly, the subiculum inputs (as depicted in Figure 2) were stimulated at 20 Hz and 50 Hz. For C-T recordings, 300 μm-thick acute coronal slices with AD/AV were obtained and the cortical inputs (as depicted in Figure 2) were electrically stimulated at 20 Hz and 50 Hz. Slices were perfused with aCSF supplemented with 100 μM picrotoxin. To isolate the effect of CTZ on short-term plasticity, we bath applied aCSF supplemented with 100 μM CTZ (Tocris). NMDAR currents were recorded as aforementioned, but at a holding potential of +40 mV and in the presence of 10 μM NBQX and 100 μM picrotoxin. Statistical significance was assessed using a two-way analysis of variance (Two-Way ANOVA) with Sidak’s multiple comparison post hoc test.

Intrinsic excitability properties were analyzed in current clamp recording mode. The recordings were done using potassium-based internal solution containing: 115mM K methanesulfonate, 20 mM KCl, 10 mM HEPES, 2.5 mM MgCl₂, 4 mM Na₂ATP, 0.4 mM Na₃GTP, 10 mM Na phosphocreatine, 0.6 mM EGTA. The osmolarity was adjusted to 295 mOsm and the pH was adjusted to pH 7.25 with KOH. In order to measure the intrinsic excitability properties, the neurons were held at ~70 mV and current was injected in 10 pA increments (from -20 to 160 pA) for 200 ms. The rheobase was determined as a minimal current injection required to elicit an action potential firing. The firing rate was determined as the number of action potentials fired at 150 pA injection. The input resistance was determined by calculating the slope of current-voltage plot (~20 to 20 pA). The firing frequency of spiking GSG1L KO neurons was determined, by calculating the frequency of fired action potentials during a 50 msec recording period.

**Fast glutamate application from outside out patches**

TetON HEK cells were plated on a coversglass coated with poly-D-lysine (37.5 μg/ml in H2O) for 15 min. Excess coating material was removed by washing in D-PBS three times. Cells were plated and incubated on a coversglass until they were adherent (typically within 12 hr) and transfected with a plasmid (DualpTREt-GluA2i(Q)+GSG1L) that DOX dependently express GluA2flip(Q) and GSG1L. After transfection, 30 μM NBQX and 5 μg/ml doxycycline (DOX) were added. Cells were used for recording 24–36 hr after induction. Ligand (1mM glutamate) was applied via theta tubing glass capillary mounted on a piezo actuator (P-830.30, Physik Instrumente) controlled...
by an LVPZT amplifier (E-505, Physik Instrumente), DAQ device (NI USB-6221, National Instruments), and LabView software (National Instruments). Recording was done using a single channel of a Multiclamp700B Amplifier (Axon Instruments) operated by pCLAMP10 software. Signals were digitized using Digidata1440A (Axon Instruments) at a sampling rate of 50 kHz and low pass filtered at 2kHz. Borosilicate glass capillaries (O.D. 1.5 mm, I.D. 0.86 mm, Sutter) were pulled to manufacture electrodes with pipette resistances of 4-5 MΩ. Internal solution was (in mM) 110 NaCl, 10 NaF, 5 EGTA, 1 CaCl₂, 1 MgCl₂, 10 Na₂ATP, 5 HEPES, adjusted to pH 7.3 with CsOH and 295 mOsm. External solution was (in mM) 145 NaCl, 2.5 KCl, 1.8 CaCl₂, 1 MgCl₂, 5 HEPES, 10 glucose, adjusted to pH 7.3 with NaOH and 301 mOsm. Standard solution without ligand was the external solution. The ligand solution contained 1mM glutamate in external solution, supplemented with 2 mM glucose and 3 mM NaCl to facilitate the visualization of the interface of the two solutions and recording liquid junction potential after breaking the patch. A protocol for four 1ms glutamate pulses at 20Hz was programmed using LabView. The 20%–80% rise time of liquid junction potential, measured after breaking the patch, was ~300 μs.

BEHAVIORAL EXPERIMENTS

Seizure behavior
To investigate seizure susceptibility behavior, GSG1L KO and WT littermates of both sexes (9-10 month old) were administered an intraperitoneal (i.p.) injection of kainic acid monohydrate (Sigma) dissolved in 0.9% saline solution at 15 mg/kg or 25 mg/kg body weight. Seizure susceptibility was determined immediately following the kainate injections. Seizure activity was video recorded for 2 hr and scored for severity using a previously described seizure severity scale in a blind manner, with the experimenter blind to the genotype during scoring (Morrison et al., 1996; Wu et al., 2005; Bateup et al., 2013). Seizure severity was assessed at a five-minute interval.

The severity of seizures was assigned based on the following chart:

0, no abnormality;
1- immobility, cessation of normal behavior;
2- rigid posture with extended tail or forelimbs;
3- repetitive behaviors including head nodding, head bobbing, twitching, or scratching;
4- Forelimb clonus with partial or intermittent rearing;
5- continuous forelimb clonus/rearing or repeated rearing and falling;
6- loss of posture, generalized tonic-clonic whole body convulsions or hyperactivity/jumping behavior;
7- mortality.

Novel object recognition task
GSG1L KO and WT littermates of both sexes (9-10 month old) were habituated for 10 min in an empty novel object recognition arena (40 × 64 × 33 cm³) on day 1. The following day, the mice were placed in the novel object recognition arena that contained two identical objects and were allowed to explore the space for 10 min. Following the habituation phase, mice were placed back into their homecage for 1 hr. During probe phase, one of the familiar (previously exposed) objects was replaced with a novel object. The exploration behavior was recorded for 10 min. Time spent exploring familiar and novel objects was scored blindly and the recognition index was identified as (time exploring novel object) - (time exploring familiar object) / total time exploring both objects.

Rotarod
Motor coordination and balance were measured by placing the mice on an accelerating rotating rod (Ugo Basile model 7650; Stoelting Co., Wood Dale, IL, USA). The initial rotations per minute (rpm) were set at 4 rpm, which gradually increased to 40 rpm. The performance of each animal was recorded over the course of 300 s. The time taken for a mouse to fall off the rod was recorded by an observer blinded to the mouse genotype. The test was conducted over the course of three days with three independent trials per day.

Elevated Zero Maze
Anxiety was measured using elevated zero maze (San Diego Instruments, CA). The device is an elevated ring-shaped platform with four equal chambers (two open arms and two closed arms). Light levels in the closed arms were at approximately 100 lux and 200 lux in open arms. Mice were placed in the center of an open arm and allowed to explore the maze for 5 min. The activity was video-recorded and analyzed using ANY-maze software (Stoelting, Wood Dale, Illinois, USA).

Grip strength
Grip strength was measured using a Grip Strength Test Apparatus (San Diego Instruments, CA). The grip strength was measured by a digital gauge over 7 independent trials. The average grip strength was calculated as an average of 7 trials (in Newtons).
Histology
GSG1L KO (P14, P21, P60, P180) rat brains with corresponding WT rat animals were anesthetized with sodium pentobarbital (Nembutal). Each animal was then perfused with normal rat Ringer solution (pH 7.4) for 2 min to ensure complete perfusion, followed by 4% paraformaldehyde in 0.1M phosphate (pH 7.4) for 4 min. Obtained brains were further fixed for 15 min. Fixed brains were subsequently sectioned to generate 300 μm-thick coronal sections using a vibratome (Leica VT 1200). The sections were then permeabilized with 0.01% Sodium deoxycholate and 0.02% Triton X-100 in PBS buffer for 2 hr. Following permeabilization, 300 μm-thick coronal sections were incubated with X-gal staining solution containing: 5 mM K₃[Fe(CN)₆], 5 mM K₄[Fe(CN)₆], 2 mM MgCl₂, 0.02% Triton X-100, and 0.1% X-gal in PBS buffer at 37°C for 8 hr in the dark. Stained brains were imaged (MULTIZOOM AZ100M, Nikon microscope) following day (1 day post staining).

Confirming the absence of targeted proteins in the KO animals
Immunoprecipitation of protein complexes were performed as previously described (Nakagawa et al., 2005). Briefly, mouse (or rat) brains were homogenized in 20 mM HEPES pH 7.4, 320 mM sucrose, 5 mM EDTA, 5 mM EGTA, 30 mM NBQX supplemented with protease inhibitors 1 mM PMSF, 10 μg/ml aprotinin, 10 μg/ml leupeptin, 1 μg/ml pepstatin, and 500 μM benzamidine. Brain homogenates were spun down at 3000 g for 15 min and the obtained supernatant was further spun at 38,400 g for 15 min resulting in a membrane pellet. The membrane pellet was resuspended in a buffer containing 20 mM HEPES pH 7.4, 1 M KI, 5 mM EDTA, 5 mM EGTA, and 30 μM NBQX. To wash off the KI, membranes were washed with wash buffer containing 20 mM HEPES pH 7.4, 5 mM EDTA, 5 mM EGTA, and 30 μM NBQX. The final membrane pellet was solubilized in a resuspension buffer containing 20 mM HEPES pH 7.4, 100 mM NaCl, 5 mM EDTA, 5 mM EGTA, 1% CHAPS, 30 μM NBQX and the protease inhibitors for two hours at 4°C. To validate GSG1L 3-loxP KO, GSG1L was immunoprecipitated with previously described anti-GSG1L Ct2 antibody (2mg/ml) and eluted with the epitope peptide CKVFEQGYREEPTFIDPEAIKYFR (Shanks et al., 2012). To validate GSG1L 1-loxP KO, GSG1L was immunoprecipitated with anti-GSG1L Ct1 antibody (2mg/ml) and eluted with the epitope peptide CRSSAHEAAELNRQCWVLGHWV (Shanks et al., 2012). Immunoprecipitated GSG1L from the brain lysates of 3-loxP KO and 1-loxP KO GSG1L was then probed in western blots with anti-GSG1L Ct1 antibody and Ct2 antibody, respectively. Of note, long and short isoforms of GSG1L transcripts are listed in the mouse genome database. The short isoform is predicted to lack the Ct2 epitope, and thus we were only able to confirm the KO of 43kDa long isoform of GSG1L. The short isoform lacks the TM4, which is essential for binding AMPAR from the cryo-EM structure (Twomey et al., 2017b), and unlikely to have a role in regulating AMPAR. The folding of TM1-TM4 is highly conserved among the tetraspanins and therefore loss of TM4 is likely to cause a substantial misfolding in the short isoform. Specifically, if the short isoform exists, it would have an extracellular C-terminal, whose topology is completely different from the long isoform with a cytoplasmic C-terminal. Moreover, our targeting design does not permit efficient transcription of the shorter isoform given that exon 4, upstream of the putative splice site, is targeted. Collectively, we suggest that the short isoform is unlikely to exist and only the long isoform is relevant in the current study. We also note that the observed differences in the non-specific band patterns in Figure S1I is a consequence of using different polyclonal antibodies for detection. Note that the band immediately below GSG1L in 1loxP KO blot is due to nonspecific binding (Figure S1I). For validating γ-2 KO, AMPAR complexes were pulled down with anti-GluA2 antibody (EGYNVYGIESVKI). Immunoprecipitated stargazin was probed with anti-stargazin antibody (Millipore, AB9876).

QUANTIFICATION AND STATISTICAL ANALYSIS
All statistical analyses were performed with Graphpad Prism. Data in the figures are represented as mean ± SEM. The information on statistical tests used, sample sizes (n) and animal subjects (N) are present in figure legends. Statistical significance for averaged data from multiple experiments was determined using Mann-Whitney U test or Two-Way Analysis of Variance (ANOVA) followed by Post hoc analysis. Error bars in all figures represent standard error of the mean. The cumulative probability difference was determined using Kolmogorov-Smirnov test.