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Convergence of Hippocampal Pathophysiology in *Syngap*^{+/-} and *Fmr1*^{-ly} Mice

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Previous studies have hypothesized that diverse genetic causes of intellectual disability (ID) and autism spectrum disorders (ASDs) converge on common cellular pathways. Testing this hypothesis requires detailed phenotypic analyses of animal models with genetic mutations that accurately reflect those seen in the human condition (i.e., have structural validity) and which produce phenotypes that mirror ID/ASDs (i.e., have face validity). We show that SynGAP haploinsufficiency, which causes ID with co-occurring ASD in humans, mimics and occludes the synaptic pathophysiology associated with deletion of the *Fmr1* gene. *Syngap*^{+/-} and *Fmr1*^{-ly} mice show increases in basal protein synthesis and metabotropic glutamate receptor (mGluR)-dependent long-term depression that, unlike in their wild-type controls, is independent of new protein synthesis. Basal levels of phosphorylated ERK1/2 are also elevated in *Syngap*^{+/-} hippocampal slices. Super-resolution microscopy reveals that *Syngap*^{+/-} and *Fmr1*^{-ly} mice show nanoscale alterations in dendritic spine morphology that predict an increase in biochemical compartmentalization. Finally, increased basal protein synthesis is rescued by negative regulators of the mGlu subtype 5 receptor and the Ras–ERK1/2 pathway, indicating that therapeutic interventions for fragile X syndrome may benefit patients with SYNGAP1 haploinsufficiency.

Key words: fragile X syndrome; long-term depression; mGluR; neurodevelopmental disorder; STED; SynGAP

Significance Statement

As the genetics of intellectual disability (ID) and autism spectrum disorders (ASDs) are unraveled, a key issue is whether genetically divergent forms of these disorders converge on common biochemical/cellular pathways and hence may be amenable to common therapeutic interventions. This study compares the pathophysiology associated with the loss of fragile X mental retardation protein (FMRP) and haploinsufficiency of synaptic GTPase-activating protein (SynGAP), two prevalent monogenic forms of ID. We show that *Syngap*^{+/-} mice phenocopy *Fmr1*^{-ly} mice in the alterations in mGluR-dependent long-term depression, basal protein synthesis, and dendritic spine morphology. Deficits in basal protein synthesis can be rescued by pharmacological interventions that reduce the mGlu₅ receptor–ERK1/2 signaling pathway, which also rescues the same deficit in *Fmr1*^{-ly} mice. Our findings support the hypothesis that phenotypes associated with genetically diverse forms of ID/ASDs result from alterations in common cellular/biochemical pathways.

Introduction

Intellectual disability (ID) refers to a group of neurodevelopmental disorders characterized by an intelligence quotient (IQ) below

70, which affects ~2–3% of the population (Huang, 2009). Large-scale genetic studies have shown that the genetic causes of ID are highly heterogeneous involving chromosomal abnormal-

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ities and single gene mutations, with several hundred ID-related genes identified (Krumm et al., 2014). Estimates of comorbidity indicate that ~30% of people with ID also present with autism spectrum disorders (ASDs), suggesting significant genetic overlap between these two disorders. For example, one-third of all boys with fragile X syndrome (FXS) also meet the diagnostic criteria for autism (Wijetunge et al., 2013).

Although the genetic causes of ID/ASDs are highly diverse, previous studies suggest that there may be convergence at the level of biochemical pathways that are affected. For example, FXS and tuberous sclerosis complex (TSC), two monogenic causes of ID with overlapping clinical symptoms, appear to be on opposite ends of a common pathophysiological axis (Auerbach et al., 2011). Thus, it is possible that rare but highly penetrant genetic mutations (e.g., FXS and TSC) may give insight into the pathophysiology of a range of IDs/ASDs (Wijetunge et al., 2013).

A core physiological consequence of the loss of fragile X mental retardation protein (FMRP) is an increase in basal protein synthesis downstream of mGlu₅ receptor activation (Qin et al., 2005; Dölen et al., 2007; Osterweil et al., 2010; Till et al., 2015). In the hippocampus, this change can be detected as an increase in metabotropic glutamate receptor (mGluR)-dependent long-term synaptic depression (LTD) that, unlike in wild type (WT), no longer requires new protein synthesis (Huber et al., 2002; Nosyreva and Huber, 2006). Reducing mGlu₅ receptor signaling, either genetically or with negative allosteric modulators of mGlu₅ receptors, rescues many of the phenotypes associated with the genetic deletion of FMRP (Dölen et al., 2007; Michalon et al., 2012; Michalon et al., 2014). In the absence of FMRP, the elevation in basal protein synthesis results from a hypersensitivity to ERK1/2-MAPK signaling, and inhibitors of this pathway also reverse cellular and behavioral effects resulting from the loss of FMRP (Osterweil et al., 2010, 2013).

In this study, we examine the pathophysiology in mice heterozygous for a null mutation in *Syngap*. Synaptic GTPase-activating protein (SynGAP) is a RasGAP encoded by a gene that gives rise to multiple isoforms, several of which have opposing effects on synaptic efficacy (McMahon et al., 2012). It is expressed throughout development with the highest levels observed in the forebrain at the peak of synaptogenesis (Porter et al., 2005; Barnett et al., 2006). In human patients, *de novo* SYNGAP1 mutations that result in a null allele lead to autosomal dominant nonsyndromic ID (NSID) and, in a minority of cases, ASDs (Hamdan et al., 2009, 2011; Berryer et al., 2013). *Syngap* heterozygous (*Syngap*^{+/-}) mice also show a range of behavioral, physiological, and anatomical abnormalities throughout development (Komiyama et al., 2002; Vazquez et al., 2004; Barnett et al., 2006; Harlow et al., 2010; Clement et al., 2012, 2013). *Fmr1*^{-/-} mice show a hypersensitivity to Ras-ERK1/2 signaling (Osterweil et al., 2010), whereas *Syngap*^{+/-} mice show elevated ERK1/2 phosphorylation in the hippocampus (Komiyama et al., 2002; Ozkan et al., 2014). The involvement of the ERK1/2 pathway led us to hypothesize that these two genetic mutations, which cause human ID, may share a common hippocampal pathophysiology downstream of mGluR-dependent ERK1/2 activation.

Materials and Methods

Animals. *Syngap*^{+/-} mutant mice, originally generated by Komiyama et al. (2002), and *Fmr1*^{-/-} mice were bred on a C57Black6/J Ola line (Harlan). For genetic occlusion experiments, *Syngap*^{+/-} males were paired with *Fmr1*^{+/-} females to obtain F1 male progeny with the following genotypes: WT (*Syngap*^{+/+}/*Fmr1*^{+/+}), *Fmr1* KO (*Syngap*^{+/+}/*Fmr1*^{-/-}), *Syngap* heterozygous (*Syngap*^{+/-}/*Fmr1*^{+/+}), and double mutants (*Syngap*^{+/-}/*Fmr1*^{-/-}). All experiments were performed with WT littermates acting as controls and with the experimenter blind to genotype. All animal work was performed in accordance with the United Kingdom Animals (Scientific Procedures) Act of 1986 under the authority of Project Licenses PPL 60/3631 and PPL 60/4290.

Hippocampal slice electrophysiology. Horizontal hippocampal slices (400 μm) were prepared from postnatal day 25–32 (P25–32) animals. Slices were collected in carbogenated (95% oxygen, 5% CO₂) ice-cold dissection buffer containing the following (in mM): 86 NaCl, 1.2 NaH₂PO₄, 25 KCl, 25 NaHCO₃, 20 glucose, 75 sucrose, 0.5 CaCl₂, and 7 MgCl₂. Slices were incubated for 30 min at ~30°C in artificial CSF (ACSF) containing the following (in mM): 124 NaCl, 1.2 NaH₂PO₄, 25 KCl, 25 NaHCO₃, 20 glucose, 2 CaCl₂, and 1 MgCl₂, bubbled with 95% oxygen and 5% CO₂. An incision was made through CA1–CA3 boundary, and slices were left to recover for a minimum of 1 h at room temperature (20–22°C) before any recordings were made. For electrophysiological recordings, slices were placed in a submersion chamber heated to 30°C and perfused with carbogenated ACSF containing picrotoxin (50 μM) at a rate of 4 ml/min.

Field EPSPs (fEPSPs) were recorded at Schaffer collateral/commissural inputs to CA1 pyramidal neurons using extracellular recording electrodes (1–3 MΩ) filled with ACSF and placed in the stratum radiatum layer of the CA1 area. Synaptic responses were evoked by applying single current pulses to the Schaffer collateral/commissural axons using a bipolar stimulating electrode. Stimuli (10–30 μA, 200 μs, delivered at a frequency of 0.033 Hz) were set to produce 30–60% of the maximal response amplitude.

Metabotropic glutamate receptor-dependent LTD (mGluR-LTD) was induced by acute application (5 min) of the group 1 mGluR agonist (*R,S*)-3,5-dihydroxyphenylglycine (DHPG; 50 μM) in the presence of the NMDAR antagonist D-AP5 (50 μM). For experiments involving anisomycin, slices were preincubated for 30 min in ACSF containing anisomycin (20 μM) before LTD induction, which remained present for the duration of the recording.

Electrophysiological traces were collected using WinLTP (University of Bristol, Bristol, UK) and exported to Microsoft Excel. The magnitude of LTD was calculated by dividing the average fEPSP slope from 40 to 60 min after DHPG application by the average fEPSP slope during the 20 min baseline before DHPG application. Statistical analysis was performed using GraphPad Prism routines (GraphPad Software). After genotyping, time-matched normalized data were averaged across experiments and expressed as means ± SE. Significant differences between the WT and mutant mice were determined using Student's *t* test or ANOVA with *p* ≤ 0.05.

Metabolic labeling. Protein synthesis levels were measured following the protocol outlined by Osterweil et al. (2010). Briefly, 500 μm transverse hippocampal slices were prepared from either *Syngap*^{+/-} or *Fmr1*^{-/-} mice with WT littermates serving as interleaved controls. Dorsal hippocampal slices were left to recover for 4 h at 30°C in preoxygenated ACSF containing the following (in mM): 124 NaCl, 1.25 NaH₂PO₄, 3 KCl, 26 NaHCO₃, 10 glucose, 2 CaCl₂, 1 MgCl₂. Slices were transferred to a chamber containing actinomycin D (25 μM) and vehicle, 2-chloro-4-((2,5-dimethyl-1-(4-(trifluoromethoxy)phenyl)-1H-imidazol-4-yl)ethynyl)pyridine (CTEP; 10 μM), 1,4-diamino-2,3-dicyano-1,4-bis(2-aminophenylthio)butadiene (U0126; 5 μM), or lovastatin (100 μM) for 30 min, and then transferred to a chamber with fresh ACSF containing 0.4 Mbq/ml ³⁵S-Met/Cys protein labeling mix (PerkinElmer) and vehicle, CTEP (10 μM), U0126 (5 μM), or lovastatin (100 μM) for 45 min. After labeling, slices were frozen on dry ice and stored at -80°C. For processing, slices were homogenized in ice-cold homogenising buffer (10 mM HEPES, pH 7.4, 2 mM EDTA, 2 mM EGTA,

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1% Triton X-100) with protease inhibitors (Roche) and phosphatase inhibitors (cocktails II and III; Sigma). Protein was precipitated with TCA (12.5% final), and total protein concentration was measured using a Bio-Rad protein assay. Samples were read using a scintillation counter, and data were expressed as the number of counts per minute per microgram of protein and normalized to the ³⁵S-Met/Cys ACSF used for incubation.

Western blotting. For Western blotting, metabolically labeled hippocampal homogenates were taken before TCA precipitation and boiled in Laemmli sample buffer, resolved on SDS polyacrylamide gels (Bio-Rad), transferred to nitrocellulose, and incubated overnight in primary antibodies for p-ERK1/2 (Thr202/Tyr204; Cell Signaling Technology), ERK1/2 (Cell Signaling Technology), and β -actin (Abcam). Blots were incubated with their respective fluorophore-conjugated secondary antibodies and imaged on an Odyssey imaging system (LiCor Biosciences).

Stimulated emission depletion microscopy and dendritic spine analysis. Quantitative morphometric analysis of dendritic spines using stimulated emission depletion (STED) microscopy on dorsal hippocampal brain sections from perfusion-fixed P35 animals was performed as reported previously (Wijetunge et al., 2014). Spines on three to four apical oblique dendritic segments within *stratum radiatum* of CA1 were acquired per animal where each dendrite was from a different cell and analyzed blind to the genotype. Confocal microscopy analysis was performed as reported previously (Till et al., 2012), except that the cells were filled in lightly postfixed (4% paraformaldehyde) acute hippocampal slices from P26–P32 animals. Statistical analyses of spine morphology were performed as described previously (Wijetunge et al., 2014).

Results

Enhanced and protein synthesis independent mGluR-LTD in *Syngap*^{+/-} hippocampal slices

To determine whether SynGAP haploinsufficiency results in a hippocampal pathophysiology similar to that observed in mouse models of FXS, we first examined whether the loss of a single copy of *Syngap* mimics the increase in mGluR-LTD reported in *Fmr1*^{-/-} mice. The application of DHPG, a selective group 1 mGluR agonist, resulted in a significant increase in the magnitude of LTD in *Syngap*^{+/-} hippocampal slices compared to WT controls (WT, 77 ± 3%, *n* = 18; *Syngap*^{+/-}, 62 ± 5%, *n* = 15; *p* = 0.012). At this developmental time point (P25–P32), the expression of this form of LTD is dependent on the translation of mRNAs at local dendritic sites (Huber et al., 2000). Preincubation of hippocampal slices in the protein synthesis inhibitor anisomycin (20 μ M) revealed that mGluR-LTD is no longer dependent on new protein synthesis in *Syngap*^{+/-} mutants (64 ± 4%, *n* = 14; Fig. 1B). In contrast, anisomycin eliminated the persistent decrease in fEPSP slope following DHPG application in WT controls (91 ± 4%, *n* = 12). Furthermore, consistent with previous observations, and under our own experimental conditions, we also observed exaggerated mGluR-LTD that was independent of protein synthesis in *Fmr1*^{-/-} mice (Fig. 1C,D; Huber et al., 2002; Nosyreva and Huber, 2006). Thus, these two ID-related mutations in *Fmr1* and *Syngap* lead to similar alterations in protein synthesis-dependent mGluR-LTD in hippocampal CA1.

Based on these LTD findings, we next wanted to examine whether mutations that cause FXS and SynGAP haploinsufficiency lead to the disruption of a common pathophysiological mechanism downstream of the mGlu₅ receptor. To test this hypothesis, we assessed whether introducing the *Syngap* mutation into the *Fmr1*^{-/-} mouse would occlude or exacerbate the LTD phenotype caused by the loss of FMRP. Figure 1, E and F, shows that in double mutant (*Syngap*^{+/-}/*Fmr1*^{-/-}) mice, the magnitude of mGluR-LTD is not significantly greater than that observed in either single mutant alone.

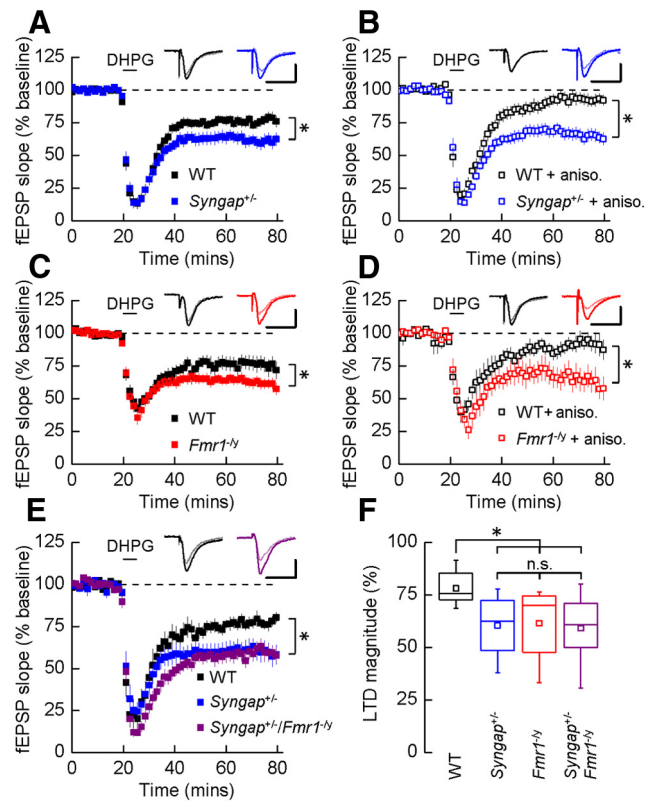


Figure 1. *Syngap*^{+/-} phenocopies the hippocampal synaptic pathophysiology observed in *Fmr1*^{-/-} mice. **A**, Application of DHPG (50 μ M) induced LTD that was significantly increased in *Syngap*^{+/-} mice (62 ± 5%, *n* = 15; *t* test, *p* = 0.01) versus WT littermate controls (77 ± 3%, *n* = 18). Representative average fEPSPs before and after DHPG application are illustrated. **B**, In the presence of protein synthesis inhibitor anisomycin (aniso., 20 μ M), DHPG-induced LTD was not sustained in WT mice (91 ± 4%, *n* = 12), while remaining intact in *Syngap*^{+/-} mice (64 ± 4%, *n* = 14). **C**, DHPG-induced mGluR-LTD was also significantly enhanced in *Fmr1*^{-/-} mice (63 ± 4%, *n* = 17; *t* test, *p* = 0.04) versus WT littermate controls (77 ± 5%, *n* = 17). **D**, In the presence of anisomycin, mGluR-LTD remained at a similar magnitude in the *Fmr1*^{-/-} (65 ± 7%, *n* = 9), whereas LTD could not be sustained in WT mice (91 ± 6%, *n* = 8). **E**, The magnitude of LTD was significantly enhanced in the *Syngap*^{+/-}/*Fmr1*^{-/-} double mutant mice (59 ± 4%, *n* = 12; ANOVA, *p* = 0.02) relative to WT littermate controls (78 ± 3%; *n* = 9). No significant differences were observed in LTD magnitude between the *Syngap*^{+/-}/*Fmr1*^{-/-} double mutant mice and either *Syngap*^{+/-} (60 ± 4%, *n* = 10) or *Fmr1*^{-/-} (62 ± 5%, *n* = 10; data not illustrated) single mutant mice. **F**, Summary of DHPG-induced LTD for each of the four genotypes generated from the *Syngap*^{+/-} × *Fmr1*^{-/-} cross. Box plots illustrate minima and maxima (whiskers), median (line), mean (square symbol), and interquartile range (box). Calibrations: **A, D**, 250 μ V, 10 ms; **B, C**, 500 μ V, 10 ms; **E**, 350 μ V, 10 ms. n.s., Not significant. **p* < 0.05.

To confirm that the occlusion of LTD in the double mutants did not arise from a “floor effect,” mGluR-LTD was induced for a second time 1 h after the initial stimulation to determine whether the first application of DHPG led to a saturating level of LTD. In agreement with previous studies (Palmer et al., 1997; Schnabel et al., 1999; Huber et al., 2001), a subsequent application of DHPG (50 μ M) significantly increased the magnitude of LTD relative to the first stimulation in slices prepared from all four genotypes (*p* < 0.05; Fig. 2A, B).

Syngap^{+/-} hippocampal slices have elevated basal protein synthesis and enhanced ERK1/2 activity

Previously, it was reported that exaggerated mGluR-LTD in *Fmr1*^{-/-} mutants correlates with an increase in basal protein synthesis (Dölen et al., 2007). Therefore, we next examined protein synthesis rates in hippocampal slices from *Syngap*^{+/-} mice em-

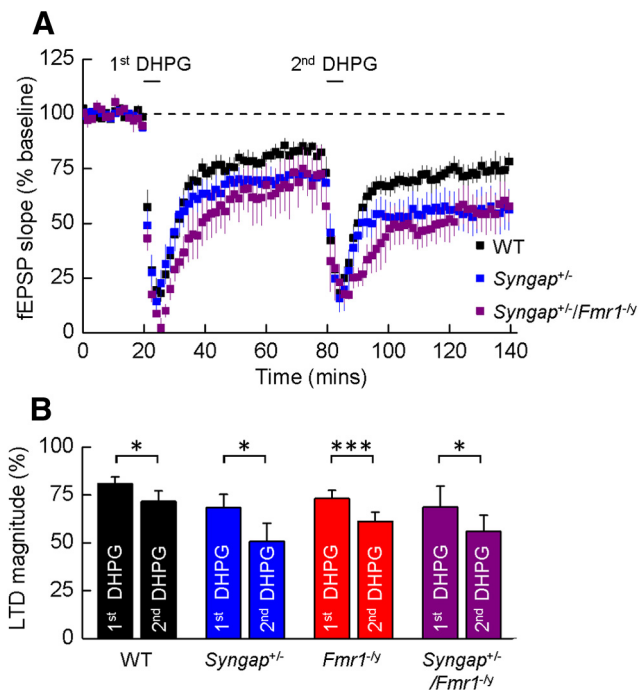


Figure 2. A single application of DHPG does not “saturate” LTD in *Syngap*^{+/-}/*Fmr1*^{-/-} double mutant mice. **A**, Mean time courses illustrating fEPSP slope in experiments on slices prepared from WT, *Syngap*^{+/-}, and *Syngap*^{+/-}/*Fmr1*^{-/-} mice where two applications of DHPG (50 μ M) were applied to determine whether the magnitude of LTD elicited by the first application could be augmented by the second. **B**, Bar graphs illustrating the magnitude of mGluR-mediated LTD produced by a first and then a second application of DHPG in slices prepared from WT (first, 81 \pm 3%; second, 71 \pm 5%; n = 8; p < 0.05, paired t test), *Syngap*^{+/-} (first, 68 \pm 7%; second, 51 \pm 9%; n = 5; p < 0.05), *Fmr1*^{-/-} (first, 73 \pm 4%; second, 61 \pm 5%; n = 3; p < 0.001), and *Syngap*^{+/-}/*Fmr1*^{-/-} (first, 69 \pm 11%; second, 56 \pm 8%; n = 4; p < 0.05) mice. * p < 0.05, *** p < 0.001.

employing a similar methodology to that described previously in *Fmr1*^{-/-} mutants (Fig. 3A; Osterweil et al., 2010). Consistent with our electrophysiological findings, we found a significant increase in ³⁵S-Met/Cys incorporation in *Syngap*^{+/-} hippocampal slices (140 \pm 12%, n = 9; Fig. 3B), indicating that SynGAP may function to suppress mRNA translation under steady-state conditions. Although the magnitude of this increase in basal protein synthesis was greater than that observed in *Fmr1*^{-/-} versus WT slices (121 \pm 7%; n = 7), our findings suggest that the proteins required for mGluR-LTD may be overtranslated in both the *Fmr1*^{+/-} and *Syngap*^{+/-} brain.

In *Fmr1*^{-/-} mice, increased protein synthesis rates are caused by a hypersensitive response to constitutive mGlu₅-Ras-ERK1/2 signaling due to the loss of the translational repression by FMRP (Osterweil et al., 2010). In the *Syngap*^{+/-} hippocampus, both Ras and ERK1/2 activity are increased (Komiya et al., 2002; Ozkan et al., 2014), leading us to hypothesize that elevated protein synthesis rates may arise from dysregulated ERK1/2 signaling due to the loss of Ras inactivation. To confirm that we observed a similar increase in ERK1/2 activity, phosphorylation levels were quantified in hippocampal slices from *Syngap*^{+/-} mice and WT controls under identical conditions to which basal protein synthesis rates were found to be enhanced. In agreement with previous studies, ERK1/2 phosphorylation was significantly increased in *Syngap*^{+/-} mutants under steady-state conditions in the absence of any detectable differences in total ERK1/2 protein (137 \pm 7%, n = 23; p = 1.5 \times 10⁻⁵; Fig. 3C).

It was shown previously that an mGluR-LTD-inducing stimulus increases protein synthesis rates in WT but not *Fmr1*^{-/-} hippocampal slices (Osterweil et al., 2010). Thus, we measured ³⁵S-Met/Cys incorporation following the acute application of DHPG (100 μ M; 5 min) in *Syngap*^{+/-} hippocampal slices. While DHPG stimulation significantly enhanced protein synthesis in WT slices, the same treatment failed to increase further the already exacerbated protein synthesis levels in *Syngap*^{+/-} mutants (Fig. 3D), suggesting that mRNA translation downstream of group 1 mGluR activation is saturated under nonstimulated conditions. ERK1/2 activation is a critical step in group 1 mGluR-mediated protein synthesis governing the expression and maintenance of mGluR-LTD (Gallagher et al., 2004; Osterweil et al., 2010). In both WT and *Syngap*^{+/-} hippocampal slices, DHPG induced a significant and robust increase in ERK1/2 phosphorylation (Fig. 3E). This result may suggest that exaggerated mGluR-LTD in *Syngap*^{+/-} hippocampal slices could arise from the dysregulation of basal, rather than stimulated, ERK1/2 signaling.

Elevated basal protein synthesis in the *Syngap*^{+/-} hippocampus can be ameliorated by negative modulators of mGlu₅ receptor and Ras-ERK1/2 signaling

To test the hypothesis that excessive mGluR-mediated protein synthesis in *Syngap*^{+/-} mice is a consequence of hyperactive Ras-ERK1/2 signaling, we examined the effects of inhibitors of this pathway on translational rates. CTEP, lovastatin, and U0126 selectively target and downregulate mGlu₅ receptors, Ras, and ERK1/2 signaling, respectively (Fig. 4A), and have been shown previously to restore exaggerated levels of protein synthesis and/or functional deficits in *Fmr1*^{-/-} mice (Osterweil et al., 2010, 2013; Michalon et al., 2012, 2014). We found that each of these compounds reduced basal protein synthesis in *Syngap*^{+/-} hippocampal slices to levels not significantly different from WT controls (Fig. 4B,Ci,Di). Importantly, none of these compounds significantly altered basal protein synthesis in WT animals, suggesting that their effect is specific to mechanisms underlying the hyperactivation of ERK1/2 in *Syngap*^{+/-} mice. To confirm that both lovastatin and U0126 elicited their effects on protein synthesis by downregulating ERK1/2 activity, we quantified the phosphorylation status of ERK1/2 in control and treated hippocampal slices from *Syngap*^{+/-} and WT mice. Lovastatin significantly reduced ERK1/2 phosphorylation in *Syngap*^{+/-} slices to WT levels without affecting ERK1/2 phosphorylation in WT treated slices (Fig. 4Cii,Ciii). In contrast, U0126 caused a dramatic reduction in ERK1/2 activity in both WT and *Syngap*^{+/-} hippocampal slices (Fig. 4Dii,Diii). Together, these findings suggest that dysregulated ERK1/2 signaling leads to exaggerated protein synthesis rates in *Syngap*^{+/-} hippocampus, and drugs that have been shown previously to ameliorate phenotypes associated with the loss of FMRP can restore translational rates in a mouse model of SYNGAP haploinsufficiency.

Altered morphology of dendritic spines of CA1 pyramidal neurons in *Syngap*^{+/-} mice

Next, we examined whether the electrophysiological and biochemical changes we observed in *Syngap*^{+/-} mice are associated with density and morphological alterations in dendritic spines, which are the main site of excitatory input to CA1 pyramidal neurons. To determine whether the reduction in SynGAP expression affected the density of dendritic spines, we dye filled cells obtained from the “partner” hemispheres of brains that had been used for electrophysiological studies. Figure 5 illustrates that no difference in spine densities were observed in *Syngap*^{+/-} mice

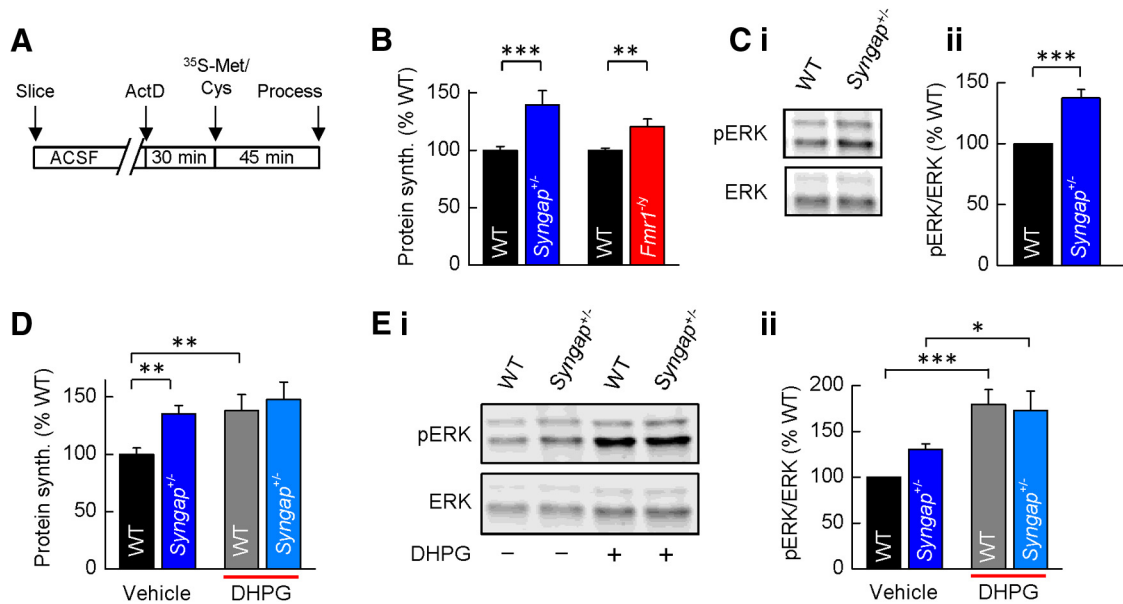


Figure 3. Basal protein synthesis and phosphorylated ERK1/2 levels are elevated in the hippocampus of *Syngap*^{+/-} mice. **A**, Schematic of the experimental timeline for [³⁵S]-Met/Cys metabolic labeling. **B**, Basal protein synthesis levels were significantly elevated in dorsal hippocampal slices from *Syngap*^{+/-} versus WT mice (WT, 100 ± 3%; *Syngap*^{+/-}, 140 ± 12%; *t* test, *p* = 0.0007; *n* = 9). Increased protein synthesis rates were also observed in the *Fmr1*^{-/-} versus WT mice (WT, 100 ± 2%; *Fmr1*^{-/-}, 121 ± 7%; *t* test, *p* = 0.01; *n* = 6). **C*i***, **C*ii***, Basal activation state of ERK1/2 in hippocampal slices was significantly increased in *Syngap*^{+/-} mice versus WT controls (phosphorylated/total ERK1/2, 137 ± 7%; *n* = 23; *p* = 1.5 × 10⁻⁵). **D**, Protein synthesis levels were significantly elevated in vehicle-treated dorsal hippocampal slices from *Syngap*^{+/-} versus WT mice (WT, 100 ± 5%; *Syngap*^{+/-}, 135 ± 7%; *n* = 6; ANOVA, *p* = 0.0038). DHPG treatment (100 μM, 5 min) significantly increased protein synthesis rates in WT slices, but did not further increase [³⁵S]-Met/Cys incorporation in *Syngap*^{+/-} slices (WT DHPG, 138 ± 14%; *Syngap*^{+/-} DHPG, 148 ± 15%; ANOVA, WT treatment, *p* = 0.0028; *n* = 6). **E*i***, **E*ii***, Western blot analysis of vehicle/DHPG-treated hippocampal slices revealed DHPG significantly increases the phosphorylation status of ERK1/2 in both *Syngap*^{+/-} and WT mice (phosphorylated/total ERK1/2, WT vehicle, 100%; *Syngap*^{+/-} vehicle, 130 ± 6%; WT DHPG, 180 ± 16%; *Syngap*^{+/-} DHPG, 173 ± 21%; *n* = 7; ANOVA, genotype × treatment, *p* < 0.05). **p* < 0.05; ***p* < 0.01; ****p* < 0.001.

and WT littermate controls (WT, 19.58 ± 1.03 spines/10 μm; *Syngap*^{+/-}, 17.34 ± 0.94 spines/10 μm; *p* = 0.1430, two-tailed independent *t* test with Welch's correction).

To accurately resolve the fine details of spine morphology we used STED microscopy, quantifying the widths and lengths of spine heads and necks at the nanoscale (Fig. 6*A*). For spine head width, no difference in the distributions between *Syngap*^{+/-} and WT neurons was observed, although spine neck lengths and widths showed significantly different distributions (Fig. 6*B–D*; WT, *n* = 393 spines; *Syngap*^{+/-}, *n* = 424 spines; head width, *p* = 0.11; neck length, *p* = 0.005; neck width, *p* = 0.0008, Kolmogorov–Smirnov test). In agreement with our STED analysis, confocal analysis of dendritic spines in the animals from which electrophysiological analyses were performed (P25–P32) also indicates that there was no difference in the mean spine head width between genotypes (WT, 0.350 ± 0.01 μm; *Syngap*^{+/-}, 0.360 ± 0.01 μm; *p* = 0.71, *t* test with Welch's correction; data not shown).

To better understand the potential functional impact of these nanoscale morphological alterations on synapse function, we calculated the compartmentalization factor (defined as *VL/A*, where *V* is the head volume, *L* is the neck length, and *A* is the cross-sectional area of the spine neck), which serves as a measure of the degree of biochemical compartmentalization of spine synapses (Tønnesen et al., 2014). We found that the compartmentalization factor was substantially increased in *Syngap*^{+/-} animals (Fig. 6*E*; *Syngap*^{+/-}, *n* = 424 spines; WT, *n* = 393 spines; compartmentalization factor, *p* = 0.0001, Kolmogorov–Smirnov test; mean compartmentalization factor, WT, 3.16 ± 0.30 μm², *n* = 4 animals; *Syngap*^{+/-}, 4.35 ± 0.37 μm², *n* = 6 animals; *p* = 0.04, two-tailed independent *t* test with Welch's correction), indi-

cating that SynGAP expression influences the biochemical coupling of spines to their parent dendrites. Interestingly, in *Fmr1*^{-/-} mice, the compartmentalization factor was also previously found to be increased relative to WT control animals (Wijetunge et al., 2014).

Discussion

De novo mutations in the *SYNGAP1* gene result in moderate to severe ID, often with co-occurring epilepsy and/or ASDs that appear to be as prevalent as FXS (Berryer et al., 2013). However, unlike FXS, little is known about the underlying mechanisms by which SynGAP reduction causes these alterations in brain function. We now show that SynGAP haploinsufficiency in mice mimics and/or occludes the deficits in synaptic structure and physiology in hippocampal CA1 pyramidal neurons associated with the loss of FMRP. Furthermore, pharmacological interventions that reduce mGlu₅ receptor signaling and rescue much of the pathophysiology associated with the loss of FMRP also rescue the protein synthesis deficits associated with loss of a single copy of *Syngap*. Together, these data suggest that these two prevalent monogenic causes of ID/ASDs share common and/or overlapping cellular mechanisms.

This is the first demonstration that *Syngap*^{+/-} mice show altered mGlu₅ receptor signaling and altered downstream protein synthesis. When considered in conjunction with previous studies that showed an increase in ERK1/2 activation following NMDA receptor activation (Vazquez et al., 2004; Rumbaugh et al., 2006), a decrease in NMDA receptor-dependent LTP (Komiyama et al., 2002; Kim et al., 2003), and a block of the induction of NMDA receptor-dependent LTD (Carlisle et al., 2008), these data suggest that SynGAP regulates two distinct glutamate receptor depen-

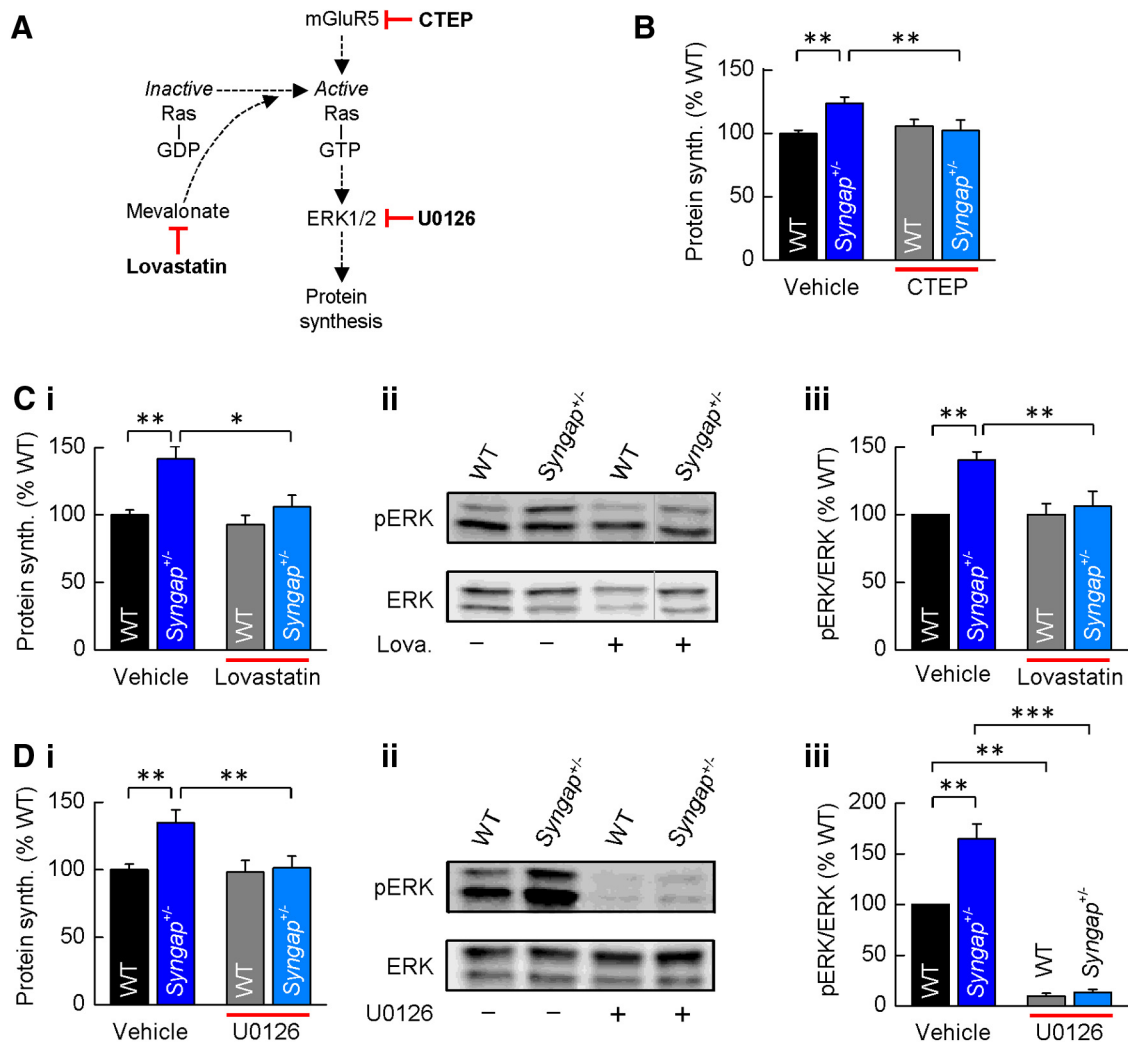


Figure 4. Elevated protein synthesis in *Syngap*^{+/-} hippocampus is corrected by inhibitors of mGluR5 and Ras–ERK1/2 signaling. **A**, schematic of the drug targets for CTEP, lovastatin, and U0126, each shown previously to reduce Ras–ERK1/2 activation and correct basal protein synthesis in *Fmr1*^{-/-} mice. **B**, CTEP (10 μ M) reverses elevated protein synthesis in *Syngap*^{+/-} mice (WT, vehicle, 100 \pm 3%; CTEP, 106 \pm 5%; *Syngap*^{+/-}, vehicle, 124 \pm 5%; CTEP, 102 \pm 8%; *n* = 9; ANOVA, genotype, *p* = 0.0007; KO treatment, *p* = 0.0014). **C–Ciii**, Lovastatin (100 μ M) reverses elevated protein synthesis (**Ci**) in *Syngap*^{+/-} mice (WT, vehicle, 100 \pm 4%; lovastatin, 93 \pm 7%; *Syngap*^{+/-}, vehicle, 142 \pm 9%; lovastatin, 106 \pm 8%; *n* = 7; ANOVA, genotype, *p* = 0.0061; KO treatment, *p* = 0.013). Example Western blot (**Cii**) and quantification of ERK1/2 activity (**Ciii**) in vehicle- and lovastatin-treated hippocampal slices revealed lovastatin significantly reduces phosphorylation levels of ERK1/2 in *Syngap*^{+/-} mice (phosphorylated/total ERK1/2, WT, vehicle, 100%; *Syngap*^{+/-}, vehicle, 140 \pm 6%; WT, lovastatin, 100 \pm 8%; *Syngap*^{+/-}, lovastatin, 106 \pm 11%; *n* = 10; ANOVA, genotype, **p* = 0.0086; KO treatment, *p* = 0.0057). The gray line to the left of the lovastatin-treated *Syngap*^{+/-} Western blot indicates that this lane was not adjacent to each of the other three lanes, but was run on the same gel, and hence was processed identically. **D**, U0126 (5 μ M) reverses elevated protein synthesis (**Di**) in *Syngap*^{+/-} mice (WT, vehicle, 100 \pm 4%; U0126, 98 \pm 8%; *Syngap*^{+/-}, vehicle, 134 \pm 9%; U0126, 102 \pm 9%; *n* = 9; ANOVA, genotype, *p* = 0.003; KO treatment, *p* = 0.004). Example Western blot (**Dii**) and quantification (**Diii**) of ERK1/2 activity in vehicle- and U0126-treated hippocampal slices show U0126 abolishes ERK1/2 activity in both WT and *Syngap*^{+/-} mice (phosphorylated/total ERK1/2, WT, vehicle, 100%; *Syngap*^{+/-}, vehicle, 165 \pm 15%; WT, lovastatin, 10 \pm 3%; *Syngap*^{+/-}, lovastatin, 13 \pm 3%; *n* = 5; ANOVA, genotype, *p* = 0.0054; WT treatment, *p* = 0.005; KO treatment, *p* = 0.0002). **p* < 0.05; ***p* < 0.01; ****p* < 0.001.

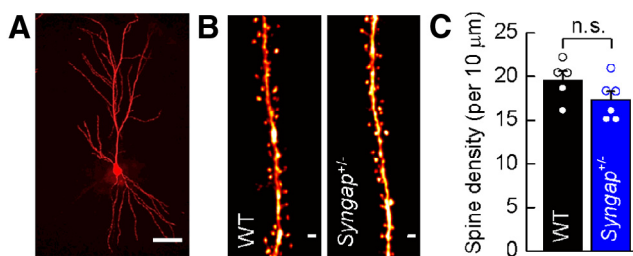


Figure 5. CA1 pyramidal neuron dendritic spine density in *Syngap*^{+/-} mice. **A**, Composite image of a representative CA1 pyramidal neuron labeled by intracellular filling with Alexa Fluor 568 in lightly fixed section through CA1 of hippocampus. Scale bar, 50 μ m. **B**, Representative composite images of apical oblique dendritic segments from WT and *Syngap*^{+/-} mice. Scale bars, 1 μ m. **C**, Mean spine density of apical dendrites for WT (*n* = 5) and *Syngap*^{+/-} (*n* = 6) animals indicating no difference in dendritic spine density (WT, 19.58 \pm 1.03 spines/10 μ m; *Syngap*^{+/-}, 17.34 \pm 0.94 spines/10 μ m; *p* = 0.1430, two-tailed independent *t* test with Welch’s correction). n.s., Not significant.

dent forms of synaptic plasticity. Intriguingly, we showed previously that the *Syngap* gene gives rise to a range of isoforms that are differentially regulated by synaptic activity and that have opposing effects on synaptic function (McMahon et al., 2012). For example, overexpression of SynGAP A α 1 results in a decrease in mEPSC frequency and amplitude, whereas SynGAP Ca2 increases mEPSC frequency and amplitude. SynGAP isoforms also demonstrate distinct activity- and age-dependent patterns of expression (McMahon et al., 2012). Therefore, the loss of a single functional allele of SynGAP may result in an imbalance of SynGAP isoforms as well as decreasing overall SynGAP levels. It remains to be determined whether the deficits seen in *Syngap*^{+/-} mice result from a reduction in total SynGAP levels or whether an imbalance of SynGAP isoform expression is responsible for the altered cellular phenotypes associated with SynGAP haploinsufficiency.

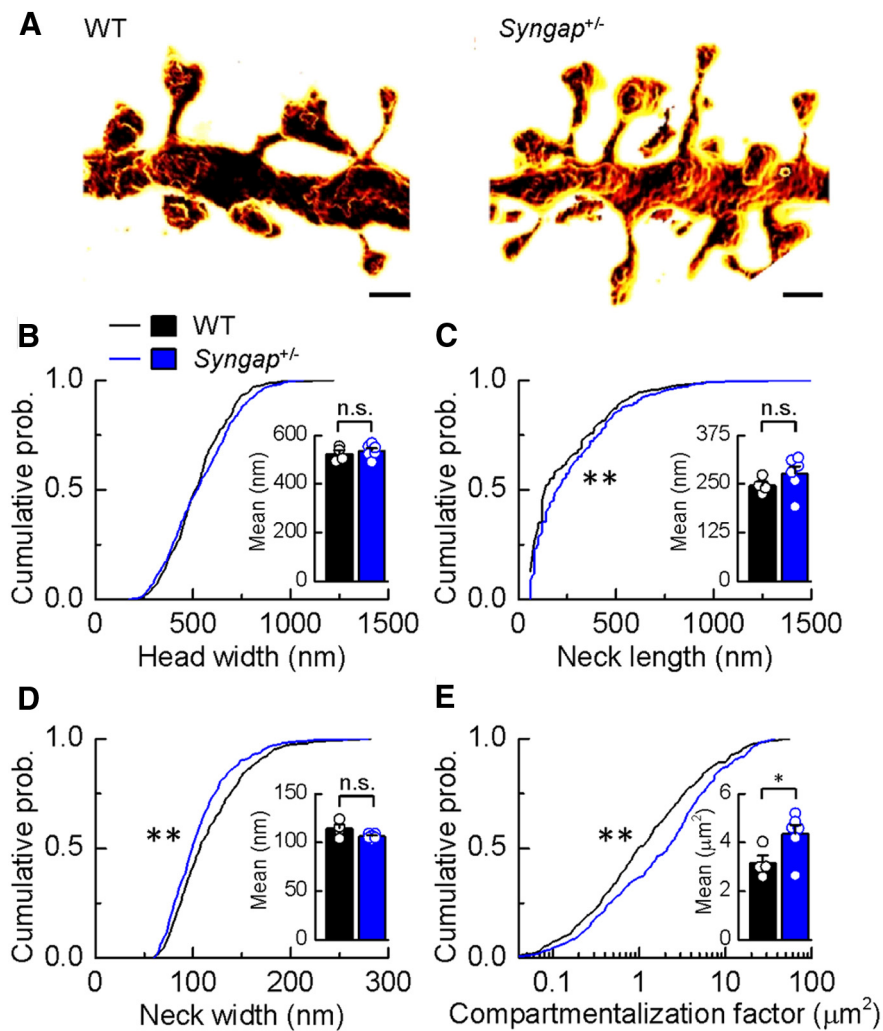


Figure 6. CA1 pyramidal neuron dendritic spine morphology in *Syngap*^{+/-} mice. **A**, Surface-rendered STED images of apical dendritic segments from pyramidal neurons in the CA1 of the hippocampus of WT and *Syngap*^{+/-} mice revealing morphological details that are invisible to conventional light microscopy. Scale bars, 500 nm. **B–D**, Kolmogorov–Smirnov tests of the cumulative frequency distributions show that the distribution profiles of spine head widths (**B**) are not significantly different between genotypes ($p = 0.11$), whereas distribution profiles of spine neck lengths and widths show that *Syngap*^{+/-} mice ($n = 424$ spines) have significantly more spines with longer (**C**; $p < 0.005$) and narrower necks (**D**; $p < 0.0008$) compared to WT mice ($n = 393$ spines). The mean values for these parameters are, however, not significantly different (WT, 523 ± 14 nm, 246 ± 10 nm, 114 ± 4 nm, $n = 4$; *Syngap*^{+/-}, 536 ± 11 nm, 276 ± 19 nm, 106 ± 1 nm, $n = 6$, for head width, neck length, and neck width, respectively). **E**, To predict the impact of these spine morphology changes on diffusional coupling, a morphological compartmentalization factor was calculated. The cumulative frequency distributions of the compartmentalization factor significantly differ between genotypes (Kolmogorov–Smirnov test, $p < 0.0001$), and this is also reflected in the mean values for each genotype (two-tailed independent t test with Welch's correction, $p = 0.04$; WT, $3.16 \pm 0.30 \mu\text{m}^2$; *Syngap*^{+/-}, $4.35 \pm 0.37 \mu\text{m}^2$). n.s., Not significant; * $p < 0.05$; ** $p < 0.01$.

Our findings are in good agreement with a previous report using acute and selective knockdown of the SynGAP $\alpha 1$ isoform in cortical cultures showing an ERK1/2-dependent increase in dendritic protein synthesis (Wang et al., 2013). Our data extend these findings by demonstrating that constitutive reduction of all SynGAP isoforms, a model that more closely reflects SYNGAP1 haploinsufficiency in humans (i.e., has excellent structural and face validity with the human condition), leads to an increase in ERK1/2 phosphorylation and basal protein synthesis. Furthermore, the latter can be corrected by inhibitors of the ERK1/2 signaling pathway. Importantly an inhibitor of ERK1/2 activator MEK1/2 (U0126) restored protein synthesis to wild-type levels, despite a dramatic reduction in ERK1/2 phosphorylation, indicating that only the exagger-

ated protein synthesis is regulated by ERK1/2. However, it seems likely that the dramatic reduction in ERK1/2 signaling could dramatically alter neuronal function and plasticity. In this respect, the finding that lovastatin restored WT levels of ERK1/2 phosphorylation and basal protein synthesis likely make it a far better candidate therapy for SYNGAP1 haploinsufficiency. Importantly, the selective effect on elevated protein synthesis in *Syngap*^{+/-} mice is similar to that observed in *Fmr1*^{-/-} mice, suggesting that dysregulation of mGlu₅-ERK1/2-dependent protein synthesis may be a common feature of a subset of neurodevelopmental disorders (Osterweil et al., 2010, 2013; Tian et al., 2015). In summary, our findings suggest that targeting the exaggerated basal protein synthesis through modulation of the ERK1/2 pathway may provide a potential therapeutic strategy in the treatment of SYNGAP haploinsufficiency.

Using STED microscopy, we reveal nanoscale changes in spine morphology in *Syngap*^{+/-} mice that indicate an increase in biochemical compartmentalization. These findings parallel those from the *Fmr1*^{-/-} mice, where changes in spine neck and head parameters decrease the magnitude of the developmental change in biochemical compartmentalization that occurs in WT animals (Wijetunge et al., 2014). As we suggested previously for the *Fmr1*^{-/-} mice (Wijetunge et al., 2014), we deem it unlikely that these subtle changes in spine morphology can, by themselves, fully account for the alterations in synaptic function and plasticity.

Previous findings from SYNGAP-deficient mice suggest that changes in synaptic function are mirrored by changes in synaptic structure (Vazquez et al., 2004; Carlisle et al., 2008; Clement et al., 2012; but see McMahan et al. 2012). More specifically, SynGAP reduction resulted in an increase in dendritic spine head size (Vazquez et al., 2004; Carlisle et al., 2008;

Clement et al., 2012). Our finding of no change in spine head width is at odds with those of Carlisle et al. (2008) and Clement et al. (2012), who examined hippocampal CA1 pyramidal neurons and hippocampal dentate gyrus cells, respectively. Although these differences may partly be due to differences in age and neuronal type that were examined [for example, Carlisle et al. (2008) examined animals up to 6 months of age, and we used animals up to 5 weeks of age], it is likely that methodological issues also explain some of the discrepancies. Both Clement et al. (2012) and Carlisle et al. (2008) categorized spines into “mushroom,” “thin,” and “stubby”; however, we demonstrated previously that spine shapes cover a wide continuum, and such categories are arbitrary (Tønnesen et al., 2014; Wijetunge et al., 2014). Carlisle et al. (2008) found small differences in the size of

“mushroom” spines and no difference in “thin” or “stubby” spines. However, without such categorization, it is unclear whether the differences between genotype would have remained. Clement et al. (2012) found only modest increases in head diameter analyzed using cumulative distributions of spine head width. They also found an increase in the relative proportion of “mushroom” to “stubby” spines in *Syngap*^{+/-} mice. However, the categorization of stubby spines may largely be a consequence of the limited resolution of regular confocal and two-photon microscopy (~200 and 350 nm, respectively) to resolve short spine necks (Tønnesen et al., 2014). For these reasons, it is not appropriate to directly compare between these studies, and the apparent differences may be simply due to differences in methodological approach. Given the excellent resolution of the STED microscope (~70 nm) and the good agreement of our findings of spine development with previous electron microscopic studies (Harris et al., 1992), we argue that our measurements are an accurate reflection of spine head morphology.

Previous studies in *Fmr1*^{-/-} mice have found a preponderance of long thin spines and have correlated this with an increase in LTD (for review, see Wijetunge et al., 2013). We propose that our recent finding of an increase in spine head size (Wijetunge et al., 2014) is more suggestive of spines that have recently undergone potentiation and hence are pre-disposed to undergo LTD. Our findings that *Syngap*^{+/-} mice have no change in head size but do show enhanced LTD indicate that these two parameters are not functionally interdependent. Notwithstanding, both models show an increase in biochemical compartmentalization, and future studies will be required to determine whether compartmentalization and excess LTD are causally linked. Finally, it should be noted that whereas an increase in head size in *Fmr1*^{-/-} mice was generally thought to reflect a premature development of spines, we demonstrated previously (also using STED microscopy) that spine head size decreases with age in cortical layer 5 and hippocampal CA1 pyramidal neurons in WT mice. This implies that maintained large spine heads in *Fmr1*^{-/-} mice actually reflects a developmental delay (Wijetunge et al., 2014).

In summary, our data demonstrate that two highly penetrant forms of ID/ASDs resulting from mutations in *FMR1* and *SYNGAP1* converge on common cellular processes. They are in agreement with studies of the *neurofibromatosis 1 (NF1)* gene, where mutations in *NF1*, encoding a RasGAP, also result in ID, and some of its associated cellular and behavioral phenotypes are alleviated by modulators of ERK1/2 signaling (Li et al., 2005). Furthermore, they support the hypothesis that alterations in synaptic plasticity resulting from dysregulation of new protein synthesis may be a common core pathophysiology underlying these disorders, and therapeutic strategies targeting modifiers of protein synthesis may be appropriate for treating genetically divergent forms of ID/ASD.

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