Interneuron Dysfunction and Inhibitory Deficits in Autism and Fragile X Syndrome

Toshihiro Nomura

Department of Neuroscience, Feinberg School of Medicine, Northwestern University, Chicago, IL 60611, USA; toshihiro.nomura@northwestern.edu

Abstract: The alteration of excitatory–inhibitory (E–I) balance has been implicated in various neurological and psychiatric diseases, including autism spectrum disorder (ASD). Fragile X syndrome (FXS) is a single-gene disorder that is the most common known cause of ASD. Understanding the molecular and physiological features of FXS is thought to enhance our knowledge of the pathophysiology of ASD. Accumulated evidence implicates deficits in the inhibitory circuits in FXS that tips E–I balance toward excitation. Deficits in interneurons, the main source of an inhibitory neurotransmitter, gamma-aminobutyric acid (GABA), have been reported in FXS, including a reduced number of cells, reduction in intrinsic cellular excitability, or weaker synaptic connectivity. Manipulating the interneuron activity ameliorated the symptoms in the FXS mouse model, which makes it reasonable to conceptualize FXS as an interneuronopathy. While it is still poorly understood how the developmental profiles of the inhibitory circuit go awry in FXS, recent works have uncovered several developmental alterations in the functional properties of interneurons. Correcting disrupted E–I balance by potentiating the inhibitory circuit by targeting interneurons may have a therapeutic potential in FXS. I will review the recent evidence about the inhibitory alterations and interneuron dysfunction in ASD and FXS and will discuss the future directions of this field.

Keywords: autism; fragile X syndrome; interneuron; GABA; E–I balance

1. Introduction

This review will discuss the pathophysiology of autism and fragile X syndrome (FXS) from a neurophysiological viewpoint. I will take FXS as a model to understand autism pathophysiology and will put this autism-related disorder at the center of the discussion. Balanced excitation and inhibition in the brain is critical to maintaining proper neuronal function. Alterations both in excitation and inhibition have been described in autism and FXS, but the inhibitory system will be the main focus in this review. I will particularly emphasize interneuron dysfunction as an important component responsible for inhibitory alterations. Taking the developmental characteristics of autism and FXS into account, how the developmental profile of the inhibitory system goes awry will also be precisely described. I will summarize and discuss the published evidence and will present future perspectives, particularly from a translational aspect, in this field.

2. Excitatory–Inhibitory (E–I) Imbalance

It is now cliché to say “Excitatory–Inhibitory (E–I) balance is disrupted in XXX disorders”. Intuitively, disrupted E–I balance has been rigorously studied in paroxysmal disorders where “more excitatory” and/or “less inhibitory” brain activity is associated with epileptic seizures. While this straightforward connection makes sense, the hypothesis that E–I imbalance is causal to a multitude of behavioral symptoms is undoubtedly too simplistic [1–3]. “XXX disorders” now, however, range from paroxysmal disorders to various other disorders, including autism [2–6]. Autism and epilepsy are commonly comorbid conditions, and the genesis for the altered E–I balance theory in autism was because of this co-occurrence [5].
In 2003, Rubenstein and Merzenich published a seminal review in which they proposed a model in which E–I balance was shifted towards excitation that was the basis for the pathological brain circuitry function in autism [7]. Alterations both in the excitatory and inhibitory systems can result in E–I imbalance. Multiple factors, including the number of neurons, intrinsic excitability of neurons, synaptic connectivity, and network activity, can all be critical components regulating E–I balance [3]. Indeed, both excitatory and inhibitory alterations have been reported in autism [8]. In this review, inhibitory alterations, particularly dysfunction of interneurons, the main source of an inhibitory neurotransmitter gamma-aminobutyric acid (GABA), will be the primary focus.

3. Autism and Fragile X Syndrome

Autism is a neurodevelopmental disorder that is conceptualized as a spectrum of disorders (autism spectrum disorder; ASD). ASD individuals have difficulties in social behaviors and communication skills as core symptoms, but comorbid symptoms such as stereotyped movements, restricted interests, anxiety, or sensory hypersensitivity vary considerably in the type and magnitude among individuals [9–11]. The heterogeneity of ASD, which likely involves multiple genetic and environmental factors, makes it challenging to clarify the pathophysiological mechanism [12,13]. Several single-gene or chromosomal disorders—for instance, FXS—have a high incidence of co-occurrence of ASD-like symptoms, and it is expected that understanding these disorders may lead to elucidating the pathophysiology for the development of new therapeutic interventions for ASD [10,14].

FXS is the most common (with a prevalence of 1 in 4000–5000 in males and 1 in 6000–8000 in females, but it varies considerably among studies [15,16]) known cause of intellectual disability and ASD [17,18]. FXS results from an epigenetic silencing of the responsible gene FMR1 caused by an abnormal expansion of a CGG repeat sequence in the 5’ untranslated region, leading to hypermethylation, transcriptional silencing, and a loss of expression of the protein product fragile X mental retardation protein (FMRP) [19–21]. FMRP is an RNA-binding protein enriched in neurons, epithelial tissues, and testes that regulates the expression of other genes as a translational repressor [22–24]. Animal models are particularly useful tools to understand the molecular, cellular, and circuit phenotypes, because they allow an accessibility to study neuronal and circuit functions, which is very difficult or practically impossible in affected human subjects. In FXS, multiple animal models are available and commonly used, including Fmr1 knock-out (KO) mice [25], Dfxr (Drosophila fragile X-related gene) null drosophila [26], Fmr1 KO zebrafish [27], and Fmr1 KO rats [28]. The “knock-in” of the expanded CGG repeat did not work well in mice to model FXS phenotypes [29–31].

As a pathological phenotype, disruption in E–I balance has been reported in FXS. The target of FMRP includes RNAs encoding synaptic proteins and trophic factors [21,32–35]. Therefore, it is not surprising that multiple morphological and functional alterations affecting E–I balance exist in FXS. Although it is not the central focus of this review, the excitatory neurotransmitter system has been intensively studied in FXS for more than two decades, and studies have revealed that glutamatergic transmission goes awry in FXS [36–38].

4. Inhibitory System Alteration

The inhibitory neurotransmitter system is a counter component to the excitatory system determining E–I balance. There are multiple lines of evidence for inhibitory alterations in individuals with FXS and in the animal models [21,39]. Circuit hyperexcitability is consistently reported in FXS. Though there is huge variability in the prevalence among the cohorts, FXS individuals have a high susceptibility to seizures, electroencephalogram (EEG) abnormalities, and epilepsy syndromes [14,39,40], which may be, in part, due to an abnormal inhibitory system. GABA is the main source of inhibitory transmission and may play an important role in dysfunction in the inhibitory system in FXS [41,42], but direct evidence for a dysfunctional GABA system in human subjects is relatively limited. A
A study using positron emission tomography (PET) in a small cohort of patients showed a significant reduction of approximately 10% in GABA<sub>A</sub> receptor availability throughout the brain, which may cause diminished GABA-mediated inhibition in FXS individuals [43]. A recent finding using transcranial magnetic stimulation (TMS) demonstrated that FXS patients had significantly increased intracortical facilitation (ICF), reduced short-interval intracortical inhibition (SICI), and increased long-interval intracortical inhibition (LICI), which are believed to represent glutamatergic excitation, GABA<sub>A</sub>-mediated inhibition, and GABA<sub>B</sub>-mediated inhibition, suggesting that cortical hyperexcitability is, in part, due to reduced GABA<sub>A</sub> inhibition in FXS individuals [44].

In FXS animal models, however, aberrant GABA signaling has been reported in multiple studies. At the behavioral level, there is evidence indicative of the altered inhibition. Fmr1 KO mice have a lower threshold for audiogenic seizures, which is one of the most robust and reliably reported behavioral phenotypes in this mouse line [45,46]. Audiogenic seizures are possibly triggered by neuronal and circuit hyperexcitability due to altered sensory processing and sensory hypersensitivity in the auditory system [21,45,47]. Moreover, a causal relationship between diminished GABA-mediated inhibition and audiogenic seizures has been established, as agonists of GABA<sub>A</sub> [48,49] and GABA<sub>B</sub> receptors can rescue enhanced audiogenic seizures in Fmr1 KO mice, and antagonists of GABA<sub>B</sub> receptors mimic the Fmr1 KO phenotype [50].

Pre-pulse inhibition (PPI) is another indicator for auditory hypersensitivity, and multiple studies have reported increased PPI in Fmr1 KO mice [52]. Increased PPI in mouse models is contradictory to the finding in FXS individuals where PPI is reported to be decreased [53]. A study using eye blink responses [54] and a recent study in young (P23–25) Fmr1 KO mice [55] have shown a significantly reduced PPI, which is somewhat reminiscent of the findings in human FXS individuals. Importantly GABA<sub>A</sub> receptor activation by its agonist 4,5,6,7-tetrahydroisoxazolo(5,4-c)pyridin-3-ol (THIP) ameliorates increased PPI in Fmr1 KO mice, suggesting a causal relationship between decreased GABA inhibition and altered PPI [52]. Alterations in PPI may also, in part, arise from sensory hypersensitivity to somatosensory stimuli, which is also one of the most commonly shared phenotypes in FXS individuals. Evidence shows a negative correlation in the GABA concentration in the somatosensory cortex to tactile sensitivity in humans [56–59]. In animal models of FXS, a recent study demonstrated that sensory hypersensitivity can be reliably monitored even in young (<P14) and in adult Fmr1 KO mice, which is associated with a deficit in an adaptation in cortical neuronal response activity in the somatosensory cortex to repetitive tactile stimuli [60].

5. ASD and FXS Are Interneuronopathy

The concept of “interneuronopathy” was originally proposed in X-linked lissencephaly with abnormal genitalia (XLAG) [61], in which deficient tangential migration and a loss of inhibitory interneurons were described [61,62]. It was later expanded to other intractable epilepsy syndromes, such as Dravet syndrome [63,64] or West syndrome [65,66], in which deficits in the development, the number, and the function of interneurons are thought to contribute to clinical manifestations such as epileptic seizures. Interneurons have local axons and innervate adjacent principal projection neurons and other interneurons [67,68]. Interneurons are composed of diverse subpopulations of neurons characterized by the expression of specific marker proteins such as parvalbumin (PV), somatostatin (SST), or vasoactive intestinal peptide (VIP) [67,68] (but there are also multiple different ways to determine interneuron subclasses, including recently proposed ways based on single-cell transcriptome or connectivity and projection pattern: please see these articles about interneuron diversity for more details [68–73]). While interneurons represent a relatively minor population (~20%) of all neurons in the brain [67,74,75], they are a source of GABA and serve as a main source of inhibitory transmission, suggesting their indispensable roles in physiological and pathological conditions.
There is now growing evidence about interneuron dysfunction in ASD and FXS. While it is not conclusive whether interneuron dysfunction is the primary pathological mechanism, it motivates researchers to consider these disorders to be interneuronopathies [8, 76, 77]. In human research, postmortem studies have consistently reported reduced cell density of PV positive fast-spiking interneurons, one of the most major classes of interneurons [67] (Figure 1), in ASD individuals [77]. In FXS individuals, however, direct evidence is limited about interneuron phenotypes. The deficit in interneurons in FXS was originally reported in the mouse model. Selby et al. first described a remarkable (~20%) decrease in the cell density of PV interneurons in the somatosensory cortex in Fmr1 KO mice [78]. Similar findings were reported in the developing auditory cortex, and perineuronal net (PNN) formation, which is the extracellular matrix associated with PV interneurons, is also impaired in the same mouse line [79]. These findings are interesting given the TMS study in human subjects showing a reduced SICI in FXS [44], because TMS is believed to act predominantly through modulating PV interneurons [80].

![Figure 1](image)

**Figure 1.** (A) PV-positive cortical interneuron filled with biocytin labeled with streptavidin-conjugated AlexaFluor 488. Alterations in this subclass of interneurons are consistently reported in ASD. Scale bar: 50 μm. (B) Typical firing pattern of PV interneurons in response to depolarizing current injection (100 pA for 500 ms). These neurons fire action potentials at a high frequency constantly for a prolonged period. The inter-spike intervals do not largely change during the prolonged spike train, i.e., fast-spiking. Calibration: 100 ms and 25 mV (from Reference [81] with edits).

Not only are the reductions in the number of cells, but also, the functional alterations in inhibitory interneurons, particularly in PV interneurons, are implicated in multiple studies in FXS animal models [77]. Early works demonstrated that, in the amygdala, there is a drastic reduction both in phasic and tonic inhibitory transmission in Fmr1 KO mice [82]. This reduced inhibition is likely caused by functional deficits in presynaptic neurons in inhibitory synaptic circuits, i.e., GABAergic interneurons, as the GABA synthesizing enzymes glutamate decarboxylase (GAD) 65/67 levels and synaptic GABA availability are reduced in Fmr1 KO mice [82]. In vivo recording results in sensory cortices agree with the reduced interneuron activity in Fmr1 KO mice [83] and Fmr1 KO rats [84]. Impaired interneuron function was also implicated in a drosophila model of FXS [85]. The optogenetic activation of interneurons fails to elicit as strong lateral inhibition in Dfmr1 KO flies as in wild-type (WT), which indirectly indicates impaired interneuron function and outputs [85]. Probably one of the most direct and important pieces of evidence about abnormal interneuron function is a recent finding in the visual cortex in Fmr1 KO mice [86]. The activity of PV interneurons was directly monitored using in vivo Ca^{2+} imaging technique during a visual discrimination task [86]. Fmr1 KO mice showed poor behavioral performances in the task,
and the interneuron activity was reduced in Fmr1 KO mice [86]. Most importantly, artificial activation of these neurons by designer receptors exclusively activated by designer drugs (DREADD) restored the impairment in the behavioral performance, which strongly suggests a causal relationship between the neuronal activity and the behavioral phenotype [86]. In addition, the poor performance in the behavioral task was fully translated in human FXS individuals [86]. Evidence for alterations in other subclasses of interneurons such as SST or VIP-positive interneurons is limited in ASD and FXS.

Although multiple studies have suggested reduced inhibition due to reduced interneuron activity in animal models of FXS, evidence in the cerebellum of Fmr1 KO mice counteracts that idea [87]. Cerebellar basket cells (BCs), a major class of cerebellar interneurons, exhibit hyperactivity in their axonal terminals, leading to an exaggerated GABAergic inhibition onto the principal neuron Purkinje cells (PCs) and reduced firing of PCs in Fmr1 KO mice [87]. This hyper-inhibition is caused by a loss of direct modulation of the voltage-gated potassium channel Kv1.2 by the N-terminal of FMRP [87]. Given the ubiquitous expression of Kv1.2 across the brain, the mechanism for this dichotomy (increased inhibition in the cerebellum vs. reduced inhibition in other brain regions) cannot be easily explained. Interestingly, exaggerated inhibitory transmission and/or the reduced firing of PCs have been reported in multiple ASD animal models, including in Tsc1 [88], PTEN [89], Shank2 [90] mutants, and BTBR [91] mouse lines. Therefore, it might be possible that these cerebellar cellular and circuitry phenotypes represent certain aspects of ASD. Given the fact that the cerebellum is implicated to be one of the key brain regions responsible for the pathophysiology of ASD [92], further studies are clearly needed to explore this regional specificity or discrepancy with other brain regions.

6. Developmental Alteration

Given the characteristics of FXS as a developmental disorder, understanding the developmental profiles of the brain circuit is undoubtedly indispensable. However, current evidence about the circuit development is relatively limited compared to the large number of studies in adult FXS animal models. Here, I will focus on evidence in the somatosensory cortex in which several studies have uncovered the abnormal development of both excitatory and inhibitory circuits. Several studies have demonstrated transient alterations in the morphology and turnover of dendritic spines [93–98], functional maturation of excitatory synapses [99,100], and the excitatory innervation pattern [101] during the somatosensory critical period (1 to 2 postnatal weeks) in Fmr1 KO mice. The closure of the critical period for synaptic plasticity was delayed in the somatosensory cortex in Fmr1 KO mice [99,100]. These series of developmental studies demonstrate that alterations in the excitatory synapses are parallel with the time course of the sensory critical period. Importantly, the timing of the critical period closure is largely determined by cortical inhibition, particularly by PV interneurons, which have been rigorously studied in the visual cortex [102].

Studies have uncovered developmental cellular and synaptic alterations in PV interneurons during the sensory critical period. The intrinsic membrane properties and characteristic fast-spiking firing patterns exhibit an immature profile in the developing (P5–P10) somatosensory cortex (layer 4) in Fmr1 KO mice [81] (Figure 2A,B). Another study demonstrated that PV interneurons show immature passive membrane properties but fire an action potential more easily (lower rheobase current) in P10 Fmr1 KO mice [103]. These studies demonstrated that the functional maturation of PV interneurons is significantly delayed in Fmr1 KO mice. The development of excitatory synapses onto PV interneurons is also significantly delayed during the critical period. The frequency of the spontaneous excitatory postsynaptic current (sEPSC) recorded from PV interneurons is lower, and the density of the synapses was decreased in <P10 Fmr1 KO mice, which suggests that the bulk excitatory synaptic input and the number of synapses are reduced in PV interneurons in Fmr1 KO mice [81] (Figure 2C–G). The synaptic connection probability and the strength of the individual synapses between excitatory stellate neurons, the principal neurons in layer 4, and PV positive fast-spiking interneurons are also reduced in 2-week-old Fmr1 KO mice [81].
Studies have uncovered developmental cellular and synaptic alterations in PV interneurons in developing (P5–P9) Fmr1 WT (top) and KO (bottom) mice. Synapses are labeled with the colocalized puncta of postsynaptic marker PSD95 (red) and presynaptic marker synaptophysin (“Synapt”: blue). The reference cube represents 1-µm thick sections. (A) Representative voltage response traces of PV interneurons in developing (postnatal day 9: P9) Fmr1 WT (top) and KO (bottom) mice. The initial and final portions of the 500-ms-long traces were magnified for clarity. Note the significantly greater ratio of the last three inter-spike intervals “b” to the first inter-spike interval “a” in KO mice compared to WT mice, which indicates that PV interneurons equip less fast-spiking properties in KO mice. Calibration: 50 ms. (B) Collective data for fast-spiking property during the development (P5–P10) in Fmr1 WT and KO mice. The spiking property was quantified by “a” divided by “b/3”. * Denotes p < 0.05. (C) Representative sEPSC traces recorded from developing (P5–P9) PV interneurons in Fmr1 WT (top) and KO (bottom) mice. Calibration: 1 s and 10 pA. (D) Collective data for sEPSC frequency. Each data point represents the mean sEPSC frequency in each cell. The frequency increases developmentally but remains lower in Fmr1 KO mice. (E,F). Three-dimensional renderings of dendritic segments (green) of WT (top) and KO (bottom) mice. Calibration: 1 s and 10 pA. (D) Collective data for the analysis of the synapse densities in Fmr1 WT and KO mice (P9–P10). The density of the synapses was significantly lower in Fmr1 KO mice (from Reference [81] with edits).

The deficits in the intrinsic and synaptic properties in PV interneurons result in antagonistic complex alterations in thalamocortical circuit activity during the critical period. Thalamocortical feed-forward inhibition (FFI) involves a di-synaptic circuit in which inhibition is mediated by PV interneurons [106,107]. The FFI is deficient (lacked in a subset of cells) in the somatosensory cortex (layer 4) in P10 Fmr1 KO mice [103]. Nomura et al. unpublished. Thalamic excitatory afferents produce action potentials in stellate cells more readily but with less precise timing upon stimuli at physiologically relevant frequencies as a result of deficiency in FFI [103]. While some cellular and synaptic measures of PV interneurons KO mice, which again suggests that the excitatory synaptic inputs onto PV interneurons are weaker in Fmr1 KO mice [103–105]. These findings indicate that PV interneurons demonstrate immature and defective activity both intrinsically and synaptically during the critical period in Fmr1 KO mice.
are normalized as the mice develop [81,104,105], as with excitatory synapses [94,99,101], there are persistent circuit-level (including FFI in layer 2/3 [83]) and behavioral-level phenotypes in adult Fmr1 KO mice beyond the sensory critical period [60,83,100,108–110]. This indicates that transient cellular and synaptic deficits in PV interneurons during the critical period can still result in persistent improper thalamocortical circuit refinement, which might warrant correcting these alterations during the developmental period is critical to treat symptoms in Fmr1 KO mice. Indeed, manipulating GABA signaling by a Cl− transporter modulator, which alters the GABA receptor polarity during the critical period, restores the abnormal whisker-evoked responses (wider receptive field and stronger signal) in the adult somatosensory cortex in Fmr1 KO mice [100]. Whether manipulating the interneuron activity during the critical period has persistent beneficial effects in Fmr1 KO mice has not been tested, but several approaches, including optogenetic or chemogenetic tools, to specifically target interneurons may help testing this critical period-interneuron hypothesis.

In addition to animal models, using human pluripotent cells such as induced pluripotent stem cells (iPS) or embryonic stem (ES) cells allows analyses of developmental profiles in vitro, which is not easy or even impossible in embryos of human patients. This approach is also useful for neurodevelopmental disorders, including ASD and FXS [111,112]. Evidence specific to interneurons in human-derived neurons is limited, but functional analyses using human ES cell-derived excitatory neurons from FXS individuals uncovered impaired action potential firing, which may be, in part, due to reduced activity in voltage-gated Na+ and K+ channels, as well as reduced synaptic excitation, which may be attributable to the reduced excitability and impaired vesicular release in the presynaptic neurons [113,114]. Similar approaches could be applied to study the interneuron pathology in human FXS, particularly developmental alterations in combination with experimental protocols, to specifically differentiate interneurons from pluripotent cells [115–118].

7. Translational Perspective

As discussed, there is now considerable evidence to believe the inhibitory deficits and interneuron dysfunction in the pathophysiology of FXS and ASD. It is theoretical and justifiable to consider the GABAergic inhibitory system as a promising therapeutic target for FXS [39,42]. However, up to now, there have been no successes in clinical trials in FXS that have tested GABA-targeting drugs. Arbaclofen, which is an agonist of GABA B receptors, was not effective in FXS individuals in phase 2 and phase 3 clinical trials, while the post hoc analyses showed several beneficial results in several measures, which might suggest younger and more severely affected individuals may have some beneficial effects [119,120]. Ganaxolone, which is an agonist of δ subunit-containing GABA A receptors, was not effective in FXS individuals in a phase 2 clinical trial, while the post hoc analyses presented some potential where individuals with higher anxiety or lower cognitive function may have more beneficial effects [121]. These clinical studies may have indicated these treatments might be effective in a subpopulation of affected individuals, depending on age, specific symptoms, or severity, for instance. However, it appears not to be game-changing.

It might be worth noting that most clinical research so far has targeted GABA receptors, the postsynaptic component of the inhibitory circuit. Given evidence is accumulating about the dysfunction of inhibitory interneurons, the presynaptic component of the inhibitory system, targeting these presynaptic neurons might be worth considering. Though manipulating a certain population of neurons is challenging, particularly in live human brains, cell type-specific proteins such as channels, receptors, or transporters specifically expressed in certain synapses or in certain circuits might be potential candidates to be considered. For instance, Kv3.1, a subfamily of voltage-gated potassium channels, is enriched in high-frequency firing neurons such as PV-positive fast-spiking GABAergic interneurons, glycinerergic medial nucleus of the trapezoid body (MNTB) neurons, and cerebellar Purkinje neurons [122]. Modulators of Kv3.1 have been shown to manipulate neuronal excitability in PV interneurons [123,124] and MNTB neurons [125]. A recent
study demonstrated that AUT2, a positive modulator of Kv3.1, restored an enhanced wave IV in the auditory brainstem response (ABR) recordings in Fmr1 KO mice, which may have suggested a therapeutic potential for sensory symptoms in FXS individuals, although the pharmacological actions for ABR rescue are likely through modulating the activity of MNTB neurons, not GABAergic interneurons [126]. Further studies for the cell type and circuit-specific analyses might be helpful for developing novel therapeutics in the future.

While the GABAergic inhibitory system looks like a reasonable therapeutic target for ASD and FXS [39,42], recent research has critically questioned if E–I imbalance particularly reduced inhibition is causal for ASD pathophysiology [83]. It has been demonstrated that multiple ASD mouse models, including Fmr1 KO mice, show E–I imbalance towards excitation and reduced interneuron activity in the somatosensory cortex (layer 2/3), but the network excitability (postsynaptic depolarization and sensory-evoked firing) is largely unchanged or even reduced in Fmr1 KO mice [83]. The authors concluded that altered E–I balance is compensatory, instead of causal, change to stabilize depolarization and spiking [83]. If this compensatory model is the case, targeting the inhibitory system to enhance the inhibition by GABA receptor activators, for instance, may amplify, instead of ameliorate, the symptoms of ASD and FXS. In addition, recent studies have shown that the frequency of basal spontaneous inhibitory postsynaptic current (sIPSC) recorded from principal neurons in the somatosensory cortex (layers 2/3) [127] and in the lateral amygdala [128] is enhanced, which may reflect the circuit hyperexcitability in Fmr1 KO mice. Synaptic plasticity in excitatory and inhibitory synapses are exaggerated and diminished, respectively, in Fmr1 KO mice [127,128], which might help to stabilize the homeostatic circuit activity. These findings may again question if correcting synaptic GABA signaling is beneficial for FXS. Further studies are clearly needed to clarify and tackle this issue.

More generally, not restricted to ASD or FXS, successful translational research, particularly in seeking new therapeutics, is extremely rare. It is known that more than 80% of the candidate compounds that demonstrated safety and efficacy in preclinical studies have no beneficial effect in human patients [129]. This discrepancy may arise from multiple complex factors, including the heterogeneity of the disorder, a lack of objective biomarkers and measures of the outcome, invalid sample size, and publication and citation biases towards positive results [129–135]. Registered reports in which research proposals are registered and reviewed before experiments are started and publication is guaranteed regardless of the outcome (whether positive or negative/novel or not novel) may reduce some of these biases and may have a potential to serve as an alternative platform in translational research, particularly in hypothesis-driven studies [136,137].

8. Conclusions

Decades of research uncovered circuit E–I imbalance as a shared feature in various brain disorders, including ASD. Single-gene disorders, including FXS, have been serving as a good model to study ASD pathophysiology together with their animal models. Evidence is accumulating about alterations in the inhibitory system, and it looks like a promising target to seek new therapeutics in ASD and FXS. Interneuron dysfunction is an important contributor to impaired inhibition, and it may also be causal for the symptoms of ASD and FXS. Evidence is accumulating about developmental alterations, but we are at an early stage of understanding. The pharmacological and modern chemogenetic manipulations of this inhibitory system ameliorate the pathological symptoms in animal models, but no successful clinical trials targeting GABA receptors are available so far. The successful translation from basic research to clinical practice is rare, which should not be overlooked. Although we are still far behind a complete picture for the satisfactory understanding of ASD and FXS, intense collaborative studies between clinical and basic researchers may promote our knowledge about these intractable disorders.

**Funding:** This research received no external funding.

**Institutional Review Board Statement:** Not applicable.
Informed Consent Statement: Not applicable.

Data Availability Statement: Not applicable.

Acknowledgments: The author thanks Jun Nishiyama (Duke-NUS), Nazim Kourdougli (UCLA), Nozomu Takata, and Anis Contractor (Northwestern) for the critical reading of the manuscript and helpful comments.

Conflicts of Interest: The author does not declare any conflict of interest.

References

1. Kofke, W.A.; Tempelhoff, R.; Dasheiff, R.M. Anesthetic implications of epilepsy, status epilepticus, and epilepsy surgery. *J. Neurosurg. Anesthesiol.* 1997, 9, 349–372. [CrossRef]

2. Eichler, S.A.; Meier, J.C. E–I balance and human diseases—From molecules to networking. *Front. Mol. Neurosci.* 2008, 1, 2. [CrossRef]

3. Tatti, R.; Haley, M.S.; Swanson, O.K.; Tselha, T.; Maffei, A. Neurophysiology and Regulation of the Balance Between Excitation and Inhibition in Neocortical Circuits. *Biol. Psychiatry* 2017, 81, 821–831. [CrossRef]

4. Bozzi, Y.; Provenzano, G.; Casarosa, S. Neurobiological bases of autism-epilepsy comorbidity: A focus on excitation/inhibition imbalance. *Eur. J. Neurosci.* 2018, 47, 534–548. [CrossRef] [PubMed]

5. Sohal, V.S.; Rubenstein, J.L.R. Excitation-inhibition balance as a framework for investigating mechanisms in neuropsychiatric disorders. *Mol. Psychiatry* 2019, 24, 1248–1257. [CrossRef] [PubMed]

6. Ghatak, S.; Talantova, M.; McKercher, S.R.; Lipton, S.A. Novel Therapeutic Approach for Excitatory/Inhibitory Imbalance in Neurodevelopmental and Neurodegenerative Diseases. *Annu. Rev. Pharmacol. Toxicol.* 2021, 61, 701–721. [CrossRef] [PubMed]

7. Rubenstein, J.L.; Merzenich, M.M. Model of autism: Increased ratio of excitation/inhibition in key neural systems. *Genes Brain Behav.* 2003, 2, 255–267. [CrossRef] [PubMed]

8. Lee, E.; Lee, J.; Kim, E. Excitation/Inhibition Imbalance in Animal Models of Autism Spectrum Disorders. *Biol. Psychiatry* 2017, 81, 838–847. [CrossRef] [PubMed]

9. Association, A.P. Diagnostic and Statistical Manual of Mental Disorders, 5th ed.; American Psychiatric Association: Arlington, VA, USA, 2013. [CrossRef]

10. Harris, J.C. The origin and natural history of autism spectrum disorders. *Nat. Neurosci.* 2016, 19, 1390–1391. [CrossRef]

11. Mottron, L.; Bzdok, D. Autism spectrum heterogeneity: Fact or artifact? *Mol. Psychiatry* 2020, 25, 3178–3185. [CrossRef] [PubMed]

12. Homberg, J.R.; Kyzar, E.J.; Scattoni, M.I.; Norton, W.H.; Pittman, J.; Gaikwad, S.; Nguyen, M.; Poudel, M.K.; Ullmann, J.F.; Diamond, D.M.; et al. Genetic and environmental modulation of neurodevelopmental disorders: Translational insights from labs to beds. *Brain Res. Bull.* 2016, 125, 79–91. [CrossRef] [PubMed]

13. Homberg, J.R.; Kyzar, E.J.; Nguyen, M.; Norton, W.H.; Pittman, J.; Poudel, M.K.; Gaikwad, S.; Nakamura, S.; Koshiba, M.; Yamanouchi, H.; et al. Understanding autism and other neurodevelopmental disorders through experimental translational neurobehavioral models. *Neurosci. Biobehav. Rev.* 2016, 65, 292–312. [CrossRef]

14. Hagerman, R.J.; Berry-Kravis, E.; Hazlett, H.C.; Bailey, D.B., Jr.; Moine, H.; Kooy, R.F.; Tassone, F.; Gantois, I.; Sonenberg, N.; Mandel, J.L.; et al. Fragile X syndrome. *Nat. Rev. Dis. Primers* 2017, 3, 17065. [CrossRef] [PubMed]

15. Hunter, J.; Rivero-Arias, O.; Angelov, A.; Kim, E.; Fotheringham, I.; Leal, J. Epidemiology of fragile X syndrome: A systematic review and meta-analysis. *Am. J. Med. Genet A* 2014, 164, 1648–1658. [CrossRef] [PubMed]

16. Riley, C.; Mailick, M.; Berry-Kravis, E.; Bolen, J. The Future of Fragile X Syndrome: CDC Stakeholder Meeting Summary. *Pediatrics* 2017, 139, S147–S152. [CrossRef] [PubMed]

17. Berry-Kravis, E.; Knox, A.; Hervey, C. Targeted treatments for fragile X syndrome. *J. Neurodev. Disord.* 2011, 3, 193–210. [CrossRef]

18. Gross, C.; Berry-Kravis, E.M.; Bassell, G.J. Therapeutic strategies in fragile X syndrome: Dysregulated mGluR signaling and beyond. *Neuropsychopharmacology* 2012, 37, 178–195. [CrossRef] [PubMed]

19. Verkerk, A.J.; Pieretti, M.; Sutcliffe, J.S.; Fu, Y.H.; Kuhl, D.P.; Pizzuti, A.; Reiner, O.; Richards, S.; Victoria, M.F.; Zhang, F.P.; et al. Identification of a gene (FMR-1) containing a CGG repeat coincident with a breakpoint cluster region exhibiting length variation in fragile X syndrome. *Cell* 1991, 65, 905–914. [CrossRef]

20. Penagarikano, O.; Mulle, J.G.; Warren, S.T. The pathophysiology of fragile x syndrome. *Annu. Rev. Genomics Hum. Genet.* 2007, 8, 109–129. [CrossRef]

21. Contractor, A.; Klyachko, V.A.; Portera-Caillau, C. Altered Neuronal and Circuit Excitability in Fragile X Syndrome. *Neuron* 2015, 87, 699–715. [CrossRef]

22. Devys, D.; Lutz, Y.; Rouyer, N.; Bellocq, J.P.; Mandel, J.L. The FMR-1 protein is cytoplasmic, most abundant in neurons and appears normal in carriers of a fragile X premutation. *Nat. Genet.* 1993, 4, 335–340. [CrossRef] [PubMed]

23. Zalfa, F.; Giorgi, M.; Primerano, B.; Moro, A.; Di Penta, A.; Reis, S.; Oostra, B.; Bagni, C. The fragile X syndrome protein FMRP associates with BC1 RNA and regulates the translation of specific mRNAs at synapses. *Cell* 2003, 112, 317–327. [CrossRef] [PubMed]

24. Bassell, G.J.; Warren, S.T. Fragile X syndrome: Loss of local mRNA regulation alters synaptic development and function. *Neuron* 2008, 60, 201–214. [CrossRef] [PubMed]
25. The Dutch-Belgian Fragile X Consortium; Bakker, C.E.; Verheij, C.; Willemsen, R.; van der Helm, R.; Oerlemans, F.; Vermeiy, M.; Bygrave, A.; Hoogeveen, A.; Oostra, B.A.; et al. Fmr1 knockout mice: A model to study fragile X mental retardation. The Dutch-Belgian Fragile X Consortium. *Cell* **1994**, *78*, 23–33.

26. Zhang, Y.Q.; Bailey, A.M.; Matthias, H.J.; Renden, R.B.; Smith, M.A.; Speese, S.D.; Rubin, G.M.; Broadie, K. Drosophila fragile X-related gene regulates the MAP1B homolog Futsch to control synaptic structure and function. *Cell* **2001**, *107*, 591–603. [CrossRef]

27. Tucker, B.; Richards, R.I.; Lardelli, M. Contribution of mGlur and Fmr1 functional pathways to neurite morphogenesis, craniofacial development and fragile X syndrome. *Hum. Mol. Genet.* **2006**, *15*, 3446–3458. [CrossRef] [PubMed]

28. Hamilton, S.M.; Green, J.R.; Veeraragavan, S.; Yuva, L.; McCoy, A.; Wu, Y.; Warren, J.; Little, L.; Ji, D.; Cui, X.; et al. Fmr1 and Ngl3 knockouts rats: Novel tools for investigating autism spectrum disorders. *Behav. Neurosci.* **2014**, *128*, 103–109. [CrossRef]

29. Bontekoe, C.J.; Bakker, C.E.; Nieuwenhuizen, I.M.; van der Linde, H.; Lans, H.; de Lange, D.; Hirst, M.C.; Oostra, B.A. Instability of a (CGG)98 repeat in the Fmr1 promoter. *Hum. Mol. Genet.* **2001**, *10*, 1693–1699. [CrossRef]

30. Van Dam, D.; Errious, V.; Kooy, R.F.; Willemsen, R.; Mientjes, E.; Oostra, B.A.; De Deyn, P.P. Cognitive decline, neuromotor and behavioural disturbances in a mouse model for fragile-X-associated tremor/ataxia syndrome (FXTAS). *Behav. Brain Res.* **2005**, *162*, 233–239. [CrossRef]

31. Brouwer, J.R.; Mientjes, E.J.; Bakker, C.E.; Nieuwenhuizen, I.M.; Severijnen, L.A.; Van der Linde, H.C.; Nelson, D.L.; Oostra, B.A.; Willemsen, R. Elevated Fmr1 mRNA levels and reduced protein expression in a mouse model with an unmodified Fragile X full mutation. *Exp. Cell Res.* **2007**, *313*, 244–253. [CrossRef]

32. Darnell, J.C.; Van Driesche, S.J.; Zhang, C.; McCoy, A.; Wu, Y.; Warren, J.; Little, L.; Ji, D.; Cui, X.; et al. FMRP stalls ribosomal translocation on mRNAs linked to synaptic function and autism. *Cell* **2011**, *146*, 247–261. [CrossRef] [PubMed]

33. Ascano, M., Jr.; Mukherjee, N.; Bandaru, P.; Miller, J.B.; Nusbaum, J.D.; Corcoran, D.L.; Langlois, C.; Munschauer, M.; Dewell, S.; Hafner, M.; et al. FMRP targets distinct mRNA sequence elements to regulate protein expression. *Nature* **2012**, *492*, 382–386. [CrossRef] [PubMed]

34. Blahak, A.L.; Dolren, G.; Bear, M.F. The pathophysiology of fragile X (and what it teaches us about synapses). *Annu. Rev. Neurosci.* **2012**, *35*, 417–443. [CrossRef] [PubMed]

35. Castren, M.L.; Castren, E. BDNF in fragile X syndrome. *Neuropharmacology* **2014**, *76 Pt C*, 729–736. [CrossRef]

36. Bear, M.F.; Huber, K.M.; Warren, S.T. The mGluR theory of fragile X mental retardation. *Trends Neurosci.* **2004**, *27*, 370–377. [CrossRef]

37. Krueger, D.D.; Bear, M.F. Toward fulfilling the promise of molecular medicine in fragile X syndrome. *Annu. Rev. Med.* **2011**, *62*, 411–429. [CrossRef]

38. Scharf, S.H.; Jaeschke, G.; Wettstein, J.G.; Lindemann, L. Metabotropic glutamate receptor 5 as drug target for Fragile X syndrome: A review of associated medical problems. *Pediatrics* **2014**, *134*, 995–1005. [CrossRef] [PubMed]

39. Van der Aa, N.; Kooy, R.F. GABAergic abnormalities in the fragile X syndrome. *Eur. J. Pediatr. Neurol.* **2020**, *24*, 100–104. [CrossRef]

40. Kidd, S.A.; Lachiewicz, A.; Barbouth, D.; Blitz, R.K.; Delahunty, C.; McBrien, D.; Visootsak, J.; Berry-Kravis, E. Fragile X syndrome: A review of associated medical problems. *Pediatrics* **2014**, *134*, 995–1005. [CrossRef] [PubMed]

41. Cellot, G.; Cherubini, E. GABAergic signaling as therapeutic target for autism spectrum disorders. *Front. Pediatr.* **2014**, *2*, 70. [CrossRef] [PubMed]

42. Braat, S.; Kooy, R.F. The GABAA Receptor as a Therapeutic Target for Neurodevelopmental Disorders. *Neuron* **2015**, *86*, 1119–1130. [CrossRef]

43. D’Hulst, C.; Heulens, I.; Van der Aa, N.; Goffin, K.; Koole, M.; Porke, K.; Van De Velde, M.; Rooms, L.; Van Paesschen, W.; Van Esch, H.; et al. Positron Emission Tomography (PET) Quantification of GABAA Receptors in the Brain of Fragile X Patients. *PLoS ONE* **2015**, *10*, e0131486. [CrossRef]

44. Morin-Parent, F.; Champigny, C.; Lacroix, A.; Corbin, F.; Lepage, J.F. Hyperexcitability and impaired intracortical inhibition in patients with fragile-X syndrome. *Transl. Psychiatry* **2019**, *9*, 312. [CrossRef]

45. Chen, L.; Toth, M. Fragile X mice develop sensory hyperreactivity to auditory stimuli. *Neuroscience* **2001**, *103*, 1043–1050. [CrossRef]

46. Gonzalez, D.; Tomasek, M.; Hays, S.; Sridhar, V.; Ammanuel, S.; Chang, C.W.; Pawlowski, K.; Huber, K.M.; Gibson, J.R. Audiogenic Seizures in the Fmr1 Knock-Out Mouse Are Induced by Fmr1 Deletion in Subcortical, VGlut2-Expressing Excitatory Neurons and Require Deletion in the Inferior Colliculus. *J. Neurosci.* **2019**, *39*, 9852–9863. [CrossRef]

47. Rotschafer, S.; Razak, K. Altered auditory processing in a mouse model of fragile X syndrome. *J. Neurosci.* **2013**, *33*, 897–905. [CrossRef] [PubMed]
52. Olmos-Serrano, J.L.; Corbin, J.G.; Burns, M.P. The GABA(A) receptor agonist THIP ameliorates specific behavioral deficits in the mouse model of fragile X syndrome. *Dev. Neurosci.* 2011, 33, 395–403. [CrossRef]

53. Frankland, P.W.; Wang, Y.; Rosner, B.; Shimizu, T.; Balleine, B.W.; Dykens, E.M.; Ornitz, E.M.; Silva, A.J. Sensorimotor gating abnormalities in young males with fragile X syndrome and Fmr1-knockout mice. *Mol. Psychiatry* 2004, 9, 417–425. [CrossRef]

54. de Vrij, F.M.; Leveugla, J.-F.; van der Linde, H.C.; Koekkooi, S.K.; De Zeeuw, C.J.; Nelson, D.L.; Oostra, B.A.; Willemens, R. Rescue of behavioral phenotype and neuronal protrusion morphology in Fmr1 KO mice. *Neurobiol. Dis.* 2008, 31, 127–132. [CrossRef]

55. Kokash, J.; Alderson, E.M.; Reinhard, S.M.; Crawford, C.A.; Binder, D.K.; Ethell, I.M.; Razak, K.A. Genetic reduction of MMP-9 in the Fmr1 KO mouse partially rescues prepulse inhibition of acoustic startle response. *Brain Res.* 2019, 1719, 24–29. [CrossRef]

56. Puts, N.A.; Edden, R.A.; Evans, C.J.; McGlone, F.; McGonigle, D.J. Regionally specific human GABA concentration correlates with tactile discrimination thresholds. *J. Neurosci.* 2011, 31, 16556–16560. [CrossRef]

57. Puts, N.A.J.; Wodka, E.L.; Harris, A.D.; Crocetti, D.; Tommerdahl, M.; Mostofsky, S.H.; Edden, R.A.E. Reduced GABA and altered somatosensory function in children with autism spectrum disorder. *Austin. Res.* 2017, 10, 608–619. [CrossRef] [PubMed]

58. Sapey-Triomphe, L.A.; Lamberton, F.; Sonie, S.; Mattout, J.; Schmitz, C. Tactile hypersensitivity and GABA concentration in the sensorimotor cortex of adults with autism. *Austin. Res.* 2019, 12, 562–575. [CrossRef]

59. Wood, E.T.; Cummings, K.K.; Jung, J.; Patterson, G.; Okada, N.; Guo, J.; O’Neill, J.; Dapretto, M.; Bookheimer, S.Y.; Green, S.A. Sensory over-responsivity is related to GABAergic inhibition in thalamocortical circuits. *Transl. Psychiatry* 2021, 11, 39. [CrossRef] [PubMed]

60. He, C.X.; Cantu, D.A.; Mantri, S.S.; Zeiger, W.A.; Goel, A.; Porteria-Cailliau, C. Tactile Defensiveness and Impaired Adaptation of Neuronal Activity in the Fmr1 Knock-Out Mouse Model of Autism. *J. Neurosci.* 2017, 37, 6475–6487. [CrossRef]

61. Kato, M.; Dobyns, W.B. X-linked lissencephaly with abnormal genitalia as a tangential migration disorder causing intractable epilepsy: Proposal for a new term, “interneuronopathy”. *J. Child Neurol.* 2005, 20, 392–397. [CrossRef] [PubMed]

62. Bonneau, D.; Toutain, A.; Laquerriere, A.; Marret, S.; Saugier-Veber, P.; Barthez, M.A.; Radi, S.; Biran-Mucignat, V.; Rodriguez, D.; Gelot, A. X-linked lissencephaly with absent corpus callosum and ambiguous genitalia (XLAG): Clinical, magnetic resonance imaging, and neuropathological findings. *Ann. Neurol.* 2002, 51, 340–349. [CrossRef]

63. Mullen, S.A.; Scheffer, I.E. Translational research in epilepsy genetics: Sodium channels in man to interneuronopathy in mouse. *Arch. Neurol.* 2009, 66, 21–26. [CrossRef]

64. Catterall, W.A. Dravet Syndrome: A Sodium Channel Interneuronopathy. *Curr. Opin. Physiol.* 2018, 2, 42–50. [CrossRef]

65. Kato, M. A new paradigm for West syndrome based on molecular and cell biology. *Epilepsy Res.* 2006, 70 (Suppl. 1), S87–S95. [CrossRef] [PubMed]

66. Ruggieri, M.; Pavone, P.; Scapagnini, G.; Romeo, L.; Lombardo, I.; Li Volti, G.; Pavone, L.; Cordella, G.; Pavone, L. The aristaless (Arx) gene: One gene for many “interneuronopathies”. *Front. Biosci. (Elite Ed)* 2010, 2, 701–710. [CrossRef]

67. Tremblay, R.; Lee, S.; Rudy, B. GABAergic Interneurons in the Neocortex: From Cellular Properties to Circuits. *Neuron* 2016, 91, 260–292. [CrossRef]

68. Fishell, G.; Kepes, A. Interneuron Types as Attractors and Controllers. *Annu. Rev. Neurosci.* 2020, 43, 1–30. [CrossRef] [PubMed]

69. Lim, L.; Mi, D.; Iturria, A.; Marin, O. Development and Functional Diversification of Cortical Interneurons. *Neurosci. 2018, 100, 294–313. [CrossRef] [PubMed]

70. Hodge, R.D.; Bakken, T.E.; Miller, J.A.; Smith, K.A.; Barkan, E.R.; Richter, T.; Close, J.; Long, B.; Johansen, N.; Penn, O.; et al. Conserved cell types with divergent features in human versus mouse cortex. *Nature* 2019, 573, 61–68. [CrossRef] [PubMed]

71. Huang, Z.J.; Paul, A. The diversity of GABAergic neurons and neural communication elements. *Nat. Rev. Neurosci.* 2019, 20, 563–572. [CrossRef] [PubMed]

72. Gouwens, N.W.; Sorensen, S.A.; Baftizadeh, F.; Budzillo, A.; Lee, B.R.; Jarsky, T.; Alifer, L.; Baker, K.; Barkan, E.; Berry, K.; et al. Integrated Morphoelectric and Transcriptomeric Classification of Cortical GABAergic Cells. *Cell 2020, 183, 935–953.e919. [CrossRef]

73. Melzer, S.; Monyer, H. Diversity and function of corticopetal and corticofugal GABAergic projection neurons. *Nat. Rev. Neurosci.* 2020, 21, 499–515. [CrossRef]

74. Bouliou, C. Numerical data on neocortical neurons in adult rat, with special reference to the GABA population. *Brain Res.* 1993, 609, 284–292. [CrossRef]

75. Jones, E.G. GABAergic neurons and their role in cortical plasticity in primates. *Cereb. Cortex 1993, 3, 361–372. [CrossRef]

76. Marin, O. Interneuron dysfunction in psychiatric disorders. *Nat. Rev. Neurosci.* 2012, 13, 107–120. [CrossRef] [PubMed]

77. Filice, F.; Janickova, L.; Henzi, T.; Bilella, A.; Schwaller, B. The Parvalbumin Hypothesis of Autism Spectrum Disorder. *Front. Cell. Neurosci.* 2020, 14, 57525. [CrossRef] [PubMed]

78. Selby, L.; Zhang, C.; Sun, Q.Q. Major defects in neocortical GABAergic inhibitory circuits in mice lacking the fragile X mental retardation protein. *Neurosci. Lett.* 2007, 422, 227–232. [CrossRef] [PubMed]

79. Wen, T.H.; Afroz, S.; Reinhard, S.M.; Palacios, A.R.; Tapia, K.; Binder, D.K.; Razak, K.A.; Ethell, L.M. Genetic Reduction of Matrix Metalloproteinase-9 Promotes Formation of Perineuronal Nets Around Parvalbumin-Expressing Interneurons and Normalizes Auditory Cortex Responses in Developing Fmr1 Knock-Out Mice. *Cereb. Cortex* 2018, 28, 3951–3964. [CrossRef]

80. Benali, A.; Tripple, J.; Weiler, E.; Mix, A.; Petrasch-Parwez, E.; Girzalsky, W.; Eysel, U.T.; Erdmann, R.; Funke, K. Theta-burst transcranial magnetic stimulation alters cortical inhibition. *J. Neurosci.* 2011, 31, 1193–1203. [CrossRef] [PubMed]
81. Nomura, T.; Musial, T.F.; Marshall, J.J.; Zhu, Y.; Remmers, C.L.; Xu, J.; Nicholson, D.A.; Contractor, A. Delayed Maturation of Fast-Spiking Interneurons Is Rectified by Activation of the TrkB Receptor in the Mouse Model of Fragile X Syndrome. J. Neurosci. 2017, 37, 11298–11310. [CrossRef]

82. Olims-Serrano, J.L.; Paluszkievicz, S.M.; Martin, B.S.; Kaufmann, W.E.; Corbin, J.G.; Huntsman, M.M. Defective GABAergic neurotransmission and pharmacological rescue of neuronal hyperexcitability in the amygdala in a mouse model of fragile X syndrome. J. Neurosci. 2010, 30, 9929–9938. [CrossRef] [PubMed]

83. Antoine, M.W.; Langberg, T.; Schnepel, P.; Feldman, D.E. Increased Excitation-Inhibition Ratio Stabilizes Synapse and Circuit Excitability in Four Autism Mouse Models. Neuron 2010, 101, 648–661.e644. [CrossRef] [PubMed]

84. Berzhanskaya, J.; Phillips, M.A.; Shen, J.; Colonnesi, M.T. Sensory hypo-excitability in a rat model of fetal development in Fragile X Syndrome. Sci. Rep. 2016, 6, 30769. [CrossRef] [PubMed]

85. Franco, L.M.; Okray, Z.; Linneweber, G.A.; Hassan, B.A.; Yaksi, E. Reduced Lateral Inhibition Impairs Olfactory Computations and Behaviors in a Drosophila Model of Fragile X Syndrome. Curr. Biol. 2017, 27, 1111–1123. [CrossRef]

86. Goel, A.; Cantu, D.A.; Guilfoyle, J.; Chaudhari, G.R.; Newadkar, A.; Todisco, B.; de Alba, D.; Kourdougli, N.; Schmitt, L.M.; Pedapati, E.; et al. Impaired perceptual learning in a mouse model of Fragile X syndrome is mediated by parvalbumin neuron dysfunction and is reversible. Nat. Neurosci. 2018, 21, 1404–1411. [CrossRef] [PubMed]

87. Yang, Y.M.; Arsenault, J.; Bah, A.; Krzeminski, M.; Fekete, A.; Chao, O.Y.; Bagni, C.; Contractor, A. Critical period plasticity is disrupted in the barrel cortex. J. Neurosci. 2001, 21, 7879–7893. [CrossRef] [PubMed]

88. Nimchinsky, E.A.; Oberlander, A.M.; Svoboda, K. Abnormal development of dendritic spines in FMR1 knock-out mice. Neuron 2001, 29, 779–789. [CrossRef] [PubMed]

89. Cruz-Martin, A.; Crespo, M.; Portera-Cailliau, C. Delayed stabilization of dendritic spines in fragile X mice. J. Neurosci. 2010, 30, 7793–7803. [CrossRef] [PubMed]

90. Cruz-Martin, A.; Crespo, M.; Portera-Cailliau, C. Glutamate induces the elongation of early dendritic protrusions via mGluRs in Purkinje cells. J. Neurophysiol. 2003, 89, 2412–2419. [CrossRef] [PubMed]

91. Wijetunge, L.S.; Angibaud, J.; Wijetunge, L.S.; Siegel, V.; Beutemans, A.; Contractor, A.; Gillingwater, T.H.; Kind, P.C. Altered excitability in the Purkinje cells of FMR1 knockout mice. J. Neurosci. 2010, 30, 385–395. [CrossRef]

92. He, Q.; Arroyo, E.D.; Smukowski, S.N.; Xu, J.; Piochon, C.; Savas, J.N.; Portera-Cailliau, C.; Contractor, A. Critical period plasticity is disrupted in the barrel cortex of FMR1 knockout mice. Neuron 2010, 65, 385–398. [CrossRef]

93. He, Q.; Arroyo, E.D.; Smukowski, S.N.; Xu, J.; Piochon, C.; Savas, J.N.; Portera-Cailliau, C.; Contractor, A. Critical period inhibition of NKCC1 rectifies synapse plasticity in the somatosensory cortex and restores adult tactile response maps in fragile X mice. Mol. Psychiatry 2019, 24, 1732–1747. [CrossRef]

94. Wijetunge, L.S.; Angibaud, J.; Frick, A.; Kind, P.C.; Nagerl, U.V. Delayed Maturation of the Primary Somatosensory Cortex in a Mouse Model of Fragile X Syndrome. J. Neurosci. 2010, 30, 11293–11304. [CrossRef] [PubMed]

95. Bureau, I.; Shepherd, G.M.; Svoboda, K. Circuit and plasticity defects in the developing somatosensory cortex of FMR1 knockout mice. J. Neurosci. 2008, 28, 5178–5188. [CrossRef]

96. Hensch, T.K. Critical period plasticity in local cortical circuits. Nat. Rev. Neurosci. 2005, 6, 877–888. [CrossRef]

97. Domanski, A.P.; Booker, S.A.; Myllie, D.J.A.; Isaac, J.T.R.; Kind, P.C. Cellular and synaptic phenotypes lead to disrupted information processing in Fmr1 KO mouse layer 4 barrel cortex. Nat. Commun. 2019, 10, 4814. [CrossRef] [PubMed]

98. Gibson, J.R.; Bartley, A.F.; Hays, S.A.; Huber, K.M. Imbalance of neocortical excitation and inhibition and altered UP states reflect network hyperexcitability in the mouse model of fragile X syndrome. J. Neurophysiol. 2008, 100, 2615–2626. [CrossRef]

99. Hensch, T.K. Critical period plasticity in local cortical circuits. Nat. Rev. Neurosci. 2005, 6, 877–888. [CrossRef]

100. Miller, K.D.; Pinto, D.J.; Simon, J.P. Processing in layer 4 of the neocortical circuit: New insights from visual and somatosensory cortex. Curr. Opin. Neurobiol. 2001, 11, 488–497. [CrossRef]
107. Feldmeyer, D.; Qi, G.; Emmenegger, V.; Staiger, J.F. Inhibitory interneurons and their circuit motifs in the many layers of the barrel cortex. *Neuroscience* **2018**, *368*, 132–151. [CrossRef] [PubMed]

108. Goncalves, J.T.; Ansley, J.E.; Golshani, P.; Portera-Cailliau, C. Circuit level defects in the developing neocortex of Fragile X mice. *Nat. Neurosci.* **2013**, *16*, 903–909. [CrossRef]

109. Arnett, M.T.; Herman, D.H.; McGee, A.W. Deficits in tactile learning in a mouse model of fragile X syndrome. *PLoS ONE* **2014**, *9*, e109116. [CrossRef]

110. Zhang, Y.; Bonnan, A.; Bony, G.; Ferezou, I.; Pietropaolo, S.; Ginger, M.; Sans, N.; Rossier, J.; Oostra, B.; LeMasson, G.; et al. Dendritic channelopathies contribute to neocortical and sensory hyperexcitability in Fmr1(-/-) mice. *Nat. Neurosci.* **2014**, *17*, 1701–1709. [CrossRef]

111. Boland, M.J.; Nazor, K.L.; Tran, H.T.; Szucs, A.; Lynch, C.L.; Paredes, R.; Tassone, F.; Sanna, P.P.; Hagerman, R.J.; Loring, J.F. Molecular analyses of neurogenic defects in a human pluripotent stem cell model of fragile X syndrome. *Brain* **2017**, *140*, 582–598. [CrossRef]

112. Vitrac, A.; Cloez-Tayarani, I. Induced pluripotent stem cells as a tool to study brain circuits in autism-related disorders. *Stem Cell Res. Ther.* **2018**, *9*, 226. [CrossRef]

113. Telias, M.; Segal, M.; Ben-Yosef, D. Neural differentiation of Fragile X human Embryonic Stem Cells reveals abnormal patterns of development despite successful neurogenesis. *Dev. Biol.* **2013**, *374*, 32–45. [CrossRef]

114. Telias, M.; Kuznitzov-Yanovsky, L.; Segal, M.; Ben-Yosef, D. Functional Deficiencies in Fragile X Neurons Derived from Human Embryonic Stem Cells. *J. Neurosci.* **2015**, *35*, 15295–15306. [CrossRef]

115. Liu, Y.; Liu, H.; Sauvey, C.; Yao, L.; Zarnowska, E.D.; Zhang, S.C. Direct differentiation of forebrain GABA interneurons from human pluripotent stem cells. *Nat. Protoc.* **2013**, *8*, 1670–1679. [CrossRef]

116. Maroof, A.M.; Keros, S.; Tyson, J.A.; Ying, S.W.; Ganat, Y.M.; Merkle, F.T.; Li, B.; Goulburn, A.; Stanley, E.G.; Elefanty, A.G.; et al. Direct differentiation and functional maturation of cortical interneurons from human embryonic stem cells. *Cell Stem Cell* **2013**, *12*, 559–572. [CrossRef]

117. Nicholas, C.R.; Chen, J.; Tang, Y.; Southwell, D.G.; Chalmers, N.; Vogt, D.; Arnold, C.M.; Chen, Y.J.; Stanley, E.G.; Elefanty, A.G.; et al. Functional maturation of hPSC-derived forebrain interneurons requires an extended timeline and mimics human neural development. *Cell Stem Cell* **2013**, *12*, 573–586. [CrossRef]

118. Kim, T.G.; Yao, R.; Monnell, T.; Cho, J.H.; Vasudevan, A.; Koh, A.; Peeyush, K.T.; Moon, M.; Datta, D.; Bolshakov, V.Y.; et al. Efficient specification of interneurons from human pluripotent stem cells by dorsoventral and rostrocaudal modulation. *Stem Cells* **2014**, *32*, 1789–1804. [CrossRef] [PubMed]

119. Berry-Kravis, E.M.; Hessl, D.; Rathmell, B.; Zarevics, P.; Cherubini, M.; Walton-Bowen, K.; Mu, Y.; Nguyen, D.V.; Gonzalez-Heydrich, J.; Wang, P.P.; et al. Effects of STX209 (arbaclofen) on neurobehavioral function in children and adults with fragile X syndrome: A randomized, controlled, phase 2 trial. *Sci. Transl. Med.* **2012**, *4*, 152ra127. [CrossRef] [PubMed]

120. Berry-Kravis, E.; Hagerman, R.; Vissootsak, J.; Budimirovic, D.; Kaufmann, W.E.; Cherubini, M.; Zarevics, P.; Walton-Bowen, K.; Wang, P.; Bear, M.F.; et al. Arbaclofen in fragile X syndrome: Results of phase 3 trials. *J. Neurodev. Disord.* **2017**, *9*, 3. [CrossRef] [PubMed]

121. Ligsay, A.; Van Dijck, A.; Nguyen, D.V.; Lozano, R.; Chen, Y.; Bickel, E.S.; Hessl, D.; Schneider, A.; Angkustsiri, K.; Tassone, F.; et al. A randomized double-blind, placebo-controlled trial of ganaxolone in children and adolescents with fragile X syndrome. *J. Neurodev. Disord.* **2017**, *9*, 26. [CrossRef] [PubMed]

122. Kaczmarek, L.K.; Zhang, Y. Kv3 Channels: Enablers of Rapid Firing, Neurotransmitter Release, and Neuronal Endurance. *J. Neurophysiol.* **2017**, *118*, 102–112. [CrossRef]

123. Kaczmarek, L.K.; Zhang, Y. Kv3 Channels: Enablers of Rapid Firing, Neurotransmitter Release, and Neuronal Endurance. *J. Neurophysiol.* **2017**, *118*, 102–112. [CrossRef]

124. Brown, M.R.; El-Hassar, L.; Zhang, Y.; Alvaro, G.; Large, C.H.; Kaczmarek, L.K. Physiological modalities of Kv3.1 channels adjust firing patterns of auditory brain stem neurons. *J. Neurophysiol.* **2016**, *116*, 106–121. [CrossRef]

125. El-Hassar, L.; Song, L.; Tan, W.J.T.; Large, C.H.; Alvaro, G.; Santos-Sacchi, J.; Kaczmarek, L.K. Modulators of Kv3 Potassium Channels Rescue the Auditory Function of Fragile X Mice. *J. Neurosci.* **2019**, *39*, 4797–4813. [CrossRef] [PubMed]

126. Cea-Del Rio, C.A.; Nunez-Parra, A.; Freedman, S.M.; Kushner, J.K.; Alexander, A.L.; Restrepo, D.; Huntsman, M.M. Disrupted inhibitory plasticity and homeostasis in Fragile X Syndrome. *Neurobiol. Dis.* **2020**, *142*, 104959. [CrossRef]

127. Svalina, M.N.; Guthman, E.M.; Cea-Del Rio, C.A.; Kushner, J.K.; Baca, S.M.; Restrepo, D.; Huntsman, M.M. Hyperexcitability and Loss of Feedforward Inhibition Contribute to Aberrant Plasticity in the Fmr1KO Amygdala. *eNeuro* **2021**, *8*. [CrossRef] [PubMed]

128. Perrin, S. Preclinical research: Make mouse studies work. *Nature* **2014**, *507*, 423–425. [CrossRef]

129. Begley, C.G.; Ellis, L.M. Drug development: Raise standards for preclinical cancer research. *Nature* **2012**, *483*, 531–533. [CrossRef] [PubMed]

130. Jannot, A.S.; Agoritsas, T.; Gayet-Ageron, A.; Feneberg, T.V. Citation bias favoring statistically significant studies was present in medical research. *J. Clin. Epidemiol.* **2013**, *66*, 296–301. [CrossRef] [PubMed]
132. Mullard, A. Fragile X disappointments upset autism ambitions. Nat. Rev. Drug Discov. 2015, 14, 151–153. [CrossRef] [PubMed]
133. Duyx, B.; Urlings, M.J.E.; Swaen, G.M.H.; Bouter, L.M.; Zeegers, M.P. Scientific citations favor positive results: A systematic review and meta-analysis. J. Clin. Epidemiol. 2017, 88, 92–101. [CrossRef] [PubMed]
134. Mlinaric, A.; Horvat, M.; Supak Smolcic, V. Dealing with the positive publication bias: Why you should really publish your negative results. Biochem. Med. (Zagreb) 2017, 27, 030201. [CrossRef]
135. Murad, M.H.; Chu, H.; Lin, L.; Wang, Z. The effect of publication bias magnitude and direction on the certainty in evidence. BMJ Evid. Based Med. 2018, 23, 84–86. [CrossRef] [PubMed]
136. Chambers, C.D.; Forstmann, B.; Pruszynski, J.A. Science in flux: Registered reports and beyond at the European Journal of Neuroscience. Eur. J. Neurosci 2019, 49, 4–5. [CrossRef]
137. Chambers, C. What’s next for Registered Reports? Nature 2019, 573, 187–189. [CrossRef]