

Gravin Orchestrates Protein Kinase A and β 2-Adrenergic Receptor Signaling Critical for Synaptic Plasticity and Memory

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A kinase-anchoring proteins (AKAPs) organize compartmentalized pools of protein kinase A (PKA) to enable localized signaling events within neurons. However, it is unclear which of the many expressed AKAPs in neurons target PKA to signaling complexes important for long-lasting forms of synaptic plasticity and memory storage. In the forebrain, the anchoring protein gravin recruits a signaling complex containing PKA, PKC, calmodulin, and PDE4D (phosphodiesterase 4D) to the β 2-adrenergic receptor. Here, we show that mice lacking the α -isoform of gravin have deficits in PKA-dependent long-lasting forms of hippocampal synaptic plasticity including β 2-adrenergic receptor-mediated plasticity, and selective impairments of long-term memory storage. Furthermore, both hippocampal β 2-adrenergic receptor phosphorylation by PKA, and learning-induced activation of ERK in the CA1 region of the hippocampus are attenuated in mice lacking gravin- α . We conclude that gravin compartmentalizes a significant pool of PKA that regulates learning-induced β 2-adrenergic receptor signaling and ERK activation in the hippocampus *in vivo*, thereby organizing molecular interactions between glutamatergic and noradrenergic signaling pathways for long-lasting synaptic plasticity, and memory storage.

Introduction

A-kinase anchoring proteins (AKAPs) are a family of functionally distinct but structurally related proteins that localize protein kinase A (PKA) and other signaling molecules to specific subcellular compartments (e.g., synapses) (Welch et al., 2010). Genetic and pharmacological inhibition of PKA anchoring impairs hippocampal synaptic plasticity and memory (Moita et al., 2002; Snyder et al., 2005; Huang et al., 2006; Nie et al., 2007; Nijholt et al., 2007), but the specific AKAPs involved in these PKA-

dependent processes, and the details of their exact functions, have not been clearly identified.

Genetic analysis of AKAP complexes involved in mediating synaptic plasticity and memory has focused on the AKAP5 gene (also known as AKAP150 in mice), because it is associated with AMPA and NMDA receptors (Colledge et al., 2000). Loss of AKAP150 disrupts hippocampal LTD, alters AMPA receptor-mediated synaptic transmission, and results in memory deficits in the Morris water maze (Tunquist et al., 2008). Surprisingly, a genetic mutation leading to a selective disruption of PKA-AKAP150 binding disrupts long-term potentiation (LTP) in adult mice (Lu et al., 2007) but does not induce a memory deficit in the Morris water maze (Weisenhaus et al., 2010). These findings suggest that either (1) other AKAP150 binding partners or (2) distinct PKA-AKAP complexes contribute to hippocampal synaptic plasticity and memory formation.

β -Adrenergic receptors (β -ARs) play a critical role in emotional arousal, metaplasticity, and long-term memory formation (McGaugh, 2000; Gibbs and Summers, 2002) through activation of the cAMP signaling pathway (O'Dell et al., 2010). Occupancy of β 2-adrenergic receptors (β 2-ARs) by their ligand noradrenaline (norepinephrine) leads to the coupling of the receptor with guanine nucleotide regulatory proteins (*G α s*), cAMP production, and the association of PKA with the β 2-AR (Shih et al., 1999; Lin et al., 2000). Phosphorylation of the β 2-AR by PKA switches the coupling from *G α s* to inhibitory guanine nucleotide regulatory protein (*G α i*), thereby activating *G α i*-linked signaling path-

Received July 28, 2012; revised Oct. 2, 2012; accepted Oct. 21, 2012.

Author contributions: R.H., G.S.B., J.D.S., and T.A. designed research; R.H., D.A.C., A.J.P., T.H., T.N., J.P.D., L.A.G., Q.G., and V.L. performed research; I.H.G. contributed unpublished reagents/analytic tools; R.H., D.A.C., A.J.P., T.H., and T.N. analyzed data; R.H. wrote the paper.

This work was supported by The Netherlands Organization for Scientific Research NWO-Rubicon Grant 825.07.029 (R.H.), the Kwanjeong Educational Foundation (A.J.P.), the Human Frontiers Science Program (T.A.), the National Institutes of Health (T.A.), National Institute on Alcohol Abuse and Alcoholism Grant R01 AA18060 (T.A.), and National Institute of General Medical Sciences Grant GM-48231 (J.D.S.). G.S.B. and J.P.D. were supported by Medical Research Council (United Kingdom) Grant G0600765 and Fondation Leducq Grant 06CVD02. We thank Dr. Jacqueline Barra for providing us with the gravin GT mouse line. We thank Drs. Seth Grant, Marcelo Coba, and members of the Abel Laboratory for helpful comments on a previous version of this manuscript, and Dr. Carlos Sindreu for technical advice.

The authors declare no competing financial interests.

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DOI:10.1523/JNEUROSCI.3612-12.2012

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ways implicated in memory such as the extracellular-regulated kinase 1/2 (ERK1/2) pathway (Daaka et al., 1997; Atkins et al., 1998; Baillie et al., 2003; Sindreu et al., 2007). One anchoring protein that recruits PKA to the β 2-AR upon receptor occupancy is the anchoring protein gravin (Tao et al., 2003). Gravin, also known as Src-suppressed C kinase substrate (SseCKS), AKAP12, and AKAP250 (Lin et al., 1996; Willoughby et al., 2006), is a multivalent scaffolding protein that binds PKA and PKC (Nauert et al., 1997), sequesters calmodulin (Lin and Gelman, 2002), and targets phosphodiesterase 4D (PDE4D) to the plasma membrane (Willoughby et al., 2006).

Although gravin modulates β 2-AR signaling complexes *in vitro*, it remains to be determined whether gravin plays a role in hippocampal synaptic plasticity and memory formation *in vivo*. We answered this important question by using a gene-trap mouse model (referred to as gravin GT mice) that lacks the α -isoform of gravin (Camus et al., 2001) and defined the role of gravin- α in β 2-AR-mediated signaling events important for learning-induced ERK1/2 activation, long-lasting forms of synaptic plasticity and long-term memory.

Materials and Methods

Animals

The gravin GT mice, also referred to as SseCKS mutants (Camus et al., 2001), were a gift from Dr. Jacqueline Barra (Institut Pasteur, Paris, France) and backcrossed to a C57BL/6J background for >10 generations. Male and female 3- to 5-month-old gravin GT mice and wild-type littermates had *ad libitum* access to food and water and were maintained on a 12 h light/dark cycle with behavioral testing occurring during the light phase. Experimentally naive mice were used for each experiment, individually housed for 1 week, and handled for 2 min for 3 consecutive days before the behavioral experiments. All experiments were conducted according to National Institutes of Health guidelines for animal care and use and were approved by the Institutional Animal Care and Use Committee of the University of Pennsylvania.

Genotyping

Tail DNA was used to genotype mice by PCR with three primers, two in the intron sequences of gravin between exon 1A2 and 1B (AGGGCTACCAGAGAGATCC and GCTGTTTGGGAGCAGAAAAG) and one at the 3' end of *LacZ* gene (GCAGAGCGAGGTATGTAGGC). The cycling parameters for PCR genotyping were 94°C, 1 min; 60°C, 1 min; and 72°C, 1 min for 40 cycles. PCR products consisted of one 562 bp band for homozygous GT mice, one 730 bp band for wild-type mice, or both 562 and 730 bp bands for heterozygous mice.

In situ hybridization

The sequence of the antisense oligonucleotide against gravin is CCGA-C AATGCCTCCAGGTCACCGACCTGGTC and designed by Sigma-Genosys. A sense probe control was accordingly designed. The hybridization was performed on 20 μ m sections of the brain of 2-month-old C57BL/6J wild-type mice as previously described (Abel et al., 1997).

Immunohistochemistry

Transcardial perfusions and immunohistochemical stainings were conducted as previously described (Havekes et al., 2006, 2007; Vecsey et al., 2009). Sections incubated with either rabbit Map2 antibody (Millipore; AB5622; 1:1000) or mouse gravin antibody (Sigma-Aldrich; JP74; 1:200) followed by incubation with the appropriate Alexa Fluor 488 antibodies (Invitrogen). Imaging was conducted on a Leica confocal microscope. In case of the phospho-ERK1/2 staining, we used rabbit anti-phospho-ERK1/2 (Cell Signaling; AB9101; 1:100), and phosphatase inhibitors were included in the fixative and all buffers (Sindreu et al., 2007). After incubation with primary antibody, sections were washed in PBS for 1 h followed by incubation with biotinylated-conjugated goat anti-rabbit IgG antibody (Jackson ImmunoResearch; 1:500) for 3 h. After washing in PBS for 1 h, sections were incubated with the avidin–biotin horseradish peroxidase complex (ABC kit; Vector Laboratories; 1:500). After 3 h of

washing in PBS, sections were processed with diaminobenzidine (0.02%) with 100 μ l of 0.1% H₂O₂ as a reaction initiator. Processing was visually monitored and stopped by rinsing with PBS. Cell counts and optical density measurements were conducted as described previously (Van der Borght et al., 2007; Hagewoud et al., 2011) using a Micromaster light microscope.

RNA isolation and cDNA synthesis

Hippocampi were dissected and stored at –80°C in 500 μ l of RNeasy Lysis Buffer (Ambion). RNA extraction and cDNA synthesis using the RETROscript (Ambion) kit were conducted as described previously (Vecsey et al., 2009).

Quantitative real-time reverse transcription-PCR

The reactions were assembled in 96-well optical reaction plates (Applied Biosystems) and covered with optical adhesive covers (Applied Biosystems). cDNA was diluted to 2 ng/ μ l with DEPC-treated H₂O and added in a quantity of 11.4 μ l per well. Each well also contained 1 μ l of 5 μ M primers and 12.4 μ l of Quantitect SYBR Green PCR master mix (QIAGEN). Five primer sets were used to assess the relative expression of gravin mRNA in the hippocampus. The *gravin- α 1a* and *gravin- α 1b* primer sets probe for transcripts spanning exon 1A1 to exon 2: primers *gravin- α 1a* (forward, 5'-GCTCAGTGGCCATGGGCCCCGAC; reverse, 5'-TGCCATTCTTTAGCTCGGTCTTTTTC), *gravin- α 1b* (forward, 5'-GCTGGAGATCCCGCTGAC; reverse, 5'-GTTTCTCTGCTCGT CCTT). *Gravin- α 2* probes for the α -isoform of gravin spanning exon 1A2 to exon 2 (forward, 5'-AAGAATGGTCAGCTGTCTGC; reverse, 5'-TGACAGTGAGTAGCTGGACG). These three primer sets all probe for the presence of the *gravin- α* transcript. The *gravin- β* primer and *LacZ* primer set probe for the β transcript (forward, 5'-AGGAGAAGGAGACTTCCTGC; reverse, 5'-TGACAGTGAGTAGCTGGACG), and *gravin-lacZ* transcript (forward, 5'-TGGCAAGAGGAAGAAGTCA; reverse, 5'-ATGTGAGCGAGTAACAACCCGTCGGATTCT), respectively. The reaction was run in the Applied Biosystems ABI Prism 7000 and began with a 2 min incubation at 50°C followed by 15 min at 90°C. Afterward, the reaction proceeded through 40 cycles of the following: (1) 95°C for 15 s; (2) 56°C for 30 s; and (3) 72°C for 30 s. Data were collected at stage 3 of each cycle and the products were examined with a dissociation curve consisting of two sets of sequential incubations at 95°C for 15 s and 60°C for 1 min. Data were normalized against actin G (*actg*), hypoxanthine phosphoribosyltransferase (*hprt*), and Tubulin (*tub4a*). Relative expression levels were calculated according to the procedure in the Applied Biosystems manual using the standard curve method.

Electrophysiology

Electrophysiological recordings were performed as described previously (Vecsey et al., 2009). LTP was induced using the following protocols: spaced four-train (four 1 s 100 Hz trains delivered 5 min apart), massed four-train (four 1 s 100 Hz trains delivered 5 s apart), theta-burst (bursts of four 100 Hz pulses delivered for a total of 3 s at 5 Hz), and one-train stimulus (one 1 s 100 Hz train of pulses). LTP was also induced by pairing a 5 Hz, 3 min stimulation with application of the 1 μ M β -adrenergic agonist isoproterenol (ISO) for 25 min. The electrical stimulation was given 15 min into ISO treatment. ISO (Sigma-Aldrich) was prepared daily as a 10 mM stock solution in Milli-Q water. Synthesis of stearated Ht31 (stHt31) (St-N-DLIEEAASRIVDAVIEQV KAAGAY-C) and pseudo-Ht31 peptide (stHt31P) (St-N-DLIEEAASRPVD AVPEQVKAAGAY-C) (Carr et al., 1992; Vijayaraghavan et al., 1997; Huang et al., 2006) was conducted by Quality Controlled Biochemicals. Lyophilized powder was resuspended at a stock concentration of 10 mM in 50 mM Tris-HCl, pH 7.0, and 0.05% DMSO and used at 10 μ M final concentration in aCSF. Forskolin was dissolved in 100% ethanol to make 50 mM stock solution and used at 50 μ M final concentration in aCSF. The β 2-AR antagonist 3-(isopropylamino)-1-[(7-methyl-4-indanyl)oxy]butan-2-ol (ICI 118551) (Sigma-Aldrich) was dissolved in H₂O at a stock concentration of 100 μ M and used at a 100 nM final concentration in aCSF.

Tissue lysates, Western blot analyses, and PKA activity assays

Proteins were separated by SDS-PAGE and transferred to PVDF membrane as described previously (Vecsey et al., 2009). The following primary antibodies were used: ERK1/2 (Cell Signaling; AB9102; 1:1000),

pERK1/2 (Cell Signaling; AB9101; 1:1000), β 2-AR (Santa Cruz; sc-569; 1:1000), p β 2-AR (Santa Cruz; sc-16718; 1:200), pGluA1 Ser845 (Millipore; AB5849; 1:1000), and GluA1 (Millipore; MAB2263; 1:50,000). After incubation with the primary antibodies, membranes were incubated with HRP-conjugated secondary antibodies for 1 h at room temperature (Santa Cruz; mouse secondary antibody, 1:1000; Santa Cruz; rabbit secondary antibody, 1:5000). PKA activity measurements were conducted on hippocampal lysates using a PKA activity ELISA kit according to the manufacturer's instructions (Arbor Assays; catalog #K027-H1).

Neuronal cultures

Primary neuronal cultures were prepared from hippocampi of embryonic day 17.5 mice, as described previously (Banker and Cowan, 1977). Dissociated neurons were plated onto poly-L-lysine-treated glass coverslips at a density of 25–50 cells/mm² and cocultured over a monolayer of astrocytes. Cells were maintained in Neurobasal medium (Invitrogen) supplemented with B27 and Glutamax. After incubation with the following primary antibodies: anti-PKA RII- α polyclonal (Santa Cruz; PKA RII- α sc-909; 1:200) and anti-gravin monoclonal (Sigma-Aldrich; G3795; 1:200), neurons were incubated with the appropriate Alexa Fluor-conjugated secondary antibodies for 2 h at room temperature. Neurons were imaged with a Zeiss confocal microscope.

Coimmunoprecipitation and RII overlays

Hippocampi were pulverized under liquid nitrogen and homogenized in HSE lysis buffer (20 mM HEPES, pH 7.2, 150 mM NaCl, 5 mM EDTA, 1% Triton X-100). Extracts were rocked for 1 h at 4°C and subsequently centrifuged at 19,400 \times g for 20 min at 4°C. Equal amounts of protein were subjected to immunoprecipitation with 5 μ g of anti-gravin (Sigma-Aldrich; G3795) by rocking overnight at 4°C. Lysates were recentrifuged at 19,400 \times g, and protein A and protein G beads (1:1 slurry) were added for 1 h. Beads were collected by centrifugation and washed four times in HSE lysis buffer. Bound proteins were eluted with 2 \times LDS sample buffer, separated on 4–12% SDS-PAGE gels (Bio-Rad) and transferred to nitrocellulose for immunoblotting. The following antibodies were used: anti-gravin monoclonal antibody (Sigma-Aldrich; G3795; 1:1000), anti-GAPDH (Sigma-Aldrich; G8795; 1:1000), anti-MAP2 (Sigma-Aldrich; M9942; 1:1000), anti- β -galactosidase (Promega; Z378A; 1:200), anti-pERK1/2 (Cell Signaling; AB9101; 1:1000), ERK1/2 (Cell Signaling; AB9102; 1:1000), and anti-AKAP150 (V088; 1:1000). Western blot data were collected using an Alpha Innotech MultiImage III with FluoroChemQ software. For the RII overlays, nitrocellulose membranes were blocked and then incubated with digoxigenin-labeled PKA RII overnight at 4°C. Binding was detected using HRP-labeled anti-digoxigenin (Abcam; ab62120).

Behavioral testing

Morris water maze. Training and testing in the hidden platform version of the water maze were performed as described previously (Abel et al., 1997). Naive mice were used in the visible platform experiment in the Morris water maze.

Object-place recognition task and novel object recognition task. Both tasks were performed as described previously (Oliveira et al., 2010).

Fear conditioning. Experiments were performed in standard conditioning chambers (San Diego Instruments) as described previously (Abel et al., 1997). Freezing behavior (defined as complete immobility except for breathing) was scored during the training and retention sessions.

Rotarod. Mice were placed on a 3-cm-diameter rotating rod elevated 16 cm and allowed to walk on it for 30 s at 4 rpm. Twenty-four hours later, mice were placed on the rotating rod again for no more than 5 min as it accelerated from 4 to 40 rpm. The time at which each mouse fell was recorded, and each was placed on the rod twice more at 1 h intervals. This procedure was repeated for 3 consecutive testing days.

Exploratory and locomotor activity. Baseline exploratory and locomotor activities were measured in a gray rectangular box (60 \times 50 \times 26 cm) built of polyvinyl chloride plastic. Mice were placed in the open field for 10 min. The time spent in peripheral and central areas and the velocity of travel were recorded using a video tracking system (Clever Systems).

Zero maze. Anxiety-related behavior was tested in the elevated zero maze as described previously (Tretter et al., 2009).

Data analysis

Data analysis was performed using SPSS. Behavioral data were analyzed using two-way ANOVAs (in some cases with repeated measures as the within-subject variable) or Student's *t* tests. Tukey–Kramer tests were used for *post hoc* analyses when necessary. Biochemical data were analyzed using independent *t* tests. Electrophysiological data were analyzed using nonparametric tests (Nie et al., 2007). The initial slope of the fEPSP at each time point was analyzed. For comparisons of the average slope over the last 20 min of the recording, the Mann–Whitney *U* test was used. Differences were considered statistically significant when *p* < 0.05. Data are plotted as mean \pm SEM.

Results

Hippocampal gravin- α but not gravin- β is lost due to gene trap insertion in gravin GT mice

In rodents, three gravin isoforms (α , β , and γ) are transcribed from a single gene locus (*gravin*). The transcription of each isoform is controlled by different promoters, yielding distinct mRNA transcripts and proteins that differ at their N termini (Streb et al., 2004). The α and β isoforms are expressed widely throughout the body, but the γ isoform is only found in the testis (Camus et al., 2001). The unique N-terminal myristoylation motif in gravin- α facilitates enrichment at the plasma membrane (Lin et al., 1996; Streb et al., 2004), where it can interact with G-protein-coupled receptors such as the β 2-AR (Shih et al., 1999). Gravin mRNA expression is observed in mouse forebrain regions (Fig. 1*A,B*), including the three major hippocampal regions [CA1, CA3, and dentate gyrus (DG)]. Moderate labeling was observed in the granule cells of the cerebellum (Fig. 1*C*). Immunohistochemical analyses confirmed the mRNA studies showing protein expression of gravin in the dorsal hippocampus (Fig. 1*D*). Higher magnification images reveal prominent gravin expression in the perinuclear region of CA1 and CA3 pyramidal cells (Fig. 1*E,F*) and dentate gyrus granular cells (Fig. 1*G*). Signals for gravin and the RII- α isoform of PKA overlap in cultured mouse hippocampal neurons (Fig. 1*H–J*), consistent with the role of gravin as an A-kinase anchoring protein.

To examine the role of this anchoring protein in hippocampal function, we used gravin GT mice because these animals give us the opportunity to exclusively assess the function of the myristoylated α -isoform of gravin. The gravin GT mouse line was generated by a gene trap insertion of the *LacZ* reporter into the genomic sequence encoding gravin between exon 1A2 and 1B (Fig. 1*K*) (Camus et al., 2001). A significant decrease in gravin- α mRNA levels was detected using three distinct primer sets: *gravin- α 1a* (both groups, *n* = 6; Student's *t* test, *p* = 0.009; Fig. 1*L*), *gravin- α 1b* (both groups, *n* = 3; Student's *t* test, *p* = 0.039; Fig. 1*L*), and *gravin- α 2* (both groups, *n* = 6; Student's *t* test, *p* = 0.013; Fig. 1*L*), all representing the *gravin- α* transcript. Gravin GT mice exhibit normal expression of the gravin- β isoform (both groups, *n* = 6; Student's *t* test, *p* = 0.444; Fig. 1*L*), suggesting that loss of the *gravin- α* transcript does not result in compensatory upregulation of the gravin- β transcript in the hippocampus.

Biochemical approaches were used to further characterize gravin GT mice. Western blot analyses indicated that the majority of full-length gravin was lost in these animals, with the remaining fraction representing the β -isoform of gravin (Fig. 1*M*, left panel). Gravin GT mice expressed a truncated form of gravin of ~150 kDa (Fig. 1*M*, left panel) that consisted of an N-terminal portion of the anchoring protein fused to lacZ (Fig. 1*M*, middle panel). The RII-overlay assay confirmed that this truncated fusion protein did not bind PKA (Fig. 1*M*, right panel). Because gravin and another anchoring protein, AKAP150, both coordinate β 2-AR-mediated signaling (Tao and Malbon, 2008), we

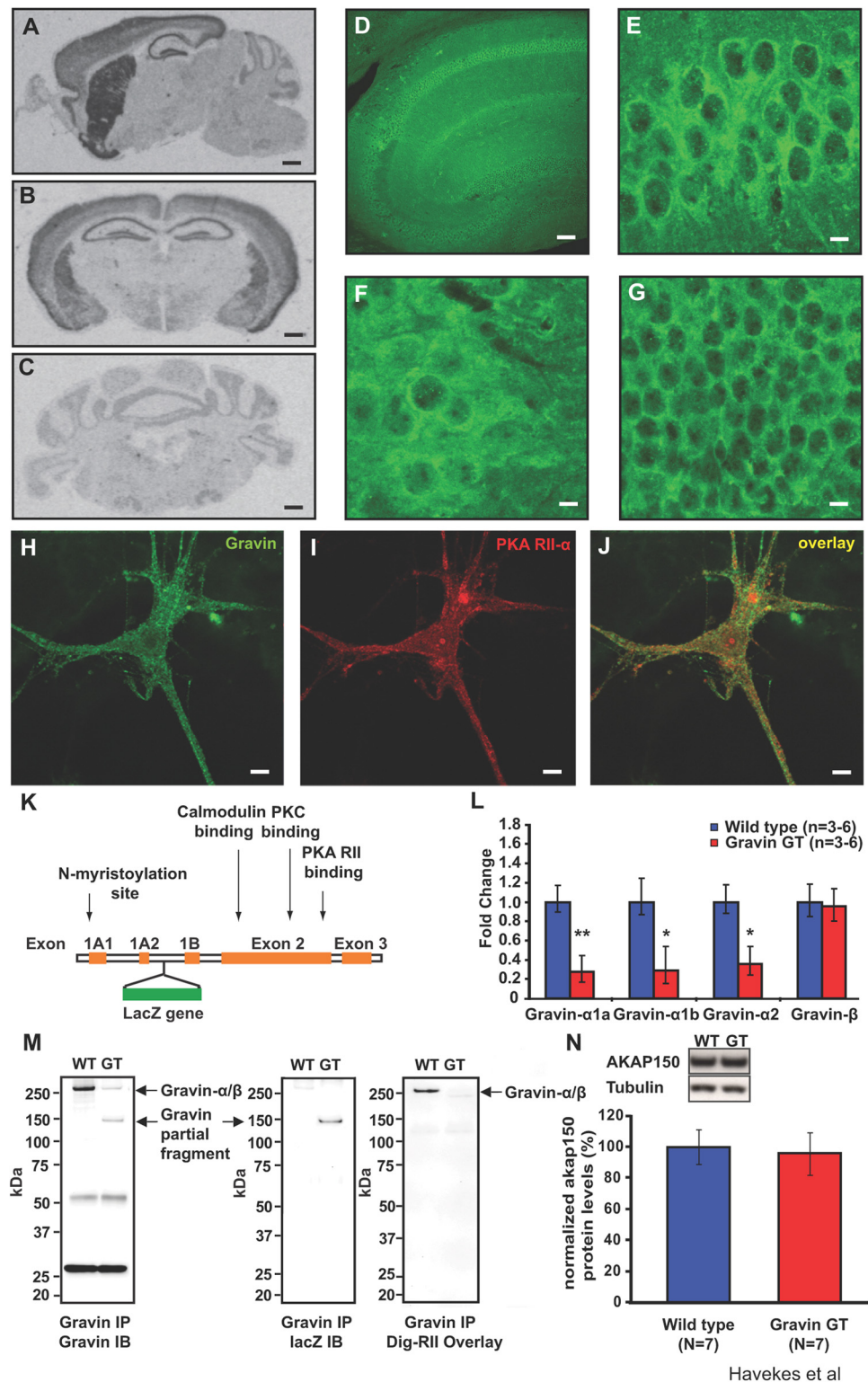


Figure 1. Expression of the α -isoform of gravin is reduced in gravin GT mice. **A**, A sagittal brain section showing gravin mRNA expression in the hippocampus, striatum, cortex, and olfactory bulb. **B**, A coronal brain section showing gravin mRNA in the hippocampus, cortex, amygdala, and striatum. **C**, Gravin mRNA is expressed at lower levels in the granular cell layer of the cerebellum. Scale bar, 800 μ m. **D**, Gravin protein expression in the dorsal hippocampus. Scale bar, 200 μ m. Higher magnification images of gravin protein expression in CA1 pyramidal cells (**E**), CA3 pyramidal cells (**F**), and granular cells of the dentate gyrus (**G**). Scale bar, 20 μ m. Gravin (**H**) and PKA RII- α (**I**) are strongly colocalized (**J**) in cultured mouse hippocampal neurons. Scale bar, 20 μ m. **K**, A schematic representation of the gravin gene. The *gravin*- α transcript consists of exons 1A1, 1A2, 2, and 3. The *gravin*- β transcript consists of exons 1B, 2, and 3. The gravin GT mice carry a LacZ insertion in the intron between exons 1A2 and 1B (Camus et al., 2001). This insertion site precedes the promoter region for the β transcript (Camus et al., 2001). **L**, Quantitative PCR experiments reveal reduced expression levels of *gravin*- α in gravin GT mice. mRNA levels for *gravin*- α 1a (Student's *t* test, $n = 6$, $p = 0.009$), *gravin*- α 1b (Student's *t* test, $n = 3$, $p = 0.039$), and *gravin*- α 2 (Student's *t* test, $n = 6$, $p = 0.014$). *Gravin*- β expression was not affected in gravin GT mice (Student's *t* test, $n = 6$, $p = 0.444$). **M**, Gravin GT mice express a truncated form of *gravin*- α of \sim 150 kDa (left panel), which contains LacZ (middle panel) but lacks an RII binding site (right panel). **N**, Hippocampal AKAP150 protein levels are not altered in gravin GT mice ($n = 7$ for both groups; Student's *t* tests, $p = 0.79$). * $p < 0.05$; ** $p < 0.01$. Error bars indicate SEM.

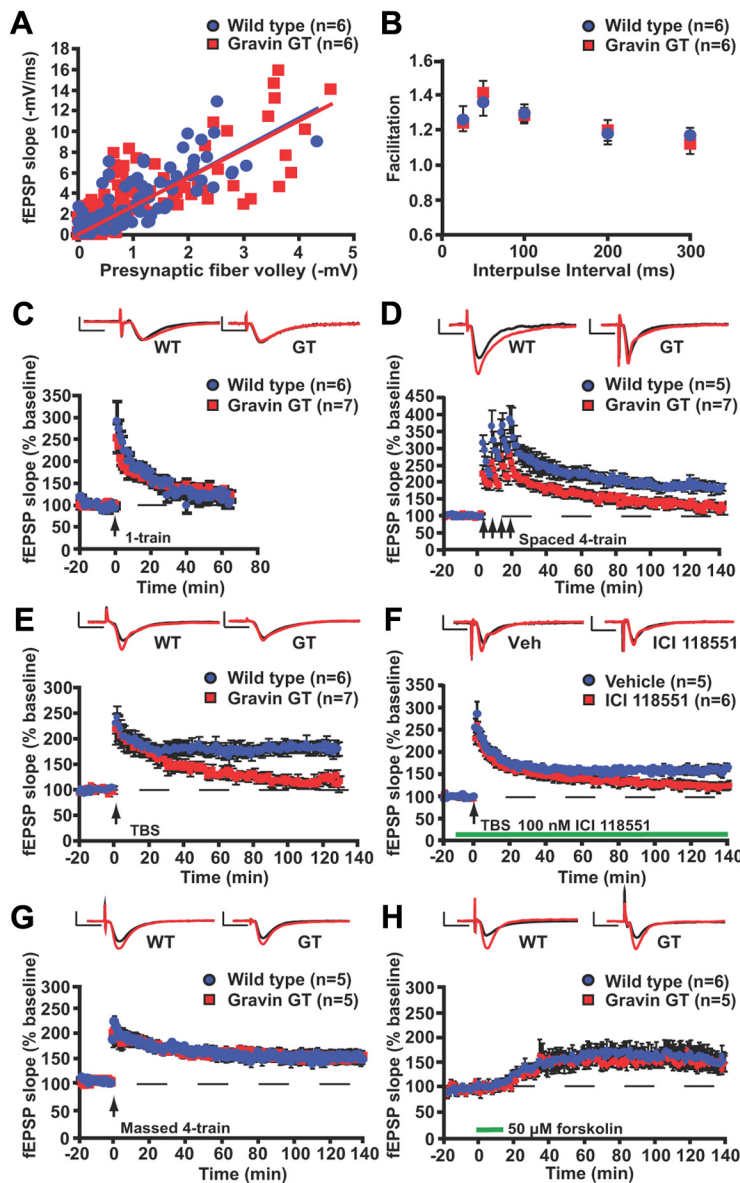


Figure 2. Long-lasting hippocampal LTP is impaired in gravin GT mice. *A*, Input–output curves relating the amplitude of the presynaptic fiber volley to the initial slope of the corresponding fEPSP at various stimulus intensities was not altered in slices from gravin GT mice (one-way ANOVA, $F_{(1,10)} = 0.128$, $p = 0.727$). *B*, Paired-pulse facilitation, a short-term form of synaptic plasticity, was not changed in slices from gravin GT mice (two-way repeated-measures ANOVA, $F_{(4,44)} = 0.415$, $p = 0.797$). *C*, One-train LTP was unaltered in hippocampal slices from gravin GT mice (Kruskal–Wallis ANOVA, $p > 0.05$). *D*, Spaced four-train LTP is impaired in slices from gravin GT mice (Kruskal–Wallis ANOVA, $p < 0.05$). *E*, A long-lasting form of LTP induced by theta-burst stimulation (15 bursts) was impaired in slices from gravin GT mice (Kruskal–Wallis ANOVA, $p < 0.05$). *F*, Bath application with the β_2 -AR antagonist ICI 118551 (100 nM) impairs a long-lasting form of LTP induced by theta-burst stimulation (15 bursts) (Kruskal–Wallis ANOVA, $p < 0.05$). *G*, Massed four-train LTP, a PKA-independent form of long-lasting LTP was not impaired in slices from gravin GT mice (Kruskal–Wallis ANOVA, $p > 0.05$). *H*, Forskolin-mediated synaptic potentiation was not impaired in slices from gravin GT mice (Kruskal–Wallis ANOVA, $p > 0.05$). In all sample sweeps, black traces indicate baseline, and red traces were acquired at 1 h after tetanus (for *C*) or ~ 2 h after stimulation (for *D–H*). Calibration: 2 mV, 10 ms. Error bars indicate SEM.

monitored AKAP150 protein levels in hippocampal lysates from GT mice and wild-type littermates. AKAP150 protein levels were not affected by the loss of gravin- α (both groups, $n = 7$; Student's t test, $p = 0.79$; Fig. 1*N*). To assess possible neuroanatomical alterations in gravin GT mice, we stained brain sections with cresyl violet and an antibody against MAP2, which is heavily expressed in neuronal dendrites (Zhong et al., 2009). No abnormalities were observed in brain architecture or dendrite structure (data not shown).

Gravin GT mice have impairments in PKA-dependent long-lasting forms of hippocampal synaptic plasticity

Initial electrophysiological characterization of Schaffer collateral–CA1 synapses in gravin GT mice involved measuring basal synaptic transmission by paired-pulse facilitation (PPF), an index of short-term plasticity. These experiments did not detect any alterations in basal synaptic transmission (WT, $n = 6$; GT, $n = 6$; one-way ANOVA, $F_{(1,11)} = 0.128$, $p > 0.05$; Fig. 2*A*) and PPF (WT, $n = 6$; GT, $n = 6$; two-way repeated-measures ANOVA, $F_{(4,44)} = 0.415$, $p > 0.05$; Fig. 2*B*) in gravin GT mice. Furthermore, maximum fEPSP slopes were not significantly different between gravin GT mice (-8.14 ± 1.73 mV/ms) and wild-type mice (-9.33 ± 2.08 mV/ms) (one-way ANOVA, $F_{(1,10)} = 0.195$, $p > 0.05$).

To determine whether gravin- α plays a role in synaptic plasticity, we examined LTP at Schaffer collateral–CA1 synapses in hippocampal slices. E-LTP is a transient form of this phenomena, induced by a single 1 s 100 Hz train, that lasts 1–2 h in hippocampal slices. E-LTP requires activation of the calcium/calmodulin-dependent kinase II α (CaMKII α), but does not require PKA or *de novo* protein synthesis (Frey et al., 1993). As expected, E-LTP was not affected by loss of gravin- α (WT, $n = 6$; GT, $n = 7$; Kruskal–Wallis ANOVA, $p > 0.05$; Fig. 2*C*).

In contrast to E-LTP, long-lasting LTP, which is induced by the spaced presentation of repeated trains of electrical stimuli, lasts for several hours, requires protein synthesis, PKA activity, and β -ARs (Frey et al., 1993; Abel et al., 1997; Kemp and Manahan-Vaughan, 2008). We have previously shown that pharmacological or genetic inhibition of PKA anchoring impairs PKA-dependent hippocampal long-lasting forms of LTP (Huang et al., 2006; Nie et al., 2007). Therefore, we reasoned that long-lasting forms of LTP might be attenuated in gravin GT mice. Hippocampal spaced four-train LTP at Schaffer collateral–CA1 synapses was impaired in slices from gravin GT mice (WT, $n = 5$; GT, $n = 7$; Kruskal–Wallis ANOVA, $p < 0.05$; Fig. 2*D*). The mean

fEPSP slope over the last 20 min of the recording was significantly reduced in gravin GT mice (WT, $185.3 \pm 7.1\%$; GT, $136.9 \pm 11.6\%$; Mann–Whitney U test, $p = 0.007$). We also examined another form of long-lasting LTP that requires PKA anchoring (Nie et al., 2007), β -ARs (Yang et al., 2002), and is induced by theta-burst stimulation, which resembles the rhythmic hippocampal electrical activity observed during spatial exploration (Buzsáki, 2005). We found that theta-burst LTP (four 100 Hz pulses delivered in bursts that are 200 ms apart for a total of 3 s) at

Schaffer collateral–CA1 synapses was impaired in slices from gravin GT mice (WT, $n = 6$; GT, $n = 7$; Kruskal–Wallis ANOVA, $p < 0.05$; Fig. 2E). The mean fEPSP slope over the last 20 min of the recording was significantly reduced in gravin GT mice (WT, $183.1 \pm 11.6\%$; GT, $114.1 \pm 7.7\%$; Mann–Whitney U test, $p = 0.004$). To determine whether theta-burst LTP requires β_2 -ARs, we repeated the experiment in wild-type mice with bath application of the β_2 -AR antagonist ICI 118551 (100 nM). Theta-burst LTP at Schaffer collateral–CA1 synapses was impaired in slices from wild-type slices treated with the β_2 -AR antagonist (WT, $n = 5$; GT, $n = 6$; Kruskal–Wallis ANOVA, $p < 0.05$; Fig. 2F). The mean fEPSP slope over the last 20 min of the recording was significantly reduced under conditions of the β_2 -AR blocker (vehicle, $156.3 \pm 10.1\%$; ICI 118551, $119.3 \pm 7.1\%$; Mann–Whitney U test, $p = 0.007$). Together, these data indicate that loss of gravin- α impairs long-lasting forms of synaptic plasticity.

LTP induced by four high-frequency trains of stimuli applied at 5 s intervals (massed four-train LTP) produces a form of long-lasting LTP that does not require cAMP/PKA signaling (Woo et al., 2003). Therefore, we anticipated that this form of LTP would not be impaired by loss of gravin- α . As expected, massed four-train LTP was not affected by loss of gravin- α (WT, $n = 5$; GT, $n = 5$; Kruskal–Wallis ANOVA, $p > 0.05$; Fig. 2G). Together, these data indicate that loss of gravin- α impairs long-lasting LTP that requires PKA anchoring without affecting PKA-independent forms of LTP.

We next wanted to establish whether loss of gravin- α impairs a form of synaptic potentiation that is induced by forskolin, a pharmacological activator of adenylyl cyclase (Huang and Kandel, 1998). The synaptic potentiation following forskolin treatment was similar in wild-type littermates and gravin GT mice (WT, $n = 6$; GT, $n = 5$; Kruskal–Wallis ANOVA, $p > 0.05$; Fig. 2H). These findings suggest that cell-wide elevation of cAMP to supraphysiological levels overcomes the mislocalization of PKA caused by the deletion of gravin- α in the mutant mice.

β_2 -Adrenergic receptor-mediated synaptic plasticity is impaired in gravin GT mice

Gravin-coordinated signaling complexes have been proposed to modulate β_2 -AR function (Tao and Malbon, 2008), but this has not been tested physiologically. Occupancy of β -ARs, when paired with distinct patterns of excitatory synaptic stimulation, can enhance hippocampal LTP (Winder et al., 1999; Gelinis et al., 2008; Qian et al., 2012). We first determined whether bath application of the β -AR agonist ISO alone induced changes in synaptic strength. Twenty-five minutes of ISO treatment did not induce any synaptic potentiation in hippocampal slices from either wild-type or gravin GT mice (data not shown). Next, we examined whether an ISO-induced form of plasticity is attenuated in the gravin GT mice. Transient low-frequency stimulation (LFS) (5 Hz for 3 min) does not cause persistent changes in synaptic strength (Qian et al., 2012). However, pairing ISO with LFS establishes long-lasting LTP (Winder et al., 1999; Gelinis et al., 2008) via the activation of PKA (Gelinis et al., 2008) and ERK1/2 (Winder et al., 1999). A recent study showed that this form of plasticity specifically requires β_2 -AR activation (Qian et al., 2012). Synaptic plasticity at Schaffer collateral–CA1 synapses induced by pairing 1 μ M ISO with LFS (5 Hz for 3 min) was impaired in slices from gravin GT mice (both groups, $n = 4$; Kruskal–Wallis ANOVA, $p < 0.05$; Fig. 3A). The mean fEPSP slope over the last 20 min of the recording was significantly reduced in gravin GT mice relative to WT mice (WT, $226.2 \pm 17.1\%$; GT, $135.3 \pm 7.9\%$; Mann–Whitney U test, $p = 0.003$).

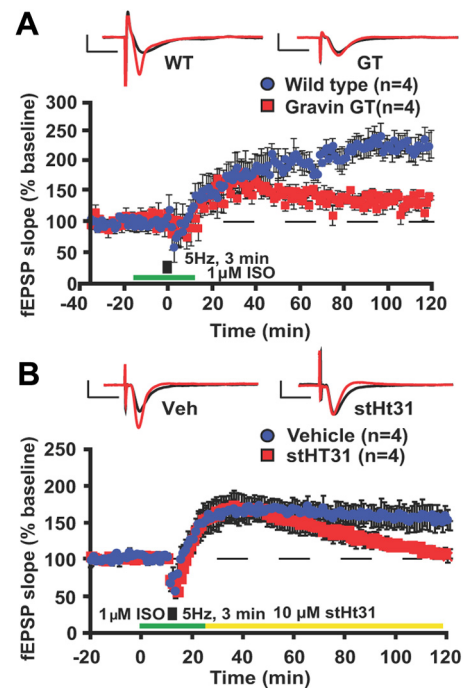


Figure 3. Hippocampal synaptic plasticity induced by β_2 -adrenergic receptor activation is impaired in gravin GT mice. **A**, Hippocampal plasticity induced by pairing 1 μ M ISO with 5 Hz train of LFS for 3 min was impaired in slices from gravin GT mice (Kruskal–Wallis ANOVA, $p < 0.05$). **B**, The maintenance of hippocampal synaptic plasticity induced by pairing 1 μ M ISO with a 5 Hz train of LFS for 3 min was impaired by disrupting PKA anchoring using membrane-permeable stearylated Ht31 after the induction of plasticity (Kruskal–Wallis ANOVA, $p < 0.05$). In all sample sweeps, black traces indicate baseline, and red traces were acquired 2 h after stimulation. Calibration: 2 mV, 10 ms. Error bars indicate SEM.

These data indicate that the loss of gravin- α impairs synaptic plasticity that requires β_2 -AR occupancy.

To determine whether PKA anchoring is required during the maintenance phase of this form of synaptic plasticity, we repeated this experiment but now treating slices from WT mice with the PKA-anchoring disrupting peptide Ht31 (Carr et al., 1992; Vijayaraghavan et al., 1997; Huang et al., 2006). Bath application of stHt31 (10 μ M) started immediately following ISO treatment and was continued throughout the recording (Fig. 3B). We found that stHt31 treatment during the maintenance phase of synaptic plasticity induced by pairing 1 μ M ISO with LFS (5 Hz for 3 min) was sufficient to impair this form of plasticity (both groups, $n = 4$; Kruskal–Wallis ANOVA, $p < 0.05$; Fig. 3B). The mean fEPSP slope over the last 20 min of the recording was significantly reduced due to treatment with stHt31 (vehicle treatment, $154.9 \pm 13.5\%$; stHt31, $98.9 \pm 4.2\%$; Mann–Whitney U test, $p = 0.029$). This reduction was not observed with bath application of stearylated pseudo-stHt31 control peptide (data not shown). These findings suggest that this form of β_2 -AR-mediated plasticity requires PKA anchoring during the maintenance phase.

Gravin GT mice exhibit impaired spatial memory in the Morris water maze

During training in the hippocampus-dependent spatial version of the Morris water maze (Schenk and Morris, 1985), mice have to learn and remember the location of a hidden platform using distal environmental cues. Because inhibition of PKA-AKAP interactions impairs memory formation in the spatial version of the Morris water maze (Nie et al., 2007; Tunquist et al., 2008), we tested whether gravin GT mice had memory deficits in this task.

Gravin GT mice ($n = 17$) and wild-type mice ($n = 18$) gradually learned to locate the hidden platform (two-way repeated-measures ANOVA, effect of day, $F_{(5,165)} = 23.20$, $p = 0.0001$; Fig. 4A). However, gravin GT mice exhibited an overall reduced acquisition rate during training (two-way repeated-measures ANOVA, effect of genotype, $F_{(1,33)} = 6.02$, $p = 0.02$; Fig. 4A). Nonetheless, they eventually reached the hidden platform with similar latency compared with wild-type littermates. Control measurements confirmed that swim speed in the gravin GT mice was similar to that of wild-type littermates (WT, 22.6 ± 1.1 cm/s; GT, 23.0 ± 1.3 cm/s; Student's t test, $p = 0.831$). Thigmotaxis was likewise not altered (WT, $15.4 \pm 5\%$; GT, $18.3 \pm 4\%$; Student's t test, $p = 0.616$).

During the probe trial 24 h after training, gravin GT mice exhibited a reduced preference for the target quadrant (two-factor ANOVA, interaction between genotype and platform, $F_{(3,99)} = 7.829$, $p = 0.0001$; Fig. 4B). *Post hoc* analyses confirmed that wild-type mice had a strong preference for the target quadrant (Tukey's test, $p < 0.001$, target vs other quadrants). In contrast, gravin GT mice extended their search for the platform to the adjacent quadrants (Tukey's test, $p > 0.1$, target vs adjacent quadrants). Wild-type mice also exhibited more platform crossings than gravin GT mice (WT, 6.4 ± 0.71 ; GT, 4.29 ± 0.63 ; Student's t test, $p = 0.036$). Together, these findings indicate that loss of gravin- α attenuates spatial memory in the Morris water maze. Gravin GT mice were not impaired in the hippocampus-independent visible platform version of the same task (WT, $n = 6$; GT, $n = 5$; two-way repeated-measures ANOVA, effect of genotype, $F_{(1,18)} = 0.056$, $p = 0.813$; interaction between genotype and day, $F_{(2,18)} = 0.240$, $p = 0.743$; Fig. 4C). To determine whether the spatial memory deficits were a consequence of alterations in locomotor behavior, we assessed motor coordination in a new batch of gravin GT mice using the rotarod test. Both wild types and gravin GT mice gradually increased the time spent on the rotating rod (two-way repeated-measures ANOVA, effect of day, $F_{(2,28)} = 15.634$, $p = 0.0001$; Fig. 4D). No effect of genotype was found (two-way repeated-measures ANOVA, effect of genotype, $F_{(1,14)} = 0.001$, $p = 0.953$). Furthermore, gravin GT mice and wild-type littermates spent a similar time in the open arms of the zero maze (Student's t test, $p = 0.694$; Fig. 4E) and a similar time in the center of the open field (Student's t test, $p = 0.388$; Fig. 4F), indicating that the spatial deficits observed in the gravin GT mice were not a consequence of alterations in exploratory or anxiety behavior.

Gravin GT mice have long-term memory impairments in the object-place recognition task, but not in the novel object recognition task

To evaluate whether other forms of hippocampus-dependent memory were impaired, we tested a naive batch of animals in the object-place recognition task, a hippocampus-dependent paradigm that examines the ability of animals to discriminate between familiar and novel spatial locations of objects (Oliveira et al., 2010). During training, both gravin GT mice ($n = 16$) and wild-type littermates ($n = 17$) gradually decreased exploration of the objects (two-way repeated-measures ANOVA, effect of genotype, $F_{(1,31)} = 4.924$, $p = 0.034$). Gravin GT mice spent considerably more time exploring the objects than wild-type mice during all three training sessions (session 1, WT, 29.1 ± 3.2 s, GT, 38.7 ± 4.0 s; session 2, WT, 17.8 ± 1.5 s, GT, 23.0 ± 2.7 s; session 3, WT, 14.1 ± 1.1 s, GT, 21.8 ± 3.1 s; two-way repeated-measures ANOVA effect of session, $F_{(2,62)} = 50.790$, $p = 0.001$). Twenty-four hours after the last training session, one of the three objects

was moved to a novel location and mice were allowed to explore the objects. A two-way ANOVA revealed a significant interaction effect between genotype and object ($F_{(1,31)} = 7.136$, $p = 0.012$; Fig. 4G). Wild-type mice significantly increased the time spent on exploring the displaced object while decreasing the time exploring the non-displaced objects (paired samples t test, $p = 0.003$; Fig. 4G). In contrast, gravin GT mice decreased the time spent exploring both displaced and non-displaced object (paired samples t test, $p = 0.518$; Fig. 4G). No difference in locomotor speed was observed between mutant and wild-type mice (WT, 51.3 ± 6.5 mm/s; GT, 47.7 ± 5.3 mm/s; Student's t test, $p = 0.651$). These findings indicate that the formation of object-place memories is impaired in the gravin GT mice.

To determine whether gravin GT mice also have a deficit in detecting object novelty, gravin GT mice ($n = 8$) and wild-type littermates ($n = 8$) were trained and tested in the novel object recognition task, a task that is based on the animal's spontaneous preference for novelty (Ennaceur and Delacour, 1988). The training conditions we used do not require an intact hippocampus (Winters et al., 2004; Oliveira et al., 2010). During training, both groups spent a similar amount of time exploring both objects (WT, 48.6 ± 8.8 s; GT, 58.4 ± 8.6 s; Student's t test, $p = 0.410$). Twenty-four hours after training, mice were reintroduced to the arena and reexposed to two objects, a familiar object and a new object. Both groups preferentially explored the novel object (two-factor ANOVA, $F_{(1,28)} = 5.806$, $p = 0.023$; Fig. 4H). No effect of genotype (ANOVA, $F_{(1,28)} = 0.006$, $p = 0.936$) or interaction effect between genotype and object (ANOVA, $F_{(1,28)} = 0.661$, $p = 0.423$) was found. These data indicate that the formation of long-term memories for objects does not require gravin- α signaling complexes.

Gravin GT mice have long-term memory impairments in contextual and tone-cued fear conditioning task

Next, we tested whether gravin GT mice showed an impairment in hippocampus-dependent contextual fear conditioning (Logue et al., 1997). During training, both groups (WT, $n = 10$; GT, $n = 8$) showed similar preshock and postshock freezing levels (Student's t test, $p > 0.2$ in both cases) (data not shown). One hour after training, both groups were reexposed to the training context and showed similar freezing levels (WT, $28.3 \pm 4.2\%$; GT, $31.2 \pm 5.4\%$; Student's t test, $p = 0.679$; Fig. 4I), indicating that gravin GT mice form normal short-term contextual fear memories. To test whether the formation of long-term contextual fear memories was attenuated, we repeated the contextual fear conditioning experiment with a new batch of mice. Both groups ($n = 8$ each) again showed similar preshock and postshock freezing levels during training (Student's t test, $p > 0.5$ in both cases) (data not shown). Twenty-four hours after training, mice were reexposed to the same context. Gravin GT mice showed reduced freezing levels ($33.2 \pm 5.8\%$) compared with wild-type littermates ($54.5 \pm 5.6\%$) (Student's t test, $p = 0.019$; Fig. 4I). Because PKA anchoring is also required for amygdala-dependent auditory fear memories (Moita et al., 2002), we subjected a third batch of mice to cued fear conditioning that critically depends on the amygdala. Both groups (WT, $n = 18$; GT, $n = 19$) showed similar freezing levels when exposed to the context, tone [the conditioned stimulus (CS)], and directly after receiving the shock (Student's t test, $p > 0.2$ for all three cases) (data not shown). Twenty-four hours after conditioning, mice were exposed to a novel context and reexposed to the tone. Exposure to the novel context induced similar freezing levels in both groups (Student's t test, $p = 0.410$; Fig. 4J). However, gravin GT mice showed reduced freezing levels

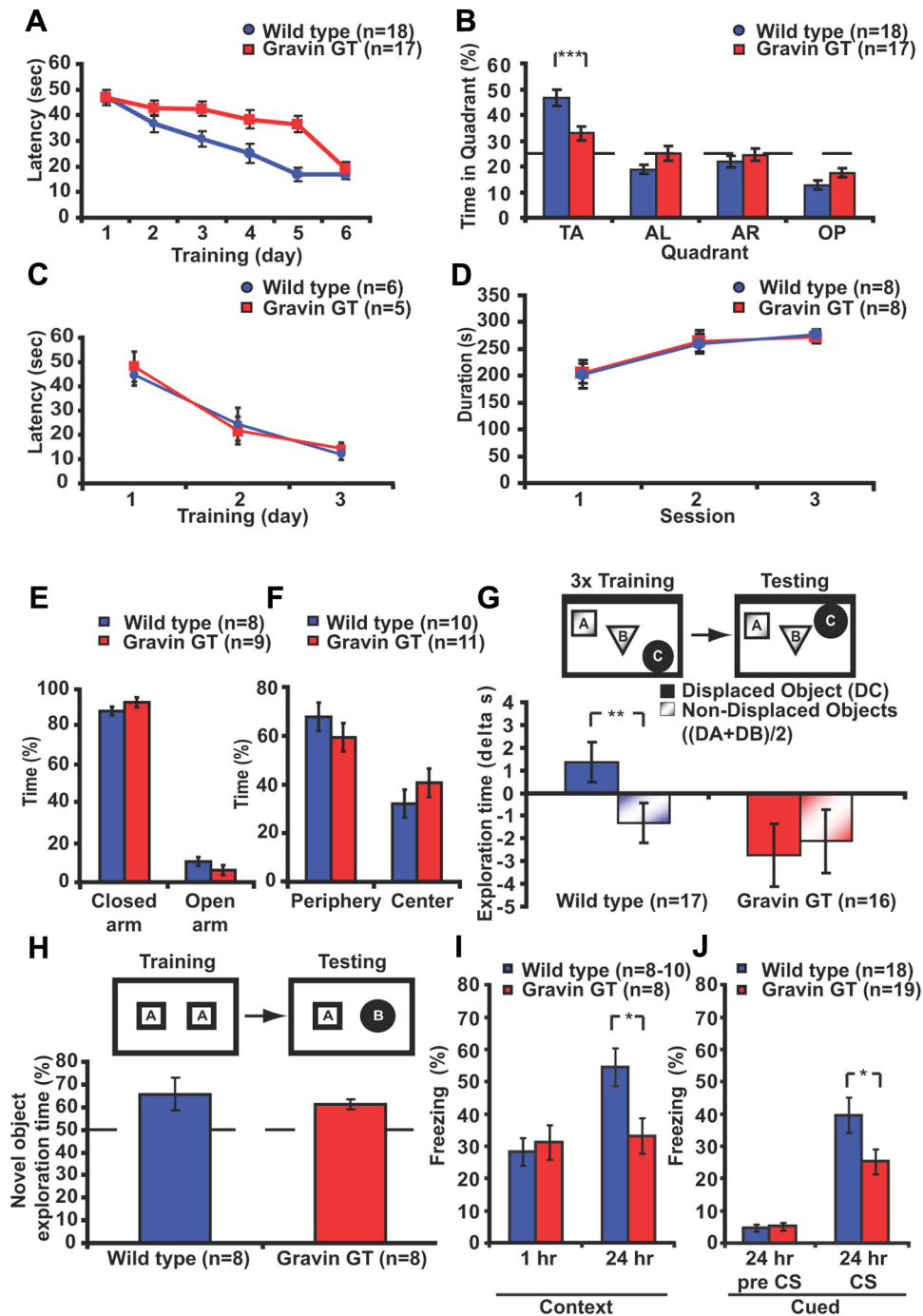


Figure 4. Long-term memory formation is impaired in gravin GT mice. **A**, In the spatial version of the water maze, gravin GT mice exhibited impairments during training (two-way repeated-measures ANOVA, effect of genotype, $F_{(1,33)} = 6.02, p = 0.02$) but eventually reached the hidden platform with a latency similar to wild-type littermates (independent-samples t test, $p = 0.523$). **B**, During the probe trial, wild-type mice preferentially searched in the target quadrant (*post hoc* Tukey's test, $p < 0.001$) in contrast to gravin GT mice (*post hoc* Tukey's test, $p > 0.1$, target vs adjacent quadrants). **C**, Gravin GT mice were not different from wild-type mice on the visible platform version of the maze (two-way repeated-measures ANOVA, $F_{(1,18)} = 0.056, p = 0.813$; interaction between genotype and day, $F_{(2,18)} = 0.240, p = 0.743$). **D**, Motor coordination as assessed by the rotarod test was not altered in the gravin GT mice. Both groups gradually increased the time spent on the rotating rod (two-way repeated-measures ANOVA, effect of day, $F_{(2,28)} = 15.634, p = 0.0001$). No effect of genotype was found (two-way repeated-measures ANOVA effect of genotype, $F_{(1,14)} = 0.001, p = 0.953$). **E**, Gravin GT mice spent a similar time in the open arms of the zero maze as wild-type mice (Student's t test, $p = 0.694$). **F**, Gravin GT mice spent a similar time in the center of the open field as wild-type mice (Student's t test, $p = 0.388$). **G**, During the test session in the object-place recognition task, wild-type mice exhibited increased exploration time for the displaced object but decreased exploration time for the non-displaced objects (paired t test, $p = 0.003$). Gravin GT mice exhibited reduced exploration times for both displaced and non-displaced objects (paired t test, $p = 0.518$). **H**, During the test session in the novel object recognition task, both wild-type and gravin GT mice preferentially explored the novel object (two-factor ANOVA $F_{(1,28)} = 5.806, p = 0.023$). No genotype or interaction effect between genotype and object was found (ANOVA, $F_{(1,28)} = 0.006, p = 0.936$; ANOVA, $F_{(1,28)} = 0.661, p = 0.423$, respectively). The dotted line indicates no preference. **I**, Mice were trained in the contextual fear conditioning paradigm. Wild-type and gravin GT mice showed similar freezing levels during the retention test 1 h after training (Student's t test, $p = 0.679$), while gravin GT mice showed reduced freezing levels during the retention test 24 h after conditioning (Student's t test, $p = 0.019$). **J**, In the cued-fear conditioning paradigm, gravin GT mice showed reduced freezing levels during the test session 24 h after training (Student's t test, $p = 0.023$). TA, Target; AL, adjacent left; AR, adjacent right; OP, opposite. * $p < 0.05$; ** $p = 0.003$; *** $p < 0.001$. Error bars indicate SEM.

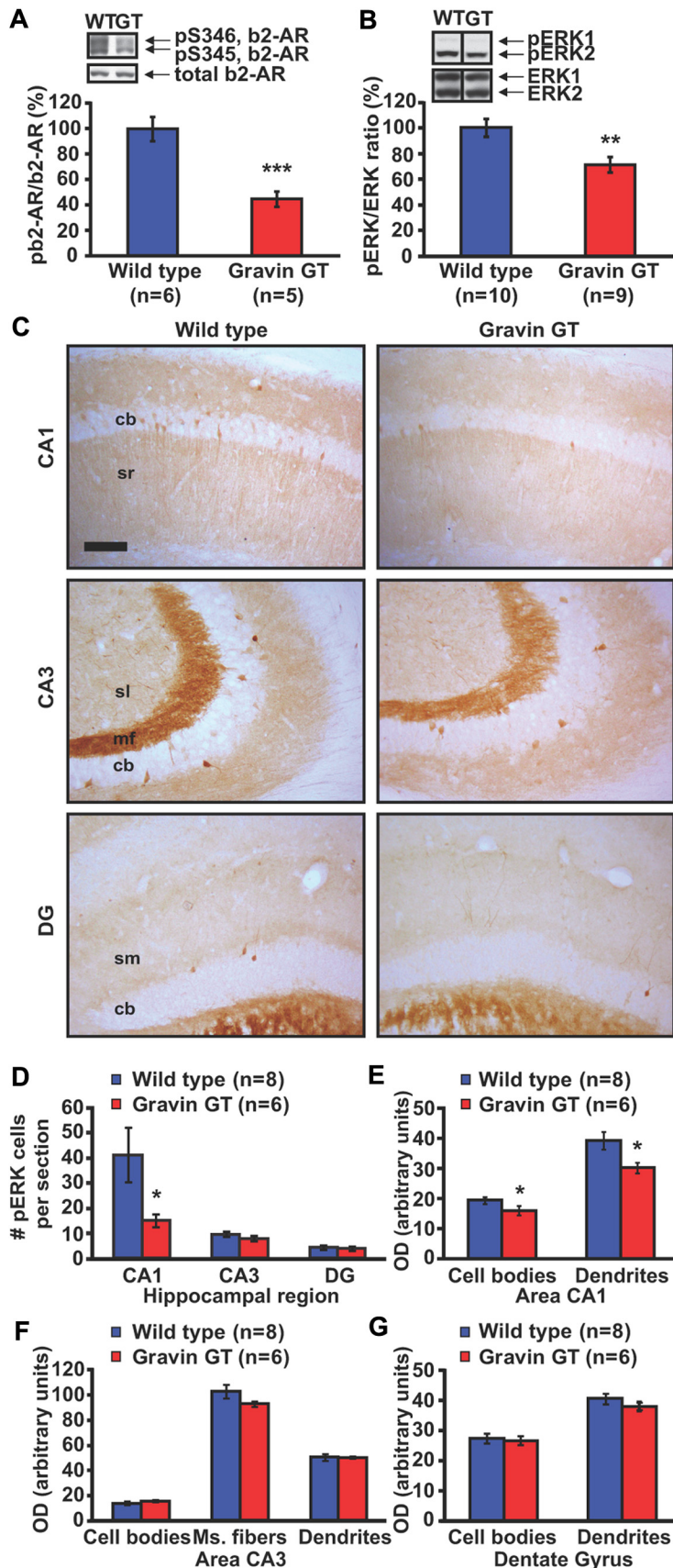


Figure 5. Phosphorylation of the β_2 -adrenergic receptor and ERK1/2 is reduced in the hippocampus of gravin GT mice 30 min after fear conditioning. **A**, Hippocampal β_2 -adrenergic receptor phosphorylation levels at the serine 345 and serine 346 sites are reduced in gravin GT mice 30 min after fear conditioning (WT, $100.0 \pm 9.6\%$; GT, $44.8 \pm 6.2\%$; Student's *t* test, $p = 0.001$). **B**, Hippocampal pERK1/2 levels are reduced in gravin GT mice 30 min after fear conditioning (WT, $100.0 \pm 7.1\%$; GT, $71.0 \pm 5.9\%$;

during the reexposure to the CS (WT, $39.6 \pm 5.5\%$; GT, $25.2 \pm 3.9\%$; Student's *t* test, $p = 0.023$; Fig. 4J), indicating that loss of gravin- α impairs cued fear conditioning.

Phosphorylation of the β_2 -AR and ERK1/2 is reduced after fear conditioning in gravin GT mice

Because gravin GT mice show deficits in the formation of hippocampus-dependent long-term memories, we next determined whether learning-induced activation of the β_2 -AR signaling pathway was altered in the gravin GT mice. Occupancy of the β_2 -AR leads to G α s-mediated activation of PKA, which in turn phosphorylates the receptor at serine 345 and serine 346 (Hausdorff et al., 1989). To determine whether the loss of gravin- α attenuates β_2 -AR phosphorylation by PKA *in vivo*, mice underwent contextual fear conditioning and hippocampal tissue was collected 30 min after training. Western blot analyses indicated that gravin GT mice showed reduced β_2 -AR phosphorylation at both serine sites (WT, $100.0 \pm 9.6\%$; GT, $44.8 \pm 6.2\%$; Student's *t* test, $p = 0.001$; Fig. 5A). Because β_2 -AR phosphorylation was not affected in experimentally naive gravin GT mice (WT, $100.0 \pm 1.3\%$; GT, $98.8 \pm 0.8\%$; Student's *t* test, $p = 0.448$), these data suggest that learning-induced β_2 -AR phosphorylation by PKA is impaired in gravin GT mice. The reduction in β_2 -AR phosphorylation by PKA 30 min after fear conditioning was also not a reflection of an overall reduction of PKA activity due to the loss of gravin- α because global PKA activity in hippocampal lysates was not at-

Student's *t* test, $p = 0.006$). **C**, Immunohistochemistry for pERK in area CA1, CA3, and DG from sections representative of wild-type mice and gravin GT mice 30 min after fear conditioning training. Note the reduced pERK1/2 labeling in area CA1. cb, Cell bodies; sr, stratum radiatum; sl, stratum lucidum; mf, mossy fibers; sm, stratum moleculare. Scale bar, 40 μ m. **D**, The number of pERK1/2-positive cells in hippocampal subregions of wild-type and gravin GT mice 30 min after fear conditioning training. The number of pERK-positive cells in area CA1 was significantly reduced in gravin GT mice (Student's *t* test, $p = 0.006$). **E**, Optical density measurements of pERK1/2 immunoreactivity in the cell body layer and stratum radiatum of area CA1 in wild-type mice and gravin GT mice. pERK1/2 immunoreactivity was significantly reduced in CA1 stratum radiatum (Student's *t* test, $p = 0.031$) and CA1 cell body layer (Student's *t* test, $p = 0.034$). **F**, Optical density measurements of pERK1/2 immunoreactivity in the cell body layer, mossy fibers, and stratum lucidum of area CA3 in wild-type mice and gravin GT mice. **G**, Optical density measurements of pERK1/2 immunoreactivity in the cell body layer and molecular layer of the DG in wild-type mice and gravin GT mice. * $p < 0.05$, ** $p < 0.01$, *** $p = 0.001$. Error bars indicate SEM.

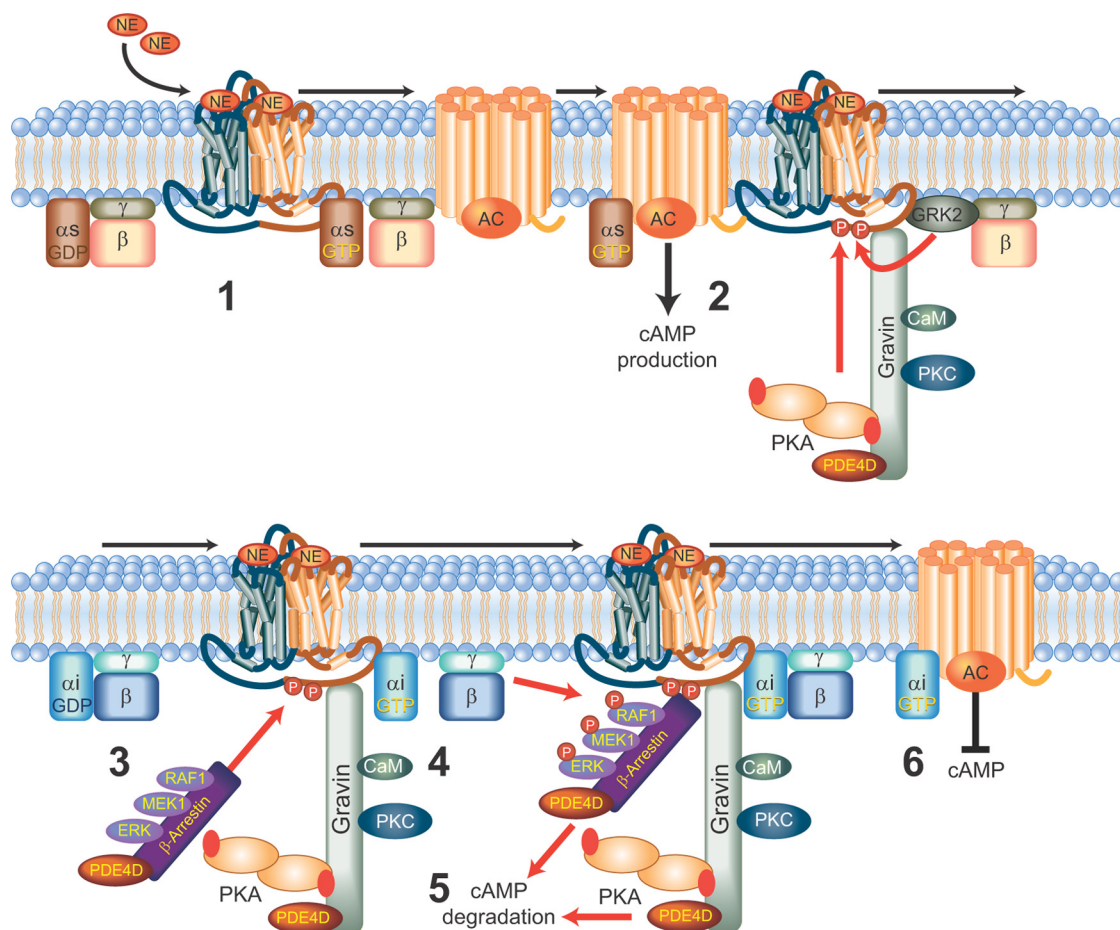


Figure 6. A potential mechanism illustrating the role of gravin complexes in β_2 -adrenergic receptor signaling. Activation of the β_2 -adrenergic receptor leads to the coupling with G_s proteins (1), the recruitment of gravin signaling complexes to the plasma membrane (Tao et al., 2003), and phosphorylation of the receptor by PKA (2) (Hausdorff et al., 1989). GRK2 initiates the desensitization process and recruitment of the β -arrestin signaling complex (Perry et al., 2002) (3). The β_2 adrenergic receptor switches coupling from G_s to G_i proteins leading to the activation of ERK1/2 (Daaka et al., 1997) (4). The β -arrestin signaling complex attenuates local cAMP signaling through the recruited PDE4D, which degrades cAMP (Baillie et al., 2003) (5), and by preventing G_s -mediated adenylate cyclase activation by the β_2 -adrenergic receptor (Perry et al., 2002) (6).

tenuated in the gravin GT mice (WT, $n = 7$, $100 \pm 5.9\%$; GT, $n = 7$, $102 \pm 8.6\%$; Student's t test, $p = 0.792$). The latter observation is in concordance with previous studies showing that loss of PKA anchoring does not affect global PKA activity in hippocampal lysates (Nie et al., 2007; Weisenhaus et al., 2010). Next, we determined whether loss of gravin affects PKA-mediated phosphorylation of the AMPA receptor GluA1 serine 845 site. We found that the phosphorylation levels of this site were not altered in hippocampal lysates from wild-type and gravin GT mice 30 min after fear conditioning (WT, $n = 7$, $100 \pm 5.9\%$; GT, $n = 7$, $99.9 \pm 5.9\%$; Student's t test, $p = 0.996$). Together, these data suggest that loss of gravin- α affects the phosphorylation of specific substrates such as the β_2 -AR rather than leading to a global reduction in PKA function. PKA-mediated β_2 -AR phosphorylation at serine 345 and serine 346 switches coupling of the β_2 -AR from G_s to $G_{\alpha i}$ -proteins leading to the activation of the ERK1/2 pathway (Daaka et al., 1997; Baillie et al., 2003). Thus, reductions in the phosphorylation of β_2 -AR phosphorylation at serine 345 and serine 346 would be predicted to reduce the activation of ERK1/2. Indeed, we found that phospho levels of ERK1/2 were significantly decreased in gravin GT mice 30 min after fear conditioning (WT, $100.0 \pm 7.1\%$; GT, $71.0 \pm 5.9\%$; Student's t test, $p = 0.006$; Fig. 5B). This decrease in hippocampal ERK1/2 phosphorylation was not observed in experimentally naive mice (WT,

$n = 15$, $100 \pm 3.8\%$; GT, $n = 14$, $114.7 \pm 8.3\%$; Student's t test, $p = 0.137$). A study by Sindreu et al. (2007) suggested that fear conditioning induces changes in pERK1/2 specifically in hippocampal area CA1. Therefore, we assessed whether the reduction in pERK1/2 levels in the gravin GT mice was restricted to area CA1. Mice were trained in the fear-conditioning task and transcardially perfused 30 min after training, and dorsal hippocampal sections were stained for pERK1/2. The largest number of pERK1/2-positive cells in the dorsal hippocampus was observed in area CA1 (Fig. 5C,D). Gravin GT mice showed significantly fewer pERK1/2-positive cells in area CA1 (WT, $41.2 \pm 10.7\%$; GT, $15.1 \pm 2.7\%$; Student's t test, $p = 0.006$; Fig. 5D). No significant changes in pERK1/2-positive cells were observed in area CA3 or in the DG. Optical density measurements in hippocampal subregions showed reduced pERK1/2 labeling in stratum radiatum (sr) of area CA1 (WT, $39.2 \pm 2.9\%$; GT, $30.3 \pm 1.7\%$; Student's t test, $p = 0.031$; Fig. 5E) and the CA1 cell body layer (WT, $19.5 \pm 1.1\%$; GT, $15.8 \pm 1.4\%$; Student's t test, $p = 0.034$; Fig. 5E). No significant changes in staining intensity were observed in CA3 cell bodies (WT, $12.4 \pm 1.7\%$; GT, $15.2 \pm 1.1\%$; Student's t test, $p = 0.422$; Fig. 5F), mossy fibers (WT, $102.6 \pm 5.19\%$; GT, $92.7 \pm 2.3\%$; Student's t test, $p = 0.117$; Fig. 5F), or stratum lucidum (WT, $50.0 \pm 2.7\%$; GT, $49.9 \pm 0.7\%$; Student's t test, $p = 0.979$; Fig. 5F). Likewise, no changes were observed in

the DG cell bodies (WT, $27.3 \pm 1.7\%$; GT, $26.7 \pm 1.6\%$; Student's *t* test, $p = 0.803$; Fig. 5G) or molecular layer (WT, $40.5 \pm 1.1\%$; GT, $38.0 \pm 1.4\%$; Student's *t* test, $p = 0.193$; Fig. 5G). Together, these results indicate that training-induced ERK1/2 activation in hippocampal area CA1 is impaired in gravin GT mice.

Discussion

Coordination of excitatory synaptic signaling events is a complex process that relies on the spatial and temporal segregation of cellular signaling pathways within the neuron. This is achieved in part by AKAPs that compartmentalize unique macromolecular complexes consisting of PKA and other signaling molecules including phosphatases, PDEs, and targets of PKA at defined synaptic locations (Welch et al., 2010). This intricate network of protein–protein interactions provides a molecular mechanism that allows different extracellular signals to engage spatially distinct pools of kinases. Although studies have implicated PKA signaling in long-lasting forms of LTP and memory, the identity of the AKAP(s) that coordinates pools of PKA critically involved in synaptic plasticity and memory formation is less well understood. Here, we show that selective loss of the α -isoform of gravin impairs long-lasting forms of hippocampal synaptic plasticity including those that require β 2-AR activation as well as the formation of long-term memories that require the hippocampus or amygdala. Furthermore, our biochemical studies reveal that the loss of gravin- α attenuates hippocampal β 2-AR phosphorylation by PKA and reduces learning-induced activation of ERK1/2 in area CA1 of the dorsal hippocampus.

Our observation that the loss of gravin- α reduces PKA-mediated β 2-AR phosphorylation in the hippocampus tallies with cell culture studies. These studies have shown that activation of the β 2-AR initiates the recruitment of gravin to plasma membrane sites (Tao et al., 2003), the phosphorylation of gravin by PKA, and association of PKA with the β 2-AR (Shih et al., 1999; Lin et al., 2000). However, we have significantly extended these biochemical studies and provide a physiological context for the findings by showing that distinct forms of hippocampal LTP that require β 2-ARs and PKA anchoring are impaired in gravin GT mice. Importantly, we have found that PKA anchoring is required for a form of synaptic plasticity that requires low-frequency electrical stimulation in combination with activation of β 2-ARs, consistent with this hypothesized role of gravin in regulating β 2-AR function during sustained activation. Thus, gravin- α is a key organizational component of PKA and β 2-AR-mediated signaling in hippocampal neurons and allowed us to propose that disruption of the β 2-AR-controlled component of hippocampal synaptic plasticity and long-term memories could be due to the observed loss of local PKA function (Fig. 6). Future studies using pharmacological reagents that modulate specific components of the β 2-AR signaling complex (Kahsai et al., 2011) are needed to further unravel the role that β 2-ARs play in hippocampal LTP and memory. Because our Ht31 experiment suggests that PKA anchoring is required during the maintenance phase of LTP, it will also be important to elucidate whether blocking PKA activity and β 2-AR function selectively during the maintenance phase is sufficient to impair long-lasting forms of LTP or behavior and memory.

As with other AKAPs, gravin is a multivalent scaffolding protein that not only binds PKA but also binds PKC and calmodulin, and recruits PDE4D to the β 2-AR (Nauert et al., 1997; Lin and Gelman, 2002; Willoughby et al., 2006), suggesting that loss of gravin- α not only reduces the localization of cAMP responsive enzymes such as PKA and PDE4D (Houslay and Baillie, 2003) but

also abrogates the efficacy of other signaling molecules that regulate β 2-AR function (Fig. 6). These include PKC, another enzyme that can phosphorylate GRK2 promoting desensitization of the β 2-ARs (Kohout and Lefkowitz, 2003) (Fig. 6), and calmodulin, which relays the increases in intracellular Ca^{2+} into activation or inhibition of downstream enzymes. The latter has been proposed to be a signal integrator for synaptic plasticity (Xia and Storm, 2005). In future experiments, it will be interesting to determine whether gravin-mediated compartmentalization of PKC and calmodulin contribute to synaptic plasticity important for memory. Nevertheless, our current study suggests that the α -isoform of gravin is particularly important for PKA signaling and for the activation of ERK1/2.

Previous studies to define the AKAP complexes involved in synaptic plasticity and memory processes have been complicated by non-mnemonic behavioral side effects and neuroanatomical abnormalities. AKAP150-null mice exhibit a reduction in motor coordination (Tunquist et al., 2008), and mice lacking the PKA binding site from MAP2B have an abnormal CA1 architecture (Khuchua et al., 2003). In contrast, loss of gravin- α impairs long-term memory formation, without affecting motor coordination, anxiety, exploratory behavior, or hippocampal architecture. This makes it unlikely that the observed memory deficits observed here are an indirect consequence of the loss of gravin- α during development. Rather, it supports the notion that gravin- α -mediated compartmentalization of signaling complexes during adulthood is required to maintain normal synaptic plasticity and memory.

ERK1/2 signaling is required for long-lasting forms of LTP (Impey et al., 1998; Patterson et al., 2001) and the formation of long-lasting memories (Davis and Laroche, 2006). More specifically, fear conditioning training activates the ERK pathway in a subset of hippocampal neurons specifically in area CA1 (Sindreu et al., 2007), and blocking ERK1/2 activity in hippocampal area CA1 during the consolidation window prevents the formation of long-term memories (Athos et al., 2002). We discovered that loss of gravin- α attenuates the formation of hippocampus-dependent long-term memories (Fig. 4) and impairs learning-induced activation of ERK1/2 pathway in area CA1 of the hippocampus (Fig. 5). These observations support the idea that gravin signaling complexes orchestrate learning-induced ERK1/2 activation in the hippocampus important for memory consolidation. β 2-AR phosphorylation by PKA is a prerequisite for activation of the ERK1/2 pathway by β 2-ARs (Daaka et al., 1997; Baillie et al., 2003). Therefore, our finding that β 2-AR phosphorylation was reduced in hippocampal neurons after fear conditioning training (Fig. 5) suggests that loss of gravin-mediated PKA phosphorylation of β 2-AR may hamper the activation of the ERK1/2 pathway (Fig. 6). However, it should be noted that β 2-ARs can also activate the ERK1/2 pathway through PKA-independent signaling mechanisms (Schmitt and Stork, 2000). Furthermore, cAMP, PKA, and ERK1/2 signaling interact in other ways (Gloerich and Bos, 2010).

This study is the first to show that gravin- α controls PKA-dependent forms of hippocampal synaptic plasticity and a β 2-AR-mediated form of synaptic plasticity that requires low-frequency electrical stimulation in combination with activation of β 2-ARs, regulates hippocampal ERK1/2 signaling, and is required for specific forms of long-term memory. Furthermore, we demonstrate that PKA anchoring is critical for the maintenance of β 2-AR-mediated synaptic plasticity. Our studies not only implicate gravin- α in the subcellular targeting of PKA but also reveal that this particular macromolecular signaling complex is critical

for the cellular and biochemical processes underlying long-term memory storage. By linking the glutamatergic and noradrenergic signaling pathways, gravin signaling complexes are able to orchestrate the formation of long-term changes in neuronal function.

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