Serotonin-1A Autoreceptors Are Necessary and Sufficient for the Normal Formation of Circuits Underlying Innate Anxiety


1Department of Psychiatry, Columbia University, New York, New York 10032; 2Department of Anesthesiology, Children’s Hospital of Philadelphia and University of Pennsylvania, Philadelphia, Pennsylvania 19132; 3Faculté Pharmacie, Université Paris Sud, Chatenay-Malabry, F-92296 France; 4Department of Radiology, University of Pennsylvania, Philadelphia, Pennsylvania 19104; 5Department of Neuroscience, Columbia University, New York, New York 10032; and 6The New York State Psychiatric Institute, New York, New York 10032

Identifying the factors contributing to the etiology of anxiety and depression is critical for the development of more efficacious therapies. Serotonin (5-HT) is intimately linked to both disorders. The inhibitory serotonin-1A (5-HT\textsubscript{1A}) receptor exists in two separate populations with distinct effects on serotonergic signaling: (1) an autoreceptor that limits 5-HT release throughout the brain and (2) a heteroreceptor that mediates inhibitory responses to released 5-HT. Traditional pharmacologic and transgenic strategies have not addressed the distinct roles of these two receptor populations. Here we use a recently developed genetic mouse system to independently manipulate 5-HT\textsubscript{1A} autoreceptor and heteroreceptor populations. We show that 5-HT\textsubscript{1A} autoreceptors act to affect anxiety-like behavior. In contrast, 5-HT\textsubscript{1A} heteroreceptors affect responses to forced swim stress, without effects on anxiety-like behavior. Together with our previously reported work, these results establish distinct roles for the two receptor populations, providing evidence that signaling through endogenous 5-HT\textsubscript{1A} autoreceptors is necessary and sufficient for the establishment of normal anxiety-like behavior.

Introduction

Anxiety and depression are highly comorbid disorders with partially overlapping genetic predisposition, environmental risk, symptom dimensions, and treatment profiles (Ressler and Nemeroff, 2000; Nemeroff, 2002). This overlap suggests that anxiety and depression share some circuitry and neurochemistry, but can be influenced by distinct factors.

Serotonin (5-HT) participates in the etiology and treatment of both anxiety and depression (Rush et al., 2006; Davidson, 2009). The most common treatments for major depressive disorder and many anxiety disorders are the selective serotonin reuptake inhibitors (SSRIs), which are thought to exert their therapeutic effects by increasing extracellular 5-HT levels (Gartside et al., 1995). However, in contrast to the beneficial effects of SSRIs in adults, evidence from rodent models suggest that SSRI treatment during early development increases anxiety or depression later in life (Caspì et al., 2003; Lià et al., 2003; Ansorge et al., 2008; Olivier et al., 2008; Oberlander et al., 2009). Thus, 5-HT may affect immature and mature mood-related circuitry differently.

Serotonin is released throughout the forebrain by axons emanating from cell bodies located in the midbrain raphe (Barnes and Sharp, 1999). The largely neuromodulatory effects of 5-HT are mediated through 14 receptor subtypes, whereas 5-HT levels are limited by two inhibitory autoreceptors expressed on 5-HT\textsubscript{1A} autoreceptor and heteroreceptor populations. We show that 5-HT\textsubscript{1A} autoreceptors act to affect anxiety-like behavior. In contrast, 5-HT\textsubscript{1A} heteroreceptors affect responses to forced swim stress, without effects on anxiety-like behavior. Together with our previously reported work, these results establish distinct roles for the two receptor populations, providing evidence that signaling through endogenous 5-HT\textsubscript{1A} autoreceptors is necessary and sufficient for the establishment of normal anxiety-like behavior.

Received Nov. 2, 2010; revised Feb. 11, 2011; accepted Feb. 23, 2011.


We thank Mathieu Araujo for technical assistance. This work was supported by NIH Grants K08 MH076083 and R01 MH091427 to E.D.L., a NARSAD Young Investigator Award to R.H., a NARSAD Distinguished Investigator Award to R.H., and a AstraZeneca Grant CUBH-0439 to R.H.

R.H. receives compensation as a consultant from PGxHealth.

Correspondence should be addressed to either Dr. E. David Leonardo or René Hen, Department of Psychiatry, Columbia University/New York State Psychiatric Institute, 1051 Riverside Drive, Box 87, New York, New York 10032.

E-mail: e0567@columbia.edu or rh95@columbia.edu.

DOI: 10.1523/JNEUROSCI.5536-10.2011

Copyright © 2011 the authors 0270-6474/11/316008-11$15.00/0

Serotonin-1A Autoreceptors Are Necessary and Sufficient for the Normal Formation of Circuits Underlying Innate Anxiety

Identifying the factors contributing to the etiology of anxiety and depression is critical for the development of more efficacious therapies. Serotonin (5-HT) is intimately linked to both disorders. The inhibitory serotonin-1A (5-HT\textsubscript{1A}) receptor exists in two separate populations with distinct effects on serotonergic signaling: (1) an autoreceptor that limits 5-HT release throughout the brain and (2) a heteroreceptor that mediates inhibitory responses to released 5-HT. Traditional pharmacologic and transgenic strategies have not addressed the distinct roles of these two receptor populations. Here we use a recently developed genetic mouse system to independently manipulate 5-HT\textsubscript{1A} autoreceptor and heteroreceptor populations. We show that 5-HT\textsubscript{1A} autoreceptors act to affect anxiety-like behavior. In contrast, 5-HT\textsubscript{1A} heteroreceptors affect responses to forced swim stress, without effects on anxiety-like behavior. Together with our previously reported work, these results establish distinct roles for the two receptor populations, providing evidence that signaling through endogenous 5-HT\textsubscript{1A} autoreceptors is necessary and sufficient for the establishment of normal anxiety-like behavior.
of-function studies have likewise suggested that the 5-HT_{1A} receptor can stably impact anxiety-like behavior during the first weeks of life (Gross et al., 2002; Bert et al., 2006). Given this evidence, 5-HT_{1A} receptors may influence anxiety and depression differentially not only between subpopulations, but also across developmental stages in an animal.

Separating both the temporal (developmental vs adult) and spatial (autoreceptor vs heteroreceptor) roles of the endogenous 5-HT_{1A} receptor in anxiety and depression has proven difficult with existing genetic or pharmacological techniques. To independently assess the functional role of endogenous 5-HT_{1A} autoreceptors and heteroreceptors, we developed a genetic mouse system with temporal and spatial specificity and tested the adult phenotype of these animals in a number of classic, mood-related paradigms.

Materials and Methods

Transgenic mice

5-HT_{1A} autoreceptor knock-out mice. Mice with suppressible 5-HT_{1A} receptors (Htr1α^{tetO}) were bred as described previously (Richardson-Jones et al., 2010). Htr1α^{tetO/mcr} mice demonstrate a pattern of 5-HT_{1A} receptors that is indistinguishable from the wild-type pattern. Mice with inducible suppression of 5-HT_{1A} autoreceptors were homozygous for the tetO insertion (Htr1α^{tetO/mcr}), possessed one copy of the raphe-specific tTS suppressor transgene, Pet1-tTS, and were maintained in the absence of doxycycline (dox) throughout life (Richardson-Jones et al., 2010).

5-HT_{1A} heteroreceptor knock-out mice. Heteroreceptor-specific suppression was achieved by placing tTS under transcriptional control of elements of the α-CaMKII promoter (Mayford et al., 1996). Briefly, an EcoRI– BamHI fragment containing the tTS coding sequence was freed from the pAct tTS plasmid (Mallo et al., 2003). The fragment was blunted and cloned into the EcoRV site of the pNN265 plasmid, flanking it with a generic 5′ intron, 3′ intron, and SV40 poly(A) sequence (Mayford et al., 1996). The tTS sequence flanked by introns was freely digested with NotI and cloned into the pMM403 plasmid, placing it under the control of α-CaMKII promoter fragments. Lastly, the promoter fragments and tTS coding sequence (CamK–tTS) were freed by Sfi/Srf digest and subjected to pronuclear injection into B6CBA/F2 hybrid zygotes. Founders were screened by PCR for tTS as described previously (Richardson-Jones et al., 2010) and by Southern blot analysis.

Seven founder lines were generated, bred to the Htr1α^{mcr/tetO} background, and screened for suppression of 5-HT_{1A} heteroreceptors. The three general patterns of suppression in these lines corresponded to the previously reported activity of these promoter fragments: (1) suppression of all receptors in the brain, (2) preferential suppression of forebrain receptors while sparing receptors in the raphe, and (3) preferential suppression of dorsal hippocampal receptors while sparing most other receptor populations (data not shown) (Gross et al., 2002; Pittenger et al., 2002). One line was identified that displayed suppression of 5-HT_{1A} heteroreceptors without affecting autoreceptors, and this line was used for subsequent experiments. Mice with inducible suppression of 5-HT_{1A} heteroreceptors were homozygous for the tetO insertion (Htr1α^{tetO/mcr}), possessed one copy of the forebrain-specific tTS suppressor transgene CamK–tTS, and were maintained in the absence of dox throughout life.

For all experiments, male Htr1α^{tetO/mcr}/Pet1–tTS^{−} and Htr1α^{tetO/mcr}/CamK–tTS^{−} mice were bred to Htr1α^{mcr/tetO} females. Thus, the tTS transgene was transmitted through the male germline, ensuring that all pups were raised by mothers of the same genotype, regardless of dox status. All experiments were conducted on male offspring. Animals were maintained on a mixed 12:12 light/dark schedule; all testing was conducted during the light period.

Animal protocols were approved by the Institutional Animal Care and Use Committee and were conducted in accordance to the NIH Guide for the Care and Use of Laboratory Animals. Care was taken to minimize the number of animals used and their suffering.

Animal husbandry

Mice were housed in groups of three to five per cage and had ad libitum access to food and water. Animals were maintained on a 12:12 light/dark schedule; all testing was conducted during the light period.

Electrophysiology

Whole-cell recordings of dorsal raphe and hippocampal CA1 pyramidal neurons were made as previously described (Beck et al., 2004; Tsetsenis et al., 2007; Crawford et al., 2010). In brief, mice were killed using decapitation. When submerged in a cold artificial CSF (aCSF) solution where NaCl had been replaced with sucrose (248 mOsm), the cortical slices were cut at the level of the dorsal hippocampus and midbrain with a VT1000s vibratome (Leica Microsystems) to generate 200 μm thick brain slices. Slices were maintained in aCSF bubbled with 95%O_{2}/5%CO_{2} at 36°C for 1 h and then at room temperature until used. Individual slices were then placed in a recording chamber and continuously perfused with 32–34°C aCSF solution bubbled with 95%O_{2}/5%CO_{2} with a solution flow rate of 1.5–2 ml/min. The composition of the aCSF was as follows (in mls): 124 NaCl, 2.5 KCl, 1.25 NaH_{2}PO_{4}, 2.0 MgSO_{4}, 2.5 CaCl_{2}, 10 dextrose, and 26 NaHCO_{3}. Neurons were visualized using Nikon E600 (Optical Apparatus) upright microscope fitted with a 40× water-immersion objective, differential interference contrast, and infrared filter. Whole-cell recording pipettes fabricated on a pipette puller (P-97; Sutter Instrument) had a resistance of 6–10 MΩ when filled with an intracellular solution containing the following (in mls): 130 K-glucionate, 10 Na-phosphocreatine, 1 MgCl_{2}, 0.02 EGTA, 10 HEPES, 2 MgATP, 0.5 Na,GTP, and 0.1% biocytin, pH 7.3. Recordings were collected online with a Multiclamp 700B amplifier, Digidata 1320 A/D converter, and Clampex 9.0 software (Molecular Devices). In current-clamp recording, resting membrane potential, action potential (AP) threshold, AP duration, after-hyperpolarization (AHP) amplitude, and the time it takes for the AHP to depolarize to one-half its peak amplitude were measured directly from traces, as previously described (Crawford et al., 2010). In voltage clamp, with a holding potential of ~60 mV, the magnitude of the outward current elicited by bath application of 100 mS 5-carboxy-amidotryptamine (5-CT) was measured. 5-CT was chosen for these experiments instead of 8-OH-DPAT as it is a high affinity full agonist, is less lipophilic, and washes out more easily from the slices. Following the experiment, slices were fixed for 2–3 h with 4% paraformaldehyde and the midbrain slices were processed for immunohistochemical detection of tryptophan hydroxylase (TPH). Identity of all recorded cells was confirmed by visualization of the biocytin-filled cells. Immunohistochemical identification of each neuron recorded in patch-clamp configuration was completed as previously described (Beck et al., 2004; Lemos et al., 2006; Kirby et al., 2008). In brief, a standard immunohistochemistry protocol was used on 200-μm-thick slices using mouse anti-TPH (1:200; Sigma) along with secondary donkey anti-mouse Alexa Fluor International) either throughout life [autoreceptor–knock-out (auto-KO) and heteroreceptor–knock-out (hetero-KO)] or beginning at postnatal day 50 (adult hetero-KO). To control for the possible effects of dox on behavior, littermate controls lacking the tTS transgene, in which dox had no effect on 5-HT_{1A} receptor expression, were also tested in baseline behavioral experiments. For all tests presented, dox exposure alone did not result in any behavioral changes (data not shown).
Intracerebral in vivo microdialysis

Extracellular 5-HT levels were measured by in vivo microdialysis as previously described (Guiard et al., 2008). Briefly, two concentric dialysis probes were implanted in the ventral hippocampus (vHPC) and prefrontal cortex (PFC) (outer diameter by active length: 0.3 × 1.6 mm and 0.3 × 2 mm, respectively) of anesthetized mice (chloral hydrate, 400 mg/kg, i.p.). Stereotaxic coordinates were as follows (in mm): PCF: anterior, 1.6; lateral, 1.3; ventral, 1.6; vHPC: anterior, −2.8; lateral, 3.0; ventral, 3.0 (Franklin and Paxinos, 1997). Animals were allowed to recover for a period of 24 h. Following recovery, probes were continuously perfused with ACSF, and dialysates were collected every 15 min for analysis by HPLC-amperometry (Guiard et al., 2008). Baseline 5-HT levels were calculated as the average of the first four samples, ± SEM. Freely moving mice were treated (t = 0) with either a challenge dose of fluoxetine (18 mg/kg; i.p.) or its vehicle, and dialysate samples were collected for a 0–120 min post-treatment period. The limit of sensitivity for 5-HT was 0.5 fmol/sample (signal-to-noise ratio, 2). Following sample collection, brains were removed and sectioned to ensure proper probe placement.

Behavioral and physiological testing

All animals used for behavioral testing were age matched within 2 weeks. Animals were initially tested at 11–13 weeks of age, at least 4 weeks after the cessation of dox in adult heteroreceptor–KO animals. Baseline anxiety tests were completed before other behavioral tests.

8-OH-DPAT-induced hyperthermia. Body temperature was assessed intracutally using a lubricated probe inserted ~2 cm and a Thermalert TH-5 thermal monitor (Physitemp). Mice were singly housed in clean cages for 10 min and three baseline body temperature measurements were taken. Ten minutes after the third baseline measurement, animals received 8-OH-DPAT intraperitoneally at the doses indicated and body temperature was monitored every 10 min for a total of 50 min. Temperatures are represented as a change from the final baseline measurement.

Stress-induced hyperthermia. The stress-induced hyperthermia paradigm measures a physiologic response to a stressful stimulus (Adriaan Bouwknecht et al., 2007). Briefly, animals in their home cages were moved to a testing room and allowed to acclimate for 1 h. One animal per cage was removed and a baseline body temperature was measured intracutally. Each animal was then placed in a novel, clean cage for 10 min, after which a second body temperature was recorded.

Open-field test. Exploration in response to a novel open field was measured as described previously (Richardson-Jones et al., 2010). Dependent measures were total path length (in cm), number, and percentage of distance in the center (distance traveled in the center divided by the total distance traveled).

Light/dark choice test. Exploration of the light/dark chamber was measured as described previously (Richardson-Jones et al., 2010). Dependent measures were total distance and percentage of time spent in the light compartment.

Modified forced-swim test. Behavioral response to forced swimming was assayed as described previously (David et al., 2007). Briefly, mice were placed into clear plastic buckets 20 cm in diameter and 23 cm deep, filled two-thirds of the way up with 26°C water and videotaped from the side. Mice had a 2 min pretest period before initiating scoring, after which two consecutive 2 min periods were scored. All animals were exposed to the swim test on two consecutive days. Scoring was done using an automated Viewpoint Videotrack software package. Dependent variables were immobility, swimming, and climbing.

Statistical analysis

In general, the effect of treatment or dose was analyzed using an ANOVA, using repeated measures where appropriate. Significant ANOVAs were followed up with Fisher protected least significant difference test for behavioral and physiological measures, and with Student–Neuman–Keuls t test for electrophysiological characterization.

Results

Conditional suppression of the 5-HT1A autoreceptors and heteroreceptors

Previous attempts to target autoreceptor versus heteroreceptor populations independently in mice relied on transgenic gain-of-function strategies using heterologous promoters that resulted in overexpression and misexpression of the receptor (Gross et al., 2002; Kussnerow et al., 2004; Bert et al., 2006; Audero et al., 2008). Such approaches fall short of establishing causal links between endogenous receptor populations and either circuit function or behavior. To independently manipulate either 5-HT1A heteroreceptors or autoreceptors while maintaining endogenous expression of the other receptor population, we used the tTS/tetO system to suppress 5-HT1A receptor transcription from the Htr1a gene locus. We previously reported the generation of mice that allow for specific suppression of 5-HT1A autoreceptors, using two engineered mouse lines: a raphe-specific TTS transgenic line and a knock-in mouse line with tetO sites inserted between the promoter and coding regions of the Htr1a locus (Richardson-Jones et al., 2010). To suppress 5-HT1A autoreceptors in the raphe throughout life (auto-KO), the previously described Htr1a<sup>(tetO<sup>cre</sup>)</sup> Pet1-tTS<sup>+</sup> mice were maintained in the absence of dox throughout life (Fig. 1C,D). Controls for these mice, in which 5-HT1A autoreceptors were not suppressed, were generated by maintaining Htr1a<sup>(tetO<sup>cre</sup>/tetO)</sup> Pet1-tTS<sup>+</sup> mice in the presence of dox throughout life (Fig. 1A,D). Quantitative receptor autoradiography revealed that <20% of control 5-HT1A receptor binding remained in the dorsal raphe of adult auto-KO mice (one-tailed t test in the dorsal raphe, t<sub>13</sub> = −15.503, p < 0.0001), whereas no changes were observed in binding of 5-HT1A in areas of high heteroreceptor expression, such as the amygdala, hippocampus, and entorhinal cortex (repeated-measures ANOVA with brain region as a within-subject factor and group as a between-subject factor, main effect of group F<sub>13,13</sub> = 1.398, p = 0.2583; region-by-group interaction, F<sub>13,13</sub> = 0.795, p = 0.46232) (Fig. 2). Attempts with this line to fully suppress the autoreceptor beginning in adulthood resulted in a modest 30% reduction in raphe 5-HT1A expression. Physiological and behavioral characterization of these animals has been reported previously (Richardson-Jones et al., 2010). The reason for more effective suppression of 5-HT1A transcription in animals maintained in the absence of dox throughout life remains unclear, although it appears that early developmental suppression at the 5-HT1A locus may be irreversible (data not shown).

To generate mice with suppression of 5-HT<sub>1A</sub> heteroreceptors without affecting autoreceptors, we first created a transgenic mouse line with TTS expression directed by promoter elements from the α-CaMKII gene (Mayford et al., 1996). One mouse line was identified that demonstrated suppression of only 5-HT1A forebrain heteroreceptors without affecting 5-HT1A raphe autoreceptors (hetero-KO) (ANOVA in dorsal raphe, F<sub>1,29</sub> = 0.429, p < 0.6635) (Figs. 1B,E; 2B,D). In hetero-KO mice, receptor expression was undetectable in the hippocampus, amygdala, or prefrontal cortex, whereas low levels of receptor remained in the oliv, entorhinal cortex, and the medial and lateral septum
Altered agonist response in mice with conditional suppression of 5-HT₁₄ autoreceptors and heteroreceptors

To directly confirm the functional impact of receptor suppression revealed by autoradiography, we performed whole-cell recordings of neurons in acute slice in response to the 5-HT₁₄ agonist 5-CT. Cellular responses were recorded in the dorsal raphe of auto-KO mice and their controls and in area CA1 of the hippocampus in hetero-KO mice and their controls. Following recording, cells were filled with biocytin. Pyramidal neurons in the hippocampus were confirmed by morphological features and raphe neurons were confirmed as serotonergic by immunohistochemistry for tryptophan hydroxylase (data not shown).

In agreement with the decreased 5-HT₁₄ autoreceptor levels indicated by autoradiography, serotonergic neurons in the dorsal raphe of auto-KO mice had a significantly decreased current response to 5-CT (two-tailed t test, \( t_{35} = 8.432, p < 0.0001 \)) (Fig. 3A–C). Indeed, we were unable to detect any response to 5-CT (defined as current < 5 pA) in 5-HT dorsal raphe neurons of auto-KO mice, consistent with a lack of functional 5-HT₁₄ raphe autoreceptors in these animals. The small amount of remaining...
receptor binding in the dorsal raphe may represent either desensitized 5-HT\textsubscript{1A} autoreceptors or receptors on non-5-HT raphe neurons (Kirby et al., 2003; Beck et al., 2004).

In pyramidal neurons in area CA1 of hetero-KO mice, we observed a large decrease in the average current response to 5-CT (one-tailed \(t\) test, \(t_{39}/H_{11005}/H_{11002} = 3.254, p/H_{11005}/H_{11002} = 0.0012\)) (Fig. 3D–F), consistent with the greatly decreased levels of heteroreceptor revealed by autoradiography.

To assess the \textit{in vivo} functional status of 5-HT\textsubscript{1A} autoreceptors in both groups of mice, we examined their hypothermic response to challenge with the 5-HT\textsubscript{1A} agonist 8-OH-DPAT or its saline vehicle. The controls for the auto-KO, hetero-KO, and adult hetero-KO mice displayed a hypothermic response to 0.5 mg/kg 8-OH-DPAT consistent with previously observed values (repeated-measures ANOVA, main effect of dose for hetero-KO controls, \(F_{(1,6)} = 46.979, p < 0.0001\); adult hetero-KO controls, \(F_{(1,9)} = 33.055, p = 0.0003\); auto-KO controls, \(F_{(1,6)} = 109.630, p < 0.0001\)) (Fig. 4Aii,Bii,Cii). Consistent with a lack of functional autoreceptor, 5-HT\textsubscript{1A} auto-KO mice displayed no detectable body temperature decrease in response to 8-OH-DPAT, with responses indistinguishable from the response to saline injection (repeated-measures ANOVA for effect of treatment, \(F_{(1,6)} = 1.38, p = 0.284\)) (Fig. 4Ci).

Conversely, 5-HT\textsubscript{1A} hetero-KO mice displayed a full hypothermic response to 8-OH-DPAT that was significantly different from the response to saline and indistinguishable from the response in their matched controls (repeated-measures ANOVA, effect of dose in hetero-KO mice, \(F_{(1,12)} = 33.10, p < 0.0001\); repeated-measures ANOVA, effect of group in 8-OH-DPAT-treated mice, \(F_{(1,12)} = 2.486, p = 0.141\)) (Fig. 4A). Likewise, 5-HT\textsubscript{1A} adult hetero-KO mice displayed a response to 8-OH-DPAT that was significantly different from the response to saline and indistinguishable from the response in their matched controls (repeated-measures ANOVA, effect of dose in adult hetero-KO mice, \(F_{(1,9)} = 36.216, p = 0.0002\); repeated-measures ANOVA effect of group in 8-OH-DPAT-treated mice, \(F_{(1,9)} = 33.055, p = 0.0003\)).

---

**Figure 2.** A, Quantitative \(^{125}\text{I}\)-MPPI autoradiography for 5-HT\textsubscript{1A} receptors revealed a significant decrease in the dorsal raphe (i) of auto-KO mice compared with their matched controls. Conversely, no difference was detected in forebrain regions (ii), such as the ventral dentate gyrus of the hippocampus (HPC), the amygdala (AMG), and the entorhinal cortex (EC). B, i, No difference was detected in binding in the dorsal raphe of hetero-KO or adult hetero-KO mice compared with their matched controls. ii, Conversely, binding was significantly decreased in dorsal CA1 of the hippocampus in both groups compared with control (\(N = 3–7/\text{group}\)). All values are mean \pm SEM. ***\(p < 0.0001\). C, D, Matched autoradiograms showing detailed expression patterns of 5-HT\textsubscript{1A} receptors binding across the rostrocaudal extent of the brain in coronal section comparing auto-KO mice and their controls (C) and hetero-KO and adult hetero-KO mice and their controls (D). Low levels of heteroreceptors remain in parts of the septum, olivary nucleus, and entorhinal cortex. E, Receptor binding in the postnatal developmental period in hetero-KO mice and their controls. Matched sections are shown at the level of the dorsal hippocampus and the dorsal raphe nucleus. Forebrain receptor suppression is complete by postnatal day (p) 14.
of group in 8-OH-DPAT-treated mice, $F_{1,10} = 0.071, p = 0.795$ (Fig. 4B). These findings are consistent with previous literature implicating 5-HT_{1A} autoreceptors, but not heteroreceptors, in the 8-OH-DPAT-induced hypothermic response in mice (Martin et al., 1992).

These results provide both a direct in vivo measurement and an in vivo confirmation of the functional selectivity of genetic suppression of 5-HT_{1A} autoreceptors and heteroreceptors, respectively, in the auto-KO, hetero-KO, and adult hetero-KO mice.

**Increased serotonergic tone in mice lacking 5-HT_{1A} autoreceptors throughout life**

The extracellular level of serotonin (5-HT_{ext}) is generally thought to be under the inhibitory control of both 5-HT_{1A} and 5-HT_{1B} autoreceptors. Here we sought to determine how specific suppression of 5-HT_{1A} autoreceptors throughout life was reflected at the neurochemical level in adult animals. We observed higher basal cortical 5-HT_{ext} in whole-life auto-KO mice compared to their wild-type littermates ($F_{1,13} = 7.79, p = 0.015$). However, differences between the two strains of mice in the vHPC were not significant ($F_{1,13} = 1.74, p = 0.21$) (Table 1).

To directly measure the effect of pharmacological inactivation of the 5-HT transporter in mice lacking 5-HT_{1A} autoreceptors, we challenged both auto-KO mice and their controls with the SSRI fluoxetine.

In the frontal cortex, a two-way ANOVA (treatment by genotype) on area under the curve (AUC) values revealed no significant effect of genotype ($F_{1,28} = 0.041, p = 0.841$), but a significant effect of treatment ($F_{1,28} = 36.984, p < 0.0001$). No genotype-by-treatment interaction was detected ($F_{1,28} = 0.02, p = 0.9672$).

In the vHPC, a two-way ANOVA (treatment by genotype) on AUC values revealed both a significant effect of genotype ($F_{1,30} = 6.82, p = 0.0139$), and a significant effect of treatment ($F_{1,30} = 58.21, p < 0.0001$), as well as a genotype-by-treatment interaction ($F_{1,30} = 6.62, p = 0.0153$).

Thus, as expected, both control and auto-KO mice displayed increased 5-HT_{ext} in response to acute treatment with the fluoxetine in the vHPC (Fig. 5A,C) and frontal cortex (Fig. 5B,D). Moreover, auto-KO mice displayed a proportionately larger increase in 5-HT_{ext} in response to fluoxetine than their controls in the vHPC, but not in the frontal cortex (post hoc for group, $p = 0.0198$) (Fig. 5C,D). Together, these results demonstrate that suppression of 5-HT_{1A} autoreceptors throughout life results in regionally distinct increases in serotonergic tone.

**Increased anxiety-like behavior in mice lacking 5-HT_{1A} autoreceptors**

Mice lacking all 5-HT_{1A} receptors throughout life display reliably increased anxiety-like behavior (Heisler et al., 1998; Parks et al., 1998; Ramboz et al., 1998). To independently assess the role of endogenous 5-HT_{1A} autoreceptors and heteroreceptors in anxiety-like behavior, we tested hetero-KO, adult hetero-KO, and auto-KO mice in conflict/anxiety paradigms: the open field, the light/dark choice test, and the elevated plus maze.

We were unable to detect any significant impact on anxiety-like behavior by suppression of 5-HT_{1A} heteroreceptors beginning in early postnatal development. Specifically, we detected no differences between 5-HT_{1A} hetero-KO mice and their controls in exploration of the center of the open field (ANOVA, $F_{1,32} = 0.025, p = 0.875$), although a trend was seen for decreased exploration of the total open field (repeated-measures ANOVA, $F_{1,32} = 3.31, p = 0.078$) (Fig. 6A). Likewise, no differences were detected between 5-HT_{1A} hetero-KO mice and their controls in total exploration (repeated-measures ANOVA, main effect of group, $F_{1,41} = 0.064, p = 0.802$), entries into the light (ANOVA, main effect of group, $F_{1,40} = 0.258, p = 0.614$), or time in the light compartment (ANOVA, main effect of group, $F_{1,41} = 0.077, p = 0.7823$) in the light/dark choice test (Fig. 6B). Finally, no effects on anxiety-like behavior were detected in the elevated plus maze (data not shown).

In adult hetero-KO mice, we did not detect changes in exploration of the center of the open field (ANOVA, $F_{1,47} = 0.686, p = 0.412$), although, similar to the results in hetero-KO mice, we observed a trend for a decrease in total exploration of the open field (repeated-measures ANOVA $F_{1,47} = 2.74, p = 0.105$) (Fig. 6C). No differences were detected between adult hetero-KO mice and their controls in total exploration (ANOVA, $F_{1,30} = 0.122, p = 0.730$), exploration of the light (ANOVA, $F_{1,30} = 2.11, p = 0.157$), or entries into the light in the light/dark choice test (ANOVA, $F_{1,30} = 2.314, p = 0.139$) (Fig. 6D). Together, these results demonstrate that suppression of 5-HT_{1A} heteroreceptors, beginning either in early postnatal development or adulthood, is not sufficient to impact classic anxiety-like behavior. However, we cannot rule out the contribution of 5-HT_{1A} heteroreceptors to exploratory behavior, as trends for decreased exploration in the open field were seen in both groups of mice that lacked 5-HT_{1A} heteroreceptors.

Having observed that suppression of 5-HT_{1A} heteroreceptors alone, either throughout life or beginning in adulthood, was not...
played decreased total exploration (ANOVA, Table 1. Mean basal serotonin levels (fmol/20 μl dialysate) ± SEM in ventral hippocampus and prefrontal cortex

<table>
<thead>
<tr>
<th>Condition</th>
<th>Ventral Hippocampus</th>
<th>Prefrontal Cortex</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>2.8 ± 0.3</td>
<td>1.9 ± 0.1</td>
</tr>
<tr>
<td>Auto-KO</td>
<td>3.7 ± 0.5</td>
<td>3.7 ± 0.78*</td>
</tr>
</tbody>
</table>

HPC, ventral hippocampus; PFC, prefrontal cortex. *Value differs from control PFC, p = 0.015.

Figure 4. A, Hypothalamic response to the 5-HT1A/5-HT1B agonist 8-OH-DPAT (DPAT) is normal in adult hetero-KO mice (i) and normal in their controls (ii). B, Hypothalamic response to the 5-HT1A agonist 8-OH-DPAT is normal in adult hetero-KO mice (i) and their controls (ii). C, Hypothalamic response to the 5-HT1A agonist 8-OH-DPAT is abolished in auto-KO mice (i) and normal in their controls (ii) (n = 3–6 mice/group/dose). Values are mean ± SEM. ***p < 0.001.

Figure 5. A, B, Extracellular serotonin levels measured by in vivo microdialysis in the HPC (A) and PFC (B) following injection of saline (Sal; t0) and 18 mg/kg fluoxetine (Flx; t105). Values are mean ± SEM for each time point. C, D, Total extracellular serotonin, measured by area under the curve analysis compared with baseline, increases in the HPC (C) and PFC (D) of both auto-KO and control mice in response to acute fluoxetine treatment. Auto-KO mice display a larger percentage increase than controls in response to fluoxetine in the HPC but not in the PFC. Values are ± SEM (n = 7–9 mice/group). *p < 0.05, **p < 0.01.

in any of the measures (data not shown). The phenotype observed in auto-KO mice is consistent with the increased anxiety-like behavior in conflict-based tasks and decreased exploration of novel environments, and is similar to what has been observed in constitutive 5-HT1A KO mice on at least three genetic backgrounds (Heisler et al., 1998; Parks et al., 1998; Ramboz et al., 1998). These data suggest that endogenous 5-HT1A autoreceptors are necessary and sufficient for the establishment of anxiety-like behavior. Furthermore, these data suggest that normal signaling though 5-HT1A heteroreceptors (with autoreceptors intact) is dispensable for the establishment of anxiety-like behavior.

Mice lacking 5-HT1A heteroreceptors throughout life demonstrate increased behavioral despair in the forced-swim test

In addition to its role in anxiety, diverse evidence has linked 5-HT1A receptors to depression and the response to stress. 5-HT1A KO mice display altered behavior in the forced-swim test (FST) and a blunted response to antidepressants as well as increased anxiety-like behavior (Ramboz et al., 1998; Santarelli et al., 2003). However, it is unclear whether the anxiety phenotype, and other non-anxiety related phenotypes like the one seen in the FST, are impacted by the same populations of 5-HT1A receptors, or whether the temporal and spatial role of 5-HT1A receptors for these two behaviors are distinct.

To test directly the role of 5-HT1A heteroreceptors and autoreceptors in mediating mood-related behavior that does not capture dimensions of anxiety, we first tested auto-KO mice in the forced swim test. In this test, which is often used as a pharmacological screen for antidepressant activity, active swimming and climbing are scored as mobility, which generally decreases over time. We observed that auto-KO mice displayed decreased mobility that was indistinguishable from their controls on the second day of testing (repeated-measures ANOVA, main effect of group, F(1,48) = 0.871, p = 0.3554; main effect of time, F(3,48) = 23.771, p < 0.0001) (Fig. 7C). This finding contrasted with our previ-
that compensation occurs when 5-HT autoreceptors (with decreased autoreceptor levels decreasing behavioral despair) (Richardson-Jones et al., 2010), and suggests that compensation occurs when 5-HT\_1\_A autoreceptors are suppressed throughout life.

As suppressing autoreceptors throughout life did not alter the response to forced-swim stress, we next tested the response of hetero-KO and adult hetero-KO mice. Although hetero-KO mice were indistinguishable from their controls on the first day of testing, they displayed markedly less mobility, or more behavioral despair, on the second day of testing (repeated-measures ANOVA, group by time interaction, $F_{(1,44)} = 2.97, p = 0.0345$; post hoc for day 2, minutes 3–4 and 5–6, $p = 0.0371$ and $p = 0.0018$, respectively) (Fig. 7A). Conversely, adult suppression of 5-HT\_1\_A autoreceptors was not sufficient to impact behavioral despair on either day of the test (repeated-measures ANOVA, main effect of group between adult hetero-KO mice and controls, $F_{(1,44)} = 2.132, p = 0.1513$; group-by-time interaction, $F_{(5,44)} = 0.570, p = 0.6357$) (Fig. 7B). This demonstrates that 5-HT\_1\_A heteroreceptors act developmentally to establish the circuitry underlying this behavior, and establishes a dissociation between the roles of 5-HT\_1\_A autoreceptors and heteroreceptors in modulating mood-related anxious and nonanxious behavior, respectively.

### Discussion

#### Independent suppression of endogenous 5-HT\_1\_A autoreceptors and heteroreceptors

The 5-HT\_1\_A receptor exists as two distinct populations in the brain (autoreceptors and heteroreceptors) that, individually, could impact the circuitry underlying mood and anxiety in both development and/or adulthood. We accomplished independent manipulation of 5-HT\_1\_A autoreceptors and heteroreceptors populations using the tTS/tetO system (Mallo et al., 2003; Richardson-Jones et al., 2010). This system provides a number of advances over classic KO and previous transgenic technology. First, it allows for spatial and temporal specificity, neither of which are possible using constitutive KO mice. Given the hypothesized difference in the roles of the 5-HT system in development and adulthood, temporal specificity was a primary goal (Ansdorfe et al., 2007, 2008). Second, this genetic strategy allows for examination of the role of endogenous receptors; unlike previous transgenic overexpression strategies, the tTS/tetO strategy relies on suppressible gene expression from the endogenous promoter elements. Moreover, we have recently demonstrated that this system can be generalized to other genes (Tanaka et al., 2010).

#### Establishment of 5-HT tone

In agreement with the observation that the firing activity of serotonergic neurons and consequently the release of 5-HT is under tonic inhibitory control by somatodendritic 5-HT\_1\_A autoreceptors (Haddjeri et al., 2004), constitutive deletion of the 5-HT\_1\_A receptor increases basal extracellular 5-HT level in some brain structures (He et al., 2001; Parsons et al., 2001). This is consistent with data obtained here in auto-KO mice (Table 1). Nevertheless, the effect observed here is region-dependent, as basal 5-HT\_2\_A was increased in the frontal cortex, but not in the vHPC of auto-KO mice. Several hypotheses may explain such differences. Regional differences in 5-HT reuptake sites in mouse brain could account for these neurochemical observations. Autoradiographic studies have reported a lower density of serotonin transporter (SERT) in

---

**Figure 6.** A, No group differences were detected between hetero-KO mice and their controls in the total exploration (i) or percentage exploration of the center (ii) of the open field ($n = 12$). B, No group differences were detected between hetero-KO mice and their controls in total exploration (i), percentage exploration of the light (ii), or entries into the light (iii) in the light/dark choice test ($n = 41$). C, No group differences were detected between adult hetero-KO mice and their controls in the total exploration (i) or percentage exploration of the center (ii) of the open field ($n = 47$). D, No group differences were detected between adult hetero-KO mice and their controls in total exploration (i), percentage exploration of the light (ii), or entries into the light (iii) in the light/dark choice test ($n = 30$). E, Auto-KO mice displayed decreased total exploration (i) and decreased percentage exploration of the center (ii) of the open field compared with their controls ($n = 50$). F, Auto-KO mice displayed less total exploration (i) and fewer entries into the light compartment (iii), but no difference in percentage exploration of the light (ii) compared with their controls in the light/dark choice test ($n = 37$). All values are mean $\pm$ SEM. *$p < 0.05$, **$p < 0.01$, ***$p < 0.001$.

**Figure 7.** A, Hetero-KO mice displayed decreased mobility compared with their controls across a 2 d forced-swim test ($N = 48$). B, C, No group differences were detected between adult hetero-KO (B) or auto-KO mice (C) and their controls across a 2 d forced-swim test ($N = 34$ and 44, respectively). All values are mean $\pm$ SEM. *$p < 0.05$, **$p < 0.01$.
the frontal cortex compared with the hippocampus in constitutive 5-HT_{1A} KO mice (Ase et al., 2001). Interestingly, SERT downregulation is also observed in double 5-HT_{1A/1B} KO mice (Guilloux et al., 2011). Thus, the increase in 5-HT_{ext} observed in the frontal cortex (but not in the vHPC) of auto-KO mice may be attributed to a specific reduction of SERT expression in the frontal cortex. A second observation to be explored with regard to the present results is the marked difference in the intrinsic organization of 5-HT innervations. The frontal cortex is primarily innervated by the dorsal raphe nucleus (DRN), whereas the hippocampus is mostly innervated by the median raphe nucleus (MRN) (Azmitia and Segal, 1978; Imai et al., 1986; McQuade and Sharp, 1997). It is thus possible that the enhancement of basal extracellular 5-HT levels specifically in the frontal cortex in auto-KO mice resulted from a greater effect of autoreceptor deletion on the release of 5-HT from DRN compared with the MRN.

The magnitude of 5-HT release in the presence of transporter blockade also displayed a region-dependent difference. As expected, upon challenge with fluoxetine, 5-HT_{ext} was increased in both control and auto-KO mice (Fig. 5). However, the magnitude of the response to SSRI in auto-KO mice was greater in the vHPC than in the frontal cortex. These neurochemical effects markedly contrast with levels observed at baseline. Because of the greater density of 5-HT uptake sites in the hippocampus in mice [compared with the frontal cortex (Ase et al., 2001)], SSRIs are expected to cause a greater increase in 5-HT_{ext} levels in the hippocampus. Indeed, this was observed in the present study. In contrast with this assumption, fluoxetine (Malagie et al., 1995) and paroxetine (Romero and Artigas, 1997) have been shown to increase 5-HT_{ext} to a comparable extent in the frontal cortex and hippocampus of wild-type mice. These data can be interpreted in light of the distinct roles that 5-HT_{1A} heteroreceptors in the frontal cortex are thought to play in the negative feedback inhibition of the raphe. Specifically, frontal cortex forebrain heteroreceptors are thought to participate in feedback via polysynaptic connections to the raphe, whereas hippocampal heteroreceptors are not (Ceci et al., 1994; Assié and Koek, 1996; Casanovas et al., 1999; Hajós et al., 1999). Our results in the auto-KO mouse suggest that autoreceptor feedback in response to fluoxetine is stronger in raphe projections to hippocampus than frontal cortex, since heteroreceptor-based negative feedback plays a substantial role solely in frontal cortex.

Alternatively, the heterogeneous elevation of 5-HT_{ext} induced by fluoxetine in auto-KO mice could be due to changes in the sensitivity of terminal 5-HT_{1A} autoreceptors. Such changes have been detected in constitutive 5-HT_{1A} KO mice (Boutrel et al., 2002).

Establishment of anxiety-like behavior

Mice lacking all 5-HT_{1A} receptors (both autoreceptors and heteroreceptors) throughout life display increased anxiety-like behavior (Ramboz et al., 1998). Subsequent gain-of-function experiments, in which 5-HT_{1A} receptors were ectopically overexpressed in forebrain areas such as the cortex and striatum (in the absence of autoreceptors), reversed the increased anxiety behavior in 5-HT_{1A} KO mice (Gross et al., 2002). These findings led to the theory that endogenous 5-HT_{1A} heteroreceptors in the forebrain control the normal establishment of anxiety-like behavior (Gross and Hen, 2004; Akimova et al., 2009; Goodfellow et al., 2009; Zhang et al., 2010). A separate body of literature has implicated serotonergic tone during postnatal development in determining anxiety-like behavior, with whole-life or early postnatal blockade of the 5-HT transporter increasing anxiety in rodents (Lira et al., 2003; Ansorge et al., 2004; Jennings et al., 2006). As one of the major modulators of serotonergic tone, the 5-HT_{1A} autoreceptors may likewise participate in the developmental establishment of anxiety. Thus, both 5-HT_{1A} autoreceptors and 5-HT_{1A} heteroreceptors are well positioned to affect the circuitry underlying anxiety-like behavior.

Here we demonstrate, using a loss-of-function approach, that suppression of endogenous 5-HT_{1A} autoreceptors throughout life is sufficient to increase anxiety in the adult. Conversely, loss of endogenous heteroreceptors beginning either in the early postnatal period or in adulthood is not sufficient to impact anxiety-like behavior. These results suggest that the anxious-like phenotype of the 5-HT_{1A} KO mouse likely results from increased serotonergic neuron excitability during development. Thus, our results suggest a new framework for interpreting the role of the 5-HT_{1A} receptor in the developmental programming of anxiety behavior that de-emphasizes the role of the heteroreceptors in favor of autoreceptor-mediated serotonergic tone.

The results presented here by no means discount the role of serotonergic signaling in the developing forebrain in establishing anxiety circuitry. Indeed, forebrain neurons undergo a shift in their response to 5-HT during this time that corresponds to differential coupling of excitatory and inhibitory receptor subtypes (Béique et al., 2004; Goodfellow et al., 2009). Thus, the serotonergic system can impact anxiety circuitry both at the level of neurotransmitter release and through the relative balance of inhibitory and excitatory response in the developing forebrain. However, using our loss-of-function approach, we conclude that, under normal conditions, endogenous 5-HT_{1A} forebrain heteroreceptors are not the primary mediators of 5-HT’s effect on developing anxiety circuitry. Whether these receptors might play a more prominent role under adverse conditions (i.e., with higher 5-HT as result of stress) is beyond the scope of these experiments but remains an intriguing hypothesis (Goodfellow et al., 2009).

Finally, our results highlight the importance of distinguishing between endogenous functions of a receptor (often most clearly revealed through loss-of-function manipulations) and the ability
of a receptor to ectopically impact circuits (often most clearly revealed through gain-of-function manipulations) (Gross et al., 2002; Kusserow et al., 2004). Recent evidence suggests that the 5-HT_{1A} receptor has great utility in pharmacogenomic experiments designed to probe circuit function, but the phenotypes conferred by such manipulations should not be confused with the endogenous role of the receptor (Tsutsenis et al., 2007; Audero et al., 2008).

**Behavioral effects of heteroreceptor suppression**

In contrast to the widely recognized antidepressant effect of increasing 5-HT in the adult, several pieces of evidence suggest that increased 5-HT tone in early development can increase adult depression as well as anxiety. Although antidepressant drugs that increase extracellular 5-HT levels reliably decrease immobility in adult rodents in the forced-swim test, mice lacking 5-HT transporter activity either throughout life or only during a postnatal developmental period show increased immobility in the this test (Lira et al., 2003; Wellman et al., 2007). Moreover, mice lacking enzymes necessary for the synthesis of 5-HT throughout life (which display lower levels of 5-HT_{ext}) show decreased immobility in the forced-swim test in adulthood (Savelieva et al., 2008). Thus, the circuitry underlying responsiveness in the forced swim test response can be influenced by extracellular 5-HT levels during development. Indeed, it appears that manipulations of extracellular 5-HT levels during postnatal development result in behaviors that are directly opposite to those caused by pharmacological manipulation of serotonin levels with SSRIs in adult animals.

Here we demonstrate that mice lacking 5-HT_{1A} autoreceptors throughout life and displaying increased anxiety-like behavior nevertheless display no differences from their controls in forced-swim behavior in adulthood. In contrast, we demonstrate that mice lacking 5-HT_{1A} heteroreceptors beginning in development display decreased mobility, or increased behavioral despair, in adulthood (Fig. 8). This phenotype was not observed when heteroreceptor suppression was initiated in adulthood, suggesting that serotonergic signaling in the forebrain during development stably impacts the circuitry underlying the behavioral response to forced-swim stress without affecting context-based anxiety measures.

Together, our findings demonstrate a functional dissociation across both spatial (raphe vs forebrain) and temporal (development vs adulthood) domains within a single receptor subtype, and may be relevant to the partially overlapping set of genes and circuitry underlying anxiety- and depression-related behavior.

**References**


Richardson-Jones et al. • 5-HT_{1A} Autoreceptors and Heteroreceptors in Anxiety in J. Neurosci., April 20, 2011 • 31(16):6008–6018 • 6017


