Serotonin Deficiency Increases Context-Dependent Fear Learning Through Modulation of Hippocampal Activity

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Brain serotonin (5-hydroxytryptamine, 5-HT) system dysfunction is implicated in exaggerated fear responses triggering various anxiety-, stress-, and trauma-related disorders. However, the underlying mechanisms are not well understood. Here, we investigated the impact of constitutively inactivated 5-HT synthesis on context-dependent fear learning and extinction using tryptophan hydroxylase 2 (Tph2) knockout mice. Fear conditioning and context-dependent fear memory extinction paradigms were combined with c-Fos imaging and electrophysiological recordings in the dorsal hippocampus (dHip). Tph2 mutant mice, completely devoid of 5-HT synthesis in brain, displayed accelerated fear memory formation and increased locomotor responses to foot shock. Furthermore, recall of context-dependent fear memory was increased. The behavioral responses were associated with increased c-Fos expression in the dHip and resistance to foot shock-induced impairment of hippocampal long-term potentiation (LTP). In conclusion, increased context-dependent fear memory resulting from brain 5-HT deficiency involves dysfunction of the hippocampal circuitry controlling contextual representation of fear-related behavioral responses.

Keywords: tryptophan hydroxylase 2, knockout, fear learning, extinction, long-term potentiation, hippocampus, immediate-early gene, serotonin deficiency

INTRODUCTION

Anxiety disorders are common and result in substantial economic costs to individuals and society (Bereza et al., 2009). The current global prevalence of anxiety disorders is approximately 7.3%, ranging from 5.4 to 10.4% (Baxter et al., 2013). Many individuals meet diagnostic criteria for multiple anxiety disorders and comorbidity with other psychiatric disorders (Kessler et al., 2005;
Miyazaki et al., 2011). A hallmark of anxiety disorders is dysfunctional acquisition and extinction of conditioned fear memories (Grillon, 2002; Milad et al., 2008).

Evidence suggests that serotonin plays an important role in control of anxiety and fear responses (Lesch et al., 1996; Lowry et al., 2005; Maier and Watkins, 2005; Maier et al., 2006; Baratta et al., 2016; Bocchio et al., 2016). Mice with a targeted inactivation of Tph2 have provided insights into the role of 5-HT in the modulation of anxiety-like behaviors. Previous studies of lifelong deficiency of brain 5-HT synthesis are consistent with the hypothesis that the brain serotonergic system plays an important role in control of anxiety-like behaviors (Mosienko et al., 2015), fear learning, and behavioral responses to stress (Gutknecht et al., 2015), effects that might be due to alterations in GABAergic transmission (Jorgensen et al., 2013; Waider et al., 2013). Furthermore, mice with defects in 5-HT system development leading to reduction of 5-HT neurons showed differential anxiety-like behaviors and fear memory (Hendricks et al., 2003; Dai et al., 2008; Schaefer et al., 2009; Kiyasova et al., 2011; Song et al., 2011; Brooks et al., 2014). Indeed, the 5-HT system is thought to play an essential role in the regulation of fear memory in rodents (Graeff and Zangrossi, 2010; Bocchio et al., 2016). Studies in animals demonstrate a direct anatomical connection between the main sources of serotonin in the brain, the brainstem dorsal and median raphe nuclei as well as forebrain limbic structures, such as the medial prefrontal cortex, hippocampus, and amygdala, that control anxiety and fear responses (Maier et al., 2006; Hale and Lowry, 2011; Fernandez et al., 2016; Muzerele et al., 2016). Of particular interest to contextual fear conditioning is the dorsal hippocampus (dHip; Bauer, 2015), which receives serotonergic projections primarily from the median raphe nucleus (Azmithia and Whitaker-Azmitia, 1995; McQuade and Sharp, 1997; Lowry, 2002).

Consistent with this hypothesis, acute administration of selective 5-HT reuptake inhibitors (SSRIs) 60 min before testing results in a decrease in contextual fear expression (Hashimoto et al., 1996; Li et al., 2001; Gravius et al., 2006), while it increases conditioned fear expression in auditory fear conditioning setting (Burghardt et al., 2007). Furthermore, peripheral administration of SSRIs decreases neuronal activity, immediate-early gene expression, and plasticity in the hippocampus (Staubli and Otaky, 1994; Igelstrom and Heyward, 2012; Ravinder et al., 2013). In addition to these effects of serotonergic signaling on fear expression, other studies provide support for a role for multiple 5-HT receptor types in the dHip in conditioned fear memory consolidation (Schmidt et al., 2017).

We previously showed that Tph2 mutant ($Tph2^{−/−}$) mice display enhanced acquisition of conditioned fear and escape-oriented behavior in response to aversive foot shock, in association with altered basolateral amygdala function (Waider et al., 2017). Here, we investigated the impact of a lifelong absence of brain 5-HT synthesis on the contextual domain of fear learning, using fear conditioning combined with an extinction paradigm, functional immunohistochemistry, and electrophysiological recordings within the hippocampal formation.

**MATERIALS AND METHODS**

**Animals**

Adult male $Tph2^{+/+}$, $Tph2^{+/-}$, and $Tph2^{−/−}$ mice on a mixed Sv129/C57BL/6N genetic background (Gutknecht et al., 2012), 2–5 months of age, were housed individually in a controlled environment (12 h/12 h light/dark cycle, light phase 7 am–7 pm, 21 ± 0.5°C room temperature, 50 ± 5% humidity) with food and water ad libitum. Mice were allowed to acclimate for 1 week before being subjected to fear conditioning experiments. Tests were performed during the light phase between 10:00 and 15:00. All experiments were performed in accordance with the European Parliament and Council Directive (2010/63/EU) and were approved by local authorities (55.2-2531.01-57/12) and (IT: 938/2017-PR).

**Fear Conditioning and Contextual Fear Extinction Training**

In order to assess context-dependent fear memory and extinction, $Tph2^{+/-}$, $Tph2^{+/-}$, and $Tph2^{−/−}$ mice ($n = 8–9$ per genotype) were exposed to a fear conditioning protocol as previously described (Waider et al., 2017) and subsequently subjected to a context-dependent extinction protocol (EXT; **Figure 1**). Briefly, on day 0, mice were placed by a blinded operator in randomized order into the fear conditioning test box (TSE Systems, Homburg, Germany), which was comprised of a transparent Perspex arena ($23 \times 23 \times 35$ cm) on a stainless steel foot shock grid (floor bars 4 mm diameter, distance rod center to rod center 8.9 mm) that was connected to a shocker-scrambler unit for delivering foot shocks of defined duration and intensity (Raab et al., 2018). The arena was placed inside in a square-shaped base frame (outer size: $31 \times 31$ cm) with integrated animal detection sensors ($XY$ and $Z$ axes featuring 16 sensors mounted 14 mm apart). All sensors were scanned with a sampling rate of up to 100 Hz to monitor the animal’s position and movement at high spatial and temporal resolution. The test box was operated in a sound-attenuating housing ($52 \times 52 \times 65$ cm) featuring a loudspeaker and two lamps in the ceiling for software-controlled application of acoustic stimuli and continuous house-light illumination (set to 100 lux in all testing phases), respectively.

Mice were allowed to freely explore for 2 min the test box before receiving three pairings of a 20 s tone cue (80 dB, 4 kHz), co-terminating with a 2 s foot shock ($FS$, 0.6 mA) at an inter-trial interval (ITI) of 1 min. Mice were removed from the chamber 2 min after the last tone-shock pairing. Starting 24 h after conditioning, mice were tested for recall and extinction of context-dependent fear on two consecutive days (day 1 and 2). Each extinction session consisted of re-exposing the animals to the conditioning context for 800 s without negative reinforcement.

Freezing was defined as no light-beam interruption for at least 2 s and expressed as percentage of time relative to total session duration (Rivero et al., 2015; Raab et al., 2018). Additionally, other indices of locomotor activity and exploratory behavior (i.e., distance moved, mean and maximum velocity, activity vs.
inactivity, rearing) were measured continuously by the light-beam detection system. Inactivity was defined as the percentage of time the animal’s speed fell below a threshold of 2 cm/s. Using the same fear conditioning system (TSE FCS 303410 series), Misane et al. (2005) have shown that there is a close linear correlation between operator-scored freezing (using a time-sampling procedure in which the animal was instantly scored as either freezing or active every 10 s) and computer-derived inactivity (activity threshold 1 cm/s) both in context- and tone-dependent memory tests. Moreover, Toth et al. (2012) have shown that there is no difference between hand-scored freezing, TSE-determined freezing (defined as no light-beam interruption for at least 3 s), and Noldus Ethovision-determined inactivity.

To determine acquisition of the conditioned fear response, freezing was measured continuously throughout the training session. The maximum movement velocity was taken as an indicator of the animal’s unconditioned reactivity to foot shock (Waider et al., 2017). During extinction training, freezing was continuously recorded in 20 s time-bins and, except for the first 20 s, averaged into blocks of 1 min to assess recall and extinction of contextual fear memory.

C-Fos and Parvalbumin Immunostaining in the Dorsal Hippocampus

Two hours after extinction on day 2, the brains of the mice that underwent context-dependent extinction training (EXT) were prepared for immunostaining of parvalbumin (PV) and c-Fos as previously described (Waider et al., 2017). The brain sections were analyzed and compared with the brains of mice that were either left undisturbed in the home cage (CTRL), exposed to the fear conditioning context and tone without foot shocks (FS-), or underwent fear conditioning with tone-signaled foot shocks (FS+) as previously described (Waider et al., 2017; Figure 1). In brief, serially cut 30 μm-thick cryostat sections were used for immunofluorescent stainings. Primary antibodies used were mouse anti-PV (1:200; Swant, Marly, Switzerland) and rabbit anti-c-Fos (1:400; Santa Cruz Biotechnology, Dallas, TX, United States). Sections were incubated in 1:400 diluted secondary antibodies, goat anti-mouse 488 and goat anti-rabbit 555 (Invitrogen, Carlsbad, CA, United States). Pictures were acquired with a motorized inverted system epifluorescence microscope IX81 (Olympus, Tokyo, Japan). Pictures were taken with 20× objective in x-y directions. Images were then processed using CellSense (Olympus, Tokyo, Japan), and corrected for contrast and brightness using ImageJ v2.0.0 (Schindelin et al., 2012). Three to five sections from −1.06 mm bregma to −2.06 mm bregma of the dHip, spaced 180 μm apart, were delineated with contours according to a stereotactic atlas of the mouse brain (Franklin and Paxinos, 1997). Immunoreactive (ir) cells were counted as c-Fos-ir only when the nucleus, according to DAPI (4′, 6-diamidino-2-phenylindole) counterstaining, showed complete fluorescent signal. When the PV signal superimposed or surrounded the cell nucleus, it was counted as c-Fos/PV-ir. The sum of counted cells of all sections per mouse were divided by the total contour area to calculate the cell densities per region of interest.

Electrophysiology

To investigate the impact of 5-HT deficiency on hippocampal plasticity under basal conditions and after contextual fear conditioning with unsignaled foot shocks, an independent cohort of mice was either left undisturbed (naive controls) or subjected to a foot shock procedure according to Dai et al. (2008). Mice were placed in the box and allowed to freely explore for 2 min before receiving five foot shocks (0.5 mA, 2 s) with ITI of 2 min. Two minutes after the last foot shock, mice were
FIGURE 2 | Increased context-dependent fear memory recall but effective contextual extinction learning due to 5-HT deficiency. (A) Time-course analysis of the conditioned fear response of Tph2+/+, Tph2+/-, and Tph2−/− mice during acquisition training showing relative freezing levels before the first pairing of a tone cue with a foot shock (pre-FS), during tone presentations and inter-trial intervals (ITI), and after the last foot shock (post-FS). (B) Unconditioned foot shock reactivity, measured as maximum movement velocity before the first pairing of a tone cue with a foot shock (pre-FS) and following the foot shock (3 × FS). (C) Time-course analysis of the freezing response during contextual fear memory recall and extinction training on day 1 and 2 after conditioning. (D) Average freezing scores in the two extinction sessions. (E) Time-course analysis of the inactivity level during acquisition training showing relative inactivity levels. (F) Positive relationship between freezing scores obtained after the last foot shock (post-FS) in the conditioning session and inactivity scores. Data are shown as means + or – SEM. *p < 0.05, **p < 0.01 and ***p < 0.001.
removed from the chamber and slices were prepared. Field excitatory postsynaptic potentials (fEPSPs) in the CA1 region of the dHip were recorded in transversal slices as previously described (Morini et al., 2011; Mlinar et al., 2015). LTP was induced by theta burst stimulation (TBS) comprised of a single train of 5 bursts of 5 stimuli (100 Hz intra-burst frequency, 5 Hz burst frequency). Stimulation intensity for baseline measurement and LTP induction was set to evoke fEPSP corresponding to 35–40% of the maximal response. Typically, more than one slice was used per mouse and the results of all determinations per genotype and treatment are shown and analyzed in order to account for the overall variability of LTP responses in the different genotypes. Mean values from replicates (2–4) in the same animal were used for genotype x treatment statistical analysis reported in results.

**Statistical Analyses**

Data were analyzed using IBM SPSS Statistics 21 (IBM Corp., Armonk, NY, United States) or GraphPad Prism version 6.00 (GraphPad Software, San Diego, CA, United States). Behavioral data were analyzed by two- or three-way mixed analysis of variance (ANOVA) with genotype as the between-subjects factor and with time and day as within-subjects factors. When appropriate, degrees of freedom were corrected using Greenhouse-Geisser estimates of sphericity.

An ANCOVA was performed in addition to the regular ANOVA to adjust for the effect of post-shock freezing (=covariate) on subsequent freezing during extinction sessions. Immunohistochemical and electrophysiological data were analyzed by two-way ANOVA with group and genotype as between-subjects factors. Unless otherwise indicated, Bonferroni post hoc tests were performed to evaluate significant main effects or simple effects following a significant interaction. Correlations between behavioral parameters and c-Fos expression were calculated using the Pearson correlation coefficient.

Results are presented as mean ± SEM unless stated otherwise. The significance level was set at \( p < 0.05 \) and \( p < 0.1 \) was highlighted as approaching statistical significance.

**RESULTS**

**Increased Acquisition of Fear Conditioning in Tph2\(^{-/-}\) Mice**

Two-way mixed ANOVA for freezing during fear acquisition training (Figure 2A) revealed a significant main effect of time \( F(2,8,64.4) = 17.54, p < 0.001 \) and genotype \( F(2,23) = 7.70, p = 0.003 \), as well as a time x genotype interaction \( F(5,6,64.4) = 2.73, p = 0.023 \). Post hoc tests showed that freezing levels were very low before the first tone-shock pairing and gradually increased thereafter over the course of training. However, Tph2\(^{-/-}\) mice acquired the conditioned fear response more rapidly than Tph2\(^{+/-}\) and Tph2\(^{+/+}\) mice, as evidenced by significantly elevated freezing levels Tph2\(^{-/-}\) mice from the second tone presentation onward. Furthermore, the reactivity to foot shock, as measured by the maximum movement velocity (Vmax), was significantly increased in Tph2\(^{-/-}\) compared to Tph2\(^{+/-}\) and Tph2\(^{+/+}\) mice [genotype effect: \( F(2,23) = 9.63, p < 0.001 \); Figure 2B].

Moreover, comparison of baseline (pre-FS) Vmax and foot shock-induced Vmax revealed a significant increase in all three genotypes (Tph2\(^{+/-}\): \( p < 0.01 \), Tph2\(^{+/-}\): \( p < 0.001 \) and Tph2\(^{-/-}\): \( p < 0.0001 \), thereby demonstrating that Tph2\(^{+/-}\) and Tph2\(^{-/-}\) mice showed a clear response to FS, which was further exaggerated in Tph2\(^{-/-}\) mice [phase x genotype interaction: \( F(2,23) = 8.223, p = 0.002 \)].

Besides freezing, we analyzed inactivity using an activity threshold of 2 cm/s. Baseline (pre-FS) inactivity (immobility/resting: \( \sim 20–25\% \)) did not significantly differ between genotypes \( (p = 0.861; \text{Figure 2E}) \). Inactivity increased linearly across tone-FS trials in all three genotypes [main effect of phase: \( F(4,2,96.2) = 55.73, p < 0.0001 \) and reached maximum levels after three tone-FS pairings (\( \sim 61\% \) in Tph2\(^{+/-}\) and Tph2\(^{+/-}\), \( \sim 76\% \) in Tph2\(^{-/-}\) mice). Similar to freezing, Tph2\(^{-/-}\) mice were significantly less active than Tph2\(^{+/-}\) and Tph2\(^{+/+}\) mice after the 2nd and 3rd tone-FS pairing [phase x genotype interaction: \( F(8,4,96.2) = 2.27, p = 0.026 \)]. Moreover, there was a close linear correlation between post-shock freezing and post-shock inactivity \( (r = 0.926, p < 0.0001) \) with similar steepness of regression slopes in Tph2\(^{+/-}\), Tph2\(^{+/-}\) and Tph2\(^{-/-}\) mice (Figure 2F).

**Increased Recall but Effective Extinction of Contextual Fear Memory in Tph2\(^{-/-}\) Mice**

Analysis of freezing during extinction training on day 1 and 2 after fear conditioning revealed a significant time-bin x genotype \( [F(9,5,149.5) = 2.02, p = 0.041; \text{Figure 2C}] \) and day x genotype \( [F(2,299) = 5.29, p = 0.013; \text{Figure 2D}] \) interaction, but no genotype main effect \( [F(2,23) = 2.09, p = 0.147] \). Post hoc tests showed that, within sessions, Tph2\(^{-/-}\) mice promptly froze upon placement into the conditioned context, while freezing onset was slightly delayed in Tph2\(^{+/-}\) and Tph2\(^{+/+}\) mice. Moreover, freezing levels of Tph2\(^{-/-}\) mice remained elevated during the first minutes of testing but were indistinguishable from the other genotypes at the end of the session due to a steeper decline of the freezing response in Tph2\(^{-/-}\) compared to Tph2\(^{+/-}\) and Tph2\(^{+/+}\) mice (Figure 2C). Accordingly, Tph2\(^{-/-}\) mice showed a significantly stronger decrease of freezing than Tph2\(^{+/-}\) and Tph2\(^{+/+}\) mice across the two extinction sessions (Figure 2D).

Since Tph2\(^{-/-}\) mice displayed significantly enhanced acquisition of conditioned fear and post-shock freezing levels in the conditioning session, we performed an analysis of covariance to control for differences in post-shock freezing levels. Three-way mixed ANCOVA confirmed a significant effect of post-shock freezing \( [F(1,2,22) = 12.56, p = 0.002] \) as well as a time-bin x genotype \( [F(9,9,138.6) = 2.46, p = 0.011] \) and day x genotype \( [F(2,286) = 2.73, p = 0.087] \) interaction on the freezing scores obtained during extinction training. Post hoc tests again showed that Tph2\(^{-/-}\) mice froze significantly more than Tph2\(^{+/-}\) and Tph2\(^{+/+}\) mice during the first 20 s of context re-exposure. However, freezing levels were indistinguishable among genotypes thereafter. Furthermore, Tph2\(^{-/-}\) mice
Increased c-Fos Activation Due to Fear Conditioning in Tph2<sup>−/−</sup> Mice

Because of the strong effect of 5-HT deficiency on post-shock freezing, an indicator of short-term memory for contextual fear (Fanselow, 1980), we analyzed c-Fos activation in the dHip and its subregions, dentate gyrus (DG), cornu ammonis area 1 (CA1), and cornu ammonis area 3 (CA3) of Tph2<sup>−/−</sup>, Tph2<sup>+/−</sup> and Tph2<sup>+/+</sup> mice that were able to extinguish the conditioned fear more efficiently than Tph2<sup>+/−</sup> and Tph2<sup>+/+</sup> mice, as evidenced by a stronger decline of the freezing response both within and across the two extinction sessions.

Taken together, these results indicate that 5-HT deficiency due to Tph2 inactivation augmented context-dependent fear memory recall and facilitated both intra- and intersession extinction learning.

**Increased c-Fos Activation Due to Fear Conditioning in Tph2<sup>−/−</sup> Mice**

An increased density of c-Fos-ir cells was found in the dHip, especially in the granule cell layer and hilus of the DG (Figure 3A) of FS+ Tph2<sup>−/−</sup> mice. Quantification confirmed a significant genotype x group interaction for the density of c-Fos-ir cells in the total dHip \([F(6,51) = 3.05, p = 0.0126; \text{Figure 3B}]\), in the DG \([F(6,51) = 2.31, p = 0.0477; \text{Figure 3C}]\), and in the CA1 area \([F(6,51) = 2.59, p = 0.0288; \text{Figure 3D}]\). Post hoc analyses showed that the number of c-Fos-ir cells was increased in the total dHip, DG and CA1 of Tph2<sup>−/−</sup> after fear conditioning relative to respective Tph2<sup>+/−</sup> and Tph2<sup>+/+</sup> FS+ controls as well as to CTRL, FS−, and EXT Tph2<sup>−/−</sup> mice \((p < 0.01)\). A similar activation
pattern was observed in the CA3 region, although only a group main effect was detected \([F_{(2,53)} = 9.59, p < 0.0001; \text{Figure 3E}]\), with a significantly increased density of c-Fos-ir cells in FS+ animals compared to CTRL, FS−, and EXT mice (all \(p < 0.001\)).

Correlation analysis of the conditioned fear response (post-shock freezing) with the c-Fos-ir cell density in FS− and FS+ animals detected a strong positive relationship specifically in \(Tph2^{+/−}\) mice \((r ≥ 0.7501, p ≤ 0.0321; \text{Table 1})\). A similar but less exclusive correlation pattern was observed for the density of c-Fos-ir cells with the unconditional response to FS (shock reactivity). Taken together, these data indicate that fear conditioning in \(Tph2^{−/−}\) mice increased hippocampal activity, while context-dependent extinction training of fear memory normalized hippocampal c-Fos expression within the dHip of \(Tph2^{−/−}\) mice.

**Foot Shock Reduces C-Fos Activation of PV-ir Cells in 5-HT-Deficient Mice**

Since PV-ir neurons in the hippocampus were previously shown to be involved in contextual memory \((Donato et al., 2013)\), we analyzed c-Fos-ir neurons in the dHip by double-immunofluorescent staining with PV, a marker of a subset of inhibitory GABAergic interneurons \((Gulyas et al., 1999)\). Because c-Fos-ir densities did not differ between CTRL and FS− animals but were highly increased in fear-conditioned FS+ \(Tph2^{−/−}\) mice, we focused our further analyses on the FS+ group compared to the CTRL group \((\text{Figure 4})\). PV-ir cells were predominantly found in the CA3 and CA1 regions of the dHip, specifically in the stratum lacunosum, pyramidal layer, and stratum oriens with a high density of c-Fos/PV double-ir neurons in the CA1 region \((\text{Figures 4B,E})\). In the total dHip, ANOVA detected an almost significant genotype × group interaction \([F_{(2,24)} = 3.14, p = 0.06; \text{Figure 4C}]\). PV-ir neurons in FS+ \(Tph2^{−/−}\) mice showed reduced c-Fos-ir cells relative to CTRL \(Tph2^{−/−}\) mice \((p = 0.053)\), while FS+ \(Tph2^{+/+}\) and \(Tph2^{+/−}\) mice displayed no alterations compared to respective CTRL mice. Within the FS+ group, \(Tph2^{+/−}\) mice showed the highest c-Fos/PV-ir density compared to \(Tph2^{+/+}\) \((p = 0.05)\) and \(Tph2^{−/−}\) \((p = 0.009)\) mice.

In CA1, a similar but significant genotype × group interaction was detected for the density of c-Fos/PV double-ir neurons \([F_{(2,24)} = 4.77, p = 0.018; \text{Figure 4E}]\). Post hoc tests revealed that CTRL \(Tph2^{−/−}\) mice showed increased c-Fos/PV double-ir cell densities relative to CTRL \(Tph2^{+/+}\) \((p = 0.047)\). However, FS+ \(Tph2^{−/−}\) mice showed reduced c-Fos/PV double-ir cell densities relative to CTRL \(Tph2^{−/−}\) as well as FS+ \(Tph2^{+/+}\) \((p < 0.05)\) and FS+ \(Tph2^{+/−}\) \((p < 0.1)\) mice. Neither differences among genotypes nor among CTRL and FS+ groups were found for the density of PV-ir neurons in dHip \((\text{Figure 4D})\) and CA1 \((\text{Figure 4F})\). Altogether these data indicate increased recruitment of PV neurons in the dorsal CA1 region of CTRL \(Tph2^{−/−}\) mice, which is absent after fear conditioning.

**LTP Impairment by Inescapable Foot Shock Is Absent in Tph2-Deficient Mice**

In the CA1 region, plasticity is modulated by endogenous 5-HT \((Mlinar et al., 2015)\). Here, we investigated the impact of 5-HT deficiency on TBS-induced LTP of fEPSP in the hippocampal CA1 region of \(Tph2\)-deficient mice \((\text{Figure 5C})\). Similar LTP of fEPSP responses were found across genotypes in naïve mice \((\text{Figure 5A, upper panels})\), indicating that basic mechanisms underlying LTP are preserved in the lifelong absence of 5-HT. Since the 5-HT system has been implicated in foot shock-induced impairment of LTP \((Dai et al., 2008)\), we compared LTP in slices obtained from animals exposed to repeated foot shock stress \((\text{Figure 5A, lower panels, B,D})\). Two-way ANOVA revealed a significant genotype × group interaction \([F_{(2,30)} = 9.37 p = 0.0484; \text{Figure 5D}]\). Post hoc analysis confirmed decreased LTP in \(Tph2^{+/+}\) \((p = 0.0011)\) and \(Tph2^{+/−}\) \((p = 0.002)\) mice after foot shock relative to naïve controls, an effect that was absent in \(Tph2^{−/−}\) mice \((\text{Figures 5A,B,D})\) indicating that presence of 5-HT during foot shock is required for the stress-induced impairment of LTP.

**DISCUSSION**

We previously showed that lifelong absence in brain 5-HT synthesis following constitutive \(Tph2\) inactivation in mice enhanced acquisition of conditioned fear and promoted unconditioned escape responses to electric foot shock \((Waider et al., 2017)\). Here, we were able to replicate and extend these findings, which indicate that

![Table 1: Pearson’s correlation of shock reactivity and post-shock freezing with c-Fos density in the dorsal hippocampus and its subregions of FS− and FS+ animals.](image-url)
FIGURE 4 | Lifelong brain 5-HT synthesis deficiency prevents activation of parvalbumin-ir neurons in fear conditioning. (A) Anti-parvalbumin (PV) (green) and c-Fos (red) immunofluorescent staining with DAPI in the dorsal hippocampus (dHip) of a fear conditioned Tph2+/− mouse. (B) Magnified with magnified area of cornu ammonis area 1 (CA1). Density of c-Fos/PV-ir cells of home cage control (CTRL) and fear-conditioned (FS+) Tph2+/−, Tph2+/−, and Tph2−/− mice in (C) dHip and (E) CA1. PV-ir cell density of CTRL and FS+ Tph2+/−, Tph2+/−, and Tph2−/− mice in (D) dHip and (F) CA1. Data are shown as means + SEM. *p < 0.05; **p < 0.01. Two-way ANOVA followed by Fisher’s LSD post hoc test. Scale bar: 100 µm.

constitutive Tph2 inactivation results in faster acquisition of conditioned fear and increased escape-like behavior in response to foot shock, as well as enhanced contextual representation of the fear memory, but functional extinction of the context. By combining fear conditioning with immediate-early gene expression, our data emphasize that lifelong 5-HT deficiency renders the dHip hyperexcitable in fear conditioning.

The dHip encodes the contextual component of fear learning during early acquisition phases (Fanselow and Dong, 2010; Maren et al., 2013). Furthermore, direct glutamatergic projections from the BLA to the dHip have been shown to be both necessary and sufficient for repeated long-term foot shock-mediated memory impairments (Rai et al., 2015), while neurons in the BLA become activated during hippocampal theta network activity or optogenetic stimulation of CA1 pyramidal neurons (Bienvenu et al., 2012; Bazelot et al., 2015).

Because compensation through a 5-HT-dependent mechanism are absent in Tph2−/− mice (Gutknecht et al., 2012; Waider et al., 2017), resilience to foot shock-mediated memory impairments may be derived directly from absence of 5-HT signaling in the dHip. Indeed, inescapable foot shock and novel aversive contexts were reported to impair contextual memory and the induction of LTP in the hippocampal CA1 region (Foy et al., 1987; Sacchetti et al., 2002; Dai et al., 2008).

Moreover, it has long been suggested that 5-HT counteracts the consolidation of stressful memories, presumably mediated by 5-HT1A receptors in the dHip, which then may lead to tolerance with chronic aversive events (Graeff et al., 1996). In contrast to Lmx1b cKO mice, which lack raphe 5-HT neurons (Dai et al., 2008), 5-HT neurons in Tph2−/− mice are unable to synthesize 5-HT, but are functionally preserved (Gutknecht et al., 2012; Montalbano et al., 2015).

Here, we chose a consolidated foot shock protocol (Dai et al., 2008) to investigate whether the absence of 5-HT influences hippocampal LTP formation to allow direct comparison of our data with those obtained using a different animal model of 5-HT system impairment. Although the protocol applied for the LTP and the behavioral experiments differed and, therefore, quantitative correlation of plasticity impairment with the behavioral effects was not possible, our results demonstrate that 5-HT is required for foot shock-induced impairment of LTP. Indeed, 5-HT has been shown to shift hippocampal activity along the longitudinal axis toward the ventral part (Mlinar and Corradetti, 2018). Thus, an absence of this mechanism in Tph2−/− mice may result in an overactivation of the hippocampal circuitry, manifested as enhanced c-Fos immunostaining and overrepresentation of the contextual
FIGURE 5 | Lack of 5-HT impedes foot shock-induced reduction in hippocampal long-term potentiation (LTP). (A) Time-course of LTP in slices obtained from naïve and foot shock-treated mice (Tph2+/+, Tph2+/−, and Tph2−/− mice; naïve: n = 8,6,5; foot shock: n = 5,6,6). Mean ± SD of fEPSP slope values normalized to the baseline mean from 5 to 0 min before theta burst stimulation (TBS). Numbers (n) in graphs indicate slices tested, including replicates per mouse. (B) Representative traces of recordings foot shock-treated Tph2+/+, Tph2+/−, and Tph2−/− mice before and after TBS. (C) Positioning of stimulation electrode (S) on Schaffer collaterals and recording electrode (R) in CA1. The dashed line indicates a cut made between cornu ammonis area 1 (CA1) and cornu ammonis area 3 (CA3) to prevent recurrent propagation of action potentials. (D) LTP is expressed as the percent change measured 40–45 min after TBS in respect to baseline between naïve and foot shock-treated Tph2+/+, Tph2+/−, and Tph2−/− mice. **p < 0.01. Two-way ANOVA, followed by Sidak’s post hoc test.
component in fear learning. In line with this, a mouse model lacking hippocampal serotonergic input showed alterations in contextual fear memory with no differences in LTP formation under baseline conditions (Fernandez et al., 2017).

However, recall of context-dependent aversive memory is exaggerated in Tph2+/− mice, supporting the view that 5-HT mediates inhibition of context-dependent aversive memories through inhibition of LTP, while under non-aversive conditions, increased endogenous 5-HT release facilitated LTP induction in the CA1 region, and may underlie the in vivo positive effects of augmented 5-HT tone on cognitive performance (Mlinar et al., 2015).

In this respect, it may be of interest in a follow up study to analyze the response of the ventral hippocampus, which is involved in emotional fear processing (Fanselow and Dong, 2010), and is modulated differentially by 5-HT along its longitudinal axis (Mlinar and Corradetti, 2018).

Formation of context-dependent memories requires changes in the expression of calcium-binding proteins of GABAergic interneurons, including hippocampal PV-ir cells (Bienvenu et al., 2012; Donato et al., 2013, 2015). In line with previous studies, 5-HT has been shown to modulate PV-specific neurons in the hippocampus (Gulyas et al., 1999). Our results hint toward an involvement of CA1 PV-ir neurons in context-dependent fear conditioning, which is prevented in Tph2+/− mice, although activity of the hippocampus is generally increased. Especially in Tph2+/− mice, fear conditioning recruited the highest number of PV-ir neurons. Thus, it seems that 5-HT-dependent activation of PV neurons may protect the dHip from overactivation. Furthermore, the increased hippocampal activity in Tph2+/− mice due to fear conditioning may prevent an increased response of the BLA, as observed in Tph2+/− mice, to inhibit flight or panic responses (Waider et al., 2017). Thus, further studies are required to investigate the differential role of 5-HT on PV neuron-dependent theta synchronization in Hip and BLA (Aamilhon et al., 2015).

Increased active coping in novel, aversive, and inescapable situations is mediated through an amygdala-ventrolateral periaqueductal gray (vlPAG) circuit, mediated through the dorsolateral PAG (Hale et al., 2012; Toyote et al., 2016). 5-HT neurons of the vlPAG inhibit panic-like responses mediated by dorsolateral PAG neurons (Pobbe et al., 2011; Spannuth et al., 2011). Indeed, Tph2+/− mice show increased post-shock freezing and increased flight responses, although the BLA did not respond to foot shock with an altered activity compared to Tph2+/+ mice (Waider et al., 2017). Thus, the increased hippocampal response in mice completely devoid of 5-HT-mediated regulatory mechanisms may influence the PAG in its role to translate inputs from the amygdala into an appropriate behavioral response, resulting in increased freezing and flight behaviors.

In conclusion, exaggerated context-dependent fear memory and shock reactivity resulting from brain 5-HT deficiency likely involves dysfunction of the raphe-hippocampal innervation controlling fear-related behavioral responses and is presumably due to the failure of 5-HT receptor-mediated inhibition of hippocampal circuitry. Furthermore, our data indicate largely effective extinction learning, during repetitive context exposure without negative reinforcement and without 5-HT functioning. Thus, context-dependent extinction training may represent a strategy to adapt behavioral therapy for patients suffering from 5-HT system dysfunction associated with anxiety-, stress-, and trauma-related disorders.

AUTHOR CONTRIBUTIONS

JW, RC, ASB, and KPL designed and supervised the study. JW, SP, BM, AM, FB, BA, ET, RK, CT, NA, ES, and ASB performed and analyzed the experiments. JW, SP, BM, RC, CL, and KPL interpreted the results and wrote the manuscript.

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REFERENCES

Aamilhon, B., Huh, C. Y., Manseau, F., Ducharme, G., Nichol, H., Adamantidis, A., et al. (2015). Parvalbumin interneurons of hippocampus tune population activity at theta frequency. *Neuron* 86, 1277–1289. doi: 10.1016/j.neuron.2015.05.027

Azmitia, E., and Whitaker-Azmitia, P. (1995). “Anatomy, cell biology, and plasticity of the serotonergic system. Neuropsychopharmacological implications for the actions of psychotropic drugs,” in *Psychopharmacology: the Fourth Generation of Progress*, eds F. E. Bloom and D. J. Kupfer (New York, NY: Raven Press), 443–449.

Baratta, M. V., Kodandaramaiah, S. B., Monahan, P. E., Yao, J., Weber, M. D., Lin, P. A., et al. (2016). Stress enables reinforcement-elicited serotonergic consolidation of fear memory. *Biol. Psychiatry* 79, 814–822. doi: 10.1016/j.biopsych.2015.06.025

Bauer, E. P. (2015). Serotonin in fear conditioning processes. *Behav. Brain Res.* 277, 68–77. doi: 10.1016/j.bbr.2014.07.028

Baxter, A. J., Scott, K. M., Vos, T., and Whiteford, H. A. (2013). Global prevalence of anxiety disorders: a systematic review and meta-regression. *Psychol. Med.* 43, 897–910. doi: 10.1017/S003329171200147X

Bazelon, M., Bocchio, M., Kasugai, Y., Fischer, D., Dodson, P. D., Ferraguti, F., et al. (2015). Hippocampal theta input to the amygdala shapes feedforward inhibition to gate heterosynaptic plasticity. *Neuron* 87, 1290–1303. doi: 10.1016/j.neuron.2015.08.024

Bereza, B. G., Machado, M., and Einarson, T. R. (2009). Systematic review and quality assessment of economic evaluations and quality-of-life studies related to...
generalized anxiety disorder. Clin. Ther. 31, 1279–1308. doi: 10.1016/j.clinthera.2009.06.004
Bienvenu, T. C., Busti, D., Magill, P. J., Ferraguti, F., and Capogna, M. (2012). Cell-type-specific recruitment of amydala interneurons to hippocampal theta rhythm and anxious stimuli in vivo. Neuron 74, 1059–1074. doi: 10.1016/j.neuron.2012.04.022
Bocchio, M., McHugh, S. B., Bannerman, D. M., Sharp, T., and Capogna, M. (2016). Serotonin, amygdala and fear: assembling the puzzle. Front. Neural Circuits 10:24. doi: 10.3389/fncir.2016.00024
Brooks, L. R., Enix, C. L., Rich, S. C., Magno, J. A., Lowry, C. A., and Tsai, P. S. (2014). Fibroblast growth factor deficiencies impact anxiety-like behavior and the serotonergic system. Behav. Brain Res. 246, 74–81. doi: 10.1016/j.bbr.2014.01.053
Burghardt, N. S., Bush, D. E. A., McEwen, B. S., and LeDoux, J. E. (2007). Acute selective serotonin reuptake inhibitors increase conditioned fear expression: blockade with a 5-HT(2C) receptor antagonist. Biol. Psychiatry 62, 1111–1118. doi: 10.1016/j.biopsych.2006.11.023
Dai, J. X., Han, H. L., Tian, M., Cao, J., Xiu, J. B., Song, N. N., et al. (2015). Enhanced freezing. Pavlov. J. Biol. Sci. 15, 177–182.
Foy, M. R., Stanton, M. E., Levine, S., and Thompson, R. F. (1987). Behavioral stress and the mode of action of antidepressants on generalized anxiety and panic disorders. Pharmacol. Biochem. Behav. 3223(02)01665-7
Fernandez, S. P., Cauli, B., Cabezas, C., Muzerelle, A., Poncer, J. C., and Gaspar, P. (2012). Impacts of brain serotonin deficiency following Tph2 inactivation on development and raphe neuron serotonergic specification. PLoS One 7:e43157. doi: 10.1371/journal.pone.0043157
Gutknecht, L., Popp, S., Waider, J., Sommerlandt, F. M., Goppner, C., Post, A., et al. (2015). Interaction of brain 5-HT synthesis deficiency, chronic stress and sex differentially impact emotional behavior in Tph2 knockout mice. Psychopharmacology 232, 2429–2441. doi: 10.1007/s00213-015-3879-0
Hale, M. W., and Lowry, C. A. (2011). Functional topography of midbrain and pontine serotonergic systems: implications for synaptic regulation of serotonergic circuits. Psychopharmacology 213, 243–264. doi: 10.1007/s00213-010-1809-z
Hale, M. W., Shekhar, A., and Lowry, C. A. (2012). Stress-related serotonergic systems: implications for symptomatology of anxiety and affective disorders. Cell Mol. Neurobiol. 32, 695–708. doi: 10.1007/s10571-012-9827-1
Hashimoto, S., Inoue, T., and Koyama, T. (1996). Serotonin reuptake inhibitors reduce conditioned fear stress-induced freezing behavior in rats. Psychopharmacology 123, 182–186. doi: 10.1007/BF02246175
Hendricks, T. J., Fyodorov, D. V., Wegman, I. J., Lelutiu, N. B., Pehek, E. A., Yamamoto, B., et al. (2003). Pet-1 ETS gene plays a critical role in 5-HT neuron development and is required for normal anxiety-like and aggressive behavior. Neuron 37, 233–247. doi: 10.1016/S0896-6273(02)01167-4
Igelström, K. M., and Heyward, P. M. (2012). Inhibition of hippocampal excitability by citalopram. Epilepsia 53, 2034–2042. doi: 10.1111/j.1528-1167.2012.03660.x
Jorgensen, C. V., Jacobsen, J. P., Caron, M. G., Klein, A. B., Knudsen, G. M., and Mikkelsen, J. D. (2013). Cerebral 5-HT2A receptor binding, but not mGluR2, is increased in tryptophan hydroxylase 2-deficient mice. Neurosci. Lett. 555, 118–122. doi: 10.1016/j.neulet.2013.08.073
Kessler, R. C., Chiu, W. T., Demler, O., Merikangas, K. R., and Walters, E. E. (2005). Prevalence, severity, and comorbidity of 12-month DSM-IV disorders in the National Comorbidity Survey Replication. Arch. Gen. Psychiatry 62, 617–627. doi: 10.1001/archpsyc.62.6.617
Kiyasova, V., Fernandez, S. P., Laine, J., Stankowski, L., Muzerelle, A., Doly, S., et al. (2011). A genetically defined morphologically and functionally unique subset of 5-HT neurons in the mouse raphe nuclei. J. Neurosci. 31, 2756–2768. doi: 10.1523/JNEUROSCI.4808-10.2011
Lesch, K. P., Bengel, D., Heils, A., Sabol, S. Z., Greenberg, B. D., Petri, S., et al. (1996). Association of anxiety-related traits with a polymorphism in the serotonin transporter gene regulatory region. Science 274, 1527–1531. doi: 10.1126/science.274.5292.1527
Li, X. R., Inoue, T., Hashimoto, S., and Koyama, T. (2001). Effect of chronic administration of fluoxetine and fluvoxamine on freezing behavior induced by conditioned fear. Eur. J. Pharmacol. 425, 43–50. doi: 10.1016/S0014-2990(01)01195-1
Lowry, C. A. (2002). Functional subsets of serotonergic neurons: implications for control of the hypothalamic-pituitary-adrenal axis. J. Neuroendocrinol. 14, 911–923. doi: 10.1046/j.1365-2826.2002.00886.x
Lowry, C. A., Johnson, P. L., Hay-Schmidt, A., Mikkelsen, J., and Shekhar, A. (2003). Modulation of anxiety circuits by serotonergic systems. Stress 8, 233–246. doi: 10.1007/s10880-003-0049-7
Maier, S. F., Amat, J., Baratta, M. V., Paul, E., and Watkins, L. R. (2006). Behavioral control, the medial prefrontal cortex, and resilience. Dialogues Clin. Neurosci. 8, 397–406.
Maier, S. F., and Watkins, L. R. (2005). Stressor controllability and learned helplessness: the roles of the dorsal raphe nucleus, serotonin, and corticotropin-releasing factor. Neurosci. Biobehav. Rev. 29, 829–841. doi: 10.1016/j.neubiorev.2005.03.021
Maren, S., Phan, K. L., and Liberzon, I. (2013). The contextual brain: implications for fear conditioning, extinction and psychopathology. Nat. Rev. Neurosci. 14, 417–428. doi: 10.1038/nrn3492
McQuade, R., and Sharp, T. (1997). Functional mapping of dorsal and median raphe 5-hydroxytryptamine pathways in forebrain of the rat using microdialysis. J. Neurochem. 69, 791–796. doi: 10.1046/j.1471-4159.1997.010-2089-z
Milad, R. M., Orr, S. P., Lasko, N. B., Chang, Y., Rauch, S. L., and Pitman, R. K. (2008). Presence and acquired origin of reduced recall for fear extinction in PTSD: results of a twin study. J. Psychiatr. Res. 42, 515–520. doi: 10.1016/j.jpsychires.2008.01.017
Mizane, I., Toyote, P., Meyer, M., Spiess, J., Ogren, S. O., and Stiedl, O. (2005). Time-dependent involvement of the dorsal hippocampus in trace fear conditioning in mice. Hippocampus 15, 418–426. doi: 10.1002/hipo.20067
Miyazaki, M., Yoshino, A., and Nomura, S. (2011). Relationships between anxiety severity, diagnosis of multiple anxiety disorders, and comorbid major
depressive disorder. Asian J. Psychiatry 4, 293–296. doi: 10.1016/j.ajp.2011.10.003

Mlinar, B., and Corradetti, R. (2018). Differential modulation of CA1 impulse flow by endogenous serotonin along the hippocampal longitudinal axis. Hippocampus 28, 217–225. doi: 10.1002/hipo.22315

Mlinar, B., Stocca, G., and Corradetti, R. (2015). Endogenous serotonin facilitates hippocampal long-term potentiation at CA3/CA1 synapses. J. Neural Transm. 122, 177–185. doi: 10.1007/s00702-014-1246-7

Montalbano, A., Waider, J., Barbieri, M., Baytas, O., Lesch, K.-P., Corradetti, R., et al. (2015). Cellular resilience: 5-HT neurons in Tph2−/− mice retain normal firing behavior despite the lack of brain 5-HT. Eur. Neuropsychopharmacol. 25, 2022–2035. doi: 10.1016/j.euroneuro.2015.08.021

Morini, R., Mlinar, B., Baccini, G., and Corradetti, R. (2011). Enhanced hippocampal long-term potentiation following repeated MDMA treatment in Dark-Agouti rats. Eur. Neuropsychopharmacol. 21, 80–91. doi: 10.1016/j.euroneuro.2010.07.007

Mosienko, V., Beis, D., Pasqualetti, M., Waider, J., Matthews, S., Qadri, F., et al. (2015). Life without brain serotonin: reevaluation of serotonin function with mice deficient in brain serotonin synthesis. Behav. Brain Res. 277, 78–88. doi: 10.1016/j.bbr.2014.06.005

Muzerelle, A., Scotto-Lomassese, S., Bernard, J. F., Soiza-Reilly, M., and Gaspar, P. (2016). Conditional anterograde tracing reveals distinct targeting of individual serotonin cell groups (B5-B9) to the forebrain and brainstem. Brain Struct. Funct. 221, 535–561. doi: 10.1007/s00429-014-0924-4

Pobbe, R. L., Zangrossi, H. Jr., Blanchard, D. C., and Blanchard, R. J. (2011). Involvement of dorsal raphe nucleus and dorsal periaqueductal gray 5-HT receptors in the modulation of mouse defensive behaviors. Eur. Neuropsychopharmacol. 21, 306–315. doi: 10.1016/j.euroneuro.2010.05.004

Raab, A., Popp, S., Lesch, K. P., Lohse, M. J., Fischer, M., Deckert, J., et al. (2018). Increased fear learning, spatial learning as well as neophobia in Rgs2(−/−) mice. Genes Brain Behav. 17:e12420. doi: 10.1111/gbb.12420

Ravinder, S., Burghardt, N. S., Brodsky, R., Bauer, E. P., and Chattarji, S. (2013). A role for the extended amygdala in the fear-enhancing effects of acute selective serotonin reuptake inhibitor treatment. Transl. Psychiatry 3:e209. doi: 10.1038/tp.2012.137

Rei, D., Mason, X., Seo, J., Graff, J., Rudenko, A., Wang, J., et al. (2015). Basolateral amygdala bidirectionally modulates stress-induced hippocampal learning and memory deficits through a p25/Cdk5-dependent pathway. Proc. Natl. Acad. Sci. U.S.A. 112, 7291–7296. doi: 10.1073/pnas.1415845112

Rivero, O., Selten, M. M., Sich, S., Popp, S., Bacmeister, L., Amendola, E., et al. (2015). Cadherin-13, a risk gene for ADHD and comorbid disorders, impacts GABAergic function in hippocampus and cognition. Transl. Psychiatry 5:e565. doi: 10.1038/tp.2015.152

Sacchetti, B., Lorenzini, C. A., Baldi, E., Bucherelli, C., Roberto, M., Tassoni, G., et al. (2002). Time-dependent inhibition of hippocampal LTP in vitro following contextual fear conditioning in the rat. Eur. J. Neurosci. 15, 143–150. doi: 10.1046/j.1460-9581.2001.01844.x

Schaefer, T. L., Vorhees, C. V., and Williams, M. T. (2009). Mouse plasmacytoma-expressed transcript 1 knock out induced 5-HT disruption results in a lack of cognitive deficits and an anxiety phenotype complicated by hypoactivity and defensiveness. Neuroscience 164, 1431–1443. doi: 10.1016/j.neuroscience.2009.09.059

Schindelin, J., Arganda-Carreras, I., Frise, E., Kaynig, V., Longair, M., Pietzsch, T., et al. (2012). Fiji: an open-source platform for biological-image analysis. Nat. Methods 9, 676–682. doi: 10.1038/nmeth.2019

Schmidt, S. D., Furini, C. R. G., Zinn, C. G., Cavalcante, L. E., Ferreira, F. F., Behling, J. A. K., et al. (2017). Modulation of the consolidation and reconsolidation of fear memory by three different serotonin receptors in hippocampus. Neurobiol. Learn. Mem. 142(Pt A), 48–54. doi: 10.1016/j.nlm.2016.12.017

Song, N. N., Xiu, J. B., Huang, Y., Chen, J. Y., Zhang, L., Gutknecht, L., et al. (2011). Adult raphe-specific deletion of LMX1B leads to central serotonin deficiency. PLoS One 6:e15998. doi: 10.1371/journal.pone.0015998

Spannuth, B. M., Hale, M. W., Evans, A. K., Lukkes, J. L., Campeau, S., and Lowery, C. A. (2011). Investigation of a central nucleus of the amygdala/dorsal raphe serotoninergic circuit implicated in fear-potentiated startle. Neuroscience 179, 104–119. doi: 10.1016/j.neuroscience.2011.01.042

Staubli, U., and Otaky, N. (1994). Serotonin controls the magnitude of LTP induced by theta bursts via an action on NMDA-receptor-mediated responses. Brain Res. 643, 10–16. doi: 10.1016/0006-8993(94)90003-5

Toth, I., Dietz, M., Peterlik, D., Huber, S. E., Fendt, M., Neumann, I. D., et al. (2012). Pharmacological interference with metabotropic glutamate receptor subtype 7 but not subtype 5 differentially affects within- and between-session extinction of Pavlovian conditioned fear. Neuropharmacology 62, 1619–1626. doi: 10.1016/j.neuropharm.2011.10.021

Tovote, P., Esposito, M. S., Botta, P., Chaudun, F., Fadok, J. P., Markovic, M., et al. (2016). Midbrain circuits for defensive behaviour. Nature 534, 206–212. doi: 10.1038/nature17996

Waider, J., Popp, S., Lange, M. D., Kern, R., Kolter, J. F., Kobler, J., et al. (2017). Genetically driven brain serotonin deficiency facilitates panic-like escape behavior in mice. Transl. Psychiatry 7:e1246. doi: 10.1038/tp.2017.209

Waider, J., Proft, F., Langhofer, G., Asan, E., Lesch, K. P., and Gutknecht, L. (2013). GABA concentration and GABAergic neuron populations in limbic areas are differentially altered by brain serotonin deficiency in Tph2 knockout mice. Histochim. Cell Biol. 139, 267–281. doi: 10.1007/s00148-012-1029-x

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