Neuroscience

Systemic pharmacological suppression of neural activity reverses learning impairment in a mouse model of Fragile X syndrome

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Abstract

The enhancement of associative synaptic plasticity often results in impaired rather than enhanced learning. Previously, we proposed that such learning impairments may result from saturation of the plasticity mechanism making it unavailable to be recruited at the appropriate synapses to support learning (Nguyen-Vu et al., 2017). This hypothesis was based on experimental results from mice lacking two class I major histocompatibility molecules, MHCI H2-K^{b} and H2-D^{b} (MHCI $K^{b}D^{b-/-}$), which have enhanced associative long-term depression at the parallel fiber-Purkinje cell synapses in the cerebellum (PF-Purkinje cell LTD). Here we extend this work by testing predictions of the saturation hypothesis in a second mouse line with enhanced PF-Purkinje cell LTD, the *Fmr1* knockout mouse model of Fragile X syndrome (FXS). Mice lacking *Fmr1* gene expression in cerebellar Purkinje cells (L7-*Fmr1* KO) were selectively impaired on an oculomotor learning task in which PF-Purkinje cell LTD has been implicated, with no impairment on an LTD-independent oculomotor learning task. Consistent with the saturation hypothesis, behavioral pre-training designed to reverse LTD at the PF-Purkinje cell synapses eliminated the oculomotor learning deficit in the L7-*Fmr1* KO mice, as previously reported in MHCI $K^{b}D^{b-/-}$mice. In addition, diazepam treatment to suppress neural activity and thereby limit the induction of associative LTD during the pre-training period also eliminated the learning deficit in L7-*Fmr1* KO mice. These results support the hypothesis that the enhancement of synaptic plasticity can lead to its saturation *in vivo* and inability to support learning, providing a novel mechanistic perspective that could inform the development of new clinical approaches for autism and other disorders of the nervous system.

eLife assessment

This **valuable** manuscript follows up on previous findings from the same lab supporting the idea that deficits in learning due to enhanced synaptic plasticity are due to saturation effects. **Convincing** evidence is presented that behavioral learning deficits associated with enhanced synaptic plasticity in a transgenic mouse model can be rescued by manipulations designed to reverse the saturation of synaptic plasticity. In particular, the finding that a previously FDA-approved therapeutic can rescue learning could provide **important** new insights for biologists, psychologists, and others studying learning and neurodevelopment.

Introduction

Since its discovery, long term synaptic plasticity has been of great interest to neuroscientists as a therapeutic target for brain disorders, especially disorders affecting learning and memory. Scientific and technological advances have provided an array of tools for enhancing synaptic plasticity. In some cases, experimental manipulations that augment plasticity have succeeded in augmenting learning (Tang et al., 1999; Van Praag et al., 1999; Lee and Silva, 2009). However, in many cases, manipulations that augment plasticity have impaired learning (Migaud et al., 1998; Uetani et al., 2000; Gu et al., 2002; Cox et al., 2003; Hayashi et al., 2004; Rutten et al., 2011; Navakkode et al., 2022). Surprisingly, there have been few attempts to reconcile these conflicting findings with a mechanistic explanation for why enhancing synaptic plasticity can have opposite effects on learning. Such mechanistic insight about how enhanced synaptic plasticity functions *in vivo* could facilitate the development of this approach as a viable clinical intervention for learning disorders, recovery from stroke or brain injury, dementia, and other neurological and psychiatric disorders.

Recently, we proposed a testable hypothesis about what can go wrong with augmented plasticity *in vivo*, based on experimental and theoretical analysis of learning in mice with enhanced associative synaptic plasticity in the cerebellum (Nguyen-Vu et al., 2017). Associative LTD at the cerebellar PF-Purkinje cell synapses (PF-Purkinje cell LTD) has been implicated in certain cerebellum-dependent learning tasks and not others, based in part on the observation of selective learning impairments in mouse lines with impaired PF-Purkinje cell LTD (reviewed in Raymond and Medina, 2018; De Zeeuw et al., 2021). Initially, we expected that mice with enhanced PF-Purkinje cell LTD would exhibit the exact opposite behavioral phenotype as mice with impaired PF-Purkinje LTD, i.e., enhancement of learning on the same tasks in which mice with impaired PF-Purkinje cell LTD exhibit impaired learning. Contrary to this expectation, double knockout of the major histocompatibility class I molecules MHCI H2-K^{b} and H2-D^{b} (MHCI $K^{b}D^{b-/-}$), which enhances PF-Purkinje cell LTD (McConnell et al., 2009), results in the very same, specific oculomotor learning impairment as observed in mice with impaired PF-Purkinje cell LTD (Nguyen-Vu et al., 2017). To explain the puzzling observation that the enhancement of a plasticity mechanism could yield the same behavioral phenotype as its impairment, we hypothesized that enhanced LTD prevents learning by allowing spontaneous activity in the circuit to saturate this form of plasticity, making it unavailable at the specific synapses where it is needed to support learning. Two key predictions of the saturation hypothesis were confirmed experimentally by previous work: optogenetic stimulation of the circuit designed to saturate PF-Purkinje cell LTD before training recapitulated in WT mice the same, specific oculomotor learning deficit observed in the MHCI $K^{b}D^{b-/-}$ mice with enhanced LTD; and a behavioral manipulation designed to reverse PF-Purkinje cell LTD before oculomotor training reversed the learning deficit in MHCI $K^{b}D^{b-/-}$ mice (Nguyen-Vu et al., 2017).

Here we further tested the hypothesis that the enhancement of associative LTD at the parallel fiber-Purkinje cell synapses can result in the saturation of this plasticity mechanism before learning can take place and thereby impair learning. First, we replicated key behavioral findings in a different line of mice with enhanced LTD at the parallel fiber-Purkinje cell synapses. Purkinje cell-specific knock out of the Fragile X gene *Fmr1* enhances PF-Purkinje cell LTD (Koekkoek et al., 2005). We show that these L7-*Fmr1* KO mice are selectively impaired on an LTD-dependent oculomotor learning task, and that this learning deficit can be reversed with behavioral pre-training designed to reverse PF-Purkinje cell LTD, as previously reported in the MHCI $K^bD^{b-/-}$ mice with enhanced PF-Purkinje cell LTD. We then test a new prediction of the saturation hypothesis about a pharmacological treatment to reverse the learning deficit in mice with enhanced associative synaptic plasticity.

Results

Selective learning impairment in mice with enhanced associative long-term depression in the cerebellum

We assessed oculomotor learning in mice lacking expression of the fragile X gene *Fmr1* in cerebellar Purkinje cells, which have been shown to have enhanced PF-Purkinje cell LTD (Koekkoek et al., 2005). Purkinje cell-specific *Fmr1* knock out mice were generated by crossing conditional *Fmr1* knockout mice (Mientjes et al., 2006) with mice expressing Cre under the control of the L7/Pcp2 promoter (Zhang et al., 2004; see Methods). We tested the ability of these L7-*Fmr1* KO mice to adaptively modify their vestibulo-ocular reflex (VOR), and compared their performance on different VOR learning tasks that have previously been shown to have different sensitivity to perturbations of PF-Purkinje cell LTD.

The VOR stabilizes images on the retina by using the vestibular sensory input caused by a head movement to drive an oppositely directed eye movement response. Learning can adjust the amplitude of this oculomotor reflex to improve the stabilization of visual images on the retina for successful navigation in the world (Gonshor and Melvill Jones, 1973; Ito et al., 1974; Miles and Fuller, 1974; Gauthier and Robinson, 1975; Batini et al., 1979; Raymond, 1998; Broussard and Kassardjian, 2004; Gittis and du Lac, 2006; Cullen, 2023). Mice were trained to adaptively increase or decrease their VOR amplitude using two types of vestibular-visual stimulus pairings (**Fig. 1**; Boyden and Raymond 2003; Boyden et al., 2004). When a vestibular stimulus (1 Hz sinusoidal rotation about an earth-vertical axis with peak velocity of ±10°/s) was paired with oppositely directed motion of a large-field visual stimulus for 30 min (**Fig. 1A**, *left*; see Methods), this induced an adaptive learned increase in the eye movement responses of wild type (WT) mice to the vestibular stimulus alone (VOR-increase learning; **Fig. 1A**, *right, black*; p<0.001, 0 vs. 30 min, Tukey). When the vestibular stimulus was instead paired with motion of a visual stimulus in the same direction as the head (**Fig. 1B**, *left*), this induced an adaptive learned decrease in the eye movement responses of WT mice to the vestibular stimulus alone (VOR-decrease learning; **Fig. 1B**, *right, black;* p<0.001, 0 vs. 30 min, Tukey).

Both VOR-increase and VOR-decrease learning are cerebellum dependent (Ito et al., 1974; Robinson, 1976; Lisberger et al., 1984, 1994; Watanabe, 1984; Michnovicz and Bennett, 1987; Koekkoek et al., 1997; McElligott et al., 1998; Rambold et al., 2002). However, manipulations that impair or enhance PF-Purkinje cell LTD have previously been found to selectively alter VOR-increase learning, with less or no effect on VOR-decrease learning (Li et al., 1995; Boyden et al., 2006; Hansel et al., 2006; Guo et al., 2014; Kimpo et al., 2014; Nguyen-Vu et al., 2017; Zhang et al., 2023). Accordingly, the L7-*Fmr1* KO mice with enhanced PF-Purkinje cell LTD were selectively and profoundly impaired on VOR-increase learning. Unlike the WT control group, L7-*Fmr1* KO mice exhibited no significant change in the amplitude of their VOR after 30 min of VOR-increase training (**Fig. 1A**, *red*; p=0.97, L7-*Fmr1* KO, 0 vs 30 min;

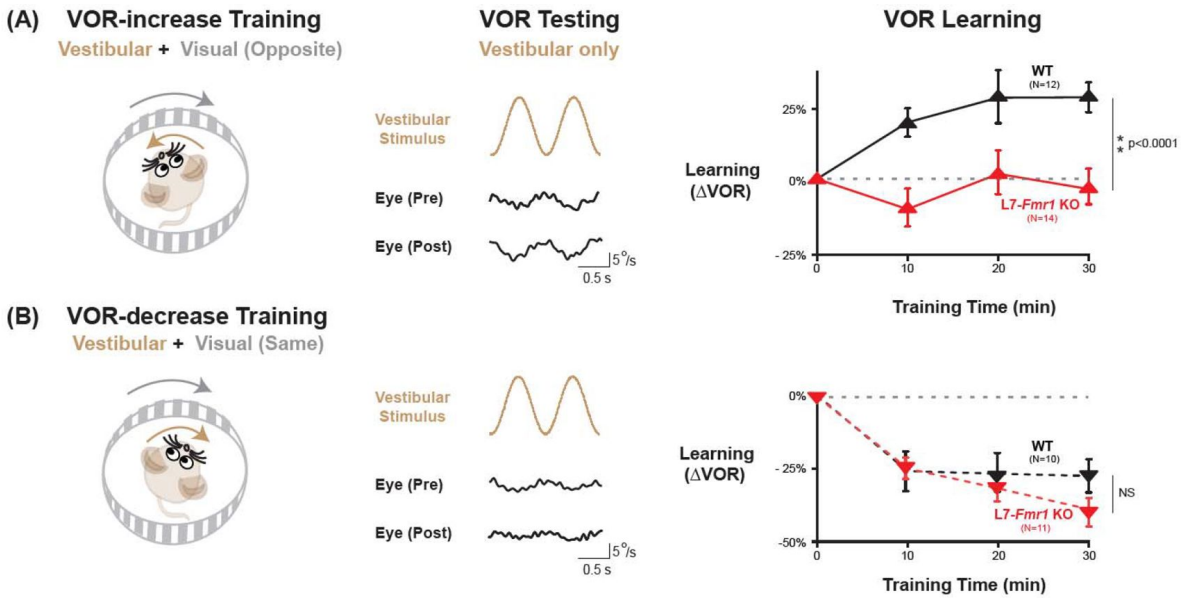

Figure 1.

VOR-increase learning is impaired in L7-*Fmr1* KO mice with enhanced cerebellar LTD.

(A) Training to increase the VOR. ***Left***, VOR-increase training paired a vestibular stimulus (1 Hz sinusoidal rotation about an earth-vertical axis, *brown*) with oppositely directed visual stimulus motion (*grey*). ***Middle***, Example raw eye velocity responses (*black*) to the vestibular stimulus alone in the dark, i.e., the VOR, measured Pre and Post VOR-increase training. ***Right***, Average learned change in the amplitude of the VOR relative to pre-training, measured in the dark (*upward triangles*) after each 10-min VOR-increase training block in the L7-*Fmr1* KO (*red*) and WT mice (*black*). **(B)** Training to decrease the VOR. ***Left***, VOR-decrease training paired a vestibular stimulus (1 Hz sinusoidal rotation) with visual stimulus motion in the same direction. ***Middle***, Example VOR responses in the dark, measured Pre and Post VOR-decrease training. ***Right***, VOR-decrease learning (*downward triangles*). NS= not significant. In this and all figures, values plotted are mean ± SEM.

p<0.001, L7-*Fmr1* KO vs. WT, 30 min; Tukey). In contrast, VOR-decrease learning in the L7-*Fmr1* KO mice was robust and indistinguishable from that of their WT littermates (**Fig. 1B**, *red*; p<0.001, L7-*Fmr1* KO, 0 vs 30 min; p= 0.09, L7-*Fmr1* KO vs. WT; Tukey). Baseline oculomotor performance of L7-*Fmr1* KO mice was normal, as were the eye movement responses to the paired presentation of visual and vestibular stimuli used for both types of training (**Fig. 1 - figure supplement 1**), suggesting that there was no deficit in the vestibular, visual or oculomotor functions required to perform the learning tasks; rather the L7-*Fmr1* KO mice have a selective deficit in learning. These results support previous findings that manipulations of PF-Purkinje cell LTD selectively affect VOR-increase learning, and that the enhancement of PF-Purkinje cell LTD impairs rather than enhances this form of learning.

Behavioral pre-training eliminates learning impairment in L7-*Fmr1* KO mice with enhanced LTD

A key question is why the enhancement of PF-Purkinje cell LTD would impair LTD-dependent learning. One potential explanation is that the enhancement of LTD allows the spontaneous activity in the cerebellar circuit to recruit and saturate this mechanism, making it unavailable to support new LTD-dependent learning. If this is the case, then manipulations that prevent or reverse excessive PF-Purkinje cell LTD before training should reset the circuit to a state compatible with new LTD-dependent learning, and thereby improve VOR-increase learning in the L7-*Fmr1* KO mice. To test this prediction, we first employed a behavioral approach designed to reverse PF-Purkinje cell LTD in the oculomotor cerebellum before training.

In wild-type mice, VOR-decrease training can rapidly reverse any behavioral evidence of prior VOR-increase learning, which suggests that VOR-decrease training can reverse any plasticity induced during VOR-increase learning, including any PF-Purkinje cell LTD (Boyden and Raymond, 2003). Accordingly, VOR-decrease pre-training was previously found to reverse the oculomotor learning deficit in *MHCI* $K^bD^{b-/-}$ mice with enhanced PF-Purkinje cell LTD (Nguyen-Vu et al., 2017). We tested whether the same behavioral pre-training intervention could also eliminate the learning deficient in L7-*Fmr1* KO mice.

L7-*Fmr1* KO and WT mice were subjected to 30 min of VOR-decrease pre-training followed by 30 min of VOR-increase training. In WT mice, there were adaptive changes in the amplitude of the VOR during both the pre-training and training periods—first a decrease and then an increase in the eye movement response to the vestibular stimulus alone (**Fig. 2B**, *black;* VOR-decrease, *dotted lines*, p<0.001, WT -30 vs 0 min; VOR-increase, *solid lines*, p=0.02, WT 0 vs 30 min; Tukey). The L7-*Fmr1* KO mice exhibited changes in the VOR during both the pre-training and training periods that were statistically indistinguishable from WT (**Fig. 2B**, *red;* VOR-decrease, *dotted lines*, p=0.18, L7-*Fmr1* KO vs. WT, 0 min, Tukey; **Fig. 2B**, *bar graphs*, p=0.17, L7-*Fmr1* KO vs. WT, VOR-increase from 0 to 30 min, t test; **Fig. 2—figure supplement 1**). Although in the absence of pre-training, VOR-increase training failed to induce any significant change in the VOR of the L7-*Fmr1* KO mice (**Fig. 2A**, *red;* p= 0.99, L7-*Fmr1* KO, 0 vs 30 min, Tukey), the same VOR-increase training procedure did induce a significant increase in VOR amplitude when delivered to the same cohort of mice after VOR-decrease pre-training (**Fig. 2B**, *red, solid lines and bar graph*, p<0.001, 0 vs 30 min, Tukey). In other words, the ability of the L7-*Fmr1* KO mice to learn in response to the VOR-increase training varied with the recent history of experience (**Fig. 2**, *compare red bars in **A** vs **B***; p< 0.05, VOR-increase learning of L7-*Fmr1* KO without vs. with pre-training, paired sample t-test). Pre-training experience did not have the same effect in WT mice. The amount of learning exhibited by WT mice in response to VOR-increase training was not enhanced after VOR-decrease pre-training (**Fig. 2**, *compare black bars in **A** vs **B***; p= 0.41, paired sample t-test). Thus, VOR-decrease pre-training had different effects on the L7-*Fmr1* KO and WT mice, putting the L7-*Fmr1* KO mice, but not the WT mice, into a state more compatible with VOR-increase learning.

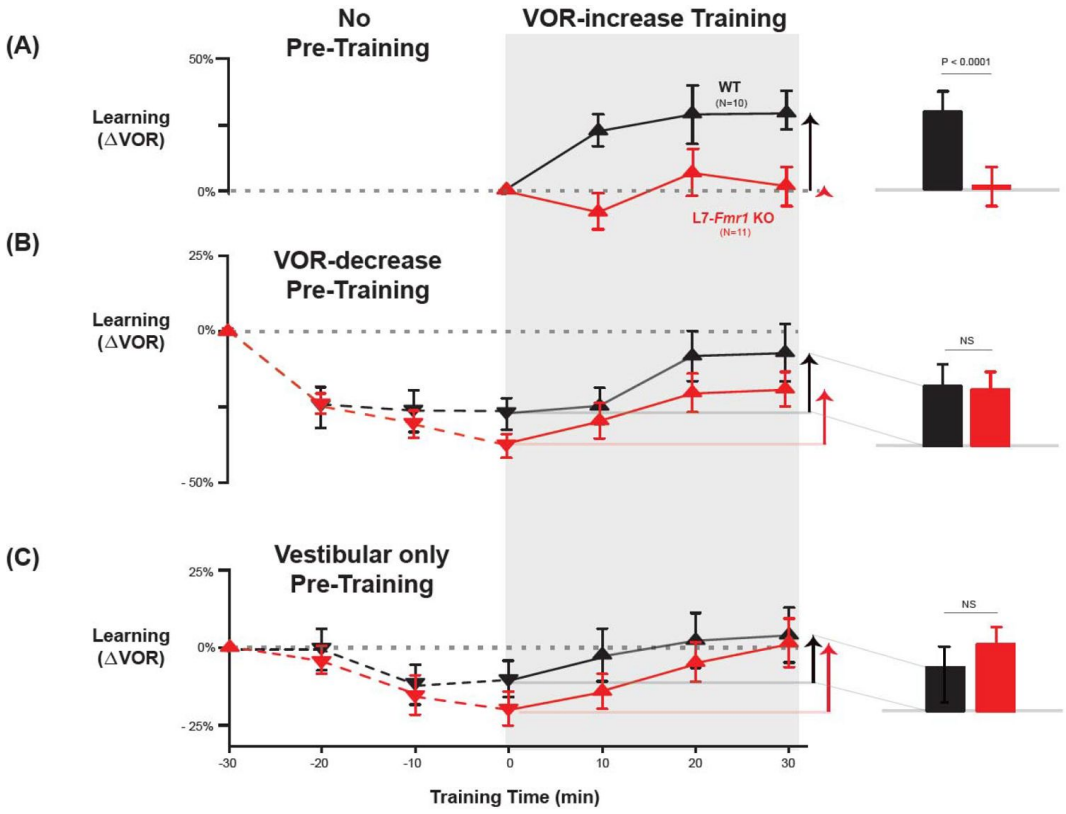

Figure 2.

Behavioral pre-training rescued learning impairment of L7-*Fmr1* KO mice with enhanced associative LTD.

Associative VOR-increase learning (*shaded area* and *bar graphs*), without pre-training (***A***), after VOR-decrease pre-training (***B***), and after Vestibular only pre-training (***C***). ***A***, learned change in the VOR response measured in the dark after each 10-min block of VOR-increase training in the subset of L7-*Fmr1* KO (*red*) and WT (*black*) mice from **Figure 1A** that were also tested after pre-training. ***B***, Changes in the VOR measured in the dark after each block of VOR-decrease pre-training *(downward triangles, dashed lines)* and then subsequent VOR-increase training *(upward triangles, solid lines)*. ***C***, Changes in the VOR measured in the dark after each block of Vestibular only pre-training *(downward triangles, dashed lines)* and then VOR-increase training *(upward triangles, solid lines)*. ***Right***, *Arrows* and *bars graphs* show the total change in the VOR induced by 30 min of VOR-increase training (training time = 30) compared with just before VOR-increase training (training time = 0).

A second behavioral pre-training procedure, habituation of the VOR, induced by presentation of the vestibular stimulus alone in complete darkness (Vestibular only pre-training), had effects similar to those of VOR-decrease pre-training on subsequent VOR-increase learning. After thirty minutes of Vestibular only pre-training, subsequent VOR-increase learning in the L7-*Fmr1* KO mice was comparable to that of their WT littermates (**Fig. 2C**, *red vs black bars;* p=0.84, L7-*Fmr1* KO vs. WT, VOR-increase from 0 to 30 min, paired sample t-test).

Pharmacological suppression of neural activity the day before training eliminates learning impairment of L7-*Fmr1* KO mice with enhanced LTD

The preceding results are consistent with the hypothesis (Nguyen-Vu et al., 2017) that spontaneous activity in the circuit can induce and saturate PF-Purkinje cell LTD in mice with enhanced LTD, and that behavioral pre-training can alter neural activity in a manner that prevents or reverses this saturation, thereby reversing the learning impairment in mice with enhanced LTD. Since PF-Purkinje cell LTD is driven by co-activation of cerebellar parallel fibers and climbing fibers (Ito and Kano, 1982; Ito et al., 1982; Linden and Connor, 1995), pharmacological suppression of neural activity should also prevent the induction and saturation of LTD during the pre-training period, and restore the capacity for subsequent LTD-dependent learning in mice with enhanced LTD. We tested this prediction by administering the benzodiazepine diazepam, a positive allosteric modulator of $GABA_A$ receptors, to enhance inhibition and suppress neural activity in the L7-*Fmr1* KO mice during the period preceding VOR-increase training. Diazepam has been shown to reduce neural firing in cerebellar neurons and neural responses to vestibular stimuli (Ryu and McCabe, 1974; Barmack, N. H. & Pettorossi, 1980). We assessed VOR learning 2 hours after diazepam administration, immediately after recovery from diazepam (18-24 hours after), and 1 week later.

The acute effect of diazepam administration was to impair learning. There was no effect of diazepam on the baseline amplitude of the VOR response measured in the dark 2 hours after diazepam (**Fig. 3 – figure supplement 1**), contrary to what has been reported in rabbit (Barmack, N. H. & Pettorossi, 1980). However, when VOR-increase training was delivered 2 hours after systemic administration of diazepam, VOR-increase learning was profoundly impaired in WT as well as L7-*Fmr1* KO mice (**Fig. 3 – figure supplement 2**).

It is not surprising that the acute effect of suppressing neural activity was to impair learning. The key question was whether this suppression of activity could reset the circuit to a state compatible with subsequent LTD-dependent learning. Therefore, VOR learning was tested after recovery from the acute effects of diazepam. Diazepam has a long half-life of ~24 hours (Riss et al., 2008), therefore mice were allowed to recover in their home cage for 18-24 hours after diazepam administration, and then VOR learning was tested after recovery from this prolonged period of pharmacological suppression of neural activity **(Fig. 3A)**. Remarkably, the L7-*Fmr1* KO mice exhibited robust VOR-increase learning, comparable to their WT littermates (**Fig. 3B**, *top, red vs black*; p=0.86, Tukey). Although the same individual L7-*Fmr1* KO had exhibited no significant learning in response to VOR-increase training in the absence of the pharmacological pre-treatment (**Fig. 2A**, *red*; p= 0.99, 0 vs 30 min, Tukey), diazepam pre-treatment eliminated this learning deficit.

The enhancement of learning by diazepam pre-treatment was temporary. When the same mice were re-tested one week after diazepam administration, the L7-*Fmr1* KOs again failed to learn in response to VOR-increase training (**Fig. 3C**, *red;* p=0.12, 0 vs 30 min, Tukey). Thus, diazepam pre-treatment could restore the VOR circuit of L7-*Fmr1* KO mice to a state compatible with VOR-increase learning, but this effect was transient.

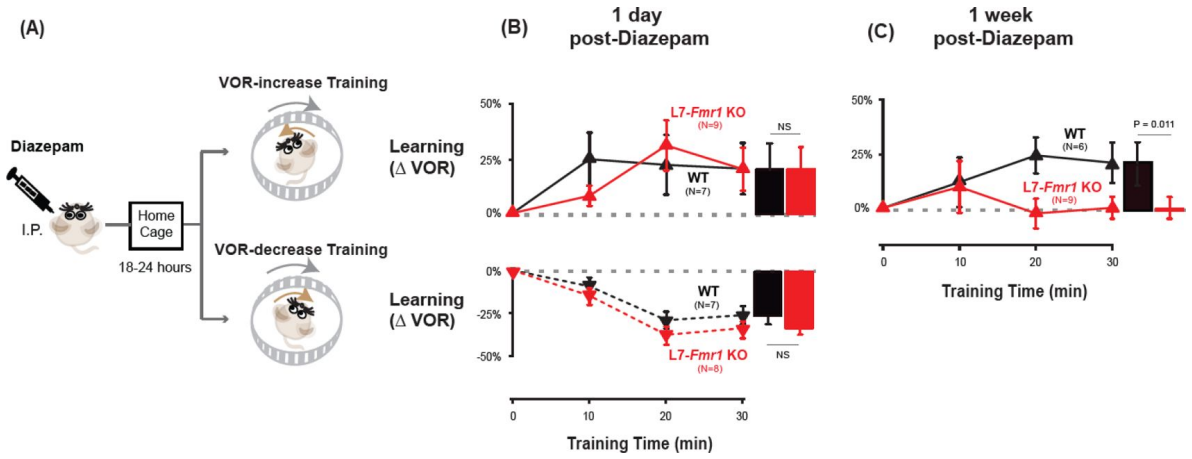

Figure 3.

Diazepam pre-treatment rescued learning impairment of L7-*Fmr1* KO mice with enhanced associative LTD.

(A) Mice were given an IP injection of diazepam (0.5 mg/kg) and then returned to the home cage for 18-24 hours, followed by VOR-increase (***top***) or VOR-decrease (***bottom***) training. **(B)** ***Top***, VOR-increase learning 1 day (18-24 hours) after diazepam administration in L7-*Fmr1* KO (*red upward triangles*) and WT mice (*black upward triangles)*. ***Bottom***, VOR-decrease learning (*downward triangles)* 1 day after diazepam. **(C)** VOR-increase learning in the same mice as in **B**, 1 week after diazepam treatment, and 18-24 hours after IP saline injection.

Specificity of pre-training effects on learning

The ability of behavioral and pharmacological pre-training interventions to enhance learning was specific to mice with enhanced PF-Purkinje cell LTD and to the type of VOR learning task. Wild type mice did not exhibit enhanced VOR-increase learning after diazepam pre-treatment (**Fig. 3B vs 3C**, *top, black*; p= 0.55, paired sample t test). Moreover, there was no effect of diazepam pre-treatment on VOR-decrease learning in either the WT or L7-*Fmr1* KO mice (*compare* **Fig. 3B**, *bottom vs* **1B**; p=0.11, L7-*Fmr1* KO vs. WT 1-day post-diazepam, VOR-decrease at 30 min, Tukey; p= 0.91, WT, diazepam vs control, VOR-decrease at 30 min paired sample t-test; p= 0.37, L7-*Fmr1* KO, diazepam vs control, VOR-decrease at 30 min, paired sample t-test). Thus, both the learning impairment in the L7-*Fmr1* KO mice and the effects of diazepam pre-treatment were selective for VOR-increase learning, consistent with previous evidence that this form of VOR learning is more dependent on PF-Purkinje cell LTD than VOR-decrease learning (Boyden et al., 2006; Nguyen-Vu et al., 2017). Previous work has also suggested a selective contribution of PF-Purkinje cell LTD to VOR learning induced with high-frequency (≥ 1 Hz) vestibular and visual stimuli, with less contribution of LTD when VOR learning is induced with low-frequency (≤0.66 Hz) vestibular and visual stimuli (Boyden et al., 2006; Nguyen Vu et al., 217). We found a trend for low-frequency (0.5 Hz) as well as high-frequency VOR-increase learning to be impaired in the L7-*Fmr1* KO mice (**Fig. 4A**, *left, red vs black;* p= 0.06, L7-*Fmr1* KO vs. WT, 30 min, Tukey). However, the low-frequency learning impairment was not reversed by the pre-training procedures that reversed the high-frequency learning impairment. Neither behavioral pre-training (**Fig. 4A**, *middle*, p= 0.47, VOR-decrease Pre-training vs. no Pre-training; **Fig. 4A**, *right*, p= 0.35, Vestibular only Pre-training vs. no Pre-training; paired sample t test), nor treatment with diazepam 18-24 hours before training (**Fig. 4B**, *compare red bars;* p= 0.66, saline vs. diazepam. paired sample t test), reversed the impairment of low-frequency VOR-increase learning in the L7-*Fmr1* KO mice, in contrast to their effectiveness at reversing the impairment of high-frequency VOR-increase learning (**Figs. 2**,**3**). This is consistent with the hypothesis that the behavioral and pharmacological pre-training manipulations selectively restore the capacity for learning tasks that depend on PF-Purkinje cell LTD.

Discussion

The results support the hypothesis proposed by Nguyen-Vu et al. (2017) that the enhancement of associative synaptic plasticity can impair learning by enabling ongoing neural activity in the circuit to saturate the plasticity mechanism. Consistent with this saturation hypothesis, L7-*Fmr1* KO mice with enhanced PF-Purkinje cell LTD exhibit impaired rather than enhanced learning on an oculomotor learning task in which PF-Purkinje cell LTD has been implicated, as previously shown for MHCI $K^bD^{b-/-}$ mice. Moreover, in both mouse lines, behavioral manipulations designed to prevent or reverse the saturation of PF-Purkinje cell LTD before training reversed the learning impairment, demonstrating that mice with enhanced LTD retain the capacity for robust LTD-dependent learning, although this capacity is influenced by the recent history of activity in the circuit. The new finding that pharmacological suppression of neural activity with diazepam can enhance subsequent learning in mice with enhanced associative plasticity provides new evidence for the saturation hypothesis and also could guide the development of novel clinical approaches.

The very similar behavioral phenotypes observed when LTD is enhanced by manipulating different molecular cascades strengthens the evidence that their shared effect of enhancing LTD is responsible for the learning impairment, rather than some other, off-target effect of the molecular manipulations. Whereas MHCI H2-K^b and H2-D^b act on MAP kinase and integrin via interaction with immune receptors such as PirB (Shatz, 2019), *Fmr1* acts by inhibiting mGluR-dependent dendritic protein translation (Huber et al., 2002). Previously, when different behavioral tasks were used to assess cerebellum-dependent learning in these two lines of mice, different phenotypes were reported–enhanced rotorod learning in the MHCI $K^bD^{b-/-}$ mice (McConnell et al.,

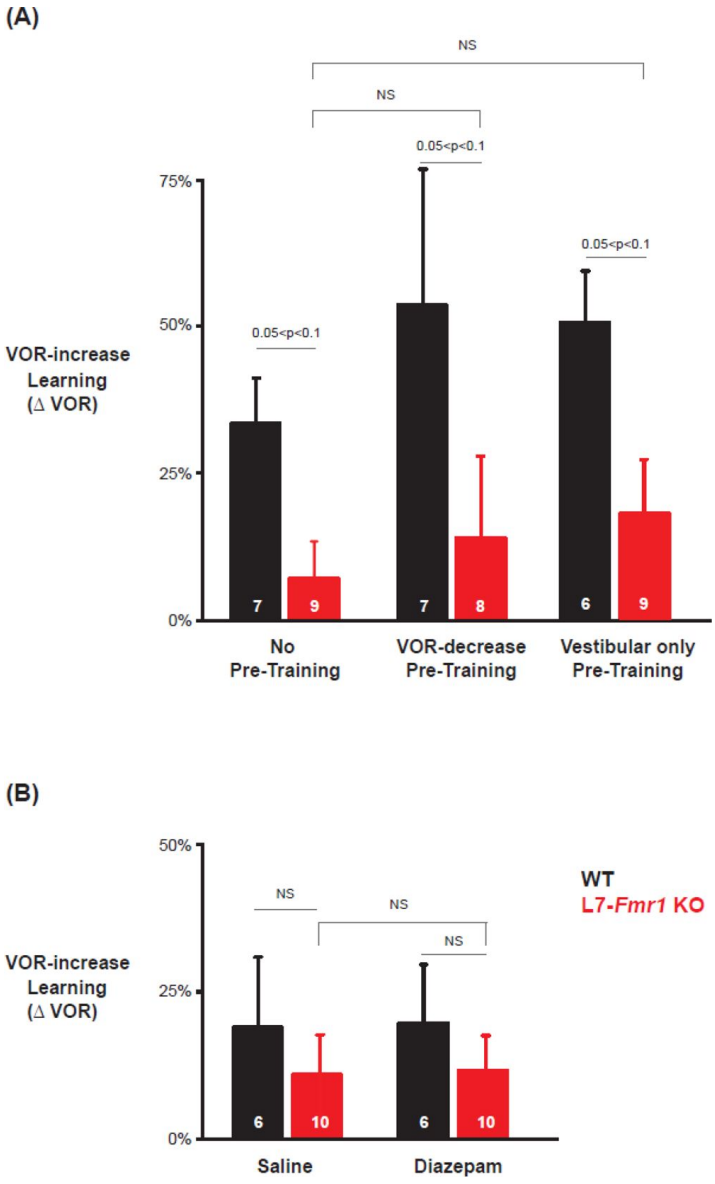

Figure 4.

Low frequency (0.5 Hz) VOR-increase learning impairment was not rescued by behavioral pre-training or diazepam pre-treatment.

(A) Low-frequency VOR-increase learning of L7-*Fmr1* KO mice (*red*) and WT mice (*black*), without pre-training (***left***), after 0.5 Hz VOR-decrease pre-training (***middle***), and after 0.5 Hz Vestibular only pre-training (***right***). **(B)** Low frequency (0.5 Hz) VOR-increase learning without diazepam pre-treatment (***left***) and 18-24 hours after IP injection of 0.5 mg/kg diazepam (***right***).

2009), but impaired eyeblink conditioning in the L7-*Fmr1* KO mice, global *Fmr1* KO mice and Fragile X patients (Koekkoek et al., 2005). This highlights the importance of the specific choice of behavioral task for assessing cerebellar learning, and the differential dependence of different cerebellar learning tasks on specific molecular and cellular processes within the cerebellum.

Although PF-Purkinje cell LTD was the main candidate mechanism of cerebellum-dependent learning for many decades, there is growing evidence that PF-Purkinje cell LTD contributes selectively to certain cerebellum-dependent learning tasks, and not others (Shibuki et al., 1996; Van Alphen and De Zeeuw, 2002; Feil et al., 2003; Boyden et al., 2004; Boyden et al., 2006; Kimpo et al., 2014; Nguyen-Vu et al., 2017; De Zeeuw et al., 2021; Zhang et al., 2023). Oculomotor learning is particularly advantageous for studying the role of PF-Purkinje cell LTD in learning because this plasticity mechanism is thought to contribute differentially to closely related oculomotor learning tasks that all depend on the same sensory and motor signaling pathways through the cerebellar flocculus, providing powerful control conditions for distinguishing observations related to PF-Purkinje cell LTD vs. other functional components within the same circuit. The VOR learning tasks examined all involve a change in the gain of the eye movement response to a vestibular stimulus that is induced by pairing vestibular and visual stimuli. Despite these commonalities, a number of experimental approaches, including *ex vivo* slice physiology (Yamaguchi et al., 2016; Shim et al., 2022), optogenetic stimulation (Nguyen-Vu et al., 2017; Rowan et al., 2018; Zhang et al., 2023) and studies of oculomotor learning in mice with impaired LTD (Boyden et al., 2006; Hansel et al., 2006; Kakegawa et al., 2018) have suggested a selective contribution of PF-Purkinje cell LTD to VOR-increase learning induced by high-frequency (≥ 1 Hz) vestibular and visual stimuli, with less or no contribution to VOR-decrease learning or VOR-increase learning induced with lower frequency vestibular and visual stimuli. Accordingly, both lines of mice with enhanced LTD (L7-*Fmr1* KO and MHCI $K^bD^{b-/-}$ mice) exhibited selective alteration of VOR-increase and not VOR-decrease learning. In MHCI $K^bD^{b-/-}$ mice the impairment of VOR-increase learning was also selective for high-frequency training (Nguyen-Vu et al., 2017). In the L7-*Fmr1* KO mice, there was a trend for low-frequency as well as high-frequency VOR-increase learning to be impaired. However, the behavioral and diazepam pre-treatments designed to reverse the saturation of LTD only improved the high-frequency and not the low-frequency VOR-increase learning, suggesting different mechanistic underpinnings of the low- and high-frequency impairments. In other words, the deletion of *Fmr1* from Purkinje cells may have two distinct effects: enhancement of PF-Purkinje cell LTD, which recapitulates the high-frequency VOR-increase learning phenotypes observed in the MHCI $K^bD^{b-/-}$ mice with enhanced LTD, plus disruption of an additional cellular mechanism that contributes to low-frequency VOR-increase learning. Overall, in the two lines of mice with enhanced PF-Purkinje cell LTD, both the learning impairments and the effects of manipulations designed to reverse or prevent the saturation of LTD before training were remarkably selective for the specific VOR learning task in which PF-Purkinje cell LTD has been most strongly implicated.

A question of central scientific and clinical importance is why the enhancement of synaptic plasticity would impair rather than enhance learning. One hypothesis is that a lower threshold for LTD induction might cause it to be over-recruited during training, at synapses that should not have undergone LTD in addition to synapses where LTD would support adaptive behavioral changes, thereby corrupting the learning process (Migaud et al., 1998; Koekkoek et al., 2005; McConnell et al., 2009; Lee et al., 2014). Our alternative, saturation hypothesis suggests that the enhancement of PF-Purkinje cell LTD allows the spontaneous activity in the cerebellar circuit to recruit and saturate this mechanism even before training begins, making it unavailable during training to support new LTD-dependent learning (**Fig. 5**). Similarly, the aberrant recruitment of LTD before training may lead, not to its saturation *per se*, but to some other kind of reduced availability, such as an increased threshold for its induction (Bienenstock, Cooper, and Munro, 1982; Leet, Bear, and Gaier, 2022). This saturation or increased threshold hypothesis differs from the over-recruitment hypothesis by suggesting that LTD is under-rather than over-recruited during the training period in mice with enhanced LTD, and that the problem with enhanced LTD

arises because of what it does to the circuit before training, rather than how it functions during training. Our finding that manipulations designed to prevent or reverse excessive PF-Purkinje cell LTD before training improve subsequent learning favor the saturation hypothesis.

Learning could be restored with two different approaches designed to prevent or reverse the saturation of LTD in the L7-*Fmr1* KO before training: behavioral pre-training (**Fig. 2**), and direct, pharmacological suppression of neural activity during the pre-training period (**Fig. 3**). In the absence of pre-training, high-frequency VOR-increase learning was not only impaired but completely absent in the L7-*Fmr1* KO mice (**Fig. 1A**, *right, red bar*). However, the same mice exhibited robust high-frequency VOR-increase learning, equal to that of WT, after behavioral pre-training, demonstrating that these mice can respond to the VOR-increase training with adaptive changes in the behavior (**Fig. 2**). At the end of those experiments, the amplitude of the VOR response (VOR gain = ratio of eye velocity to head velocity; see Methods) achieved in both L7-*Fmr1* KO and WT was roughly to the level of the VOR baseline value before pre-training. The additional experiments using pharmacological suppression of activity with diazepam during the pre-training period extends this work by providing the additional demonstration that the L7-*Fmr1* KO are capable of learning a VOR gain higher than baseline (**Fig. 3**). The ability of both behavioral and pharmacological pre-treatments to rescue learning shows that mice with enhanced LTD are not incapable of LTD-dependent learning. Rather, the enhancement of associative plasticity in the context of ongoing neural activity in the circuit appears to create a state in which LTD is saturated or otherwise unavailable to support learning. However, this saturated state can be reversed when the patterns of neural activity that create it are eliminated.

The striking specificity of the effects of the pre-training manipulations is consistent with the pre-training selectively reversing limitations caused by enhanced LTD, rather than generally enhancing cerebellum-dependent learning. Neither behavioral nor pharmacological pre-training enhanced learning in WT mice. Moreover, in the L7-*Fmr1* KO and MHCI $K^bD^{b-/-}$ mice, the pre-training treatments selectively enhanced high-frequency VOR-increase learning, with no significant effect on other forms of VOR learning (VOR-decrease learning or low-frequency VOR-increase learning), which previous research suggests are less dependent on PF-Purkinje cell LTD (Aiba et al., 1994; Li et al., 1995; Feil et al., 2003; Boyden et al., 2006; Hansel et al., 2006; Titley et al., 2010; Schonewille et al., 2011; Nguyen Vu et al., 2017; Zhang et al., 2023).

The present findings offer a scientific perspective that could guide the development of new clinical approaches for Fragile X syndrome and a range of other neurological and psychiatric conditions with enhanced associative plasticity. The oculomotor learning impairment in L7-*Fmr1* KO mice could be reversed with diazepam, which is an FDA-approved drug. The specificity of the effects of diazepam enhances its therapeutic potential. Pre-treatment with diazepam restored the capacity for high-frequency VOR-increase learning in the L7-*Fmr1* KO mice without compromising the normal performance of these mice on other oculomotor learning tasks. This was true even for VOR-decrease learning, which may depend on LTP of the same population of PF-Purkinje cell synapses that undergo LTD during VOR-increase learning (Shim et al., 2022). In other words, diazepam rescued the learning impairment with no apparent side effects on other, closely related functions of the same neural circuitry. At the same time, there is reason to expect that this approach of suppressing neural activity during a pre-training period may be generally applicable to any learning task, motor or cognitive, that is impaired by enhanced associative LTD (Rochefort et al., 2013; Badura et al., 2018; Ashburn et al., 2019; Frontera et al., 2020; Stoodley and Tsai, 2021; Hwang et al., 2022). Pharmacological suppression of neural activity should suppress PF-Purkinje cell LTD throughout the cerebellum, and hence may have the general effect of restoring all regions of the cerebellum to a state compatible with new LTD-dependent learning. This generality of the pharmacological approach stands in contrast to the behavioral pre-training approach, which would require extensive additional knowledge and experimentation to design the appropriate behavioral pre-treatment to reset each functional regional of the cerebellum to a state compatible with LTD-dependent learning, and different behavioral pre-treatments would be

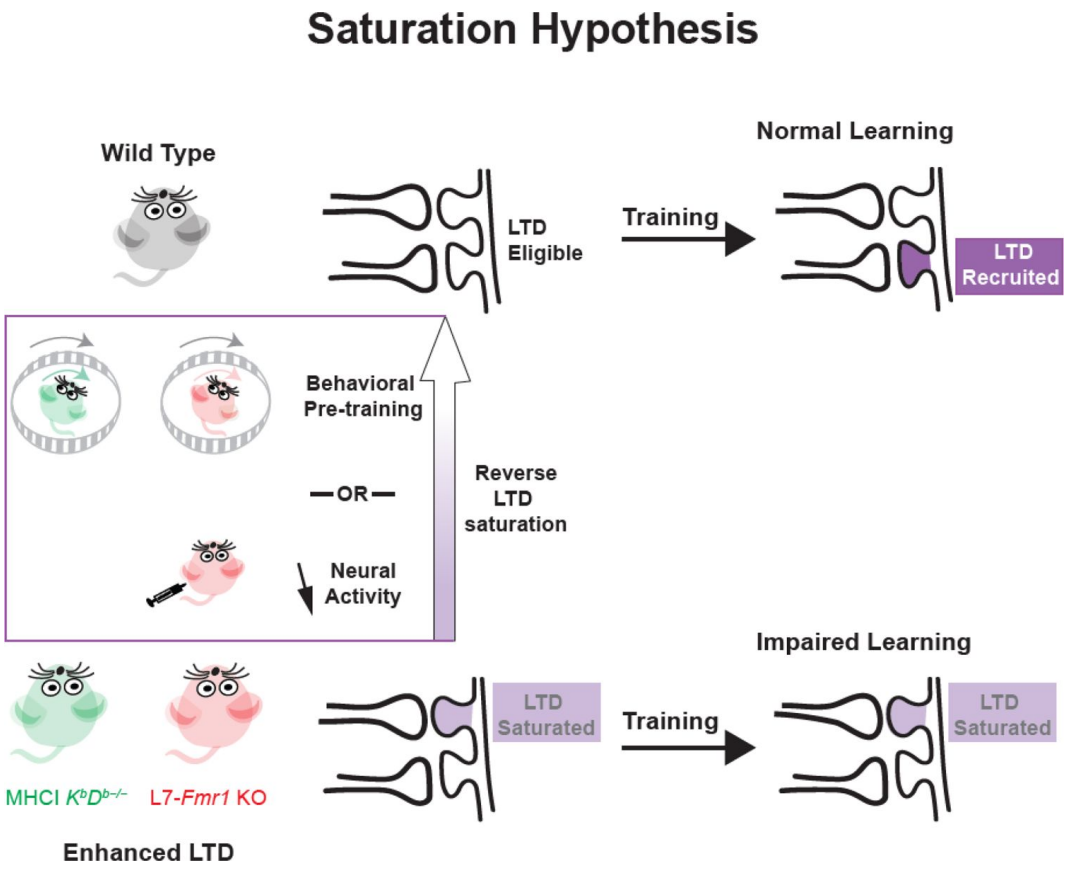

Figure 5.

Saturation hypothesis for how enhanced plasticity could impair learning. *Top*, In naïve wild type mice, synapses are eligible to undergo associative synaptic plasticity (LTD; *dark violet*) in response to training, thereby supporting normal learning.

Bottom, In mice with enhanced LTD, such as L7-*Fmr1* KO (*pink*) and MHCI $K^bD^{b-/-}$ (*green*), the lower threshold for induction of LTD allows it to be aberrantly recruited by spontaneous activity in the circuit (*light violet*), saturating the capacity for LTD and reducing its availability to be recruited during training at the synapses where it is needed to support learning, and thus impairing learning. Behavioral training that can reverse the LTD or drugs that reduce neural activity to reduce LTD induction can reset the synapses to an LTD-eligible state (*upward arrow*), restoring normal learning capacity.

required to target each of the many functional regions supporting the myriad motor and cognitive functions of the cerebellum. Thus, from a practical standpoint, pharmacological pre-treatment to prevent or reverse the saturation of LTD before training is likely to be a more feasible and general approach to restoring the capacity for PF-Purkinje cell LTD-dependent learning. Alternative approaches for manipulating neural activity, such as transcranial magnetic stimulation (TMS) and transcranial direct current stimulation (tDCS), may also hold promise (Tan et al., 2013; Gschwind and Seeck, 2016; Biabani et al., 2017; Miterko et al., 2019; Denoyer et al., 2020).

Conclusion

We leveraged the relatively simple and well understood physiology and function of the cerebellum and oculomotor system to develop and test a new hypothesis to explain why enhanced plasticity often impairs rather than enhances learning. The current results, along with the previous work by Nguyen-Vu and colleagues (2017) provide convergent evidence that a lower threshold for synaptic plasticity can result in its saturation and hence the impairment of learning. This saturation hypothesis may be useful in considering the impact of enhanced plasticity not only in the cerebellum, but in other brain areas as well. Moreover, the approach of limiting neural activity during a period before training may be broadly applicable for reversing maladaptive plasticity and resetting neural circuits to a state compatible with adaptive plasticity and new learning. For example, suppression of neural activity in the retina has been employed to reset the visual circuitry and enable recovery from amblyopia (Fong et al., 2021). The suppression of neural activity may be an especially useful approach if plasticity is pathologically enhanced in areas like the cerebellum or basal ganglia, with a high level of spontaneous spiking activity. More generally, the present results highlight the principle that synaptic properties do not control learning in isolation but interact with the patterns of neural activity in the corresponding circuits to control the capacity for new learning. The implication is that learning deficits associated with abnormal plasticity are not necessarily permanent, but in some cases can be remedied with appropriate reset of the circuit, opening up the possibility for therapeutic approaches targeting neural activity as well as the plasticity mechanisms themselves.

Materials and Methods

All experimental procedures were approved by the Administrative Panel on Laboratory Animal Care at Stanford University.

Mice

Mice with the *Fmr1* gene knocked out selectively from cerebellar Purkinje cells were generated through the following breeding strategy. First, homozygous female mice whose *Fmr1* gene, located on the X-chromosome, was floxed (*Fmr1* conditional knockout, cKO; Mientjes et al., 2006) were crossed with male mice expressing L7/Pcp2-Cre on an autosome (L7/Pcp2-Cre *Jdhu*; The Jackson Laboratory, Stock No. 010536; Zhang et al., 2004;). The L7/Pcp2-Cre *Jdhu* line expresses Cre-recombinase in a manner that is highly selective for Purkinje cells. Male offspring from this first cross were mated with females homozygous for the *Fmr1 c*KO allele to generate offspring homozygous for *Fmr1* cKO, with some mice L7/Pcp2-Cre-positive and some L7/Pcp2-Cre-negative. Cre-positive offspring of this second cross are referred to as L7-*Fmr1* KO, and their Cre-negative littermates were used as controls and referred to as wild type (WT). Genotyping was performed by Transnetyx Inc on ear-clipped samples to confirm the presence of the floxed *Fmr1* allele in all mice and the presence or absence of Cre using RT-qPCR.

Mice were kept on a reversed 12-h light/12-h dark cycle, and behavioral experiments were conducted during the dark cycle of the mice. After surgical implantation (see below), mice were housed individually in standard cages and provided food and water *ad libidum*. Male and female mice 8-22 weeks old were used in the behavioral experiments. Similar learning deficits were observed in male and female L7-*Fmr1* KO mice (p=0.30, VOR-increase learning in 7 male vs 7 female L7-*Fmr1* KO mice, **Fig. 1A**), therefore results were pooled across sex.

Surgery

Mice underwent surgery between 8-12 weeks of age to implant hardware for restraining the head and measuring eye movements, as described previously (Payne and Raymond, 2017; Nguyen-Vu et al., 2017). Mice were anesthetized with 1.5-2.5% isoflurane. An incision was made in the scalp and a custom-made head post (Shapeways Inc) was attached to the top of the skull using dental acrylic (Relyx Unicem Self-Adhesive Universal Resin Cement, Aplicap Capsule Refills-3M). Two stacked neodymium magnets with a total size of 0.75 x 2 mm (grade N50, axially magnetized, *SuperMagnetMan.com*) were implanted beneath the conjunctiva on the temporal side of the left eye. An angular magnetic field sensor (HMC1512, Honeywell Inc.) was soldered to an 8-pin connector and attached to the skull above the eye using dental acrylic, in a plane parallel to horizontal (nasal-temporal) eye movements. Eye movements were measured by detecting changes in the magnetic field created by movements of the magnet implanted on the eye (Payne and Raymond, 2017). Mice recovered from surgery for at least five days before experiments were performed.

Behavioral experiments

Mice were acclimatized to the laboratory for at least 20 min after transport from the animal care facility before the start of an experiment. Experiments were conducted in a light-proof, sound-attenuated chamber (IAC acoustics). The head of the mouse was secured by attaching its head post to a restrainer, which was then attached to a vestibular turntable controlled by a Carco Model 823 rate table and Model 405D controller. The turntable delivered vestibular stimuli to the mouse by rotation about a yaw (earth-vertical) axis centered on the head of the mouse. An optokinetic drum controlled by a Yaskawa AC-Servo SGMCS-02B3B11 motor provided visual stimulation by rotation about an earth-vertical axis aligned with that of the vestibular turntable. The drum was made of translucent white plastic, and had alternating black and white stripes, with each stripe subtending approximately 7.5° of the visual field, illuminated by an LED light strip attached to the rim of the drum. Eye movements were recorded using the method described in Payne & Raymond (2017).

Experiments to assess VOR learning consisted of testing blocks and training blocks. Testing blocks consisted of three 45 second tests of the eye movement response to the vestibular stimulus delivered alone in complete darkness, i.e., the VOR. The vestibular stimulus was sinusoidal vestibular turntable rotation at 1 Hz or 0.5 Hz with a peak velocity of ±10°/s. The three 45 s VOR tests in a block were separated by 10 s with the turntable stationary. Training blocks were ten minutes long, and were repeated three times for a total of 30 min training, with a testing block following each training block. For VOR-increase training, the vestibular stimulus used for testing the VOR (1 Hz or 0.5 Hz) was paired with oppositely directed motion of the illuminated optokinetic drum with the same peak velocity (±10°/s). For VOR-decrease training, the vestibular stimulus used for testing was paired with motion of the optokinetic drum in the same direction with the same velocity, so that the drum was stationary relative to the head of the mouse. In behavioral pre-training experiments, the pre-training consisted of three 10-min blocks of either VOR-decrease training or delivery of the vestibular stimulus alone in the dark (Vestibular only), with a testing block before each training block. Calibration of the signals from the magnetic sensor used to record eye movements was performed after the experiment, as described in Payne & Raymond (2017).

Prior to some experiments (**Figures 3**, **3-figure supplement 1** and **2** and 4B,), mice received a single IP injection of 0.4, 0.5, or 2.5 mg/kg diazepam (in saline) or saline control. After diazepam or saline administration, mice were returned to the home cage, and then behavioral testing was performed either 2 hours, 18-24 hours, and/or 1 week later.

Each mouse underwent multiple behavioral experiments, with at least two days between successive experiments. The same cohort of mice was used for the experiments shown in **Figures 1** and **2**, with the order of the experiments randomized. A subset of the same cohort was then used for the diazepam experiments shown in **Figure 3**. A separate cohort of mice was used for the low-frequency training experiments shown in **Figure 4**, with the order of randomized for the behavioral pre-training conditions shown in **Fig. 4A** (no pretraining, VOR-decrease pre-training and Vestibular only pre-training) followed by the diazepam pre-treatment experiments in **Fig. 4B**, with randomized order for drug and saline conditions.

Analysis of eye movement measurements

Signals from the magnetic sensor related to eye position were fourth-order low-pass (15 Hz) Butterworth filtered and then digitally differentiated to obtain eye velocity using a windowed (30 ms) Savitzky-Golay filter. Eye velocity data from each VOR test were fit with a 1 Hz or 0.5 Hz sinusoid. Values deviating from the sinusoidal fit by more than 31°/s were identified as saccades or movement artifacts and excluded, along with data from 50 ms before and after. Segments of data less than 10 ms in duration were also excluded. The entire 45 s VOR test was excluded if more than 45% of the data points were excluded. The amplitude of the sinusoidal fit provided the measure of the amplitude of the eye movement response, and values from the three VOR tests in a block were averaged. VOR learning (ΔVOR) was calculated as the percentage change in the VOR amplitude following each 10 min block of training relative to the baseline VOR amplitude measured before training. Eye movement gain was calculated at the ratio of eye movement amplitude to vestibular stimulus amplitude.

Statistical analysis

Data were analyzed with a Shapiro-Wilk test of normality, followed by a two-factor repeated measures ANOVA with posthoc Tukey or by a two-sample or paired sample t-test, executed in OriginPro 2022 software. A value of p less than 0.05 was considered significant. Data are plotted as mean ± SEM.

Code

All code used for data acquisition (*https://github.com/RaymondLab/Code/tree/Master/Experiment%20Protocols*) and analysis (*https://github.com/RaymondLab/Code/tree/Master/Tools/VOR_Analysis*) is available at *github.com/RaymondLab/Code*.

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In preparing this manuscript, we made a conscious effort to address citation bias. Following the approach outlined in Dworkin et al., (2020), we used open source code to assess the gender balance of our citations based on the first names of the first and last authors (Zhou et al., 2020). Excluding self-citations, our article includes citations as follows: 52.80% man/man, 12.50%

man/woman, 25.0% woman/man, and 9.7% woman/woman citations. For comparison, the proportions obtained from articles in the top five neuroscience journals (Dworkin et al., 2020) are as follows: 58.4% man/man, 9.4% man/woman, 25.5% woman/man, and 6.7% woman/woman.

Declaration of interest

The authors declare no competing interests.

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Author contributions

A.S. designed the study, performed the experiments, analyzed, and interpreted experimental data, and wrote the manuscript. J.G.F. and A.B.N. conducted experiments. J.B. conducted experiments and wrote the first draft of the methods section. J.L.R. designed experiments, interpreted data, and wrote the manuscript.

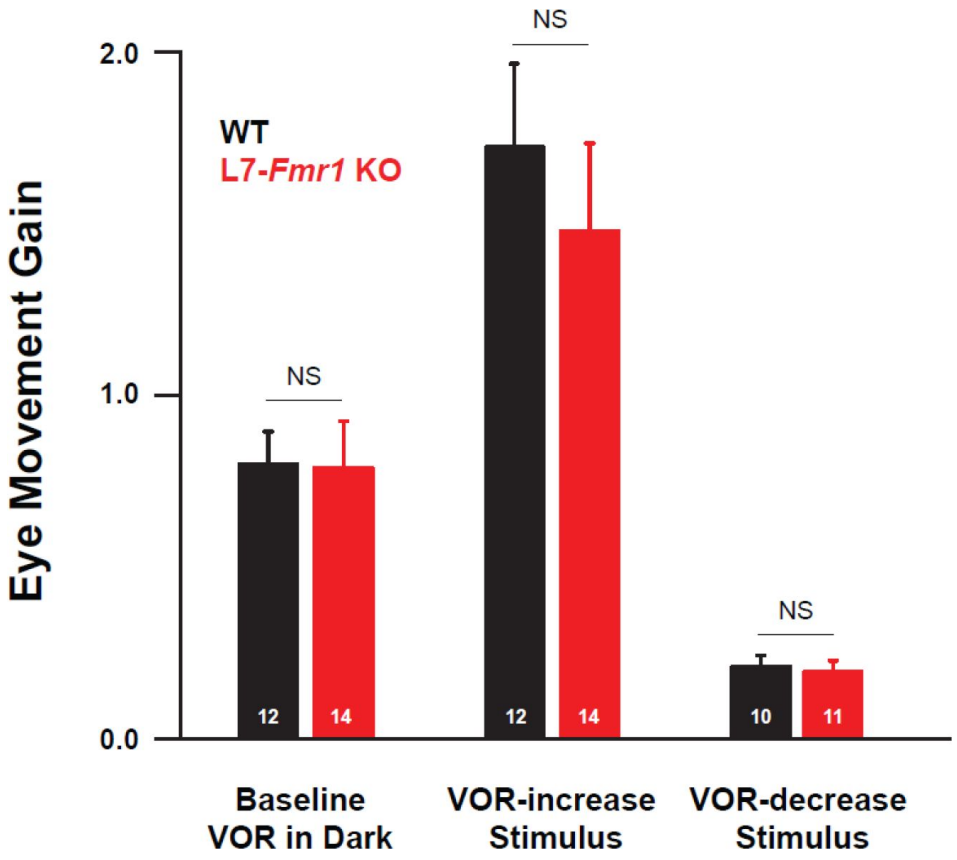

Figure 1-figure supplement 1.

Baseline oculomotor performance of L7-*Fmr1* KO mice was indistinguishable from WT.

The gain of the eye movement responses (ratio of eye movement amplitude to vestibular stimulus amplitude; see Methods) of L7-*Fmr1* KO mice (*red*) was not significantly different from that of WT mice (*black*) during baseline tests of the VOR in the dark before training (***left***; p= 0.95, two sample t-test) or during the first 45 sec of the paired presentation of visual and vestibular stimuli used for VOR-increase training (***middle***; p= 0.50, two sample t-test) or for VOR-decrease training (***right***; p= 0.76, two sample t-test). Number of mice tested is indicated in each bar.

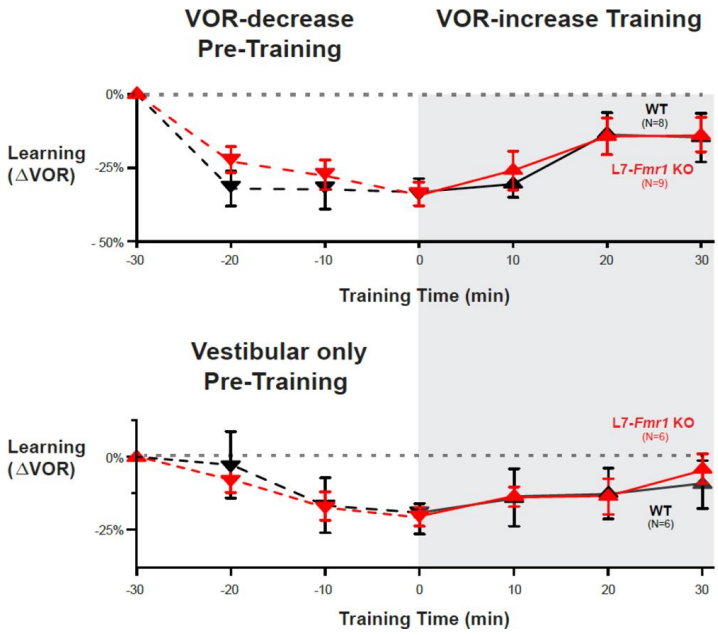

Figure 2-figure supplement 1.

Data from Figure 2 were subsampled to compare VOR-increase learning in subpopulations of mice matched for the mean learned decrease in the VOR during pre-training.

Subsampling was done by eliminating the WT mice (*black*) with the smallest decrease and L7-*Fmr1* KO mice (*red*) with the largest decrease in the VOR measured after 30 min of pre-training (just before the start of VOR-increase training), until the mean values in the two populations were within 2%. In these sub-sampled populations, the amount of VOR-increase learning was not significantly different between the L7-*Fmr1* KO and WT mice after VOR-decrease pre-training (*top;* p=0.74, L7-*Fmr1* KO mice vs. WT, 30 min, Tukey) or after Vestibular only pre-training (*bottom;* p=0.40, L7-*Fmr1* KO mice vs. WT, 30 min, Tukey), as also observed in the full samples.

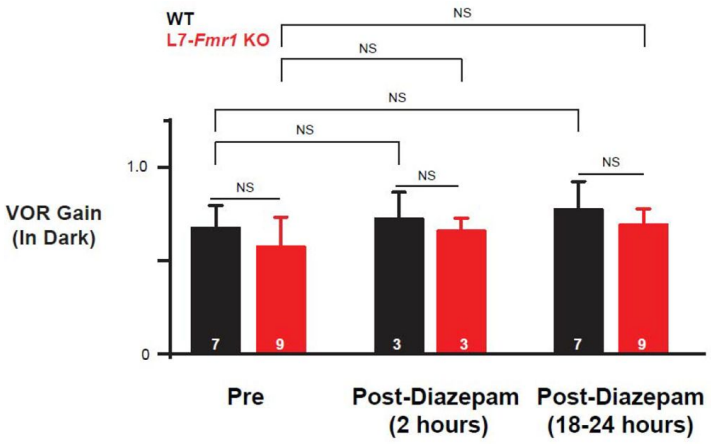

Figure 3- figure supplement 1.

Diazepam did not affect baseline VOR performance.

The gain of the VOR (ratio of eye velocity to vestibular stimulus velocity) was measured in the dark in L7-*Fmr1* KO (*red*) and WT (*black*) mice before (*Pre*), 2 hours after (*Post-Diazepam (2 hours)*) and 18-24 hours after (*Post-Diazepam (18-24 hours)*) an IP injection of diazepam (0.5 mg/kg). There was no effect of diazepam on the gain of the VOR in L7-Fmr1 KO mice (*red;* p=0.72, Pre vs 2 hours Post Diazepam; p= 0.77, Pre vs 18-14 hours Post-Diazepam; Tukey) or WT mice (*black*; p=0.99, Pre vs 2 hours Post Diazepam; p= 0.36, Pre vs 18-14 hours Post-Diazepam; Tukey). Moreover, the gain of the VOR of L7-*Fmr1* KO mice was not significantly different from that of WT mice during baseline tests of the VOR in the dark before diazepam administration Pre (***left***; p= 0.79, Tukey), ***Post-Diazepam (2 hours)*** (***middle***; p= 0.77, Tukey) and ***Post-Diazepam (18-24 hours)*** (***right***; p= 0.97, Tukey). The 2-hour and 18-24-hour VOR performance measurements were made just before the VOR-increase training sessions (training time = 0) shown in **Fig. 3-figure supplement 2B**, and **Fig. 3B** ***top***, respectively. The Pre VOR-performance measurements were made just before the VOR-increase training sessions shown in **Fig. 1A**, ***right*** for the subset of mice that were also tested 1 day after diazepam administration.

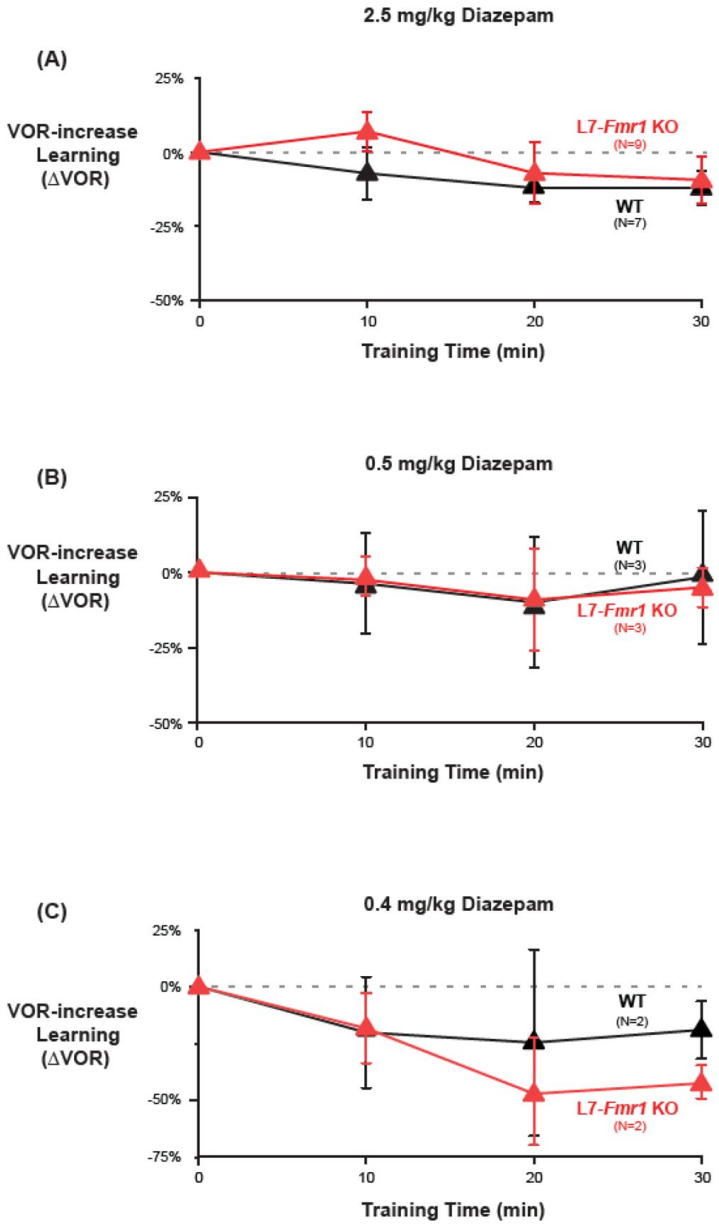

Figure 3- figure supplement 2

The acute effect of diazepam was inhibition of VOR-increase learning. When VOR-increase training was delivered two hours after IP injection of 0.5 mg/kg diazepam **(A)**, 0.4 mg/kg diazepam **(B)**, or 2.5 mg/kg diazepam **(C)**, no learned increase in VOR amplitude was observed in L7-*Fmr1* KO or WT mice.

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Reviewer #1 (Public Review):

Summary:
Shakhawat et al., investigated how enhancement of plasticity and impairment could result in the same behavioral phenotype. The authors tested the hypothesis that learning impairments result from saturation of plasticity mechanisms and had previously tested this hypothesis using mice lacking two class I major histocompatibility molecules. The current study extends this work by testing the saturation hypothesis in a Purkinje-cell (L7) specific Fmr1 knockout mouse mice, which have enhanced parallel fiber-Purkinje cell LTD. The authors found that L7-Fmr1 knockout mice are impaired on an oculomotor learning task and both pre-training, to reverse LTD, and diazepam, to suppress neural activity, eliminated the deficit when compared to controls.

Strengths:

This study tests the "saturation hypothesis" to understand plasticity in learning using a well-known behavior task, VOR, and an additional genetic mouse line with a cerebellar cell-specific target, L7-Fmr1 KO. This hypothesis is of interest to the community as it evokes a novel inquisition into LTD that has not been examined previously.

Utilizing a cell-specific mouse line that has been previously used as a genetic model to study Fragile X syndrome is a unique way to study the role of Purkinje cells and the Fmr1 gene. This increases the understanding in the field in regards to Fragile X syndrome and LTD.

The VOR task is a classic behavior task that is well understood, therefore using this metric is very reliable for testing new animal models and treatment strategies. The effects of pretraining are clearly robust and this analysis technique could be applied across different behavior data sets.

The rescue shown using diazepam is very interesting as this is a therapeutic that could be used in clinical populations as it is already approved.

There was a proper use of controls and all animal information was described. The statistical analysis and figures are clear and well describe the results.

Weaknesses:
While the proposed hypothesis is tested using genetic animal models and the VOR task, LTD itself is not measured. This study would have benefited from a direct analysis of LTD in the cerebellar cortex in the proposed circuits.

Diazepam was shown to rescue learning in L7-Fmr1 KO mice, but this drug is a benzodiazepine and can cause a physical dependence. While the concentrations used in this study were quite low and animals were dosed acutely, potential side-effects of the drug were not examined, including any possible withdrawal. This drug is not specific to Purkinje cells or cerebellar circuits, so the action of the drug on cerebellar circuitry is not well understood for the study presented.

It was not mentioned if L7-Fmr1 KO mice have behavior impairments that worsen with age or if Purkinje cells and the cerebellar microcircuit are intact throughout the lifespan. Connections between Purkinje cells and interneurons could also influence the behavior results found.

While males and females were both used for the current study, only 7 of each sex were analyzed, which could be underpowered. While it might be justified to combine sexes for this particular study, it would be worth understanding this model in more detail.

Training was only shown up to 30 minutes and learning did not seem to plateau in most cases. What would happen if training continued beyond the 30 minutes? Would L7-Fmr1 KO mice catch-up to WT littermates?

The pathway discussed as the main focus for VOR in this learning paradigm was connections between parallel fibers (PF) and Purkinje cells, but the possibility of other local or downstream circuitry being involved was not discussed. PF-Purkinje cell circuits were not directly analyzed, which makes this claim difficult to assess.

The authors mostly achieved their aim and the results support their conclusion and proposed hypothesis. This work will be impactful on the field as it uses a new Purkinje-cell specific mouse model to study a classic cerebellar task. The use of diazepam could be further analyzed in other genetic models of neurodevelopmental disorders to understand if effects on LTD can rescue other pathways and behavior outcomes.

Reviewer #2 (Public Review):

This manuscript explores the seemingly paradoxical observation that enhanced synaptic plasticity impairs (rather than enhances) certain forms of learning and memory. The central hypothesis is that such impairments arise due to saturation of synaptic plasticity, such that the synaptic plasticity required for learning can no longer be induced. A prior study provided evidence for this hypothesis using transgenic mice that lack major histocompatibility class 1 molecules and show enhanced long-term depression (LTD) at synapses between granule cells and Purkinje cells of the cerebellum. The study found that a form of LTD-dependent motor learning-increasing the gain of the vestibulo-ocular reflex (VOR)-is impaired in these mice and can be rescued by manipulations designed to "unsaturate" LTD. The present study extends this line of investigation to another transgenic mouse line with enhanced LTD, namely, mice with the Fragile X gene knocked out. The main findings are that VOR gain increased learning is selectively impaired in these mice but can be rescued by specific manipulations of visuomotor experience known to reverse cerebellar LTD. Additionally, the authors show that a transient global enhancement of neuronal inhibition also selectively rescues gain increases learning. This latter finding has potential clinical relevance since the drug used to boost inhibition, diazepam, is FDA-approved and commonly used in the clinic. The evidence provided for the saturation is somewhat indirect because directly measuring synaptic strength in vivo is technically difficult. Nevertheless, the experimental results are solid. In particular, the specificity of the effects to forms of plasticity previously shown to require LTD is remarkable. The authors should consider including a brief discussion of some of the important untested assumptions of the saturation hypothesis, including the requirement that cerebellar LTD depends not only on pre- and postsynaptic activity (as is typically assumed) but also on the prior history of synaptic activation.