Mdm2 mediates FMRP- and Gp1 mGluR-dependent protein translation and neural network activity

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Abstract
Activating Group 1 (Gp1) metabotropic glutamate receptors (mGluRs), including mGluR1 and mGluR5, elicits translation-dependent neural plasticity mechanisms that are crucial to animal behavior and circuit development. Dysregulated Gp1 mGluR signaling has been observed in numerous neurological and psychiatric disorders. However, the molecular pathways underlying Gp1 mGluR-dependent plasticity mechanisms are complex and have been elusive. In this study, we identified a novel mechanism through which Gp1 mGluR mediates protein translation and neural plasticity. Using a multi-electrode array (MEA) recording system, we showed that activating Gp1 mGluR elevates neural network activity, as demonstrated by increased spontaneous spike frequency and burst activity. Importantly, we validated that elevating neural network activity requires protein translation and is dependent on fragile X mental retardation protein (FMRP), the protein that is deficient in the most common inherited form of mental retardation and autism, fragile X syndrome (FXS). In an effort to determine the mechanism by which FMRP mediates protein translation and neural network activity, we demonstrated that a ubiquitin E3 ligase, murine double minute-2 (Mdm2), is required for Gp1 mGluR-induced translation and neural network activity. Our data showed that Mdm2 acts as a translation suppressor, and FMRP is required for its ubiquitination and down-regulation upon Gp1 mGluR activation. These data revealed a novel mechanism by which Gp1 mGluR and FMRP mediate protein translation and neural network activity, potentially through de-repressing Mdm2. Our results also introduce an alternative way for understanding altered protein translation and brain circuit excitability associated with Gp1 mGluR in neurological diseases such as FXS.

Introduction
The ability of living organisms to adapt to constantly changing environments depends on the plasticity of their nervous systems. Neural plasticity often requires activity-dependent translation to rapidly supply specific proteins (1–4). One well-known mechanism underlying activity-mediated translation occurs through Group 1 metabotropic glutamate receptors (Gp1 mGluRs) (5,6). Activating Gp1 mGluRs, including mGluR1 and
mGluR5, elicits multiple translation-dependent neural plasticity mechanisms (7–9). Dysregulated Gp1 mGluR signaling is observed in both neurological and psychiatric disorders, including fragile X syndrome (FXS) (10), autism spectrum disorders (ASDs) (11), Alzheimer’s disease (12), and epilepsy (13). However, because of the complexity of Gp1 mGluR signaling, pharmacological correction of Gp1 mGluR signaling in animal models of such diseases has been challenging. In this study, we aim to characterize novel and promising signaling pathways that underlie Gp1 mGluR-dependent protein translation and neural plasticity with the intention of revealing novel therapeutic targets.

Activating Gp1 mGluRs leads to elevated Phosphatidylinositol-4,5-bisphosphate 3-kinase (PI3K) or Extracellular Signal-regulated Kinase-1/2 (ERK1/2) signaling (14–16), which in turn activates eIF4E and translation initiation (17–21). Several recent studies have also demonstrated that translation elongation is facilitated upon mGluR activation through the phosphorylation of eukaryotic elongation factor 2 (eEF2) (22–26). These mechanisms belong to general translational regulation and are often difficult to target for therapeutic intervention. To begin to search for other promising targets underlying Gp1 mGluR-mediated protein translation, we are studying murine double minute-2 (Mdm2), a ubiquitin E3 ligase which was recently shown to be dysregulated in FXS mouse model, Fmr1 knockout (KO) mice (27,28). Mdm2 directly interacts with multiple ribosomal proteins, including ribosomal protein 11 and ribosomal protein 5 (29–31). Studies indicate that interactions between Mdm2 and ribosomal proteins prevent Mdm2 from interacting with Mdm2’s substrates (29,32,33). Although Mdm2 did not appear to regulate translation in cancer cells (34), it remains unknown whether Mdm2 can participate in translational regulation under physiological conditions in normal cells. In our current study, we revealed strong association between Mdm2 and active protein translation has an effect on the synchrony of spontaneous spikes (i.e. action potentials). To determine whether this elevation requires protein translation, we employed a translation inhibitor, cycloheximide (CHX, 60 μM). While CHX alone did not have an effect, pre-application of CHX for 10 min attenuates DHPG-elevated spontaneous spike frequency. The average spontaneous spike amplitude did not differ between all 4 groups (Fig. 1B). These data indicate that Gp1 mGluR-elevated spontaneous spike frequency requires protein translation.

Results

Activation of Gp1 mGluR leads to translation-dependent elevation of neural network activity

To study Gp1 mGluR-mediated neural network activity, we employed a multi-electrode array (MEA) recording system to record extracellular spontaneous spikes in primary cortical neuron cultures (36). Because this system allows recording and analysis from the same cultures before and after treatment, the variability between cultures is greatly diminished. As shown in Figure 1A, activating Gp1 mGluR using an agonist, (RS)-3,5-Dihydroxyphenylglycine (DHPG, 100 μM, 30 min), significantly elevates the frequency of spontaneous spikes (i.e. action potentials). To determine whether this elevation requires protein translation, we employed a translation inhibitor, cycloheximide (CHX, 60 μM). While CHX alone did not have an effect, pre-application of CHX for 10 min attenuates DHPG-elevated spontaneous spike frequency. The average spontaneous spike amplitude did not differ between all 4 groups (Fig. 1B). These data indicate that Gp1 mGluR-induced spontaneous spike frequency requires protein translation.

In addition to elevated spontaneous spike frequency, we also asked whether the pattern of spontaneous spikes is affected by Gp1 mGluR activation. If so, we further investigated whether or not it is also dependent on protein translation. We first determine the burst activity, which is a series of spontaneous spikes occurred within a short period of time. We defined a minimum of 5 spontaneous spikes with a maximum inter-spike interval of 0.1 second as a burst, as previously reported (36), and confirmed that activating Gp1 mGluR using applications of DHPG (100 μM, 15 min) elevates burst activity, as demonstrated by increased burst duration (Fig. 2A and B) and average number of spikes per burst (Fig. 2C). Remarkably, inhibiting translation by CHX attenuates DHPG-triggered burst activity, suggesting protein translation is also required for Gp1 mGluR-induced burst activity.

Our previous study demonstrated synchronization of spontaneous spikes across a culture when the neurons are hyperexcitable (36). To determine whether activating Gp1 mGluR also affects the synchrony of spontaneous spikes, we measured the synchrony index before and after DHPG treatment. As shown in Figure 2D, neither activation of Gp1 mGluR nor inhibition of translation has an effect on the synchrony of spontaneous spikes. Together, our data showed that Gp1 mGluR mediates the translation-dependent elevation of spontaneous spike frequency and burst activity.

FMRP is required for Gp1 mGluR-induced neural network activity

Gp1 mGluR-dependent protein translation is co-regulated by FMRP (37–39). In the absence of FMRP, Gp1 mGluR signaling is dysregulated and leads to many mGluR-associated deficits in FXS (40–45). To determine whether FMRP participates in the elevation of neural network activity upon Gp1 mGluR activation, we employed Fmr1 KO mice, in whom the gene encoding FMRP has been deleted. As shown in Figure 3A, compared to the cortical neuron cultures made from wild type (WT) mice (Fig. 3A1 and 3A3), Fmr1 KO cortical neuron cultures did not exhibit elevated spontaneous spike frequency after DHPG treatment (Fig. 3A2 and A3). The spontaneous spike amplitude did not differ between vehicle or DHPG treatment in either WT or Fmr1 KO cultures (Fig. 3B). We also asked whether elevated burst activity...
activity upon Gp1 mGluR activation requires FMRP. As shown in Figure 3C, DHPG fails to increase burst duration or the number of spikes per burst in $Fmr1$ KO cultures. Together, these data suggest that Gp1 mGluR-induced elevation of neural network activity, demonstrated by increased spontaneous spike frequency and burst activity, requires FMRP.

Mdm2 is required for Gp1 mGluR-induced neural network activity

We next sought to identify the signaling pathway by which Gp1 mGluR and FMRP mediate neural network activity. Multiple recent studies have identified a connection between FMRP and Mdm2 in the neuronal and synaptic deficits observed in $Fmr1$ KO mice (27,28). To determine the role of Mdm2 in Gp1 mGluR-induced neural network activity, we employed a conditional Mdm2 knockdown mouse model by crossing Mdm2 floxed mice ($Mdm2^{f/f}$) with $Emx1$-Cre mice to obtain $Mdm2^{f/+}$-Emx-Cre+ and $Mdm2^{f/+}$-Emx-Cre- mice. $Emx1$-Cre can confer Mdm2 reduction in the cortex and hippocampus beginning as early as embryonic day 10.5 (E10.5) (46,47). We used heterozygous mice ($Mdm2^{f/+}$) to avoid potential apoptosis caused by complete Mdm2 knockout (48,49). As shown in Figure 4A, we observed approximately 40% knockdown of Mdm2 in the $Mdm2^{f/+}$-Emx-Cre+ cortical neuron cultures in comparison to the $Mdm2^{f/+}$-Emx-Cre- cultures at days-in-vitro (DIV 14) when the cultures were prepared on postnatal day 0.

We next employed primary cortical neuron cultures made from $Mdm2^{f/+}$-Emx-Cre+ or $Mdm2^{f/+}$-Emx-Cre- mice to determine whether and how Mdm2 is involved in regulating neural network activity. As shown in Figure 4B, DHPG elevated

![Figure 1. Activation of Gp1 mGluR induces protein translation-dependent elevation of spontaneous spike frequency. (A) Representative raster plots of spontaneous spikes from 1-min recordings of the same MEA dishes before and after drug treatments as labeled in the Figure (A). Quantification of relative spontaneous spike frequency by comparing ‘after treatment’ to ‘before treatment’, and the dot plot showing the distribution of data points were on the bottom (A) ($n=7–9$). (B) Representative average traces of 1-min recording from the same single electrodes before and after drug treatments as labeled in the figure (B). Quantification of average spontaneous spike amplitude by comparing ‘after treatment’ to ‘before treatment’, and the dot plot were on the bottom (B) ($n=7–9$). For both figures, a two-way ANOVA with Tukey test was used. Data are represented as mean ± SEM with *$P<0.05$, ns: non-significant.](https://humanmoleculargenetics.oxfordjournals.org/content/17/12/3240/F1.large.jpg)
Data are represented as mean comparing ‘after treatment’ to ‘before treatment’, and the dot plot were on the right (PSD-95 were indeed elevated in (PSD-95) (28,50), we tested whether knocking down Mdm2 affects the ubiquitination of post-synaptic density protein 95 To answer this question, we first determined whether knocking down Mdm2 affects basal synaptic transmission or Gp1 mGluR signaling. Despite the connection between FMRP and Mdm2 that was identified in the previous studies (27,28), it remains unknown whether and how Mdm2 participates in Gp1 mGluR signaling. To answer this question, we first determined whether knocking down Mdm2 affects basal synaptic activity. Because Mdm2 mediates the ubiquitination of post-synaptic density protein 95 (PSD-95) (28,50), we tested whether knocking down Mdm2 affects the levels of PSD-95. As shown in Figure 5A, the levels of PSD-95 were indeed elevated in Mdm2$^{f/-}$ cultures but not in the Mdm2$^{f/-}$ cultures (Fig. 4D). These data introduced a novel function of Mdm2 in neuroplasticity and demonstrated the necessity of Mdm2 in Gp1 mGluR-mediated elevation of neural network activity.

Knocking down Mdm2 does not affect basal synaptic transmission or Gp1 mGluR signaling

Despite the connection between FMRP and Mdm2 that was identified in the previous studies (27,28), it remains unknown whether and how Mdm2 participates in Gp1 mGluR signaling. To answer this question, we first determined whether knocking down Mdm2 affects basal synaptic activity. Because Mdm2 mediates the ubiquitination of post-synaptic density protein 95 (PSD-95) (28,50), we tested whether knocking down Mdm2 affects the levels of PSD-95. As shown in Figure 5A, the levels of PSD-95 were indeed elevated in Mdm2$^{f/-}$ cultures but not in the Mdm2$^{f/-}$ cultures (Fig. 4D). These data introduced a novel function of Mdm2 in neuroplasticity and demonstrated the necessity of Mdm2 in Gp1 mGluR-mediated elevation of neural network activity.

Mdm2 mediates Gp1 mGluR-induced protein translation

Because Mdm2 directly interacts with multiple ribosomal proteins (29–31), we next asked whether Mdm2 functions to mediate Gp1 mGluR-induced protein translation. We first determined whether Mdm2 associates with intact ribosomes. Using sucrose density ultracentrifugation to isolate ribosomes from the brains of 3-week-old mice, we found that Mdm2 is highly enriched in the ribosome pellet (Fig. 6A), suggesting Mdm2 associates with ribosomes. Interestingly, the distribution of Mdm2 in ribosomes did not differ between WT and Fmr1 KO brains, suggesting the association between Mdm2 and ribosomes is independent of FMRP. To determine whether Mdm2 associates more with monosomes or polysomes, we utilized polysome profiling by sucrose gradient sedimentation using mouse brain homogenates. As shown in Figure 6B and C, Mdm2 is detected in the lighter fractions, likely with monosomes or individual ribosomal subunits, as compared to PABPC1 which is heavily associated with polyribosomes. This phenomenon is consistent between WT and Fmr1 KO brains. Furthermore, EDTA, a ribosome-disrupting agent, successfully collapsed the polysome gradient and shifted both Mdm2 and PABPC1 to lighter fractions. These data indicate that Mdm2 might be involved in Gp1 mGluR-mediated protein translation by interacting with ribosomes and ribosomal components.

We next asked whether and how Mdm2 affects protein translation. We employed rabbit reticulocyte lysate system to perform in vitro translation with total RNA extracted from WT cortical neuron cultures. As shown in Figure 7A, the global translation efficiency in the presence of recombinant Mdm2 (lane 2), as measured by puromycin incorporation, was significantly lower than in reaction mixtures without Mdm2 (lane 1) or with control Glutathione S-Transferase (GST) protein (lane 3). These results suggest that Mdm2 functions as a translational
suppressor. To determine whether such suppressive activity requires Mdm2’s association with ribosomal proteins, we utilized two mutant Mdm2s, D294A-Mdm2 and T306A-S307A-Mdm2, which exert reduced affinity to interact with ribosomal proteins (53). When we expressed WT or either of the mutant Mdm2s in HEK 293 cells and used puromycin to label newly synthesized proteins, we found that while WT-Mdm2 reduces global translation, both of the mutants fail to exhibit significant activity toward translational suppression (Fig. 7B). These data suggest that Mdm2 represses translation likely through its association with ribosomal proteins.

To further validate the role of Mdm2 in Gp1 mGluR-mediated protein translation, we employed primary cortical neuron cultures prepared from Mdm2f/+–Emx1-Cre- or Mdm2f/+–Emx1-Cre+ mice. Using puromycin to label newly synthesized proteins in cultures, we found that the Mdm2f/+–Emx1-Cre+ cultures showed basally elevated protein translation in comparison to the Mdm2f/+–Emx1-Cre- cultures (Fig. 7C, comparing lanes 1 and 3). When we applied DHPG to activate Gp1 mGluR signaling and translation, we found that elevated protein translation is occluded in Mdm2f/+–Emx1-Cre+ cultures (Fig. 7C, comparing lanes 3 and 4). Because Gp1 mGluR-induced translation often occurs to FMRP target mRNAs (54), we tested the expression of one of the known FMRP targets, protocadherin-7 (55), upon DHPG treatment for 30 min. As shown in Figure 7D, the levels of Pcdh7 protein were moderately elevated, and further elevation upon DHPG treatment was occluded in Mdm2f/+–Emx1-Cre+ cultures. Altogether, our data suggest that Mdm2 functions as a novel translational suppressor.
suppressor and is required for Gp1 mGluR-induced translation activation.

FMRP mediates Gp1 mGluR-induced Mdm2 ubiquitination and down-regulation

Because Mdm2 inhibits protein translation and is required for Gp1 mGluR-induced translation activation (Fig. 7), we hypothesized that activating Gp1 mGluR will de-repress Mdm2’s function so as to permit protein translation. To test this idea, we evaluated the protein level of Mdm2 before and after DHPG treatment. As shown in Figure 8A, DHPG treatment significantly reduced Mdm2 levels. Because this reduction was observed within 30 min, we hypothesized that the process involves protein degradation. To this end, we measured Mdm2 ubiquitination by performing immunoprecipitation of Mdm2 followed by western blotting with the anti-ubiquitin antibody. We found that DHPG treatment elevates Mdm2 ubiquitination (Fig. 8B), which is consistent with the reduction of total Mdm2 levels (Fig. 8A). Furthermore, the downregulation of Mdm2 upon DHPG treatment was inhibited by a proteasome inhibitor MG132 (10 μM) (Fig. 8C), conforming that the downregulation is resulted from proteasome-mediated degradation. These data suggest that activating Gp1 mGluR-induced protein translation likely occurs in part through de-repression of Mdm2’s function in translational suppression.

Because Gp1 mGluR-induced protein translation (37–39) and neural network activity (Fig. 3) requires FMRP, we next asked whether Gp1 mGluR-induced Mdm2 ubiquitination and down-regulation requires FMRP. As shown in Figure 8D and E, Fmr1 KO cortical neuron cultures exhibited neither ubiquitination nor down-regulation of Mdm2 upon DHPG treatment. In fact, Mdm2 ubiquitination was significantly reduced (Fig. 8E). These results indicate that FMRP is required for Gp1 mGluR-induced Mdm2 ubiquitination and down-regulation. They also suggested a novel signaling process by which FMRP and Mdm2 mediate protein translation and neural network activity upon Gp1 mGluR activation (Fig. 9).
Discussion

A novel mechanism by which FMRP regulates translation

FMRP is an RNA-binding protein with nearly 1,000 target genes identified through high throughput screening (55). It is well accepted that FMRP usually acts as a translational suppressor by hindering ribosomes from moving along its target mRNAs (55). However, because of the numerous targets regulated by FMRP, the mechanisms by which FMRP affects protein translation are complex and involve many indirect signaling pathways, leading to a global effect on translation. Our current study suggests a novel signaling pathway by which FMRP mediates protein ubiquitination and down-regulation to affect protein translation.
through de-repression of Mdm2 (Fig. 9). It is currently unknown what molecular pathway mediates Mdm2 degradation upon Gp1 mGluR activation and how FMRP participates in this mechanism. Our previous study showed that FMRP functions to limit phosphorylation of Mdm2. It is therefore possible that the phosphorylation status of Mdm2 is crucial to permitting its degradation upon Gp1 mGluR activation. Future work is necessary to test this possibility.

**Figure 6.** Mdm2 associates with ribosomes. (A) Representative western blots of Mdm2, poly(A) binding protein cytoplasmic 1 (PABPC1) and Gapdh after sucrose density ultracentrifugation of WT or Fmr1 KO whole brains. PABPC1 and Gapdh serve as positive and negative controls, respectively, for ribosomes. Quantification is shown on the right (n = 4). (B, C) Representative western blots of Mdm2 and PABPC1, and polysome analysis from WT (B) or Fmr1 KO (C) brains. EDTA was used as a control to validate the fractionation by disrupting ribosomes.
Although both Mdm2 and FMRP associate with ribosomal components, due to the lack of known RNA-binding motifs, Mdm2 likely provides a more general translation suppressive activity than FMRP. Because our data showed that Mdm2 is involved in FMRP-associated translation potentially through its association with ribosomal proteins, and also because FMRP regulates the expression of Mdm2 through direct binding to Mdm2 mRNA (27), it is possible that FMRP might exert other functions to mediate global translation. Further dissecting these functions would offer new molecular targets for designing new therapies to ameliorate translation-associated neuronal deficits in FXS.

Gp1 mGluR-mediated neural network activity

Activating Gp1 mGluR has multiple effects on neuroplasticity. We focus on neural network activity because it is relevant to brain circuit excitability, and hyperexcitability is seen in multiple mGluR-associated neurological disorders including epilepsy, FXS, and autism spectrum disorders (56–58). Using a MEA system, our study found that activation of Gp1 mGluR elevates spontaneous spike frequency and burst activity, two indicators of hyperexcitability. Although Gp1 mGluR signaling was basally activated in FXS, and elevated burst activity was observed in Fmr1 KO mice (35), we were unable to appropriately compare burst activity basally between WT and Fmr1 KO cultures in MEA recordings due to high variability between cultures and the necessity to compare activity internally within the same culture. Instead, we focused on the neural network and burst activity upon Gp1 mGluR activation, which was clearly dysregulated in Fmr1 KO cortical neuron cultures. Importantly, we revealed that this phenomenon is mediated by Mdm2. A previous study demonstrated that Gp1 mGluR activation elevates neuronal excitability through downregulation of dendritic h-channels (59), while another study showed that such elevation of neuronal excitability depends on protein translation (13). Because we found that Gp1 mGluR activation leads to the elevation of translation by down-regulating Mdm2 in this study, and because FMRP also undergoes degradation upon Gp1 mGluR activation (60), it would be important to determine whether Mdm2, FMRP and h-channels undergo the same degradation pathway, and whether downregulation of h-channels contributes to the elevation of network activity that we observed. Furthermore, our previous studies showed that chronic neuronal activity stimulation triggers Mdm2-mediated degradation of p53, and this degradation homeostatically down-regulates the synchronization of spontaneous network activity without affecting spontaneous spike frequency or burst activity (35). This study, together with our current study, suggests that Mdm2 is a multifaceted regulator that mediates different properties of neural network activity, depending on the stimulation paradigms. It again shows the complexity of mGluR signaling and emphasizes the necessity of elucidating downstream mechanisms to identify better and more specific therapeutic targets for mGluR-associated diseases.

Figure 7. Mdm2 mediates Gp1 mGluR-induced protein translation. (A) Representative western blots of puromycin after in vitro translation using rabbit reticulocyte lysate and purified total RNA from WT cortical neuron cultures. Quantification (n = 5) and the purity of recombinant Mdm2 and GST were shown on the right. (B) Representative western blots of puromycin, Mdm2 and Gapdh after 30-min labeling of puromycin in HEK 293 cells transfected with HA-WT-Mdm2, HA-D294A-Mdm2 or HA-T306A-S307A-Mdm2. Quantification is shown on the right (n = 12). (C) Representative western blots of puromycin and Gapdh, and quantification after 30-min labeling of puromycin in Mdm2<sup>fl/fl</sup>-Emx-Cre<sup>+</sup> or Mdm2<sup>fl/fl</sup>-Emx-Cre<sup>-</sup> cortical neuron cultures at DIV 14 (n = 6). (D) Representative western blots of Pcdh7 and Gapdh, and quantification after vehicle or DHPG treatment for 30 min in Mdm2<sup>fl/fl</sup>-Emx-Cre<sup>+</sup> or Mdm2<sup>fl/fl</sup>-Emx-Cre<sup>-</sup> cortical neuron cultures at DIV 14 (n = 12). For the quantification above, Student t-test (A, B) or a two-way ANOVA with Tukey test (C, D) was used. Data are represented as mean ± SEM with *P < 0.05, ns: non-significant.
Other mGluR-dependent neuronal plasticity mechanisms

Another neuroplasticity mechanism that is well-known to be mediated by mGluR is long-term synaptic depression (LTD) (61). LTD describes a long-lasting attenuation of excitatory synaptic strength (62). LTD is critical to properly establishing activity- or experience-dependent synaptic connections during circuit development (63) and to learning and memory consolidation in the mature brain (64). Inducing mGluR-LTD, as with mGluR-regulated neural network activity (Figs 1 and 2), requires de novo protein translation. Because Mdm2 mediates Gp1 mGluR-mediated translation, it is anticipated that Mdm2 also participates in mGluR-LTD and will be an important future direction. mGluR-LTD is often associated with local protein translation. Local protein translation is thought to efficiently provide specific proteins, such as activity-regulated cytoskeleton-associated protein (Arc; also known as Arg3.1) (54,65), at close proximity to the synapses, thereby allowing for rapid weakening of the synapses. Mdm2 can be found in dendrites and postsynaptic terminals (28) and therefore can potentially contribute to local protein translation. A broader effect of Mdm2 is waiting to be discovered.

Figure 8. FMRP mediates Gp1 mGluR-induced ubiquitination and down-regulation of Mdm2. (A) Representative western blots of Mdm2 and Gapdh from WT cortical neuron cultures after vehicle or DHPG treatment for 30 min (n = 4). (B) Representative western blots of Ubiquitin and Mdm2 from WT cortical neuron cultures after immunoprecipitation with anti-Mdm2 antibody (n = 4). (C) Representative western blots of Mdm2 and Gapdh from WT cortical neuron cultures after vehicle, MG132, DHPG or DHPG + MG132 treatment for 30 min. The cells in all four conditions were pre-treated with translation inhibitor cycloheximide (60 μM, 10 mins) to avoid potential interference from translation after Gp1 mGluR activation (n = 8). (D) Representative western blots of Mdm2 and Gapdh from Fmr1 KO cortical neuron cultures after vehicle or DHPG treatment for 30 min (n = 4). (E) Representative western blots of Ubiquitin and Mdm2 from Fmr1 KO cortical neuron cultures after immunoprecipitation with anti-Mdm2 antibody (n = 4). For the quantification above, a one-sample t-test was used (A, B and D, E) to compare DHPG-treated groups to vehicle-treated groups while a two-way ANOVA with Tukey test was used for (C). Data are represented as mean ± SEM with *P < 0.05, ns: non-significant.
Mdm2 in FXS and in cancer biology

Mdm2 is a ubiquitin E3 ligase that has been well-characterized in the field of cancer biology, and the tumor suppressor p53 is one of its major substrates (66). Mdm2-mediated p53 ubiquitination leads to inactivation of p53-dependent transcriptional regulation and ultimately alters the cell cycle (67–69). Unbiased screening of direct p53 target mRNAs has revealed many brain-enriched genes (70). However, the function and regulation of Mdm2-p53 signaling in neuronal cells is largely unclear. Recently, several studies have begun to uncover the functions of Mdm2-p53 signaling in neurodevelopment and neuroplasticity (27,36,71). However, it remains unknown whether or not p53 is also involved in Mdm2-mediated protein translation. If this is the case, it would open another entirely new avenue for studying mGluR-dependent plasticity. Given that a number of small molecules related to Mdm2-p53 signaling are currently available, and some of them are currently on clinical trials, exploring if these small molecules could affect Mdm2-Gp1 mGluR signaling could open the possibility of developing new potential therapies for treating neurological diseases associated with dysregulated Gp1 mGluR signaling.

Materials and Methods

All experimental protocols involving mice were performed in accordance with the guidelines and regulations set forth by the Institutional Animal Care and Use Committee at the University of Illinois at Urbana-Champaign.

Reagents

Cycloheximide (CHX) and picrotoxin was from Santa Cruz Biotechnology. Dimethyl sulfoxide (DMSO) was from Fisher Scientific. Tetrodotoxin was from Cayman Chemical. DHPG was from Abcam. MG132 was from Selleck Chemical. Puromycin was from MP Biomedicals. DMSO was used as vehicles in this study. The antibodies used in this study were purchased from Santa Cruz Biotechnology (anti-Mdm2), GenScript Corporation (anti-Gapdh), Millipore (anti-puromycin), Enzo Life Sciences (anti-Akt), Bioss (anti-Pcdh7), Abcam (anti-PABPC1) and Cell Signaling (anti-ubiquitin, anti-p-Akt, anti-ERK and anti-p-ERK). Recombinant Mdm2 protein was from Abcam and GST was from Sigma. HRP-conjugated secondary antibodies were from Abcam.

In vitro translation, transfection, immunoprecipitation and western blotting

In vitro translation was conducted using rabbit reticulocyte lysate (RRL, Promega). In brief, 15 μl of RRL was mixed with 100 ng of total RNA extracted from WT cortical neuron cultures at DIV 14. After initial reaction for 5 min, puromycin (10 μg/ml) was added for another 10 min to label actively translating proteins. At the end of the experiment, SDS buffer was added to stop the reaction followed by standard western blotting.

For western blotting after SDS-PAGE, the gel was transferred onto a polyvinylidene fluoride (PVDF) membrane. After blocking with 1% Bovine Serum Albumins in TBST buffer (20 mM Tris pH 7.4, 120 mM NaCl, 0.5% Nonidet P-40) for 3 times subjecting to western blotting.

For western blotting after SDS-PAGE, the gel was transferred onto a polyvinylidene fluoride (PVDF) membrane. After blocking with 1% Bovine Serum Albumins in TBST buffer (20 mM Tris pH 7.4, 150 mM NaCl, 0.1% Tween-20), the membrane was incubated with the a primary antibody for overnight at 4 °C, followed by a 30-min wash with TBS. The membrane was then incubated with an HRP-conjugated secondary antibody for 1 h, followed by another 30-min wash. The membrane was then developed with an ECL (enhanced chemiluminescence) solution as described (65).

MEA recording

All the multi-electrode array recordings were done using an Axion Muse 64-channel system in single well MEAs (M64-GL1-30Pt200, Axion Biosystems) inside a 5% CO2,3 7°C incubator. Field potentials (voltage) at each electrode relative to the ground electrode were recorded with a sampling rate of 25 kHz. After 30 min of recording the baseline (before), drug(s) indicated in each experiment was added, and the MEA dish was immediately put back into the recording device for another 30 min of recording (after). Due to changes in network activity caused by physical movement of the MEA, only the last 15 min of each recording were used in data analyses as performed previously (36). AxIS software (Axion Biosystems) was used for the extraction of spikes (i.e. action potentials) from the raw electrical signal obtained from the Axion Muse system. After filtering, a threshold of ±6 standard deviations was independently set for each channel; activity exceeding this threshold was counted as a spike. The settings for burst detection were a minimum of 5 spikes with a maximum inter-spike interval of 0.1 s. Synchrony index was computed through AxIS software, based on a previously published algorithm (73), by taking the cross-correlation between two spike trains, removing the portions of the cross-correlogram that are contributed by the auto-correlations of each spike train, and reducing the distribution to a single metric. A value of 0 corresponds to no synchrony and a value of 1 corresponds to perfect synchrony.
Patch-clamp electrophysiology

Whole-cell patch-clamp recordings were made at room temperature (23–25 °C) in a submersion chamber continuously perfused with artificial cerebrospinal fluid (aCSF) containing (in mM): 119 NaCl, 2.5 KCl, 4 CaCl2, 4 MgCl2, 1 Na2HPO4, 26 NaHCO3, and 11 D-Glucose, saturated with 95% O2/5% CO2 (pH 7.4, 310 mOsm). aCSF was supplemented with 1 µM TTX and 100 µM picrotoxin for mEPSC measurements to block action potential-dependent EPSCs and GABA<sub>A</sub> receptors, respectively. Whole cell recording pipettes (~4–6 MΩ) were filled with intracellular solution containing (in mM): 130 K-gluconate, 6 KCl, 3 NaCl, 10 HEPES, 0.2 EGTA, 4 Mg-ATP, 0.4 Na-GTP, 14 Tris-phosphocreatine, 2 QX-314 (pH 7.25, 285 mOsm). Neurons at DIV 11–14 were used for electrophysiological analyses. Membrane potential was clamped at ~60 mV. All recordings were performed with Clampex 10.6 and Multiclamp 700B amplifier interfaced with Digidata 1440A data acquisition system (Molecular Devices). Recordings were filtered at 1 kHz and digitized at 10 kHz. Miniature excitatory postsynaptic currents (mEPSCs) were analyzed with Mini Analysis Program (Synaptosoft) with a 5-pA threshold level.

Isolation of total ribosomes

Whole brains were extracted, immediately frozen in liquid nitrogen, and stored at −80 °C.Brains were pulverized under liquid nitrogen using a previously cooled, RNAse free mortar and pestle. The powder obtained was transferred to a 10-cm plate on dry ice while subsequent brains were pulverized. Afterward, 1 ml of lysis buffer (10 mM Tris-HCl at pH 8.0, 150 mM NaCl, 5 mM MgCl2, 1 mg/ml heparin, 1% Nonidet-P40, 5% deoxycholate, 40 mM dithiothreitol, 1 U/ml SUPERaseIn RNAse inhibitor [Thermo Fisher], and 150 µg/ml cycloheximide) was added to the tissue powder. Next, re-suspended powder was scraped from the plate and transferred to a microcentrifuge tube with pipetting 10 times to lyse the cells. The cell nuclei and cell debris were removed by centrifugation (2,000g, 10 min, at 4 °C). The supernatant was transferred to a fresh tube and then centrifuged again at (16,000g, 7.5 min, at 4 °C). 400 μl of supernatant was layered onto a 10 ml linear sucrose gradient (15–45% sucrose [w/v] made using a Biocorp Gradient Master) and centrifuged in a SW41T rotor (Beckman) for 125 min at 38,000 rpm and 4 °C, with the brake off. Polysome profiles were recorded using a UA-6 absorbance (ISCO) detector at 254 nm and fractions were collected along the gradient. Fractions were used for subsequent western blot analysis.

Statistical analysis

For multiple comparisons, a two-way ANOVA and Tukey post-hoc test were performed. Student t-test was used in paired condition (Figs 5B, 6A and 7A). One sample t-tests were used when experimental group is normalized to control groups (Fig. 8). In all figures, error bars represent SEM and *P < 0.05, **P < 0.01.

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Conflict of Interest statement. None declared.

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