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# Human genetic variants disrupt RGS14 nuclear shuttling and regulation of LTP in hippocampal neurons

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The human genome contains vast genetic diversity as naturally occurring coding variants, yet the impact of these variants on protein function and physiology is poorly understood. RGS14 is a multifunctional signaling protein that suppresses synaptic plasticity in dendritic spines of hippocampal neurons. RGS14 also is a nucleocytoplasmic shuttling protein, suggesting that balanced nuclear import/export and dendritic spine localization are essential for RGS14 functions. We identified genetic variants L505R (LR) and R507Q (RQ) located within the nuclear export sequence (NES) of human RGS14. Here we report that RGS14 encoding LR or RQ profoundly impacts protein functions in hippocampal neurons. RGS14 membrane localization is regulated by binding Gai-GDP, whereas RGS14 nuclear export is regulated by Exportin 1 (XPO1). Remarkably, LR and RQ variants disrupt RGS14 binding to Gai1-GDP and XPO1, nucleocytoplasmic equilibrium, and capacity to inhibit long-term potentiation (LTP). Variant LR accumulates irreversibly in the nucleus, preventing RGS14 binding to Gai1, localization to dendritic spines, and inhibitory actions on LTP induction, while variant RQ exhibits a mixed phenotype. When introduced into mice by CRISPR/Cas9, RGS14-LR protein expression was detected predominantly in the nuclei of neurons within hippocampus, central amygdala, piriform cortex, and striatum, brain regions associated with learning and synaptic plasticity. Whereas mice completely lacking RGS14 exhibit enhanced spatial learning, mice carrying variant LR exhibit normal spatial learning, suggesting that RGS14 may have distinct functions in the nucleus independent from those in dendrites and spines. These findings show that naturally occurring genetic variants can profoundly alter normal protein function, impacting physiology in unexpected ways.

Despite 99.9% similarity across all human genomes, the human population nevertheless exhibits vast genetic diversity responsible for disease predisposition and distinct human

traits. This genetic diversity is not captured by canonical reference protein sequences and, despite selective pressure, missense variations within exonic coding regions frequently arise in modular protein domains that can impact protein functions. While many studies focus on monogenetic variations in the context of severe diseases and phenotypes, rare variants may actually play a more important role in susceptibility, particularly for complex diseases (1–8). Many of these rare variants are predicted to alter protein function, which could impact linked physiology. Thus, exploration of rare missense human variants in cell and animal models can serve as a useful tool to reveal novel protein functions relevant to human physiology.

The Regulators of G protein Signaling (RGS) proteins modulate GPCR/G protein signaling (9–11), which controls many aspects of neurobiology and cognition (12), including synaptic plasticity and learning (13, 14). RGS proteins catalyze the off-rate of active G $\alpha$ -GTP and thus also control many aspects of synaptic plasticity in both physiology and disease states (15, 16). All RGS proteins (20 classical family members) share a conserved ~120 amino acid RGS domain, and many contain accessory domains that act to regulate different aspects of cellular function (9, 10), such as MAP kinase signaling (17–19). Although canonical reference sequences are used to define and study RGS and other proteins, our recent report identified considerable genetic diversity within each human RGS protein family member, with many encoding missense variants (4). These missense variants can have potentially deleterious effects on RGS protein function with physiological consequences (4, 20).

In our report (4), RGS14 emerged as one family member with considerable genetic variance. RGS14 is a dynamically regulated multifunctional signaling protein that suppresses synaptic plasticity in hippocampal neurons (21, 22). In addition to an RGS domain, RGS14 also contains other signaling domains including tandem Ras/Rap binding domains (R1 and R2), which bind active H-Ras-GTP and Rap2A-GTP to regulate MAP kinase signaling (17–19), and a G protein regulatory (GPR) motif, which binds inactive Gai1/3-GDP at the plasma

This article contains [supporting information](#).

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## Genetic variants disrupt RGS14 regulation of hippocampal LTP

membrane to govern RGS14 subcellular localization (23, 24) and capacity to engage downstream effectors (25, 26). Of note, RGS14 is also a nucleocytoplasmic shuttling protein whose subcellular movement is regulated by a nuclear localization sequence (NLS) and a nuclear export sequence (NES) (26, 27). The NES is encoded within the GPR motif and dictates RGS14 nuclear export (26, 28). Thus, the GPR motif and the NES within are critically important for RGS14 subcellular localization, distribution, and cellular functions.

We previously reported that, in mouse brain, RGS14 is highly expressed within the CA2 subregion of the hippocampus, but not in the neighboring CA1 or CA3 subregions (21, 29). While CA1 is known for its robust long-term potentiation (LTP), a form of synaptic plasticity linked to learning and memory (30), LTP is absent from CA2 neurons where RGS14 is naturally expressed (31). RGS14 localizes predominately to spines and dendrites in CA2 neurons, making it well positioned to modulate LTP and synaptic plasticity. In support of this idea, genetic ablation of RGS14 resulted in robust LTP in CA2 (mirroring that of CA1), with no changes in CA1, and introduction of RGS14 into CA1 neurons blocked LTP there (21, 22). Further, loss of RGS14 conferred an enhancement in hippocampal-based spatial learning (21), demonstrating that RGS14 is a natural suppressor of LTP linked to spatial learning. However, whether these findings extended to primates, specifically humans, remained an open question.

We recently reported the regional and cellular localization of RGS14 in the primate brain (32), observing that RGS14 is expressed not only in hippocampus, as expected (29), but also in amygdala and striatum. There, we observed native expression of RGS14 in the nuclei of striatal neurons in primates, consistent with reports of RGS14 as a dynamically regulated nucleocytoplasmic shuttling protein (26, 27). These observations raised numerous questions including: 1) how is RGS14 nuclear localization controlled, 2) what is the impact of nuclear localization on RGS14 cellular functions, and 3) do naturally occurring human variants shift nucleocytoplasmic shuttling?

To explore these questions, we took advantage of human genetic variant data collected from over 130,000 individuals to examine the contribution of naturally occurring mutations on RGS14 functions (33, 34). Human RGS14 contains many genetic variants observed throughout the gene. Here we report that two human genetic variants, specifically L505R (LR) and R507Q (RQ) located within the NES of RGS14, profoundly impact RGS14 functions in mouse hippocampal neurons and brain. We provide evidence that the GPR motif of RGS14 and the NES encoded within are critically important for RGS14 subcellular localization and capacity to prevent LTP induction. Variants LR and RQ disrupt RGS14 binding to Gai1-GDP and XPO1, nucleocytoplasmic equilibrium, and capacity to inhibit LTP in hippocampal neurons. We introduced RGS14 variant LR into mice by CRISPR/Cas9 and found this variant to direct RGS14 protein to the nuclei of neurons in the hippocampus, central amygdala (CeA), and striatum, brain regions rich in synaptic plasticity. Whereas mice lacking RGS14 are reported to exhibit enhanced spatial learning (21), those expressing variant LR exhibit normal spatial learning, suggesting that

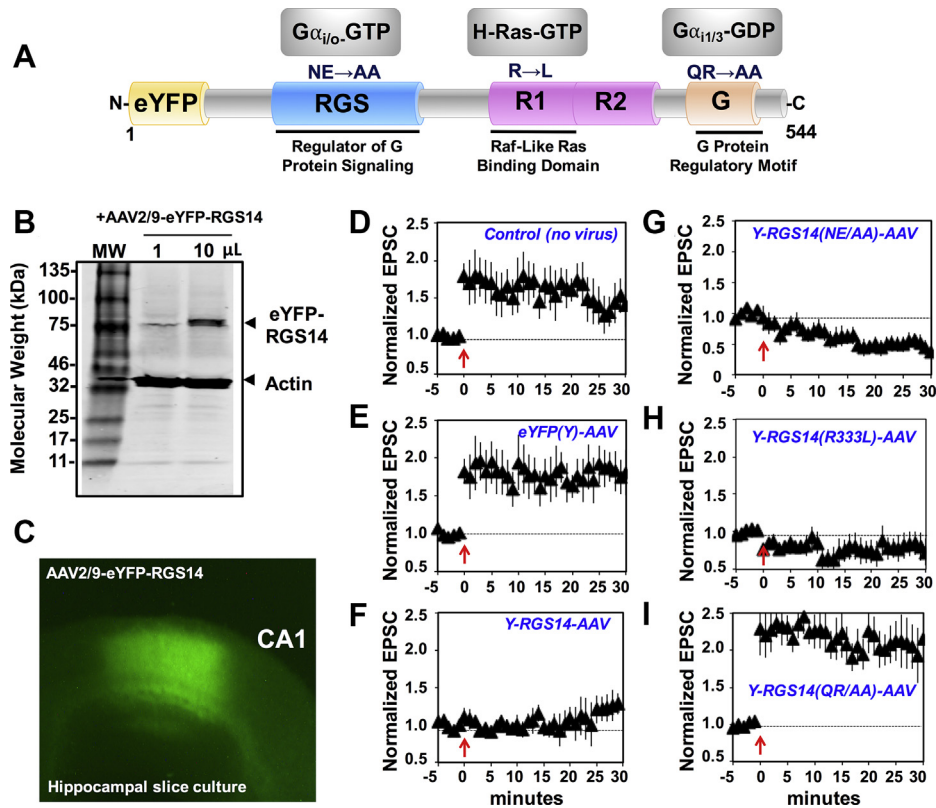
RGS14 has distinct functions in the nucleus *versus* dendrites and postsynaptic dendritic spines. These findings demonstrate that naturally occurring genetic variants can alter normal protein functions in unexpected ways and, in the case of RGS14, with profound physiological consequences in the hippocampus and likely other brain regions.

### Results

#### *The GPR motif is critical for RGS14 capacity to inhibit synaptic plasticity in hippocampal neurons*

Loss of RGS14 (RGS14 KO) restores LTP in CA2 neurons in acute mouse hippocampal slices (21, 22), and exogenous introduction of RGS14 into area CA1 completely blocks synaptic plasticity there (22), suggesting that RGS14 can engage postsynaptic signaling pathways shared by both CA1 and CA2 neurons to inhibit synaptic plasticity. Based on these observations, we sought to isolate which structural region(s) of RGS14 is/are most important for regulating LTP in hippocampal neurons (Fig. 1).

RGS14 is a multifunctional protein containing: 1) an RGS domain, which binds active Gai/o-GTP; 2) tandem Ras-binding domains (R1 and R2), which bind active H-Ras-GTP and Rap2A-GTP; and 3) a GPR (G) motif, which binds inactive Gai1/3-GDP (17–19, 23–26) (Fig. 1A). We initiated studies to determine which domain(s) and protein interaction(s) are most critical for RGS14 regulation of LTP. Because RGS14 blocks LTP in CA1 neurons when introduced by ectopic expression (22), we used this system to examine the effects of targeted loss-of-function mutations (Fig. S1) of RGS14 on LTP in CA1 neurons. Each of these mutations allowed for specific RGS14 functions to be blocked while leaving others intact. We generated adeno-associated viruses (AAV) expressing YFP and wild-type YFP-RGS14 as negative and positive controls (Fig. S1). We also generated AAV expressing: 1) YFP-RGS14 E92A/N93A (EN/AA) in the RGS domain, which blocks Gai/o-GTP binding (35); 2) YFP-RGS14-R333L in the R1 domain, which blocks H-Ras-GTP binding (18); and 3) YFP-RGS14-Q515A/R516A (QR/AA; GPRm) in the GPR motif, which blocks Gai1/3-GDP binding (25, 35) (Fig. S1 and Fig. 1A). Virus was expressed in CA1 of hippocampal slice cultures (Fig. 1, B–C), and excitatory postsynaptic currents (EPSCs) were measured by whole-cell voltage clamp recordings as performed previously on acute slices (21, 22, 31). While control and YFP-only recordings show robust LTP in response to a pairing protocol, expression of wild-type YFP-RGS14 fully blocks LTP when ectopically expressed in CA1 hippocampal neurons (Fig. 1, D–F) as expected (22), demonstrating that RGS14 is sufficient to suppress LTP. Inactivation of the RGS domain or R1 domain did not alter RGS14 capacity to suppress LTP (Fig. 1, G–H). However, inactivation of the GPR (G) motif was able to block RGS14 function and allow LTP in CA1 neurons (Fig. 1I). These results indicate that the GPR motif, a regulator of RGS14 localization within the cell (25, 35), is critical for RGS14 suppression of synaptic plasticity, prompting the notion that spatial dynamics are a critical mediator of RGS14 function.



**Figure 1. The GPR motif is critical for RGS14 suppression of LTP.** A, RGS14 is a multifunctional signaling protein, which contains an RGS domain that binds active Gai/o-GTP, an R1 domain that binds H-Ras-GTP, and the GPR motif (G) that binds inactive Gai1/3-GDP. Amino acid substitutions are indicated for each domain, which confer a loss-of-function phenotype for that specific domain (see Fig. S1, A–C). Each eYFP-RGS14 construct, wild-type and mutant, was placed in AAV expression vector and made into live AAV2/9 virus (Fig. S1D). B, AAV2/9-eYFP-RGS14 was added, either as 1  $\mu$ L or as 10  $\mu$ L, to cultured primary neurons (35 mm dish), and after 24 h, infected cells were harvested and cell lysates were subjected to SDS-PAGE and western immunoblot. Expressed YFP-RGS14 and actin were detected with mouse monoclonal anti-RGS14 and rabbit polyclonal antiactin sera, respectively. C, eYFP-RGS14 is expressed in area CA1 of cultured hippocampal slices following infection with 5 nL AAV2/9-eYFP-RGS14 ( $3.14 \times 10^{13}$  VG/mL) by glass pipette. D–F, long-term potentiation (LTP) was performed by pairing Schaffer collateral input with CA1 neuron depolarization (time indicated with the arrow) in cultured hippocampal slices expressing AAV-eYFP-RGS14 following infection with virus as in C. Following the LTP-pairing protocol, EPSCs were assessed for potentiation. Control CA1 neurons ( $n = 12$ ) and those expressing YFP alone ( $n = 6$ ) showed robust LTP, while eYFP-RGS14 ( $n = 5$ ) suppressed induction of LTP. G–I, although the eYFP-RGS14 NE/AA (RGS domain knockout;  $n = 6$ ) and eYFP-RGS14 R/L (R1 domain knockout;  $n = 5$ ) suppressed LTP similar to WT RGS14, the eYFP-RGS14 GPR motif knockout QR/AA ( $n = 6$ ) failed to suppress LTP, suggesting that the GPR motif is a critical mediator of RGS14 function. Data is presented as mean  $\pm$  SD.

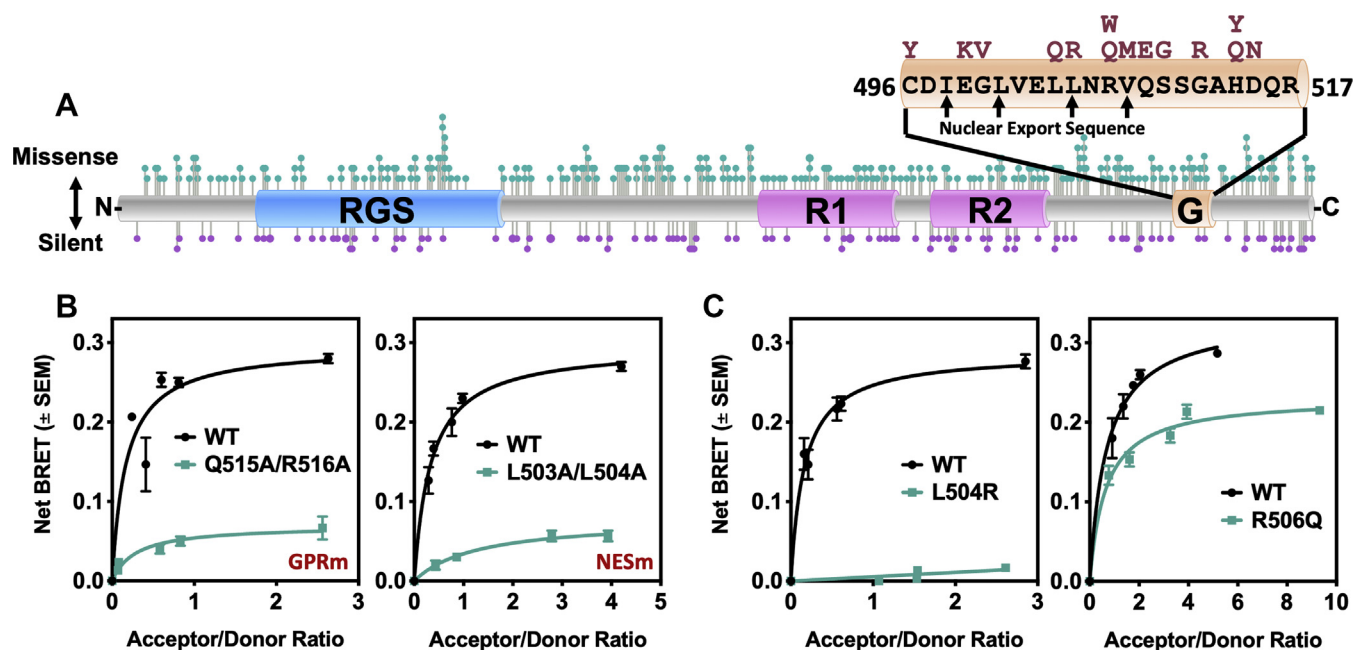
### Naturally occurring rare human genetic variants within the GPR motif disrupt RGS14 interactions with Gai1-GDP

The mutations used earlier to disrupt the RGS14 GPR motif function were rationally designed within the context of the canonical RGS14 sequence. Based on the newly appreciated importance of the GPR motif for RGS14 function in LTP (Fig. 1), we next explored whether naturally occurring human variants existed within the GPR motif that might impact RGS14 protein function. We recently reported the existence of numerous missense variants in the human RGS14 sequence, including many within the GPR motif (4) that are predicted to disrupt function. For this, we accessed the Genome Aggregation Database (GnomAD, version 2.0) for all RGS14 missense (amino acid change) and silent (DNA change but no amino acid change) variants. We observed a relatively even distribution of missense and silent variants throughout the RGS14 gene (Fig. 2A, missense on top, silent on bottom). Germane to RGS14 regulation of G protein signaling, multiple naturally occurring rare variants were found within the GPR motif

(Fig. 2A, GPR motif and sequence expanded with variants shown on top in red).

The GPR motif binds inactive Gai1-GDP and serves to recruit and stabilize RGS14 at the plasma membrane (18, 25). RGS14:Gai1-GDP interactions are readily detectable by bioluminescence resonance energy transfer (BRET) and are reflective of functional RGS14 trafficking between the cytosol and the plasma membrane (25). Wild-type RGS14-Luc binds Gai1-YFP and saturates the net BRET signal in a concentration-dependent manner (Fig. 2B, black line). Importantly, the GPR motif has embedded within it a nuclear export sequence (NES, Fig. 2A), which shuttles nuclear RGS14 to the cytoplasm (27). The NES is therefore also an important mediator of RGS14 subcellular localization, as it provides a cytoplasmic pool of RGS14 for Gai1-GDP recruitment and colocalization at the plasma membrane. When the GPR motif is mutated (Q515A/R516A) such that it cannot bind Gai1-GDP (GPRm) (25, 36) (Fig. S1) or when the NES is mutated (L503A/L504A) such that RGS14 is incapable of nuclear

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**Figure 2. Rare human genetic variants within the GPR and NES motifs of RGS14 disrupt its association with Gai1-GDP, suggestive of aberrant cellular trafficking.** A, RGS14 missense and silent variants were obtained from GnomAD (Broad Institute) and plotted onto the sequence of RGS14 using Lollipop (https://github.com/pbnjay/lollipop) as described previously (4). Missense variants are plotted on top in teal, with silent variants on the bottom in purple. The GPR motif, which contains an embedded NES, has human variants as shown (red). B–C, BRET measurement of RGS14-Luc interaction with Gai1-eYFP-GDP. RGS14-Luc is recruited from the cytoplasm to the plasma membrane by inactive Gai1-eYFP-GDP (25). Inactivation of the GPR in RGS14-Luc (Q515A/R516A; GPRm; n = 3) prevents Gai1-eYFP-GDP binding and recruitment to the plasma membrane. Similarly, inactivation of the NES (L503A/L504A; NESm; n = 3) prevents RGS14 from being accessible for recruitment in the cytoplasm and therefore cannot associate with Gai1-GDP at the plasma membrane. Variants L504R or R506Q were placed in RGS14-Luc and tested for their capacity to associate with Gai1-eYFP-GDP by BRET, indicative of proper recruitment to the plasma membrane. L504R (n = 3) completely abolished Gai1-GDP association, while R506Q (n = 3) exhibited a partial reduction in Gai1-GDP association. Data is presented as mean  $\pm$  SEM.

export (NESm) (26, 27), RGS14-Luc ceases to bind Gai1-YFP as measured by BRET, indicating aberrant subcellular trafficking (Fig. 2B, teal lines). Together, these findings confirm that this screening assay is a sensitive measure of Gai1-GDP binding and RGS14 subcellular spatial dynamics.

We identified 14 human variants in the RGS14 GPR motif (Fig. 2A). Each of these variants was measured for their interactions with Gai1-GDP by BRET, and we found that, whereas the majority did not affect the Gai1-YFP BRET signal (Fig. S2), several did. Two rare variants (defined here as <2% frequency) emerged as especially interesting based on their effect and population frequency. Human RGS14 variant L505R (L504R in rat sequence), which is very rare (found in 0.006% of East Asian population [GnomAD v2.0]), completely abolished interaction with Gai1-YFP (Fig. 2C). Human RGS14 variant R507Q (R506Q in rat sequence, found in 1.25% of Ashkenazi Jewish population, among other populations [GnomAD v2.0]) has a submaximal reduction in Gai1-YFP interaction and BRET signal.

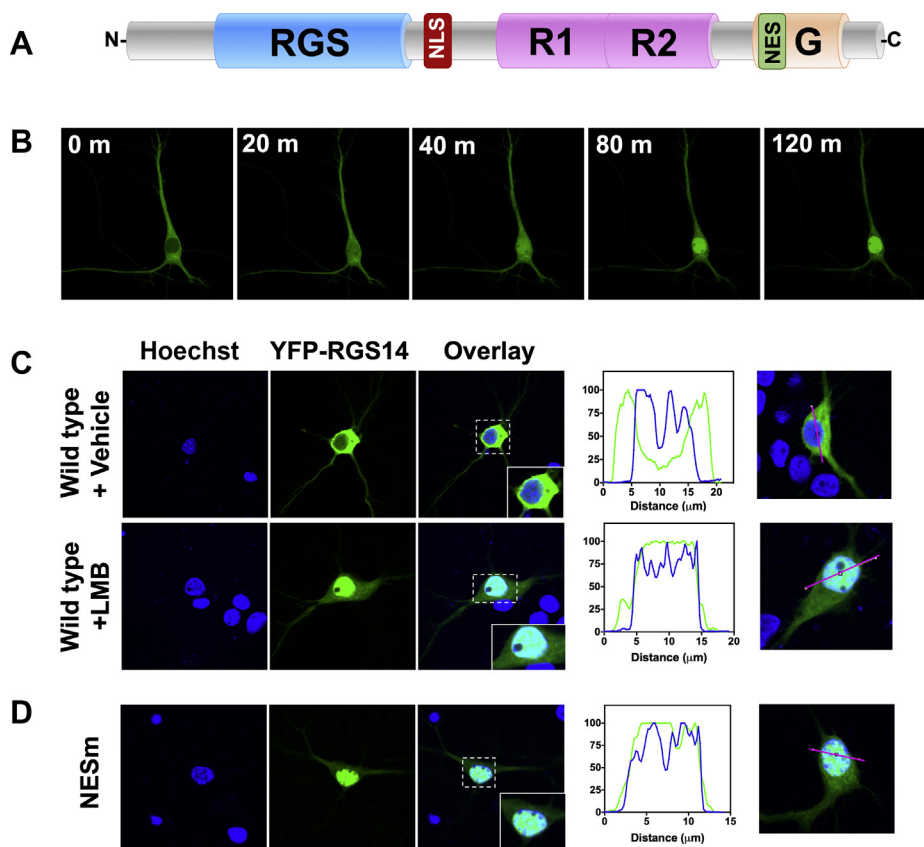
### RGS14 is a nuclear shuttling protein in primary hippocampal neurons

We and others have reported that recombinant RGS14 shuttles into and out of the nucleus in cultured cell lines (26–28) and neurons (37); however, the dynamic subcellular behavior of RGS14 in neurons has not been well described. As noted, RGS14 contains a functional NLS between the RGS

domain and the first Ras-binding domain (R1) and a functional NES embedded within the GPR motif of RGS14 (26, 27) (Fig. 3A).

Neurons (DIV 8) were transfected with AAV-hSyn-YFP-RGS14 for 18 h and treated with the nuclear export inhibitor Leptomycin B (LMB) over two and a half hours, concurrent with imaging by live cell confocal microscopy. We found that cytoplasmic RGS14 translocated to the nucleus within 40 min and was maximally nuclear by 2 h (Fig. 3B), suggesting that cytoplasmic pools of RGS14 are constantly targeted for nuclear import/export with an equilibrium favoring the cytosol. These results are consistent with our recently published report demonstrating that nuclear import of RGS14 is regulated by 14-3-3 $\gamma$  in neurons (37).

We next verified colocalization of RGS14 with a nuclear marker, Hoechst, as well as validated the functional status of the RGS14 NES motif in neurons. DIV 8 neurons were again transfected with AAV-YFP-RGS14 WT or the NES mutant (NESm) incapable of being shuttled out of the nucleus (26, 27). Neurons were treated with vehicle or LMB for 2 h, then fixed and stained with Hoechst, and YFP signal was assessed by confocal microscopy. Vehicle-treated neurons exhibited robust YFP-RGS14 expression that filled the soma, dendrites, and spines of the neurons, but did not overlap with the Hoechst-stained nucleus. In contrast, YFP-RGS14 WT colocalized entirely with nuclear Hoechst in LMB-treated neurons (Fig. 3C). Immunoblot analysis verified that full-length



**Figure 3. RGS14 is a nucleocytoplasmic shuttling protein in hippocampal neurons.** *A*, RGS14 rapidly shuttles into and out of the nucleus in dividing cells directed by a nuclear localization sequence (NLS) and nuclear export sequence (NES). Studies validated that RGS14 followed the same cellular dynamics in primary hippocampal neurons. *B*, neurons were infected with AAV2/9-eYFP-RGS14 for 18 h and then imaged using live cell confocal microscopy. Leptomycin B (LMB) was added (final concentration 20 nM) and YFP signal was measured. At time zero, YFP-RGS14 is entirely cytosolic. After 20 min, YFP-RGS14 was detectable in the nucleus, at 40 min YFP-RGS14 was mostly nuclear, and by 80 to 120 min YFP-RGS14 was entirely nuclear. *C*, wild-type YFP-RGS14 does not colocalize with Hoechst in vehicle-treated conditions, but colocalizes with Hoechst entirely under LMB conditions. *D*, mutation of the NES (L503A/L504A, NESm) in RGS14 sequesters RGS14 entirely in the nucleus under vehicle-treated conditions. Approximately ten images were collected per condition and representative images are shown.

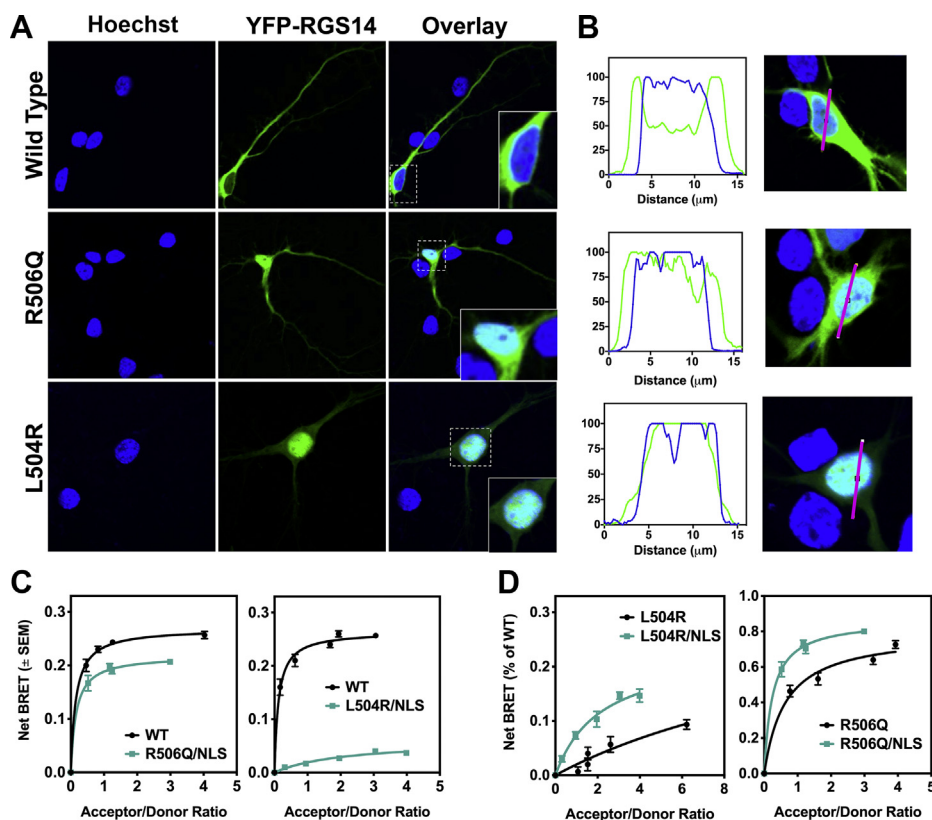
YFP-RGS14 was expressed (Fig. S3), confirming this nuclear localization was not an effect of posttranslationally modified (e.g., truncated) RGS14. LMB strongly and specifically inhibits nuclear export receptor Exportin 1 (XPO1, also known as CRM1) (38). The XPO1-binding motif is a leucine-rich sequence (39). We and others previously showed that mutating two leucines (L503A/L504A) as mentioned earlier (Fig. 2B) in the RGS14 GPR motif (LL/AA) inhibited nuclear export of RGS14 in immortal cells (26, 27). Thus, we examined in complimentary experiments whether RGS14 nuclear shuttling in primary hippocampal neurons occurred through the same mechanism. AAV-hSyn-YFP-RGS14 LL/AA (NESm) was transfected into DIV 8 neurons and stained with Hoechst as before. Again, we verified expression of a single band corresponding to full-length YFP-RGS14 NESm by immunoblot (Fig. S3). We found that under vehicle-treated conditions, YFP-RGS14 NESm sequestered in the nuclei and colocalized with Hoechst, indistinguishable from the LMB-treated WT RGS14 (Fig. 3D). These findings confirm that RGS14 nucleocytoplasmic shuttling behaves similarly in primary hippocampal neurons as it does in other cells and that RGS14 nuclear export is mediated by XPO1.

#### Human variants LR and RQ mislocalize RGS14 to the nucleus in primary hippocampal neurons

As illustrated in Figure 2, RGS14 genetic variants LR and RQ exhibit reduced Gai1-YFP binding. Based on this observation, we hypothesized that RGS14 variants might also exhibit aberrant cellular trafficking, *via* either reduced plasma membrane recruitment or reduced nuclear shuttling. To distinguish between these possibilities, we transfected DIV 8 neurons with AAV-hSyn-YFP-RGS14 WT, AAV-hSyn-YFP-RGS14 RQ, or AAV-hSyn-YFP-RGS14 LR for 18 h, then fixed and stained them with Hoechst as before. Confocal imaging captured the subcellular localization of wild-type or variant RGS14 within the neuron. Wild-type RGS14 localized to the soma, dendrites, and spines. In stark contrast, RGS14 LR concentrated within the nucleus while the RQ variant exhibited a mixed phenotype (Fig. 4, A–B).

Given that the RGS14-binding motifs for XPO1 and Gai1-GDP overlap, coupled with previous studies showing that Gai1-GDP interactions can affect nuclear localization (26), we next investigated whether the reduction in RGS14-Luc:Gai1-YFP binding observed in Figure 2 was due to RGS14 sequestration in the nucleus, or if there was an additional inhibitory

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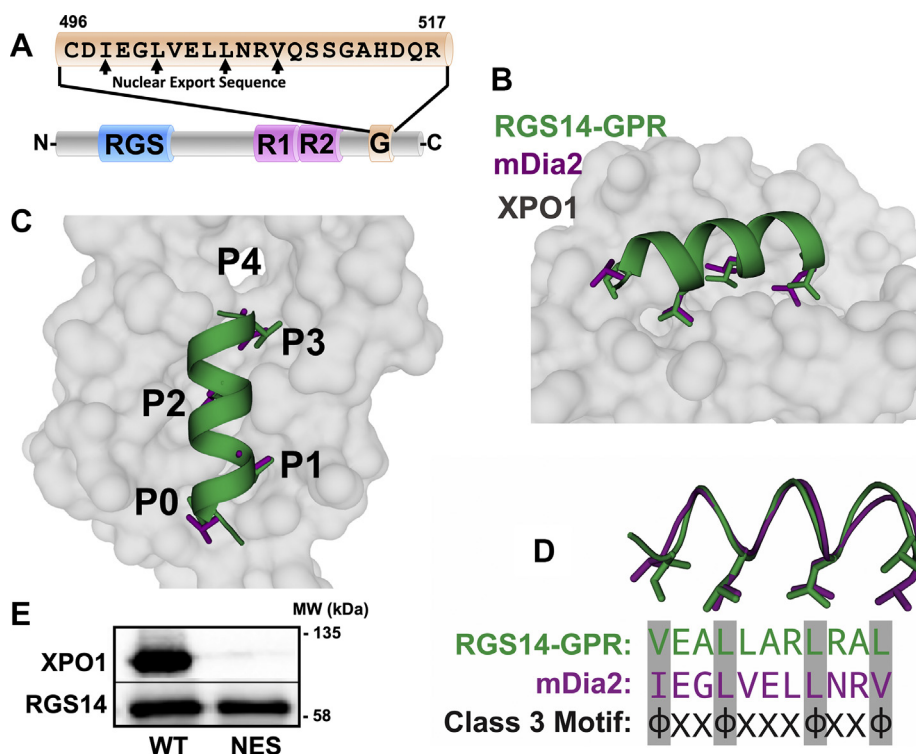
**Figure 4. Human variants LR and RQ disrupt RGS14 nucleocytoplasmic equilibrium to favor the nucleus over the cytoplasm.** Primary hippocampal neurons were infected with AAV-eYFP-RGS14 or RGS14 carrying variants LR or RQ for 18 h and then imaged using live cell confocal microscopy. *A*, wild-type RGS14 does not colocalize with Hoechst in neurons under unstimulated conditions. The RQ variant occupies both the nucleus and the cytoplasm, while the LR variant localizes entirely to the nucleus. *B*, quantification of YFP (RGS14) and Hoechst overlay for WT RGS14 and each of the variants provides a measure of colocalization. Approximately ten images were collected per condition and representative images are shown. *C*, RGS14-Luc and Gai1-eYFP-GDP interactions in live HEK cells were measured as static net BRET. Since the RGS14 NES motif overlaps with the GPR binding site of Gai1-GDP, an NLS mutation (NLSm) was introduced to prevent nuclear localization. Thus, any change in net BRET from wild-type is due to effects of Gai1-GDP binding, independent of mislocalization to the nucleus and interaction with XPO1. Net BRET for both RGS14 RQ ( $n = 3$ ) and RGS14 LR ( $n = 3$ ) is partially (although not fully) restored when an NLS mutation is introduced. *D*, measure of RGS14-Luc and Gai1-eYFP-GDP Net BRET comparing L504R and L504R/NLSm and R506Q and R506Q/NLSm (L504R and R506Q data from Fig. 2). Adding the NLS mutation enhances Gai1-GDP association, indicating that XPO1 and Gai1-GDP interactions are both impacted by the L504R and R506Q mutations.

effect on the RGS14:Gai1 binding interaction itself. Results show that these variants mislocalize RGS14 to the nucleus, similar to RGS14 NESm. RGS14 NESm has poor Gai1 binding as measured by BRET. Thus, determining whether reduced BRET activity is due to loss of RGS14 binding to Gai1 or loss of an available cytosolic pool of RGS14 presents a challenge. To tease apart these two possibilities, we generated mutations that disrupt the NLS motif (R208A/K209A/K210A; NLSm) (Fig. S4) in combination with our RGS14-Luc LR and RGS14-Luc RQ constructs to make RGS14-Luc LR/NLS and RGS14-Luc RQ/NLS, respectively. These double mutant constructs are unable to translocate to the nucleus (Fig. S4). Therefore, contributions of loss of XPO1 binding and nuclear sequestration should be eliminated, allowing a measure of the effect of these human variants on RGS14-Luc:Gai1-YFP binding alone. We found that RGS14 LR/NLS and RGS14 RQ/NLS partially, but not fully, restored Gai1-YFP binding (Fig. 4C). Comparing LR with LR/NLS and RQ to RQ/NLS as a percentage of wild-type (Fig. 4D) using the data generated in Figure 2C shows an incomplete rescue of RGS14 binding to Gai1-YFP in both the LR and RQ variants when an NLS mutation was added (Fig. 4C), suggesting at least a partial

disruption of RGS14 binding to Gai1-GDP unrelated to XPO1 binding. These results also suggest that any RGS14 LR or RGS14 RQ that is translocated from the nucleus will be an unlikely target of Gai1-GDP, thus further disrupting plasma membrane recruitment and proper cellular trafficking necessary for inhibitory actions on LTP (Fig. 1). While loss of both XPO1 and Gai1 binding contributes to overall inhibition of RGS14 localization, variant RGS14 must first overcome nuclear sequestration, and thus we interpret the predominant phenotype to be aberrant nuclear export. Together, our BRET data (Fig. 2) and imaging data (Fig. 4) indicate that naturally occurring rare human variants LR and RQ confer aberrant cellular trafficking properties to RGS14.

### The XPO1 binding motif in RGS14 fits a class 3 model

RGS14 is a nucleocytoplasmic shuttling protein in neurons that utilizes XPO1 as its export receptor (Fig. 3), and variants LR and RQ disrupt this interaction (Fig. 4). To explore the molecular basis for this, we next created a structural model of the RGS14 NES motif bound to XPO1 (Fig. 5). XPO1, also known as CRM1, recognizes a diverse range of NES motifs that can be grouped into ten classes according to their unique



**Figure 5. The GPR motif of RGS14 contains an embedded class 3 nuclear export sequence that binds XPO1.** Structural modeling of the RGS14 GPR/NES motif binding to Exportin 1 (XPO1). *A*, the NES of RGS14 is embedded within the GPR motif and follows a class 3 pattern:  $\Phi_1XX\Phi_2XXX\Phi_3XX\Phi_4$ . *B*, the crystallized GPR peptide of RGS14 (PDB 1KJY) was structurally aligned with a similar class 3 NES motif (mDia2 PDB 5UWP) crystallized with XPO1. *C–D*, the RGS14 NES fits a class 3 binding motif, occupying pockets 0 to 3, but not pocket 4, as reported previously (40). *E*, RGS14 WT, but not RGS14 NES, coimmunoprecipitates with Flag-XPO1. Flag-XPO1 and RGS14 WT or NES were transfected into HEK cells and Flag was immunoprecipitated (Flag affinity gel). RGS14 (1:1000) and XPO1 (1:300) protein were measured by immunoblot.

spacing of hydrophobic residues (40). The RGS14 NES motif embedded within the GPR motif forms an all  $\alpha$ -helical secondary structure that matches a class 3 spacing motif:  $\Phi_1XX\Phi_2XXX\Phi_3XX\Phi_4$  where  $\Phi$  is L, V, I, F, or M, and X is any amino acid (Fig. 5A). We therefore hypothesized that the RGS14 NES would fit a class 3 binding model to XPO1. The RGS14 GPR peptide, which contains the full NES, was crystallized previously in complex with G $\alpha$ i1-GDP (36), and a class 3 NES (mDia2) was recently crystallized with XPO1 (40). Using this information, we generated an RGS14-NES/XPO1 model by structurally aligning RGS14's NES with the crystal structure of the mDia2 NES–XPO1 complex (Fig. 5B). The alignment of the two NES-containing helices reveals optimal positioning of the hydrophobic residues ( $\Phi$ ) necessary for interaction with XPO1 (Fig. 5C). These  $\Phi$  residues stabilize the RGS14–XPO1 interaction through hydrophobic interactions with XPO1 pockets P0–P3, mirroring the previously reported mDia2–XPO1 complex (Fig. 5D) (40). When coexpressed in HEK293 cells, coimmunoprecipitation of RGS14 with Flag-XPO1 confirmed that RGS14, but not RGS14 NES, is a binding partner of XPO1 (Fig. 5E).

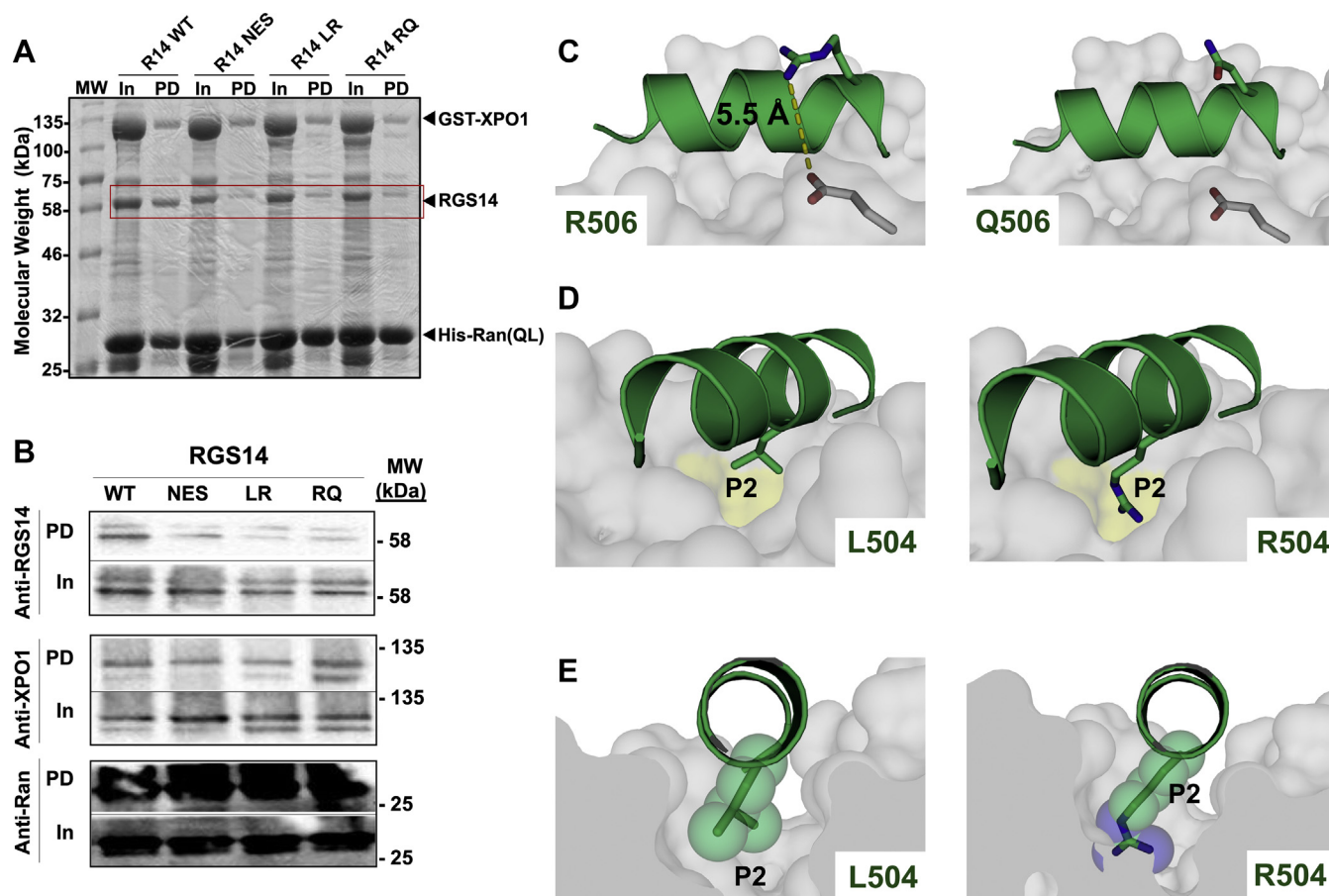
#### Human variants LR and RQ disrupt RGS14 interactions with XPO1

As shown earlier, RGS14 is a nuclear shuttling protein that utilizes XPO1 as a nuclear export receptor, and human variants RQ and LR cause RGS14 to mislocalize and accumulate in the

nucleus. Based on this, we hypothesized that these variants disrupt RGS14 binding with the XPO1. To test this idea, we examined whether these variants altered RGS14 complex formation with purified XPO1 and its obligate binding partner, Ran-GTP. Purified bacterially expressed recombinant RGS14 WT, RGS14 NES, RGS14 LR, or RGS14 RQ was mixed with purified XPO1-GST and constitutively active His-Ran Q69L-GTP. After mixing for 2 h for complex formation to occur, His-Ran QL-GTP was recovered by Ni-NTA affinity pull-down along with XPO1-GST and/or RGS14 in complex. As expected, RGS14 WT was precipitated in complex with XPO1. In contrast, RGS14 NES, RGS14 LR, or RGS14 RQ did not bind XPO1 (Fig. 6, A–B). We next utilized the RGS14-NES/XPO1 model described earlier (Fig. 5) to further explore the mechanism by which these variants disrupt RGS14 binding to XPO1. As the reported RGS14 GPR crystal structure utilized rat RGS14 (36), the human variants R507Q and L505R correspond to rat residue number R506Q and L504R. Based on our model, arginine 506 (R506) in wild-type RGS14 contains a positively charged nitrogen atom that is 5.5 Å away from a negatively charged oxygen atom of glutamic acid in XPO1, which is in line with a possible ionic interaction. We then mutated this residue in our model to the RQ variant, glutamine 506 (Q506), in order to provide mechanistic insight for the decreased nuclear export present with the RQ variant (Fig. 6C). The variant (Q506) disrupts the ionic interaction that was present between R506 and glutamic acid in XPO1. The distance between Q506 and the



## Genetic variants disrupt RGS14 regulation of hippocampal LTP



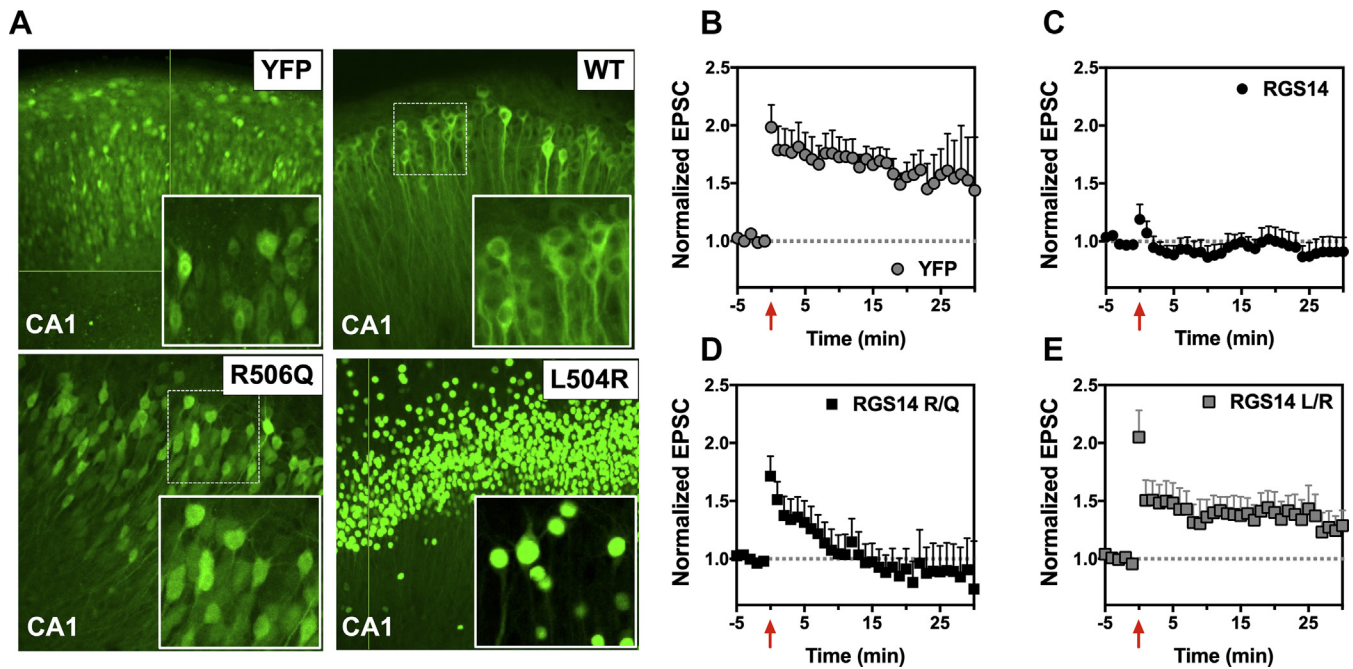
**Figure 6. Human variants LR and RQ disrupt RGS14 binding to Exportin 1 (XPO1) through loss of side-chain interactions.** A–B, recombinant RGS14 WT and variants LR and RQ were purified from *E. coli* and mixed with purified GST-XPO1 and constitutively active His-Ran (GV)-GTP. His-Ran (GV)-GTP was captured by Ni-NTA affinity pull-down and complexes were assessed by Coomassie (A) and immunoblot (B). WT RGS14 was recovered in complex with XPO1, but RGS14 NES, RGS14 LR, and RGS14 RQ did not bind XPO1. Blots are representative of five independent experiments. C, structural modeling of the R506Q variant binding to XPO1. Left panel is the wild-type amino acid, arginine, which makes a salt bridge with a nearby glutamate near the binding pocket on XPO1. Mutation to a glutamine (right panel) removes this salt bridge, presumably destabilizing the protein–protein interaction. D–E, structural modeling of the L504R variant onto XPO1. The leucine interacts directly with the hydrophobic P2 pocket (left panel). Mutation of this amino acid to an arginine (right panel) introduces a charged side chain into a hydrophobic pocket. These data mechanistically explain the difference in phenotype (R506Q being a subtler phenotype compared with L504R).

glutamic acid in XPO1 is too great for hydrogen bonding, suggesting this variant will reduce affinity of RGS14 for XPO1 (Fig. 6C) and thus explains the mixed phenotype. Although RGS14 RQ can bind XPO1, it does so less efficiently than WT RGS14. We next wanted to visualize the impact of the LR variant, arginine 504 (R504). In wild-type RGS14, leucine 504 (L504) fits nicely within the P2 hydrophobic pocket of XPO1, stabilizing the complex through hydrophobic interactions. The LR variant (R504) places a charged arginine within the P2 hydrophobic pocket of XPO1 causing a polar incompatibility that will greatly disrupt the RGS14–XPO1 interaction (Fig. 6, D–E). This severe mutation within the binding interface of RGS14 and XPO1 suggests that the LR variant is highly unlikely to interact with XPO1 and thus explains why RGS14 LR becomes trapped within the nucleus.

### Human variants LR and RQ disrupt RGS14 suppression of LTP in hippocampal slices

The profound disruption of RGS14 binding to Gai1-GDP and XPO1 observed with the LR and RQ human variants,

which results in their subcellular mislocalization, predicts that these variants will negatively impact RGS14 suppression of synaptic plasticity. To examine this, we utilized the experimental system of RGS14 ectopic expression in CA1 hippocampal slices as shown in Figure 1 and as described previously (22). We generated wild-type (WT) AAV-YFP-RGS14 and AAV-YFP as positive and negative controls and compared these with AAV-YFP-RGS14 LR and AAV-YFP-RGS14 RQ. Each of these AAV constructs was exogenously expressed in area CA1 of cultured hippocampal slices (Fig. 7A). We found that wild-type YFP-RGS14 was detected throughout the soma, dendrites, and spines (Fig. 7A), consistent with expression patterns in dissociated hippocampal neurons (Fig. 4A). YFP-RGS14 RQ filled the entire cell, occupying both the nucleus and cytoplasm, while YFP-RGS14 LR localized to the nucleus (Fig. 7A), consistent with our findings in dissociated neurons (Fig. 4A). We recorded EPSCs at baseline and following an LTP induction protocol, as before (Fig. 1 and (21)). Neurons from slices with YFP expression alone showed robust LTP (Fig. 7B), whereas WT YFP-RGS14 expression completely



**Figure 7. Human variants LR and RQ that disrupt RGS14 nuclear-cytoplasmic equilibrium also block RGS14 capacity to inhibit long-term potentiation in hippocampal slices.** AAV-eYFP-RGS14 was used to make AAV2/9 viruses, including wild-type RGS14 as well as human variants RQ and LR. Furthermore, a stop codon was generated as follows AAV-YFP-STOP-RGS14, to make a truncated YFP virus lacking RGS14. Virus was injected into mouse CA1 slice cultures and incubated for 1 week, at which point electrophysiological recordings were made to assess LTP. A, expression of YFP-RGS14 in CA1 brain slices. To improve visualization of individual neurons, insets are shown derived from cutout white dashed boxes (WT and R506Q) or as 20 $\times$  images where individual cell resolution in the cutout was not sufficient (YFP and L504R). The horizontal and vertical green lines in images for YFP and L504R represent the interface where composite confocal images were spliced together. WT RGS14 fills the cytoplasm of neurons. RQ fills both the cytoplasm and nucleus, while LR localizes predominantly to the nucleus, entirely consistent with our data in dissociated neurons. Images are representative of 6 to 9 experiments. B–E, long-term potentiation (LTP) was induced by an LTP pairing protocol in CA1. YFP alone ( $n = 8$ ) had no effect on LTP, but in contrast, WT RGS14 ( $n = 6$ ) completely inhibits LTP in CA1, a hippocampal region where RGS14 is not natively found. Potentiation lasting only about 10 min was induced in RQ variant-expressing neurons ( $n = 9$ ), whereas the LR variant ( $n = 9$ ) failed to suppress LTP. ANOVA with Tukey's post-hoc analysis demonstrated a statistically significant difference between YFP alone (control) and WT RGS14 at 5 ( $p = 0.03$ ) and 10 ( $p = 0.006$ ) min poststimulation, while RGS14 LR was not statistically different from YFP alone at either time point. RGS14 RQ was statistically different from YFP alone at 10 min poststimulation ( $p = 0.018$ ), but not 5 min. This suggests that the degree of mislocalization by these variants correlates with ablation of RGS14 function in CA1 neurons. Data is presented as mean  $\pm$  SEM.

blocked LTP, as expected (Fig. 7C). Consistent with our other data here, the YFP-RGS14 RQ variant allowed induction of a short-lasting potentiation (Fig. 7D), while the YFP-RGS14 LR variant blocked RGS14's capacity to inhibit LTP (Fig. 7E). ANOVA with Tukey's post-hoc comparisons revealed a statistically significant difference between YFP alone and WT RGS14 at 5 min ( $p = 0.03$ ) and 10 min ( $p = 0.006$ ) poststimulation. RGS14 LR was not statistically different (defined as  $p < 0.05$ ) from YFP alone at either time point, while RGS14 RQ was statistically different from YFP alone only at 10 min poststimulation ( $p = 0.018$ ). Together, these findings demonstrate that: 1) RGS14 mislocalization is associated with impaired postsynaptic plasticity functions; and 2) the degree of nuclear accumulation is correlated with the degree of RGS14 dysfunction as a regulator of postsynaptic plasticity.

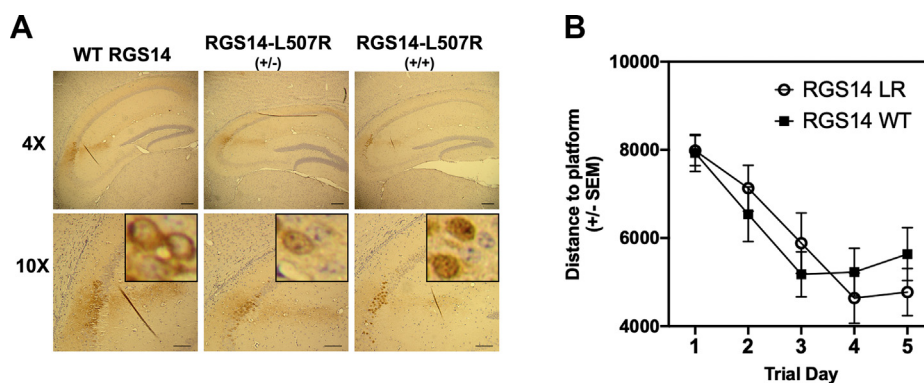
#### **Variant LR disassociates RGS14 actions on synaptic plasticity from spatial learning in mice**

We previously reported that genetic loss of RGS14 (RGS14 KO mice) restores LTP in hippocampal area CA2 that is correlated with enhanced spatial learning as measured by the Morris Water Maze test (21). Based on our observation that the LR variant completely ablated RGS14-dependent

suppression of LTP (Fig. 7), we hypothesized that RGS14 LR may behave as a functional knockout and recapitulate the enhanced spatial learning phenotype of the RGS14 KO mice (21). To test this idea, we generated a novel mouse line carrying the RGS14 LR variant using CRISPR/Cas9 on a C57Bl/6J background. To confirm that RGS14 LR variant mice exhibited the characteristic aberrant localization of RGS14 in hippocampal CA2 neurons, we performed immunohistochemical staining of hippocampal slices. We observed that, like our findings with cultured hippocampal slices (Fig. 7) and neurons (Fig. 4), homozygously expressed RGS14 LR was localized to the nuclei of CA2 neurons as compared with RGS14 WT, which exhibited typical diffuse staining throughout the soma and neurites of CA2 neurons but was absent from nuclei (Fig. 8A). Heterozygously expressed RGS14 LR showed a mixed expression pattern present in soma, neurites, and the nuclei of CA2 neurons. These findings with whole animal are completely consistent with our observations in cultured hippocampal slices (Fig. 7) and primary neurons (Fig. 4).

We took advantage of these animals to examine RGS14 protein expression in brain regions outside of the hippocampus, particularly brain regions recently reported in monkey and human to express RGS14 (29, 32). Consistent with RGS14

## Genetic variants disrupt RGS14 regulation of hippocampal LTP



**Figure 8. The RGS14 variant LR localizes to the nuclei of hippocampal neurons but does not alter hippocampal-based spatial learning in CRISPR/Cas9 mice.** Mice carrying the LR variant were generated by CRISPR/Cas9 (RGS14 LR mice). *A*, expression and localization of RGS14 LR in CA2 hippocampal neurons were confirmed by immunohistochemical staining. Consistent with our slice electrophysiology data, RGS14-LR was predominantly nuclear, while RGS14 WT was localized to soma, dendrites, and spines but excluded from the nucleus of CA2 hippocampal neurons. The heterozygous mice exhibited a mixed nuclear-cytoplasmic phenotype. *B*, RGS14-LR mice ( $n = 20$ ) and RGS14 WT ( $n = 20$ ) littermates were subjected to the Morris Water Maze test. RGS14-LR mice do not exhibit enhanced spatial learning, suggesting a unique nuclear role of RGS14 on suppression of spatial learning.

protein expression patterns in primates, we observed RGS14 protein expression in the CeA, striatum, and piriform cortex of the RGS14 LR mouse (Fig. S5). RGS14 expression had not been reported previously in the striatum and amygdala of rodents, due to low mRNA expression (Allen Brain Atlas: <https://portal.brain-map.org/>). In the RGS14 LR animals, RGS14 is concentrated in the nuclei resulting in a bright punctate staining pattern, greatly improving detection and visualization of RGS14. The concentration of RGS14 LR in the nucleus allows for effortless distinction between cells that express RGS14 and cells that do not, but which are contacted by RGS14-containing neurites.

To determine the effects of nuclear-bound RGS14 LR on spatial learning, we subjected RGS14 WT and RGS14 LR mice to the Morris Water Maze test (Fig. 8B). We expected the LR mice to mimic the RGS14 KO mice and exhibit enhanced spatial learning as before (21). However, quite unexpectedly, RGS14 LR mice did not exhibit an enhancement of the spatial learning phenotype and behaved similarly to WT littermates (Fig. 8B). In summary, RGS14 LR mimicked one aspect of genetically ablated RGS14 (RGS14 KO), *i.e.*, a release on the block of LTP in hippocampal neurons, but failed to recapitulate the other phenotype of RGS14 KO mice, *i.e.*, enhancement of spatial learning (21). Taken together, these findings suggest that nuclear RGS14 may serve a distinct cellular role from RGS14 localized to postsynaptic dendritic spines.

Although RGS14 is a well-established nucleocytoplasmic shuttling protein, nuclear roles for RGS14 remain unknown. In an initial attempt to define a nuclear role for RGS14, we examined if RGS14 nuclear shuttling and localization impacted mRNA expression levels or patterns in hippocampal neurons using RNA-seq methodology (41). Cultured primary hippocampal neurons were infected with AAV expressing RGS14 with normal nucleocytoplasmic shuttling (WT), RGS14 excluded from the nucleus (NLSm), or RGS14 targeted exclusively to the nucleus (NESm) (Fig. S4). After 2 weeks, neurons were harvested, mRNA was isolated and converted to a sequencing library for analysis. No differences in the gene expression patterns or total number of genes expressed

(10,324 of 17,324 transcripts) were observed in hippocampal neurons expressing either RGS14 exclusively in the cytosol (NLSm) or the nucleus (NESm) or both (WT) (Fig. S6).

### Discussion

Here, we provide evidence for the interdependence of RGS14 subcellular localization and protein function in host neurons, specifically driven by genetic diversity. Our findings suggest distinct roles for RGS14 in the nucleus and dendritic spines of hippocampal neurons and highlight the underappreciated fact that rare human genetic variants can alter the function of affected proteins in unexpected ways to markedly impact cellular physiology. We identified the GPR motif, and the embedded NES, as being critical for RGS14 functions in hippocampal neurons. We demonstrate that genetic variants within the GPR and NES motifs disrupt RGS14 localization to dendritic spines and capacity to inhibit LTP, serving to highlight the relationship between spatial localization and function for RGS14.

One surprising result from our studies is that the nuclear-bound RGS14 LR variant presented a discordant behavioral phenotype compared with the RGS14 KO phenotype in mice, which exhibit unrestricted LTP in hippocampal slices that is correlated with an enhancement in spatial learning (21). Although hippocampal LTP and learning/memory are not necessarily causally linked, they are tightly correlated (42), and thus we expected the RGS14 LR mice to behave identically to the RGS14 KO mice in the behavioral assessment. Whereas the RGS14 LR mice exhibited a release of the brake on LTP in hippocampal slices, they did not exhibit altered spatial learning. Reasons for this distinct phenotype are unclear. Early LTP lasts 1 to 2 h and is dependent on local signaling and local protein synthesis at the spines and dendrites (43, 44). Both the KO and LR mice lack RGS14 in dendritic spines, resulting in unrestricted early LTP. By contrast, late LTP and linked memory last much longer and are reliant on synthesis of *new* mRNA/protein from the nucleus, as well as synaptic shuttling of those newly formed mRNA/proteins (45–47). In the RGS14 KO mice, RGS14 is neither at the spines nor in the nucleus,

whereas RGS14 LR exists primarily in the nucleus. In the case of RGS14 LR mice, any enhancement of early LTP at the spines may not be properly consolidated in late (nuclear-dependent) LTP processes. Conversely, RGS14 KO mice have a release of molecular brakes at the spines and dendrites, and *RGS14 is absent from the nucleus*, allowing for unabated inhibition of plasticity, thereby suggesting a possible distinct nuclear role of RGS14.

Specific roles for RGS14 in the nucleus remain a mystery. RGS14 lacks a DNA-binding motif but may nonetheless regulate the expression of synaptic proteins critical for late-term LTP and memory in neurons. This could occur through RGS14 interactions with accessory proteins, chromatin remodeling, mRNA splicing machinery, or other nuclear processes, as is the case for other closely related RGS proteins (48–52). Consistent with this idea, native RGS14 has been observed to exist at intranuclear membrane channels and within both chromatin-poor and chromatin-rich regions of a neuronal cell line (28). In addition, a subset of nuclear RGS14 localizes adjacent to active RNA polymerase II. Together, these findings suggest potential roles for RGS14 in transcriptional regulation (28, 53, 54). Even so, our pilot studies here with cultured hippocampal neurons showed no differences in the number or variety of mRNA transcripts in neurons expressing either normal RGS14, or RGS14 excluded from the nucleus or RGS14 trapped in the nucleus (Fig. S6). Of note, the neurons in this study were at rest and in culture. Future studies examining mRNA expression in CA2 punches from WT and LR mice could elucidate potential regulated transcripts as well as differential splice variants. Though speculative, a possibility is that synaptic activation could redirect a subpopulation of RGS14 and/or other regulatory factors to the nucleus to modulate transcription. As we show here, human variants target RGS14 to the nucleus constitutively, which would disrupt any temporal regulation or necessary posttranslational modification of RGS14 necessary for nuclear functions and would confer a different phenotype than would be exhibited by normal nuclear shuttling of RGS14. Ideally, changes in mRNA would be examined in WT or LR hippocampal slices following an LTP pairing protocol; however, gene dysregulation is a known issue following slice preparation (55) and thus would complicate result interpretation.

Another possible nuclear role for RGS14 may involve synapse-nucleus signaling, which is important for spatial memory (56) and has been postulated to facilitate synapse-specific targeting of newly synthesized mRNAs or proteins following an LTP stimulus (57–60) as mentioned earlier. Signals from the activated synapse must reach the nucleus, and newly synthesized mRNA must reach the activated synapse within a relatively short window in order to support synaptic strengthening and maturation (58). How the neuron coordinates this has become generally accepted under the synaptic tagging and capture (STC) hypothesis (58, 61). RGS14 could play a role in either retrograde or anterograde trafficking, coordinating signals from the spine to the nucleus or coordinating cargo delivery back to the spine. In support of this idea, RGS14 has been found to colocalize with cytoskeletal

elements, including microtubules, and influences tubulin polymerization (28, 62, 63), which enables trafficking of cargo to the tagged synapse (64, 65). As a multifunctional scaffold, RGS14 integrates G protein, Ca<sup>++</sup>/CaM/CaMKII, H-Ras/ERK, and 14-3-3 $\gamma$  signaling (17, 18, 22, 37, 66, 67), all critical mediators of both early- and late-stage LTP and learning (68–71), and thus could be acting as a scaffold to coordinate signaling between synapse and nucleus. Importantly, while some studies suggest that proteins at the activated synapse translocate to the soma/nucleus, others indicate that action potentials stimulate nuclear import of somatic molecules to mediate transcription (57), suggesting that a nuclear/somatic pool of RGS14 could have distinct functions from a spino-dendritic pool of RGS14.

The simplest explanation for RGS14 targeting to the nucleus is to redirect and sequester the protein away from the spine as a mechanism by the neuron to remove the brake on LTP, with no specific nuclear function (*i.e.*, time-out). RGS14 prepositioning at the spine allows for immediate participation in, and regulation of, signaling events that drive LTP. Removing the available pool of RGS14 from spines may be one way in which the cell can allow for differential regulation of these LTP-linked pathways. The discordant phenotype of nuclear-bound RGS14 LR *versus* RGS14 KO suggests that RGS14 is not simply passively hiding out in the nucleus, but likely confers an expression- and localization-dependent outcome on nuclear processes necessary for spatial learning. Ongoing and future studies will explore these possibilities.

In addition to the hippocampus, RGS14 is natively expressed in the piriform cortex, caudate, putamen, globus pallidus, substantia nigra, and amygdala of primate brain (32). Our studies here confirm for the first time in adult rodents the expression of RGS14 in brain regions outside of the hippocampus, including the striatum and amygdala, and raise questions about RGS14 roles beyond the hippocampus. One feature these brain regions all share is experience-based synaptic plasticity (72–74), suggesting that RGS14 may serve as a broad regulator of plasticity in these brain regions. Within the striatum, RGS14 could be regulating changes in learned behavior linked to reward and addiction (75). Other RGS proteins have been implicated in various measures of drug seeking and drug reward behavior (76). For example, both RGS9-2 and RGS7 in the striatum modulate locomotor responses to cocaine (77, 78). Within the amygdala, RGS14 could be regulating fear memory and stress responses (74). Consistent with this idea, a recent report showed that a global loss of RGS14 (RGS14 KO) enhances certain fear behaviors in female mice (79). Although the report demonstrates that this behavioral phenotype is linked to the known RGS14-rich CA2 region of the hippocampus, the findings do not rule out contributions of RGS14 within the CeA, a region long known to mediate behaviors associated with fear. These questions motivate current and ongoing studies.

Our findings here were made possible by the study of natural genetic variation within the human RGS14 gene. Genes/proteins are typically studied as canonical reference sequences (defined as normal) that do not reflect the rich genetic diversity in the human population, which can have profound

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impacts on protein function and linked physiology. Regulatory proteins, such as those in the RGS family, that contribute to complex signaling cascades (15, 80, 81) are more likely to contribute to complex polygenic diseases or physiological traits, particularly in combination with variants in other proteins. The variants chosen for study were selected from a data set of human variants within the RGS protein family (4). While these variants were derived from a “healthy” population, they may nevertheless impart important functional consequences in human carriers. As an example, genetic variants within the noncoding region of RGS16 are linked with self-reported “morning people” (82, 83), a human trait not captured by studies focused exclusively on disease-linked SNPs. RGS14 protein harboring rare variants LR or RQ could lead to aberrant signaling in synaptic plasticity in the striatum or amygdala, for example, contributing to a predisposition to fear learning or addiction, among others. Within the hippocampus, variants LR and RQ tip the nucleo-cytosolic-membrane shuttling balance in favor of the nucleus, leading to functional consequences in RGS14 modulation of plasticity. Indeed, protein mislocalization due to a coding variant has been shown to contribute to protein dysfunction in other complex diseases (84), underscoring the importance of signaling proteins being in the right place, at the right time, for the right amount of time for proper function.

In summary, we show that RGS14 harboring human genetic variants exhibits aberrant subcellular trafficking and mislocalization that results in loss of function in dendritic spines. While inactivating the RGS or R1 domain individually did not indicate these domains were required, *per se*, for function, removing the entire protein from the spines by mislocalization suggests potential redundancy or complexity in how these domains coordinate inhibitory signaling on LTP. As the precise mechanism for how RGS14 inhibits LTP is not known, we propose a working model suggesting that RGS14 regulates multiple processes for both early and (potentially) late LTP at the spines and nucleus, respectively (Fig. 9). At postsynaptic spines, RGS14 regulates GPCR/G protein, H-Ras/ERK,  $\text{Ca}^{++}/\text{CaM}$ , and other signaling events to suppress LTP (15, 18, 19, 85) (Fig. 9A). Mislocalization and nuclear accumulation of RGS14 due to genetic variant LR (or variant RQ, to a lesser extent) leads to absence of RGS14 from spines and dendrites, rendering the protein unable to modulate early LTP signaling (Fig. 9B). Overall, our findings here highlight the importance of dynamic spatial regulation of signaling proteins in the context of synaptic plasticity and demonstrate the contribution of naturally occurring rare genetic variation to RGS14 compartmental equilibrium. These data underscore the importance of considering not only the function of proteins derived from “canonical” sequences, but also those representing diverse human populations.

### Experimental procedures

#### Constructs and reagents

Leptomycin B was obtained from Sigma, and cells were treated at 20 nM for 2 h. Rat RGS14 was cloned into pRLucN2 as described previously (17), pLic-GST as described

previously (32), and pAAV-hSyn-YFP using restriction sites AgeI and HindIII. pET-His-hRan-Q69L was acquired from AddGene (catalog #42048), and pGEX-TEV-hCRM1 (XPO1) was kindly provided by Dr YuhMin Chook (UT Southwestern) (86). Flag-hCRM1 (XPO1) was obtained from AddGene (catalog # 17647). pET-Ran(Q69L) was obtained from Add Gene (catalog #42048). Isopropyl  $\beta$ -D-1-thiogalactopyranoside (IPTG) was purchased from Fisher Scientific. Primers used to make mutant constructs are included in Table S1. Anti-RGS14 was obtained from Proteintech. Anti-XPO1 was purchased from Sigma.

#### Purification of recombinant protein and measurement of pure protein-protein interactions

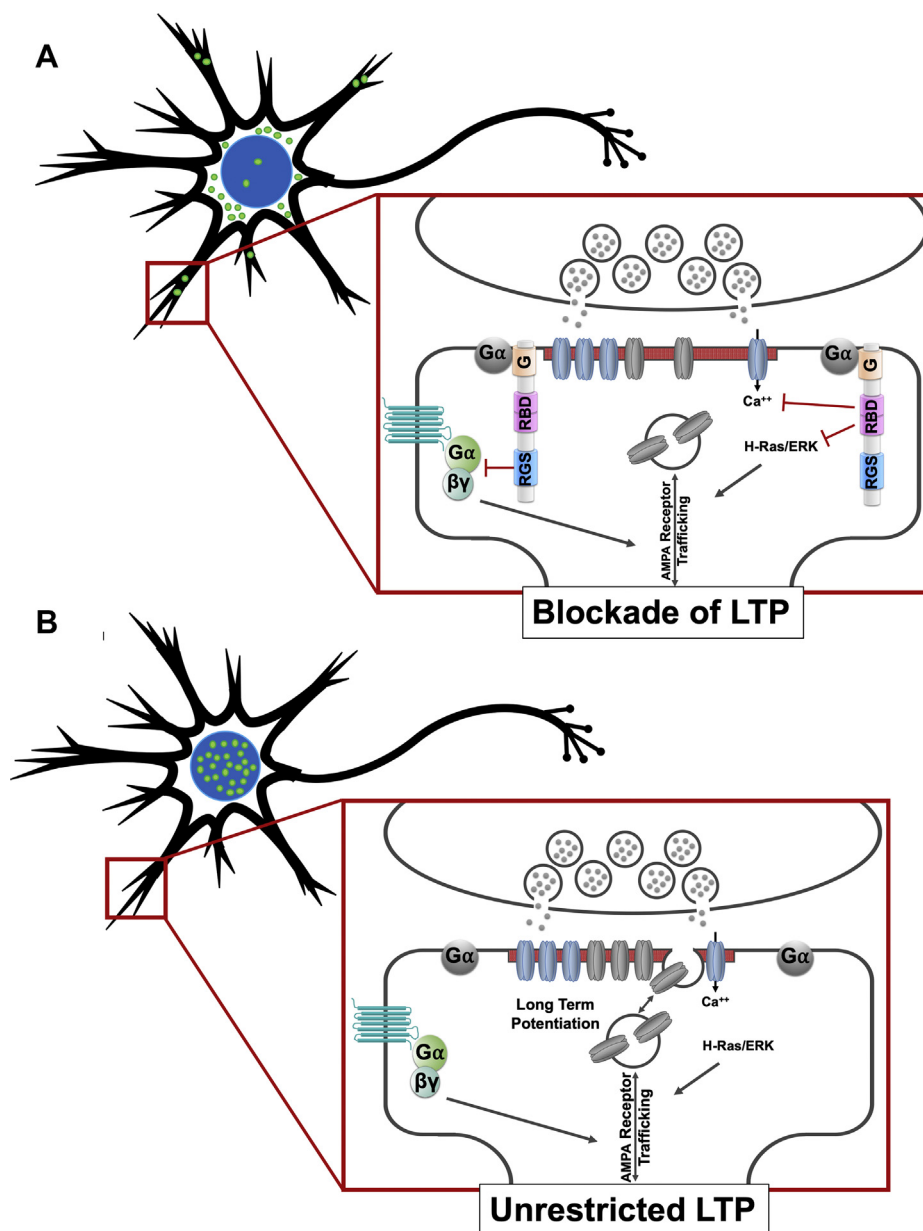
RGS14 was purified as described previously (25). Briefly, MBP-TEV-RGS14 was expressed in BL21 (DE3) *Escherichia coli* (*E. Coli*), pellets were resuspended in 50 mM HEPES, 200 mM NaCl, 10% glycerol, and 2 mM BME. Bacterial lysate was passed over a glutathione affinity column, washed with phosphate buffered saline (PBS), eluted with 10 mM reduced glutathione in 50 mM Tris (pH 8), and cleaved overnight with TEV protease. Purity was verified by coomassie. His-Ran Q69L (Ran QL) was purified as follows. BL21 bacterial cells were transformed with His-Ran QL, and protein expression was induced with IPTG at 37 °C for 2 h. Bacterial pellets were resuspended in 50 mM HEPES, 150 mM NaCl, 10% glycerol, 10 mM GTP, 2 mM BME, 20 mM imidazole, 2 mM  $\text{MgCl}_2$ , and PMSF. Lysates were then passed over a  $\text{Ni}^{2+}$  affinity column, washed with resuspension buffer and finally eluted with resuspension buffer containing 200 mM imidazole. Purity was verified by coomassie. GST-CRM1 (Exportin 1, or XPO1) was purified as described by the Chook Lab (86) as follows. BL21 bacterial cells expressing GST-CRM1 were induced with IPTG overnight at 25 °C. Lysate (40 mM HEPES pH 7.5, 2 mM  $\text{MgSO}_4$ , 200 mM NaCl, 5 mM dithiothreitol (DTT), 10% glycerol, and protease inhibitors) was then passed over a glutathione affinity column (Sepharose 4B, GE Healthcare), washed with buffer (40 mM HEPES, 2 mM  $\text{MgSO}_4$ , 150 mM NaCl, 2 mM DTT), and beads were resuspended in PBS. CRM1/XPO1 was cleaved from the beads by incubating with TEV protease overnight at 4 °C. Purity was verified by coomassie. Immunoprecipitation with FLAG and immunoblotting were performed as described previously (26, 37).

#### Structural modeling

The model depicting RGS14-NES/XPO1 interactions was generated in PyMol (The PyMOL Molecular Graphics System, Version 2.0 Schrödinger, LLC.) by structurally aligning the RGS14 GPR peptide/Gai1 structure (PDB code 1KJY) with the XPO1/mDia2 structure (PDB code 5UWP). Figures were constructed using PyMol.

#### Human genetic variants and lollipop plot

Human variant information was obtained from the Genome Aggregation Database (GnomAD version 2.0) at the Broad Institute (<http://gnomad.broadinstitute.org>) (33). Data was



**Figure 9. Proposed working model of RGS14 spatial-dependent inhibition of LTP.** *A*, wild-type RGS14 (green dot) is dispersed throughout the neuron, occupying multiple compartments including the nucleus, the cytoplasm, and the dendrites and spines (red box). Within a dendritic spine (expanded red box), Gα-GDP interaction with the G protein regulatory motif (G) brings RGS14 to the microcompartment necessary for inhibition of GPCR signaling by the RGS domain, inhibition of H-Ras/ERK signaling by the Ras binding domains (RBD), and inhibition of Ca<sup>++</sup> signaling, all of which support LTP in the absence of RGS14. *B*, variant forms of RGS14 (green dots) are sequestered in the nucleus and cannot reach dendrites (red box). Due to their mislocalization, they cannot interact with, and be recruited by, Gα-GDP to dendritic compartments (expanded red box). The lack of RGS14 at these microcompartments allows for unrestricted GPCR, H-Ras/ERK, and Ca<sup>++</sup> signaling, removing the block on LTP.

sorted by missense and synonymous annotation. The lollipop plot was then generated by open source code available on GitHub (87). Human variants were generated in rat RGS14 (of which the GPR motif shares 100% identity compared with human, but the amino acid number is less 1) and are reported in this article as such. Primers used to make the human variants as well as functional mutations are listed in Table S1.

#### **Bioluminescence resonance energy transfer (BRET)**

BRET was performed as described previously (17, 25, 88). Briefly, HEK293 cells were maintained in 1x Dulbecco's Modified Eagle Medium (Mediatech, Inc) without phenol red

and supplemented with 10% fetal bovine serum (FBS), 2 mM L-Glutamine, and 1% penicillin/streptomycin (VWR, Calbiochem, and Invitrogen, respectively). Cells were transiently transfected with polyethyleneimine and RGS14-Luciferase(Luc) plus Gαi1-YFP (89) for 40 h. Cells were resuspended in Tyrode's Solution (140 mM NaCl, 5 mM KCl, 1 mM MgCl<sub>2</sub>, 1 mM CaCl<sub>2</sub>, 0.37 mM NaH<sub>2</sub>PO<sub>4</sub>, 24 mM NaHCO<sub>3</sub>, 10 mM HEPES, and 0.1% glucose, pH 7.4) and plated at 10<sup>5</sup> cells per well in a 96-well Optiplate (PerkinElmer Life Sciences). YFP expression was quantified using a TriStar LB 941 plate reader (Berthold Technologies) at 485 nm excitation and 530 nm emission. Next, 5 μM coelenterazine H

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(Nanolight Technologies) was incubated for 2 min, and BRET measurements were taken at 485 nm (Luc emission) and 530 nm (YFP emission). The Net BRET ratio was quantified as follows: (530 nm signal/485 nm signal) – (485 nm signal from Luc alone). Acceptor/donor ratios were calculated as follows: (530 nm signal from YFP measurement)/(485 nm signal from BRET measurement). Each experiment was repeated three times.

### Dissociated hippocampal neuronal culture

Our protocol for culturing neurons was adapted from Beaudoin *et al.*, 2012 (90). Brains were removed from E18 to 19 embryos obtained from a timed pregnant Sprague-Dawley rat (Charles River). The meninges were removed, and the hippocampi were isolated in calcium and magnesium-free HBSS (Invitrogen) supplemented with 1x sodium pyruvate (Invitrogen), 0.1% glucose (Sigma-Aldrich), and 10 mM HEPES (Sigma-Aldrich) pH 7.3. Isolated hippocampi were washed with the HBSS solution, then dissociated using the same buffer containing 0.25% trypsin (Worthington) for 8 min at 37 °C. Trypsinized hippocampi were washed two times with the same HBSS buffer before being triturated 5 to 6 times with a fire-polished glass Pasteur pipette in BME (Invitrogen) supplemented with 10% FBS (VWR), 0.45% glucose (Sigma-Aldrich), 1x sodium pyruvate (Invitrogen), 1x Glutamax (Invitrogen), and 1x penicillin/streptomycin (HyClone). Neurons were counted and plated at a density of 80,000 cells/cm<sup>2</sup> in the BME-based buffer on coverslips that had been etched with 70% nitric acid (Sigma-Aldrich) before being coated with 1 mg/ml poly-L-lysine (Sigma-Aldrich) in borate buffer. Cells were allowed to adhere for 1 to 3 h before media was changed to Neurobasal (Invitrogen) supplemented with 1x B27 (Invitrogen) and 1x Glutamax (Invitrogen). Neurons were kept in a 37 °C, 5% CO<sub>2</sub> incubator, and half of the media was replaced with new Neurobasal every 3 to 4 days until neurons were used for experiments.

### Confocal microscopy imaging

All reactions were carried out at room temperature. Cells were fixed with 4% paraformaldehyde for 10 min, then quenched with 0.75% glycine in 200 mM Tris pH 7.4, for 5 min. Cells were then permeabilized for 10 min in 0.1% Triton-X in PBS and stained with Hoechst (1:12,500 in PBS) for 4 min. Cells were finally washed three times in PBS and mounted onto slides with ProLong Diamond Antifade mountant (Thermo Fisher). Images were taken on an Olympus FV1000 confocal microscope at 60x, then processed using ImageJ software. Approximately ten images per condition were obtained and representative images are shown.

### Hippocampal slice preparation and electrophysiological recordings

Hippocampal slices were prepared from wild-type C57Bl/6J mice (postnatal 5–8 days). Mouse pups were briefly placed in 70% ethanol, rapidly sacrificed by decapitation, and then brains were rapidly removed. Recovered brains were sectioned into

coronal slices (250 μm thick) by vibratome (Leica VT1200 S) in ice-cold cutting solution (MEM (GIBCO cat # 61100), NaHCO<sub>3</sub> 26 mM, HEPES 25 mM, Tris 15 mM, Glucose 10 mM, MgCl<sub>2</sub> 3 mM). Under a dissecting microscope, hippocampal slices were removed and then carefully placed in a 0.4-μm pore size membrane insert (Transwell 3450-Clear, Corning Incorporated) 6-well plate. Slices (1–3 slices per well) were cultured at 37 °C and 5% CO<sub>2</sub> with the medium changed twice per week. Once cultured slices were stable, AAV-YFP-RGS14 (WT or mutant) was injected into CA1 and incubated for 10 to 14 days.

For electrophysiological recordings, slices were transferred to a recording chamber including ACSF containing (in mM): 124 NaCl, 2.5 KCl, 2 MgCl<sub>2</sub>, 2 CaCl<sub>2</sub>, 1.25 NaH<sub>2</sub>PO<sub>4</sub>, 26 NaHCO<sub>3</sub>, and 15 glucose. For the pairing-induced LTP (pairing postsynaptic depolarization with presynaptic stimulation) experiments, the internal solution contained the following (in mM): 115 Cs-methanesulfonate, 20 CsCl, 2.5 MgCl<sub>2</sub>, 0.6 EGTA, 10 HEPES, 4 Na<sub>2</sub>-ATP, 0.4 Na<sub>2</sub>-GTP, and 10 phosphocreatine disodium salt. LTP was induced in whole-cell voltage clamp mode by pairing a postsynaptic depolarization to 0 mV with presynaptic stimulation delivered at 3 Hz (270 pulses over 1.5 min).

For imaging of slice cultures, slices were transferred to 4% paraformaldehyde (PFA) in PBS for fixation at room temperature for 2 h and then stored in PBS at 4 °C until use. Twenty minutes prior to imaging, slices were incubated in 60% 2,2'-thiodiethanol to clear the tissue (91) and then imaged on a Zeiss 880 laser scanning confocal microscope (Carl Zeiss AG, Oberkochen, Germany).

### Generation of RGS14 (L507R) CRISPR mouse line

To generate the RGS14 point mutation, a CRISPR gRNA (CTGGTGGAGCTGCTGAATCGGG) targeting exon 15 along with a donor oligo (5'-CAGCAGAATGGTCAAGCAT TGGTGTAGGTGCTTTGGCCATATCGGCCCTGACTCTG TGGCTCCCTCCAGGCCTGGTGGAGCTGCGGAATCGAG TGCAGAGCAGCGGGGCCACGATCAGAGAGGACTTCT TCGCAAAGAGGACCTGGTCCCTTCCAGAATTTCTGCAG CTTCCCTTCCCAA-3'; underlined bases are engineered) was designed to generate a CTG-to-CGG change creating the L507R (equivalent to L505R in human sequence and L504R in rat sequence) point mutation as well as G-to-A SNP conversion to create a PAM blocking mutation and G-to-A silent changes to create a novel TaqI restriction site in Rgs14 mouse that could be used for screening.

To generate the RGS14 (L507R) mice, the gRNA (20 ng/μl), oligo donor (20 ng/μl), and CRISPR protein (20 ng/μl) were injected into 1-cell C57Bl/6J zygotes and subsequently transplanted at the 2-cell stage into C57Bl/6J pseudopregnant females by the Emory Mouse Transgenic and Gene Targeting Core. Genomic DNA from toes was amplified *via* PCR using primers: FW 5'-GTGGCATCAGAGAGGCTG-3' and RV 5'-GTTACACAGATGCCAGAGGAC-3'. PCR bands at 387 bp (wild-type), 188 bp, and/or 199 (mutant) were produced. Of the 95 pups born, nine of them (approx. 10%) were positive for the mutation. Two positive animals were selected initially for

continued breeding. To dilute out/eliminate potential off-target mutations due to CRISPR/Cas9 gene editing, each animal was back-crossed (Het-Het) against a fresh wild-type C57Bl/6J background for three generations, and then a single line was continued for back-crossing for five generations prior to behavioral tests.

### Morris water maze behavioral test

All animal studies were approved by the Emory University IACUC. Morris Water Maze was conducted as described previously (21). Adult wild-type RGS14 and homozygous RGS14 LR mice (littermates ages 2–6 months) were used. The water maze consisted of a circular swim arena (diameter of 116 cm, height of 75 cm) surrounded by extramaze visual cues that remained in the same position for the duration of training and filled to cover the platform by 1 cm at 22 °C. Water was made opaque with nontoxic, white tempera paint. The escape platform was a circular, nonskid surface (area 127 cm<sup>2</sup>) placed in the NW quadrant of the maze. Acquisition training consisted of 5 test days with four daily trials. Mice entered the maze facing the wall and began each trial at a different entry point in a semirandom order. Trials lasted 60 s or until the animal mounted the platform with a 15-min intertrial interval. A probe trial was conducted on day 6 wherein the platform was removed, and the animal swam for 60 s, and the time spent in the target quadrant (NW) *versus* the adjacent and opposite quadrants was recorded. A video camera mounted above the swim arena and linked to TopScan software recorded swim distance, swim speed, and time to platform and was used for tracking and analysis. Statistics were ANOVA and post hoc Dunnett's test.

### RGS14 LR CRISPR mouse immunofluorescence

Male RGS14 WT (n = 3) and RGS14 LR (n = 3) mice were euthanized with an overdose of sodium pentobarbital (Fatal Plus, 150 mg/kg, i.p.; Med-Vet International, Mettawa, IL) and transcardially perfused with 4% paraformaldehyde dissolved in 0.01 M PBS. After extraction, brains were postfixed for 24 h in 4% paraformaldehyde at 4 °C and subsequently transferred to 30% sucrose/PBS solution for 72 h at 4 °C. Brains were embedded in OCT medium (Tissue-Tek; Sakura, Torrance, CA) and serially sectioned by cryostat (Leica) into 40- $\mu$ m coronal slices. Brain sections were stored in 0.01 M PBS (0.02% sodium azide) at 4 °C before IHC.

Immunohistochemistry (IHC) was performed to visualize RGS14 protein in brain sections at the level of the hippocampus and striatum. Brain sections from mice of both genotypes were subjected to 3 min antigen retrieval at 100 °C in 10 mM sodium citrate buffer. Following antigen retrieval, brain sections were blocked for 1 h at room temperature in 5% normal goat serum (NGS; Vector Laboratories, Burlingame, CA) diluted in 0.01 M PBS/0.1% Triton-X permeabilization buffer. Sections were then incubated for 24 h at 4 °C in this same NGS blocking/permeabilization buffer now including a mouse primary antibody raised against RGS14 (NeuroMab; clone N133/21) at 1:500 dilution. After washing, sections were

then incubated for 2 h in blocking/permeabilization buffer with goat antimouse AlexaFluor 568 (Invitrogen, Carlsbad, CA) at 1:500 dilution. After washing, the sections were mounted onto Superfrost Plus slides (Thermo Fisher Scientific, Norcross, GA) and coverslipped with Fluoromount-G plus DAPI (Southern Biotech, Birmingham, AL).

Fluorescent micrographs of immunostained sections were acquired on a Leica DM6000B epifluorescent upright microscope at 5, 10, or 20x magnification. Micrographs were acquired for dorsal hippocampal Cornu Ammonis fields CA1 and CA2, piriform cortex (Pir Ctx), CeA, dorsal striatum (DS), and nucleus accumbens (NAc). For each regional comparison of RGS14 immunostaining between genotypes, uniform exposure parameters were used. All images were processed with ImageJ (NIH) to reduce background and enhance contrast.

### Next-generation RNA-sequencing and data analysis

Cultured rat hippocampal neurons (DIV18) were infected with AAV2/9 virus expressing either RGS14 WT (n = 4) or RGS14 carrying mutations RKK/AAA (NLSm) (n = 3) or RGS14 LL/AA (NESm) (n = 3). Infected neurons then were harvested and RNA was collected (Qiagen RNeasy Mini Kit). Purified RNA (500 ng) was converted to sequencing libraries using the KAPA mRNA HyperPrep kit. To perform RNA-seq, libraries were pooled at equimolar ratio and sequenced on a NextSeq500 using 75 bp paired-end sequencing. Data processing involved raw fastq reads mapped to the rn6 version of the rat genome. High percentage of reads were mapped (>90%) and were unique (orange bar), indicating high-quality libraries. Fastq files were mapped to the RGSC 6.0/rn6 genome using STAR (92) and gene coverage for all exons annotated using GenomicRanges (93). To identify detected transcripts, genes are filtered for detection based on being expressed at > 3 reads per million in at least three samples (first blue dotted line) resulting in the detection of 10,324 out of 17,324 transcripts. To analyze samples for differential gene expression and sample variation, all detected genes were used as input for edgeR v3.24.3 (94) to identify differentially expressed genes. Data sets were subjected to principal component analysis of variation using the princomp function in R v3.5.2 between samples to examine interclustering and differences between the groups.

### Data availability

All data are contained within the article.

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**Abbreviations**—The abbreviations used are: BRET, bioluminescence resonance energy transfer; CeA, central amygdala; DS, dorsal striatum; EPSCs, excitatory postsynaptic currents; FBS, fetal bovine serum; GPR, G protein regulatory; IHC, Immunohistochemistry; IPTG, Isopropyl  $\beta$ -D-1-thiogalactopyranoside; LMB, Leptomycin B; LTP, long-term potentiation; NAc, nucleus accumbens; NES, nuclear export sequence; NLS, nuclear localization sequence; PBS, phosphate buffered saline; PFA, paraformaldehyde; RGS, regulator of G protein signaling; STC, synaptic tagging and capture.

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